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Method and apparatus for neuroenhancement to enhance emotional response

Abstract

A method of transplanting a desired emotional state from a donor to a recipient, comprising determining an emotional state of the donor, recording neural correlates of the emotional state of the donor who is in the desired emotional state; analyzing neural correlates of the emotional state of the donor to decode at least one of a temporal and a spatial pattern corresponding to the desirable emotional state; converting said at least one of a temporal and a spatial pattern corresponding to the desirable emotional state into a neurostimulation pattern; storing the neurostimulation pattern in the nonvolatile memory; retrieving the neurostimulation pattern from the nonvolatile memory; stimulating the recipients brain with at least one stimulus modulated with the neurostimulation pattern to induce the desired emotional state in the recipient.

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6322515	12/2000	Goor et al.	N/A	N/A
6325475	12/2000	Hayes et al.	N/A	N/A
6325761	12/2000	Jay	N/A	N/A
6331164	12/2000	Shaw et al.	N/A	N/A
6332087	12/2000	Svenson et al.	N/A	N/A
6338713	12/2001	Chamoun et al.	N/A	N/A
6339725	12/2001	Naritoku et al.	N/A	N/A
6341236	12/2001	Osorio et al.	N/A	N/A
6343229	12/2001	Siebler et al.	N/A	N/A
6354087	12/2001	Nakahara et al.	N/A	N/A
6354299	12/2001	Fischell et al.	N/A	N/A
6356079	12/2001	Mizoguchi et al.	N/A	N/A
6356781	12/2001	Lee et al.	N/A	N/A
6356788	12/2001	Boveja	N/A	N/A
6358201	12/2001	Childre et al.	N/A	N/A
6364845	12/2001	Duffy et al.	N/A	N/A
6366813	12/2001	DiLorenzo	N/A	N/A
6366814	12/2001	Boveja et al.	N/A	N/A
6370414	12/2001	Robinson	N/A	N/A
6370423	12/2001	Guerrero et al.	N/A	N/A
6374131	12/2001	Tomita et al.	N/A	N/A
6375614	12/2001	Braun et al.	N/A	N/A
6377833	12/2001	Albert	N/A	N/A
6385479	12/2001	Sibbitt et al.	N/A	N/A
6385486	12/2001	John et al.	N/A	N/A
6390979	12/2001	Njemanze	N/A	N/A
6393363	12/2001	Wilt et al.	N/A	N/A
6394963	12/2001	Blazey et al.	N/A	N/A
6402520	12/2001	Freer	N/A	N/A
6402689	12/2001	Scarantino et al.	N/A	N/A
6408107	12/2001	Miller et al.	N/A	N/A
6418344	12/2001	Rezai et al.	N/A	N/A
6419629	12/2001	Balkin et al.	N/A	N/A
6427086	12/2001	Fischell et al.	N/A	N/A
6428490	12/2001	Kramer et al.	N/A	N/A
6430443	12/2001	Karell	N/A	N/A
6435878	12/2001	Reynolds et al.	N/A	N/A
6442421	12/2001	Le Van Quyen et al.	N/A	N/A
6442948	12/2001	Takeda	N/A	N/A
6466816	12/2001	Granger et al.	N/A	N/A
6470220	12/2001	Kraus, Jr. et al.	N/A	N/A
6475163	12/2001	Smits et al.	N/A	N/A
6482165	12/2001	Patton et al.	N/A	N/A
6487441	12/2001	Swanson et al.	N/A	N/A
6488617	12/2001	Katz	N/A	N/A
6490472	12/2001	Li et al.	N/A	N/A
6493577	12/2001	Williams	N/A	N/A
6496724	12/2001	Levendowski et al.	N/A	N/A
6497658	12/2001	Roizen et al.	N/A	N/A
6497699	12/2001	Ludvig et al.	N/A	N/A
6503085	12/2002	Elkind	N/A	N/A
6507754	12/2002	Le Van Quyen et al.	N/A	N/A
6510340	12/2002	Jordan	N/A	N/A
6511424	12/2002	Moore-Ede et al.	N/A	N/A
6516246	12/2002	Derakhshan	N/A	N/A
6520905	12/2002	Surve et al.	N/A	N/A
6520921	12/2002	Patton et al.	N/A	N/A
6522906	12/2002	Salisbury, Jr. et al.	N/A	N/A
6524249	12/2002	Moehring et al.	N/A	N/A

6526297	12/2002	Merilainen	N/A	N/A
6526415	12/2002	Smith et al.	N/A	N/A
6527715	12/2002	Balkin et al.	N/A	N/A
6527730	12/2002	Blazey et al.	N/A	N/A
6529759	12/2002	Tucker et al.	N/A	N/A
6529773	12/2002	Dewan	N/A	N/A
6530884	12/2002	Balkin et al.	N/A	N/A
6534986	12/2002	Nichols	N/A	N/A
6538436	12/2002	Simola et al.	N/A	N/A
6539245	12/2002	Tsukada et al.	N/A	N/A
6539263	12/2002	Schiff et al.	N/A	N/A
6544170	12/2002	Kajihara et al.	N/A	N/A
6546378	12/2002	Cook	N/A	N/A
6547736	12/2002	Moehring et al.	N/A	N/A
6547746	12/2002	Marino	N/A	N/A
6549804	12/2002	Osorio et al.	N/A	N/A
6551243	12/2002	Bocionek et al.	N/A	N/A
6553252	12/2002	Balkin et al.	N/A	N/A
6556695	12/2002	Packer et al.	N/A	N/A
6556861	12/2002	Pritchep	N/A	N/A
6556868	12/2002	Naritoku et al.	N/A	N/A
6557558	12/2002	Tajima et al.	N/A	N/A
6560486	12/2002	Osorio et al.	N/A	N/A
6565518	12/2002	Blazey et al.	N/A	N/A
6574573	12/2002	Asano	N/A	N/A
6587727	12/2002	Osorio et al.	N/A	N/A
6587729	12/2002	O'Loughlin et al.	N/A	N/A
6591132	12/2002	Gotman et al.	N/A	N/A
6591137	12/2002	Fischell et al.	N/A	N/A
6594524	12/2002	Esteller et al.	N/A	N/A
6597954	12/2002	Pless et al.	N/A	N/A
6602202	12/2002	John et al.	N/A	N/A
6603502	12/2002	Martin et al.	N/A	N/A
6609030	12/2002	Rezai et al.	N/A	N/A
6611698	12/2002	Yamashita et al.	N/A	N/A
6615158	12/2002	Wenzel et al.	N/A	N/A
6616611	12/2002	Moehring	N/A	N/A
6622036	12/2002	Suffin	N/A	N/A
6622047	12/2002	Barrett et al.	N/A	N/A
6625485	12/2002	Levendowski et al.	N/A	N/A
6626676	12/2002	Freer	N/A	N/A
6633686	12/2002	Bakircioglu et al.	N/A	N/A
6644976	12/2002	Kullok et al.	N/A	N/A
6648822	12/2002	Hamamoto et al.	N/A	N/A
6648880	12/2002	Chauvet et al.	N/A	N/A
6650917	12/2002	Diab et al.	N/A	N/A
6652458	12/2002	Blazey et al.	N/A	N/A
6652470	12/2002	Patton et al.	N/A	N/A
6654632	12/2002	Lange et al.	N/A	N/A
6654729	12/2002	Hickman et al.	N/A	N/A
6656137	12/2002	Tyldsley et al.	N/A	N/A
6658287	12/2002	Litt et al.	N/A	N/A
6663571	12/2002	Njemanze	N/A	N/A
6665552	12/2002	Yokosawa et al.	N/A	N/A
6665553	12/2002	Kandori et al.	N/A	N/A
6665562	12/2002	Gluckman et al.	N/A	N/A
6671555	12/2002	Gielen et al.	N/A	N/A
6671556	12/2002	Osorio et al.	N/A	N/A
6678548	12/2003	Echaz et al.	N/A	N/A
6684098	12/2003	Oshio et al.	N/A	N/A
6684105	12/2003	Cohen et al.	N/A	N/A
6687525	12/2003	Llinas et al.	N/A	N/A
6695761	12/2003	Oschman et al.	N/A	N/A
6697660	12/2003	Robinson	N/A	N/A
RE38476	12/2003	Diab et al.	N/A	N/A
6699194	12/2003	Diab et al.	N/A	N/A
6701173	12/2003	Nowinski et al.	N/A	N/A
6703838	12/2003	Conti	N/A	N/A
6708051	12/2003	Durousseau	N/A	N/A
6708064	12/2003	Rezai	N/A	N/A
6708184	12/2003	Smith et al.	N/A	N/A
6709399	12/2003	Shen et al.	N/A	N/A
6725080	12/2003	Melkent et al.	N/A	N/A
6726624	12/2003	Keirsbilck et al.	N/A	N/A
6728424	12/2003	Zhu et al.	N/A	N/A

6728564	12/2003	Lahteenmaki	N/A	N/A
6731975	12/2003	Viertio-Oja et al.	N/A	N/A
6735460	12/2003	Tsukada et al.	N/A	N/A
6735467	12/2003	Wilson	N/A	N/A
6735475	12/2003	Whitehurst et al.	N/A	N/A
6740032	12/2003	Balkin et al.	N/A	N/A
6743167	12/2003	Balkin et al.	N/A	N/A
6743182	12/2003	Miller et al.	N/A	N/A
6745060	12/2003	Diab et al.	N/A	N/A
6745156	12/2003	Cook	N/A	N/A
6746409	12/2003	Keirsbilck et al.	N/A	N/A
6751499	12/2003	Lange et al.	N/A	N/A
6758813	12/2003	Meadows	N/A	N/A
6768920	12/2003	Lange et al.	N/A	N/A
6773400	12/2003	Njemanze	N/A	N/A
6774929	12/2003	Kopp	N/A	N/A
6775405	12/2003	Zhu	N/A	N/A
6782292	12/2003	Whitehurst	N/A	N/A
6785409	12/2003	Suri	N/A	N/A
6788975	12/2003	Whitehurst et al.	N/A	N/A
6791331	12/2003	Conti	N/A	N/A
6795724	12/2003	Hogan	N/A	N/A
6798898	12/2003	Fedorovskaya et al.	N/A	N/A
6801648	12/2003	Cheng	N/A	N/A
6801803	12/2003	Viertio-Oja	N/A	N/A
6804558	12/2003	Haller et al.	N/A	N/A
6804661	12/2003	Cook	N/A	N/A
6815949	12/2003	Kandori et al.	N/A	N/A
6816744	12/2003	Garfield et al.	N/A	N/A
6819956	12/2003	DiLorenzo	N/A	N/A
6826426	12/2003	Lange et al.	N/A	N/A
6843774	12/2004	Foust et al.	N/A	N/A
6853186	12/2004	Li	N/A	N/A
6856830	12/2004	He	N/A	N/A
6863127	12/2004	Clark et al.	N/A	N/A
6865494	12/2004	Duensing et al.	N/A	N/A
6873872	12/2004	Gluckman et al.	N/A	N/A
6875174	12/2004	Braun et al.	N/A	N/A
6876196	12/2004	Taulu et al.	N/A	N/A
6879859	12/2004	Boveja	N/A	N/A
6882881	12/2004	Lesser et al.	N/A	N/A
6885192	12/2004	Clarke et al.	N/A	N/A
6885886	12/2004	Bauch et al.	N/A	N/A
6886964	12/2004	Gardiner et al.	N/A	N/A
6893407	12/2004	Brooks et al.	N/A	N/A
6896655	12/2004	Patton et al.	N/A	N/A
RE38749	12/2004	Dardik	N/A	N/A
6907280	12/2004	Becerra et al.	N/A	N/A
6915241	12/2004	Kohlmorgen et al.	N/A	N/A
6920357	12/2004	Osorio et al.	N/A	N/A
6926921	12/2004	Stasiak et al.	N/A	N/A
6928354	12/2004	Ryu et al.	N/A	N/A
6931274	12/2004	Williams	N/A	N/A
6931275	12/2004	Collura	N/A	N/A
6936012	12/2004	Wells	N/A	N/A
6947790	12/2004	Gevins et al.	N/A	N/A
6950697	12/2004	Jordan	N/A	N/A
6950698	12/2004	Sarkela et al.	N/A	N/A
6959215	12/2004	Gliner et al.	N/A	N/A
6961618	12/2004	Osorio et al.	N/A	N/A
6963770	12/2004	Scarantino et al.	N/A	N/A
6963771	12/2004	Scarantino et al.	N/A	N/A
6978179	12/2004	Flagg et al.	N/A	N/A
6980863	12/2004	van Venrooij et al.	N/A	N/A
6981947	12/2005	Melker	N/A	N/A
6983184	12/2005	Price	N/A	N/A
6983264	12/2005	Shimizu	N/A	N/A
6985769	12/2005	Jordan	N/A	N/A
6988056	12/2005	Cook	N/A	N/A
6990377	12/2005	Gliner et al.	N/A	N/A
6993380	12/2005	Modarres	N/A	N/A
6996261	12/2005	deCharms	N/A	N/A
6996549	12/2005	Zhang et al.	N/A	N/A
7003352	12/2005	Whitehurst	N/A	N/A
7006872	12/2005	Gielen et al.	N/A	N/A

7010340	12/2005	Scarantino et al.	N/A	N/A
7010351	12/2005	Firlik et al.	N/A	N/A
7011410	12/2005	Bolger et al.	N/A	N/A
7011814	12/2005	Suddarth et al.	N/A	N/A
7014613	12/2005	John et al.	N/A	N/A
7016722	12/2005	Prichep	N/A	N/A
7022083	12/2005	Tanaka et al.	N/A	N/A
7023206	12/2005	Viehland et al.	N/A	N/A
7024247	12/2005	Gliner et al.	N/A	N/A
7030617	12/2005	Conti	N/A	N/A
7035686	12/2005	Hogan	N/A	N/A
7037260	12/2005	Keirsbilck et al.	N/A	N/A
7038450	12/2005	Romalis et al.	N/A	N/A
7039266	12/2005	Doty	N/A	N/A
7039547	12/2005	Wilson	N/A	N/A
7043293	12/2005	Baura	N/A	N/A
7053610	12/2005	Clarke et al.	N/A	N/A
7054454	12/2005	Causevic et al.	N/A	N/A
7062391	12/2005	Wilson	N/A	N/A
7063535	12/2005	Stamm et al.	N/A	N/A
7070571	12/2005	Kramer et al.	N/A	N/A
7079977	12/2005	Osorio et al.	N/A	N/A
7089927	12/2005	John et al.	N/A	N/A
7092748	12/2005	Valdes Sosa et al.	N/A	N/A
7099714	12/2005	Houben	N/A	N/A
7104947	12/2005	Riehl	N/A	N/A
7104963	12/2005	Melker et al.	N/A	N/A
7105824	12/2005	Stoddart et al.	N/A	N/A
7107090	12/2005	Salisbury, Jr. et al.	N/A	N/A
7116102	12/2005	Clarke et al.	N/A	N/A
7117026	12/2005	Shao et al.	N/A	N/A
7119553	12/2005	Yang et al.	N/A	N/A
7120486	12/2005	Leuthardt et al.	N/A	N/A
7123955	12/2005	Gao et al.	N/A	N/A
7127100	12/2005	Wenzel et al.	N/A	N/A
7128713	12/2005	Moehring et al.	N/A	N/A
7130673	12/2005	Tolvanen-Laakso et al.	N/A	N/A
7130675	12/2005	Ewing et al.	N/A	N/A
7130691	12/2005	Falci	N/A	N/A
7145333	12/2005	Romalis et al.	N/A	N/A
7146211	12/2005	Frei et al.	N/A	N/A
7146217	12/2005	Firlik et al.	N/A	N/A
7146218	12/2005	Esteller et al.	N/A	N/A
7149572	12/2005	Frei et al.	N/A	N/A
7149773	12/2005	Haller et al.	N/A	N/A
7150710	12/2005	Haber et al.	N/A	N/A
7150715	12/2005	Collura et al.	N/A	N/A
7150717	12/2005	Katura et al.	N/A	N/A
7150718	12/2005	Okada et al.	N/A	N/A
7151961	12/2005	Whitehurst et al.	N/A	N/A
7155279	12/2005	Whitehurst et al.	N/A	N/A
7163512	12/2006	Childre et al.	N/A	N/A
7164941	12/2006	Misczynski et al.	N/A	N/A
7167751	12/2006	Whitehurst et al.	N/A	N/A
7170294	12/2006	Kasevich	N/A	N/A
7171252	12/2006	Scarantino et al.	N/A	N/A
7171339	12/2006	Repucci et al.	N/A	N/A
7174206	12/2006	Frei et al.	N/A	N/A
7176680	12/2006	Veryaskin	N/A	N/A
7177675	12/2006	Suffin et al.	N/A	N/A
7177678	12/2006	Osorio et al.	N/A	N/A
7181505	12/2006	Haller et al.	N/A	N/A
7183381	12/2006	Varadhachary et al.	N/A	N/A
7184837	12/2006	Goetz	N/A	N/A
7186209	12/2006	Jacobson et al.	N/A	N/A
7187169	12/2006	Clarke et al.	N/A	N/A
7190826	12/2006	Russell et al.	N/A	N/A
7190995	12/2006	Chervin et al.	N/A	N/A
7193413	12/2006	Kandori et al.	N/A	N/A
7196514	12/2006	Li	N/A	N/A
7197352	12/2006	Gott et al.	N/A	N/A
7199708	12/2006	Terauchi et al.	N/A	N/A
7203548	12/2006	Whitehurst et al.	N/A	N/A
7207948	12/2006	Coyle	N/A	N/A
7209787	12/2006	DiLorenzo	N/A	N/A

7209788	12/2006	Nicolelis et al.	N/A	N/A
7212851	12/2006	Donoghue et al.	N/A	N/A
7215986	12/2006	Diab et al.	N/A	N/A
7215994	12/2006	Huiku	N/A	N/A
7218104	12/2006	Clarke et al.	N/A	N/A
7221981	12/2006	Gliner	N/A	N/A
7222964	12/2006	Gotze et al.	N/A	N/A
7224282	12/2006	Terauchi et al.	N/A	N/A
7225013	12/2006	Geva et al.	N/A	N/A
7228167	12/2006	Kara et al.	N/A	N/A
7228169	12/2006	Viertio-Oja et al.	N/A	N/A
7228171	12/2006	Lesser et al.	N/A	N/A
7228178	12/2006	Carroll et al.	N/A	N/A
7231245	12/2006	Greenwald et al.	N/A	N/A
7231254	12/2006	DiLorenzo	N/A	N/A
7236830	12/2006	Gliner	N/A	N/A
7236831	12/2006	Firlik et al.	N/A	N/A
7239731	12/2006	Semenov et al.	N/A	N/A
7239926	12/2006	Goetz	N/A	N/A
7242983	12/2006	Frei et al.	N/A	N/A
7242984	12/2006	DiLorenzo	N/A	N/A
7252090	12/2006	Goetz	N/A	N/A
7254433	12/2006	Diab et al.	N/A	N/A
7254439	12/2006	Misczynski et al.	N/A	N/A
7254500	12/2006	Makeig et al.	N/A	N/A
7257439	12/2006	Llinas	N/A	N/A
7258659	12/2006	Anninou et al.	N/A	N/A
7260430	12/2006	Wu et al.	N/A	N/A
7267644	12/2006	Thomas et al.	N/A	N/A
7267652	12/2006	Coyle et al.	N/A	N/A
7269455	12/2006	Pineda	N/A	N/A
7269456	12/2006	Collura	N/A	N/A
7269516	12/2006	Brunner et al.	N/A	N/A
7276916	12/2006	Hammer	N/A	N/A
7277758	12/2006	DiLorenzo	N/A	N/A
7278966	12/2006	Hjelt et al.	N/A	N/A
7280861	12/2006	Thomas et al.	N/A	N/A
7280867	12/2006	Frei et al.	N/A	N/A
7280870	12/2006	Nurmikko et al.	N/A	N/A
7282030	12/2006	Frei et al.	N/A	N/A
7283861	12/2006	Bystritsky	N/A	N/A
7286871	12/2006	Cohen	N/A	N/A
7288066	12/2006	Drew	N/A	N/A
7292890	12/2006	Whitehurst et al.	N/A	N/A
7295019	12/2006	Yang et al.	N/A	N/A
7297110	12/2006	Goyal et al.	N/A	N/A
7299088	12/2006	Thakor et al.	N/A	N/A
7299096	12/2006	Balzer et al.	N/A	N/A
7302298	12/2006	Lowry et al.	N/A	N/A
7305268	12/2006	Gliner et al.	N/A	N/A
7309315	12/2006	Kullok et al.	N/A	N/A
7313442	12/2006	Velasco et al.	N/A	N/A
7321837	12/2007	Osorio et al.	N/A	N/A
7324845	12/2007	Mietus et al.	N/A	N/A
7324851	12/2007	DiLorenzo	N/A	N/A
7328053	12/2007	Diab et al.	N/A	N/A
7330032	12/2007	Donnangelo	N/A	N/A
7333619	12/2007	Causevic et al.	N/A	N/A
7333851	12/2007	Echaz et al.	N/A	N/A
7334892	12/2007	Goodall et al.	N/A	N/A
7338171	12/2007	Hsieh et al.	N/A	N/A
7338455	12/2007	White et al.	N/A	N/A
7340125	12/2007	Doty	N/A	N/A
7340289	12/2007	Kandori et al.	N/A	N/A
7343198	12/2007	Behbehani et al.	N/A	N/A
7346382	12/2007	McIntyre et al.	N/A	N/A
7346395	12/2007	Lozano et al.	N/A	N/A
7353064	12/2007	Gliner et al.	N/A	N/A
7353065	12/2007	Morrell	N/A	N/A
7355597	12/2007	Laidlaw et al.	N/A	N/A
7359837	12/2007	Drew	N/A	N/A
7363164	12/2007	Little et al.	N/A	N/A
7366571	12/2007	Armstrong	N/A	N/A
7367807	12/2007	Pennebaker	N/A	N/A
7367949	12/2007	Korhonen et al.	N/A	N/A

7369896	12/2007	Gesotti	N/A	N/A
7371365	12/2007	Poduslo et al.	N/A	N/A
7373198	12/2007	Bibian et al.	N/A	N/A
7376453	12/2007	Diab et al.	N/A	N/A
7376459	12/2007	Rosenfeld	N/A	N/A
7378056	12/2007	Black	N/A	N/A
7381185	12/2007	Zhirnov et al.	N/A	N/A
7383070	12/2007	Diab et al.	N/A	N/A
7383237	12/2007	Zhang et al.	N/A	N/A
7386347	12/2007	Chung et al.	N/A	N/A
7389144	12/2007	Osorio et al.	N/A	N/A
7392079	12/2007	Donoghue et al.	N/A	N/A
7394246	12/2007	Chieh et al.	N/A	N/A
7395292	12/2007	Johnson	N/A	N/A
7396333	12/2007	Stahmann et al.	N/A	N/A
7399282	12/2007	John et al.	N/A	N/A
7400984	12/2007	Kandori et al.	N/A	N/A
7403809	12/2007	Tsukada et al.	N/A	N/A
7403814	12/2007	Cox et al.	N/A	N/A
7403815	12/2007	Katz et al.	N/A	N/A
7403820	12/2007	DiLorenzo	N/A	N/A
7407485	12/2007	Huiku	N/A	N/A
7409321	12/2007	Repucci et al.	N/A	N/A
7418290	12/2007	Devlin et al.	N/A	N/A
7420033	12/2007	Varadhachary et al.	N/A	N/A
7422555	12/2007	Zabara	N/A	N/A
7429247	12/2007	Okada et al.	N/A	N/A
7437196	12/2007	Wyler et al.	N/A	N/A
7440789	12/2007	Hannula et al.	N/A	N/A
7440806	12/2007	Whitehurst et al.	N/A	N/A
7444184	12/2007	Boveja et al.	N/A	N/A
7450986	12/2007	Nguyen et al.	N/A	N/A
7453263	12/2007	Kim et al.	N/A	N/A
7454240	12/2007	Diab et al.	N/A	N/A
7454243	12/2007	Silberstein	N/A	N/A
7454245	12/2007	Armstrong et al.	N/A	N/A
7454387	12/2007	Abercrombie et al.	N/A	N/A
7457653	12/2007	Fujimaki	N/A	N/A
7457665	12/2007	Osorio et al.	N/A	N/A
7461045	12/2007	Chaovalitwongse et al.	N/A	N/A
7462151	12/2007	Childre et al.	N/A	N/A
7462155	12/2007	England	N/A	N/A
7463024	12/2007	Simola et al.	N/A	N/A
7463142	12/2007	Lindsay	N/A	N/A
7463927	12/2007	Chaouat	N/A	N/A
7466132	12/2007	Clarke et al.	N/A	N/A
7468040	12/2007	Hartley et al.	N/A	N/A
7468350	12/2007	Gong et al.	N/A	N/A
7469697	12/2007	Lee et al.	N/A	N/A
7471971	12/2007	Diab et al.	N/A	N/A
7471978	12/2007	John et al.	N/A	N/A
7478108	12/2008	Townsend et al.	N/A	N/A
7482298	12/2008	Nepela	N/A	N/A
7483747	12/2008	Gliner et al.	N/A	N/A
7486986	12/2008	Osorio et al.	N/A	N/A
7488294	12/2008	Torch	N/A	N/A
7489958	12/2008	Diab et al.	N/A	N/A
7489964	12/2008	Suffin et al.	N/A	N/A
7490085	12/2008	Walker et al.	N/A	N/A
7491173	12/2008	Heim	N/A	N/A
7493171	12/2008	Whitehurst et al.	N/A	N/A
7493172	12/2008	Whitehurst et al.	N/A	N/A
7496393	12/2008	Diab et al.	N/A	N/A
7497828	12/2008	Wilk et al.	N/A	N/A
7499741	12/2008	Diab et al.	N/A	N/A
7499745	12/2008	Littrup et al.	N/A	N/A
7499752	12/2008	Maschino et al.	N/A	N/A
7499894	12/2008	Marom et al.	N/A	N/A
7502720	12/2008	Taulu	N/A	N/A
7509154	12/2008	Diab et al.	N/A	N/A
7509161	12/2008	Viertio-Oja	N/A	N/A
7509163	12/2008	Luo et al.	N/A	N/A
7510531	12/2008	Lee et al.	N/A	N/A
7510699	12/2008	Black et al.	N/A	N/A
7515054	12/2008	Torch	N/A	N/A

7530955	12/2008	Diab et al.	N/A	N/A
7537568	12/2008	Moehring	N/A	N/A
7539528	12/2008	Xiong et al.	N/A	N/A
7539532	12/2008	Tran	N/A	N/A
7539533	12/2008	Tran	N/A	N/A
7539543	12/2008	Schiff et al.	N/A	N/A
7547284	12/2008	Brainard, II	N/A	N/A
7553810	12/2008	Gong et al.	N/A	N/A
7558622	12/2008	Tran	N/A	N/A
7559903	12/2008	Moussavi et al.	N/A	N/A
7561918	12/2008	Armstrong et al.	N/A	N/A
7565193	12/2008	Laken	N/A	N/A
7565199	12/2008	Sheffield et al.	N/A	N/A
7565200	12/2008	Wylér et al.	N/A	N/A
7565809	12/2008	Takeda	N/A	N/A
7567693	12/2008	deCharms	N/A	N/A
7570054	12/2008	Lin	N/A	N/A
7570991	12/2008	Milgramm et al.	N/A	N/A
7572225	12/2008	Stahmann et al.	N/A	N/A
7573264	12/2008	Xu et al.	N/A	N/A
7573268	12/2008	Volegov et al.	N/A	N/A
7574007	12/2008	Shaw et al.	N/A	N/A
7574254	12/2008	Milgramm et al.	N/A	N/A
7577472	12/2008	Li et al.	N/A	N/A
7577481	12/2008	Firlik et al.	N/A	N/A
7580798	12/2008	Brunner et al.	N/A	N/A
7582062	12/2008	Magill et al.	N/A	N/A
7583857	12/2008	Xu et al.	N/A	N/A
7593767	12/2008	Modarres	N/A	N/A
7594122	12/2008	Milgramm et al.	N/A	N/A
7594889	12/2008	St. Ores et al.	N/A	N/A
7596535	12/2008	de Voir et al.	N/A	N/A
7597665	12/2008	Wilk et al.	N/A	N/A
7603168	12/2008	Bibian et al.	N/A	N/A
7603174	12/2008	De Ridder	N/A	N/A
7604603	12/2008	Sackner et al.	N/A	N/A
7606405	12/2008	Sawyer et al.	N/A	N/A
7608579	12/2008	Gong et al.	N/A	N/A
7610083	12/2008	Drew et al.	N/A	N/A
7610094	12/2008	Stahmann et al.	N/A	N/A
7610096	12/2008	McDonald, III	N/A	N/A
7610100	12/2008	Jaax et al.	N/A	N/A
7613502	12/2008	Yamamoto et al.	N/A	N/A
7613519	12/2008	De Ridder	N/A	N/A
7613520	12/2008	De Ridder	N/A	N/A
7617002	12/2008	Goetz	N/A	N/A
7618381	12/2008	Krebs et al.	N/A	N/A
7620455	12/2008	Maschino	N/A	N/A
7620456	12/2008	Gliner et al.	N/A	N/A
7623912	12/2008	Akselrod et al.	N/A	N/A
7623927	12/2008	Rezai	N/A	N/A
7623928	12/2008	DiLorenzo	N/A	N/A
7624293	12/2008	Osorio et al.	N/A	N/A
7625340	12/2008	Sarkela	N/A	N/A
7627370	12/2008	Marks	N/A	N/A
7629889	12/2008	Sachanandani et al.	N/A	N/A
7630757	12/2008	Dorfmeister et al.	N/A	N/A
7634317	12/2008	Ben-David et al.	N/A	N/A
7640055	12/2008	Geva et al.	N/A	N/A
7643655	12/2009	Liang et al.	N/A	N/A
7643881	12/2009	Armstrong	N/A	N/A
7647097	12/2009	Flaherty et al.	N/A	N/A
7647098	12/2009	Pritchep	N/A	N/A
7648498	12/2009	Hempel	N/A	N/A
7649351	12/2009	Kajola et al.	N/A	N/A
7653433	12/2009	Lozano et al.	N/A	N/A
7654948	12/2009	Kaplan et al.	N/A	N/A
7657316	12/2009	Jaax et al.	N/A	N/A
7668579	12/2009	Lynn	N/A	N/A
7668591	12/2009	Lee et al.	N/A	N/A
7670838	12/2009	Deisseroth et al.	N/A	N/A
7672707	12/2009	Takeda	N/A	N/A
7672717	12/2009	Zikov et al.	N/A	N/A
7672730	12/2009	Firlik et al.	N/A	N/A
7676263	12/2009	Harris et al.	N/A	N/A

7678047	12/2009	Shiomi et al.	N/A	N/A
7678061	12/2009	Lee et al.	N/A	N/A
7678767	12/2009	Gong et al.	N/A	N/A
7680526	12/2009	McIntyre et al.	N/A	N/A
7680540	12/2009	Jensen et al.	N/A	N/A
7684856	12/2009	Virtanen et al.	N/A	N/A
7684858	12/2009	He et al.	N/A	N/A
7684866	12/2009	Fowler et al.	N/A	N/A
7684867	12/2009	Jaax et al.	N/A	N/A
7697979	12/2009	Martinerie et al.	N/A	N/A
7702387	12/2009	Stevenson et al.	N/A	N/A
7702502	12/2009	Ricci et al.	N/A	N/A
7706871	12/2009	Devlin et al.	N/A	N/A
7706992	12/2009	Ricci et al.	N/A	N/A
7711417	12/2009	John et al.	N/A	N/A
7711432	12/2009	Thimineur et al.	N/A	N/A
7714936	12/2009	Martin et al.	N/A	N/A
7715894	12/2009	Dunseath et al.	N/A	N/A
7715910	12/2009	Hargrove et al.	N/A	N/A
7715919	12/2009	Osorio et al.	N/A	N/A
7720519	12/2009	Ruohonen	N/A	N/A
7720530	12/2009	Causevic	N/A	N/A
7725174	12/2009	Kern et al.	N/A	N/A
7725192	12/2009	Eskandar et al.	N/A	N/A
7727161	12/2009	Coyle et al.	N/A	N/A
7729740	12/2009	Kraus, Jr. et al.	N/A	N/A
7729753	12/2009	Kremliovsky et al.	N/A	N/A
7729755	12/2009	Laken	N/A	N/A
7729773	12/2009	Sloan	N/A	N/A
7733224	12/2009	Tran	N/A	N/A
7733973	12/2009	Moriya et al.	N/A	N/A
7734334	12/2009	Mietus et al.	N/A	N/A
7734340	12/2009	De Ridder	N/A	N/A
7734355	12/2009	Cohen et al.	N/A	N/A
7736382	12/2009	Webb et al.	N/A	N/A
7737687	12/2009	Na et al.	N/A	N/A
7738683	12/2009	Cahill et al.	N/A	N/A
7740592	12/2009	Graham et al.	N/A	N/A
7742820	12/2009	Wylar et al.	N/A	N/A
7746979	12/2009	Dilmanian et al.	N/A	N/A
7747318	12/2009	John et al.	N/A	N/A
7747325	12/2009	Dilorenzo	N/A	N/A
7747326	12/2009	Velasco et al.	N/A	N/A
7747551	12/2009	Snyder	N/A	N/A
7749155	12/2009	Anderson et al.	N/A	N/A
7751877	12/2009	Flaherty et al.	N/A	N/A
7751878	12/2009	Merkle et al.	N/A	N/A
7753836	12/2009	Peterchev	N/A	N/A
7754190	12/2009	Suffin	N/A	N/A
7756564	12/2009	Matsui et al.	N/A	N/A
7756568	12/2009	Scarantino et al.	N/A	N/A
7756584	12/2009	Sheffield et al.	N/A	N/A
7757690	12/2009	Stahmann et al.	N/A	N/A
7758503	12/2009	Lynn et al.	N/A	N/A
7763588	12/2009	van Praag et al.	N/A	N/A
7764987	12/2009	Dorr et al.	N/A	N/A
7765088	12/2009	Drew	N/A	N/A
7766827	12/2009	Balkin et al.	N/A	N/A
7769424	12/2009	Sato	N/A	N/A
7769431	12/2009	Scarantino et al.	N/A	N/A
7769461	12/2009	Whitehurst et al.	N/A	N/A
7769464	12/2009	Gerber et al.	N/A	N/A
7771341	12/2009	Rogers	N/A	N/A
7771364	12/2009	Arbel et al.	N/A	N/A
7774052	12/2009	Burton et al.	N/A	N/A
7774064	12/2009	Meyer et al.	N/A	N/A
7775993	12/2009	Heruth et al.	N/A	N/A
7778490	12/2009	Quist	N/A	N/A
7778692	12/2009	Scarantino et al.	N/A	N/A
7778693	12/2009	Barbour et al.	N/A	N/A
7783362	12/2009	Whitehurst et al.	N/A	N/A
7787937	12/2009	Scarantino et al.	N/A	N/A
7787946	12/2009	Stahmann et al.	N/A	N/A
7792575	12/2009	Fujimaki et al.	N/A	N/A
7794403	12/2009	Schaafsma	N/A	N/A

7794406	12/2009	Reisfeld et al.	N/A	N/A
7797040	12/2009	Pesaran et al.	N/A	N/A
7800493	12/2009	Terauchi et al.	N/A	N/A
7801591	12/2009	Shusterman	N/A	N/A
7801592	12/2009	Shan et al.	N/A	N/A
7801593	12/2009	Behbehani et al.	N/A	N/A
7801601	12/2009	Maschino et al.	N/A	N/A
7801686	12/2009	Hyde et al.	N/A	N/A
7803118	12/2009	Reisfeld et al.	N/A	N/A
7803119	12/2009	Reisfeld	N/A	N/A
7804441	12/2009	DeChiaro, Jr.	N/A	N/A
7805203	12/2009	Ben-David et al.	N/A	N/A
7809433	12/2009	Keenan	N/A	N/A
7809434	12/2009	Kofol et al.	N/A	N/A
7811279	12/2009	John	N/A	N/A
7819794	12/2009	Becker	N/A	N/A
7819812	12/2009	John et al.	N/A	N/A
7822481	12/2009	Gerber et al.	N/A	N/A
D627476	12/2009	Gaw et al.	N/A	N/A
7829562	12/2009	Shamloo et al.	N/A	N/A
7831302	12/2009	Thomas	N/A	N/A
7831305	12/2009	Gliner	N/A	N/A
7834627	12/2009	Sakai et al.	N/A	N/A
7835787	12/2009	Sajda et al.	N/A	N/A
7840039	12/2009	Fuchs	N/A	N/A
7840248	12/2009	Fuchs et al.	N/A	N/A
7840250	12/2009	Tucker	N/A	N/A
7840257	12/2009	Chance	N/A	N/A
7840280	12/2009	Parnis et al.	N/A	N/A
7841986	12/2009	He et al.	N/A	N/A
7844324	12/2009	Sarkela et al.	N/A	N/A
7848803	12/2009	Jaax et al.	N/A	N/A
7852087	12/2009	Wilt et al.	N/A	N/A
7853321	12/2009	Jaax et al.	N/A	N/A
7853322	12/2009	Bourget et al.	N/A	N/A
7853323	12/2009	Goetz	N/A	N/A
7853329	12/2009	DiLorenzo	N/A	N/A
7856264	12/2009	Firlik et al.	N/A	N/A
7860548	12/2009	McIntyre et al.	N/A	N/A
7860552	12/2009	Borsook et al.	N/A	N/A
7860561	12/2009	Modarres	N/A	N/A
7860570	12/2009	Whitehurst et al.	N/A	N/A
7863272	12/2010	Oksenberg et al.	N/A	N/A
7865234	12/2010	Modarres	N/A	N/A
7865235	12/2010	Le et al.	N/A	N/A
7865244	12/2010	Giftakis et al.	N/A	N/A
7869867	12/2010	Armstrong et al.	N/A	N/A
7869884	12/2010	Scott et al.	N/A	N/A
7869885	12/2010	Begnaud et al.	N/A	N/A
7872235	12/2010	Rousso et al.	N/A	N/A
7873411	12/2010	Eda et al.	N/A	N/A
7876938	12/2010	Huang et al.	N/A	N/A
7878965	12/2010	Haber et al.	N/A	N/A
7879043	12/2010	Meneghini et al.	N/A	N/A
7881760	12/2010	Matsui et al.	N/A	N/A
7881770	12/2010	Melkent et al.	N/A	N/A
7881780	12/2010	Flaherty	N/A	N/A
7882135	12/2010	Brunner et al.	N/A	N/A
7884101	12/2010	Teegarden et al.	N/A	N/A
7887493	12/2010	Stahmann et al.	N/A	N/A
7890155	12/2010	Burns et al.	N/A	N/A
7890176	12/2010	Jaax et al.	N/A	N/A
7890185	12/2010	Cohen et al.	N/A	N/A
7891814	12/2010	Harada et al.	N/A	N/A
7892764	12/2010	Xiong et al.	N/A	N/A
7894890	12/2010	Sun et al.	N/A	N/A
7894903	12/2010	John	N/A	N/A
7895033	12/2010	Joublin et al.	N/A	N/A
7896807	12/2010	Clancy et al.	N/A	N/A
7899524	12/2010	Kozel	N/A	N/A
7899525	12/2010	John et al.	N/A	N/A
7899539	12/2010	Whitehurst et al.	N/A	N/A
7899545	12/2010	John	N/A	N/A
7901211	12/2010	Pennebaker	N/A	N/A
7904134	12/2010	McIntyre et al.	N/A	N/A

7904139	12/2010	Chance	N/A	N/A
7904144	12/2010	Causevic et al.	N/A	N/A
7904151	12/2010	Ben-David et al.	N/A	N/A
7904175	12/2010	Scott et al.	N/A	N/A
7904507	12/2010	Jung et al.	N/A	N/A
7907994	12/2010	Stolarski et al.	N/A	N/A
7907998	12/2010	Arad	N/A	N/A
7908008	12/2010	Ben-David et al.	N/A	N/A
7908009	12/2010	Wylér et al.	N/A	N/A
7909771	12/2010	Meyer et al.	N/A	N/A
7912530	12/2010	Seki et al.	N/A	N/A
7917199	12/2010	Drew et al.	N/A	N/A
7917206	12/2010	Frei et al.	N/A	N/A
7917221	12/2010	Tass	N/A	N/A
7917225	12/2010	Wylér et al.	N/A	N/A
7918779	12/2010	Haber et al.	N/A	N/A
7920914	12/2010	Shieh et al.	N/A	N/A
7920915	12/2010	Mann et al.	N/A	N/A
7920916	12/2010	Johnson et al.	N/A	N/A
7925353	12/2010	Whitehurst et al.	N/A	N/A
7929693	12/2010	Terauchi et al.	N/A	N/A
7930035	12/2010	DiLorenzo	N/A	N/A
7932225	12/2010	Gong et al.	N/A	N/A
7933645	12/2010	Strychacz et al.	N/A	N/A
7933646	12/2010	Frei et al.	N/A	N/A
7933727	12/2010	Taulu et al.	N/A	N/A
7937138	12/2010	Liley	N/A	N/A
7937152	12/2010	Lozano	N/A	N/A
7937222	12/2010	Donadille et al.	N/A	N/A
7938782	12/2010	Stahmann et al.	N/A	N/A
7938785	12/2010	Aguilar et al.	N/A	N/A
7941209	12/2010	Hughes et al.	N/A	N/A
7942824	12/2010	Kayyali et al.	N/A	N/A
7944551	12/2010	Addison et al.	N/A	N/A
7945304	12/2010	Feinberg	N/A	N/A
7945316	12/2010	Giftakis et al.	N/A	N/A
7945330	12/2010	Gliner et al.	N/A	N/A
7957796	12/2010	Maschino	N/A	N/A
7957797	12/2010	Bourget et al.	N/A	N/A
7957806	12/2010	Stevenson et al.	N/A	N/A
7957809	12/2010	Bourget et al.	N/A	N/A
7961922	12/2010	Spence et al.	N/A	N/A
7962204	12/2010	Suffin et al.	N/A	N/A
7962214	12/2010	Byerman et al.	N/A	N/A
7962219	12/2010	Jaax et al.	N/A	N/A
7962220	12/2010	Kolafa et al.	N/A	N/A
7970734	12/2010	Townsend et al.	N/A	N/A
7972278	12/2010	Graham et al.	N/A	N/A
7974688	12/2010	Armstrong et al.	N/A	N/A
7974693	12/2010	Ben-David et al.	N/A	N/A
7974696	12/2010	DiLorenzo	N/A	N/A
7974697	12/2010	Maschino et al.	N/A	N/A
7974701	12/2010	Armstrong	N/A	N/A
7974787	12/2010	Hyde et al.	N/A	N/A
7976465	12/2010	Frei et al.	N/A	N/A
7983740	12/2010	Culver et al.	N/A	N/A
7983741	12/2010	Chance	N/A	N/A
7983757	12/2010	Miyazawa et al.	N/A	N/A
7983762	12/2010	Gliner et al.	N/A	N/A
7986991	12/2010	Pritchep	N/A	N/A
7988613	12/2010	Becker	N/A	N/A
7988969	12/2010	Poduslo et al.	N/A	N/A
7991461	12/2010	Flaherty et al.	N/A	N/A
7991477	12/2010	McDonald, III	N/A	N/A
7993279	12/2010	Hartley et al.	N/A	N/A
7996075	12/2010	Korzinov et al.	N/A	N/A
7996079	12/2010	Armstrong	N/A	N/A
8000767	12/2010	Eden et al.	N/A	N/A
8000773	12/2010	Rouso et al.	N/A	N/A
8000788	12/2010	Giftakis et al.	N/A	N/A
8000793	12/2010	Libbus	N/A	N/A
8000794	12/2010	Lozano	N/A	N/A
8000795	12/2010	Lozano	N/A	N/A
8001179	12/2010	Jung et al.	N/A	N/A
8002553	12/2010	Hatlestad et al.	N/A	N/A

8005534	12/2010	Greenwald et al.	N/A	N/A
8005624	12/2010	Starr	N/A	N/A
8005894	12/2010	Jung et al.	N/A	N/A
8010178	12/2010	Seki et al.	N/A	N/A
8010347	12/2010	Ricci et al.	N/A	N/A
8012107	12/2010	Einav et al.	N/A	N/A
8014847	12/2010	Shastri et al.	N/A	N/A
8014870	12/2010	Seidman	N/A	N/A
8016597	12/2010	Becker et al.	N/A	N/A
8019400	12/2010	Diab et al.	N/A	N/A
8019410	12/2010	Bharmi et al.	N/A	N/A
8024029	12/2010	Drew et al.	N/A	N/A
8024032	12/2010	Osorio et al.	N/A	N/A
8025404	12/2010	Bolger et al.	N/A	N/A
8027730	12/2010	John	N/A	N/A
8029553	12/2010	Nemenov	N/A	N/A
8031076	12/2010	Sachanandani et al.	N/A	N/A
8032209	12/2010	He et al.	N/A	N/A
8032229	12/2010	Gerber et al.	N/A	N/A
8032486	12/2010	Townsend et al.	N/A	N/A
8033996	12/2010	Behar	N/A	N/A
8036434	12/2010	Hewett et al.	N/A	N/A
8036728	12/2010	Diab et al.	N/A	N/A
8036736	12/2010	Snyder et al.	N/A	N/A
8036745	12/2010	Ben-David et al.	N/A	N/A
8041136	12/2010	Causevic	N/A	N/A
8041418	12/2010	Giftakis et al.	N/A	N/A
8041419	12/2010	Giftakis et al.	N/A	N/A
8046041	12/2010	Diab et al.	N/A	N/A
8046042	12/2010	Diab et al.	N/A	N/A
8046076	12/2010	Whitehurst et al.	N/A	N/A
8050768	12/2010	Firlik et al.	N/A	N/A
8055348	12/2010	Heruth et al.	N/A	N/A
8055591	12/2010	Jung et al.	N/A	N/A
8059879	12/2010	Tsukimoto	N/A	N/A
8060181	12/2010	Rodriguez Ponce et al.	N/A	N/A
8060194	12/2010	Flaherty	N/A	N/A
8064994	12/2010	Pardo et al.	N/A	N/A
8065011	12/2010	Echaz et al.	N/A	N/A
8065012	12/2010	Firlik et al.	N/A	N/A
8065017	12/2010	Cornejo Cruz et al.	N/A	N/A
8065240	12/2010	Jung et al.	N/A	N/A
8065360	12/2010	Jung et al.	N/A	N/A
8066637	12/2010	Childre et al.	N/A	N/A
8066647	12/2010	Armitstead	N/A	N/A
8068904	12/2010	Sun et al.	N/A	N/A
8068911	12/2010	Giftakis et al.	N/A	N/A
8069125	12/2010	Jung et al.	N/A	N/A
8073534	12/2010	Low	N/A	N/A
8073546	12/2010	Sheffield et al.	N/A	N/A
8073631	12/2010	Wilber et al.	N/A	N/A
8075499	12/2010	Nathan et al.	N/A	N/A
8079953	12/2010	Braun et al.	N/A	N/A
8082031	12/2010	Ochs	N/A	N/A
8082033	12/2010	Rezai et al.	N/A	N/A
8082215	12/2010	Jung et al.	N/A	N/A
8083786	12/2010	Gafni et al.	N/A	N/A
8086294	12/2010	Echaz et al.	N/A	N/A
8086296	12/2010	Bystritsky	N/A	N/A
8086563	12/2010	Jung et al.	N/A	N/A
8088057	12/2011	Honeycutt et al.	N/A	N/A
8089283	12/2011	Kaplan et al.	N/A	N/A
8090164	12/2011	Bullitt et al.	N/A	N/A
8092549	12/2011	Hillis et al.	N/A	N/A
8095209	12/2011	Flaherty	N/A	N/A
8095210	12/2011	Burdick et al.	N/A	N/A
8097926	12/2011	De Graff et al.	N/A	N/A
8099299	12/2011	Sirohey et al.	N/A	N/A
8103333	12/2011	Tran	N/A	N/A
8108033	12/2011	Drew et al.	N/A	N/A
8108036	12/2011	Tran	N/A	N/A
8108038	12/2011	Giftakis et al.	N/A	N/A
8108039	12/2011	Saliga et al.	N/A	N/A
8108042	12/2011	Johnson et al.	N/A	N/A
8112148	12/2011	Giftakis et al.	N/A	N/A

8112153	12/2011	Giftakis et al.	N/A	N/A
8114021	12/2011	Robertson et al.	N/A	N/A
8116874	12/2011	Tass	N/A	N/A
8116877	12/2011	Lozano	N/A	N/A
8116883	12/2011	Williams et al.	N/A	N/A
8121361	12/2011	Ernst et al.	N/A	N/A
8121673	12/2011	Tran	N/A	N/A
8121694	12/2011	Molnar et al.	N/A	N/A
8121695	12/2011	Gliner et al.	N/A	N/A
8126228	12/2011	Fueyo et al.	N/A	N/A
8126243	12/2011	Hamada et al.	N/A	N/A
8126528	12/2011	Diab et al.	N/A	N/A
8126542	12/2011	Grey	N/A	N/A
8126567	12/2011	Gerber et al.	N/A	N/A
8126568	12/2011	Gliner	N/A	N/A
8128572	12/2011	Diab et al.	N/A	N/A
8131354	12/2011	Arad	N/A	N/A
8131526	12/2011	Neville	N/A	N/A
8133172	12/2011	Shachar et al.	N/A	N/A
8135472	12/2011	Fowler et al.	N/A	N/A
8135957	12/2011	Dinges et al.	N/A	N/A
8137269	12/2011	Sheikhzadeh-Nadjar et al.	N/A	N/A
8137270	12/2011	Keenan et al.	N/A	N/A
8140152	12/2011	John et al.	N/A	N/A
8145295	12/2011	Boyden et al.	N/A	N/A
8145310	12/2011	Dong et al.	N/A	N/A
8148417	12/2011	Teegarden et al.	N/A	N/A
8148418	12/2011	Teegarden et al.	N/A	N/A
8150508	12/2011	Craig	N/A	N/A
8150523	12/2011	Schiff et al.	N/A	N/A
8150524	12/2011	Maschino et al.	N/A	N/A
8150796	12/2011	Jung et al.	N/A	N/A
8152732	12/2011	Lynn et al.	N/A	N/A
8155726	12/2011	Seki et al.	N/A	N/A
8155736	12/2011	Sullivan et al.	N/A	N/A
8160273	12/2011	Visser et al.	N/A	N/A
8160317	12/2011	Amunts et al.	N/A	N/A
8160680	12/2011	Boyden et al.	N/A	N/A
8160689	12/2011	Jadidi	N/A	N/A
8160696	12/2011	Bendett et al.	N/A	N/A
8165687	12/2011	Cornejo Cruz et al.	N/A	N/A
8167784	12/2011	Honeycutt et al.	N/A	N/A
8167826	12/2011	Oohashi et al.	N/A	N/A
8170315	12/2011	Mistretta et al.	N/A	N/A
8170347	12/2011	Ancelin	N/A	N/A
8172759	12/2011	Bukhman	N/A	N/A
8172766	12/2011	Kayyali et al.	N/A	N/A
8174430	12/2011	DeChiaro, Jr.	N/A	N/A
8175359	12/2011	O'Halloran et al.	N/A	N/A
8175360	12/2011	Razifar et al.	N/A	N/A
8175686	12/2011	Utsugi et al.	N/A	N/A
8175696	12/2011	Liley et al.	N/A	N/A
8175700	12/2011	Johnson et al.	N/A	N/A
8177724	12/2011	Derchak et al.	N/A	N/A
8177726	12/2011	John	N/A	N/A
8177727	12/2011	Kwak	N/A	N/A
8180125	12/2011	Avinash et al.	N/A	N/A
8180148	12/2011	Cover et al.	N/A	N/A
8180420	12/2011	Diab et al.	N/A	N/A
8180436	12/2011	Boyden et al.	N/A	N/A
8180601	12/2011	Butson et al.	N/A	N/A
8185186	12/2011	Ross et al.	N/A	N/A
8185207	12/2011	Molnar et al.	N/A	N/A
8185382	12/2011	Joublin et al.	N/A	N/A
8187181	12/2011	Osorio et al.	N/A	N/A
8187201	12/2011	Lynn	N/A	N/A
8188749	12/2011	Wilt et al.	N/A	N/A
8190227	12/2011	Diab et al.	N/A	N/A
8190248	12/2011	Besio et al.	N/A	N/A
8190249	12/2011	Gharieb et al.	N/A	N/A
8190251	12/2011	Molnar et al.	N/A	N/A
8190264	12/2011	Lozano et al.	N/A	N/A
8195295	12/2011	Stevenson et al.	N/A	N/A
8195298	12/2011	Lozano	N/A	N/A
8195300	12/2011	Gliner et al.	N/A	N/A

8195593	12/2011	Jung et al.	N/A	N/A
8197395	12/2011	Jassemidis et al.	N/A	N/A
8197437	12/2011	Kalafut et al.	N/A	N/A
8199982	12/2011	Fueyo et al.	N/A	N/A
8199985	12/2011	Jakobsson et al.	N/A	N/A
8200319	12/2011	Pu et al.	N/A	N/A
8200340	12/2011	Skelton et al.	N/A	N/A
8204583	12/2011	Sackellares et al.	N/A	N/A
8204603	12/2011	Maschino	N/A	N/A
8209009	12/2011	Giftakis et al.	N/A	N/A
8209018	12/2011	Osorio et al.	N/A	N/A
8209019	12/2011	Giftakis et al.	N/A	N/A
8209224	12/2011	Pradeep et al.	N/A	N/A
8211035	12/2011	Melker et al.	N/A	N/A
8212556	12/2011	Schwindt et al.	N/A	N/A
8213670	12/2011	Lai	N/A	N/A
8214007	12/2011	Baker et al.	N/A	N/A
8214035	12/2011	Giftakis et al.	N/A	N/A
8219188	12/2011	Craig	N/A	N/A
8221330	12/2011	Sarkela et al.	N/A	N/A
8222378	12/2011	Masure	N/A	N/A
8223023	12/2011	Sachanandani et al.	N/A	N/A
8224431	12/2011	Drew	N/A	N/A
8224433	12/2011	Suffin et al.	N/A	N/A
8224444	12/2011	Ben-David et al.	N/A	N/A
8224451	12/2011	Jaax et al.	N/A	N/A
8229540	12/2011	Sami et al.	N/A	N/A
8229559	12/2011	Westendorp et al.	N/A	N/A
8233682	12/2011	Fessler et al.	N/A	N/A
8233689	12/2011	Razifar et al.	N/A	N/A
8233965	12/2011	Bjornerud et al.	N/A	N/A
8233990	12/2011	Goetz	N/A	N/A
8235907	12/2011	Wilk et al.	N/A	N/A
8236005	12/2011	Meneghini et al.	N/A	N/A
8236038	12/2011	Nofzinger	N/A	N/A
8239014	12/2011	Ochs	N/A	N/A
8239028	12/2011	Scott	N/A	N/A
8239029	12/2011	De Ridder	N/A	N/A
8239030	12/2011	Hagedorn et al.	N/A	N/A
8241213	12/2011	Lynn et al.	N/A	N/A
8244340	12/2011	Wu et al.	N/A	N/A
8244341	12/2011	Hinrikus et al.	N/A	N/A
8244347	12/2011	Lozano	N/A	N/A
8244475	12/2011	Aguilar et al.	N/A	N/A
8244552	12/2011	Firminger et al.	N/A	N/A
8244553	12/2011	Firminger et al.	N/A	N/A
8248069	12/2011	Buracas	N/A	N/A
8249316	12/2011	Hu et al.	N/A	N/A
8249698	12/2011	Mugler et al.	N/A	N/A
8249718	12/2011	Skelton et al.	N/A	N/A
8249815	12/2011	Taylor	N/A	N/A
8260426	12/2011	Armstrong et al.	N/A	N/A
8262714	12/2011	Hulvershorn et al.	N/A	N/A
8263574	12/2011	Schaller et al.	N/A	N/A
8267851	12/2011	Kroll	N/A	N/A
8270814	12/2011	Pradeep et al.	N/A	N/A
8271077	12/2011	Rotenberg	N/A	N/A
8280502	12/2011	Hargrove et al.	N/A	N/A
8280503	12/2011	Linderman	N/A	N/A
8280505	12/2011	Craig	N/A	N/A
8280514	12/2011	Lozano et al.	N/A	N/A
8280517	12/2011	Skelton et al.	N/A	N/A
8285351	12/2011	Johnson et al.	N/A	N/A
8285368	12/2011	Chen et al.	N/A	N/A
8290575	12/2011	Tarassenko et al.	N/A	N/A
8290596	12/2011	Wei et al.	N/A	N/A
8295914	12/2011	Kalafut et al.	N/A	N/A
8295934	12/2011	Leyde	N/A	N/A
8295935	12/2011	Okun et al.	N/A	N/A
8296108	12/2011	Tanaka	N/A	N/A
8298078	12/2011	Sutton et al.	N/A	N/A
8298140	12/2011	Beck-Nielsen et al.	N/A	N/A
8301222	12/2011	Rongen et al.	N/A	N/A
8301232	12/2011	Albert et al.	N/A	N/A
8301233	12/2011	Zhang et al.	N/A	N/A

8301257	12/2011	Hsu et al.	N/A	N/A
8303636	12/2011	Schiffer	N/A	N/A
8304246	12/2011	Cook et al.	N/A	N/A
8305078	12/2011	Savukov et al.	N/A	N/A
8306607	12/2011	Levi et al.	N/A	N/A
8306610	12/2011	Mirow	N/A	N/A
8306627	12/2011	Armstrong	N/A	N/A
8308646	12/2011	Belohlavek et al.	N/A	N/A
8308661	12/2011	Miesel et al.	N/A	N/A
8311622	12/2011	Snyder et al.	N/A	N/A
8311747	12/2011	Taylor	N/A	N/A
8311748	12/2011	Taylor et al.	N/A	N/A
8311750	12/2011	Taylor	N/A	N/A
8313441	12/2011	Dalton	N/A	N/A
8314707	12/2011	Kobetski et al.	N/A	N/A
8315703	12/2011	Lozano	N/A	N/A
8315704	12/2011	Jaax et al.	N/A	N/A
8315710	12/2011	Skelton et al.	N/A	N/A
8315812	12/2011	Taylor	N/A	N/A
8315813	12/2011	Taylor et al.	N/A	N/A
8315814	12/2011	Taylor et al.	N/A	N/A
8315962	12/2011	Horne	N/A	N/A
8315970	12/2011	Zalay et al.	N/A	N/A
8320649	12/2011	Shahaf et al.	N/A	N/A
8321150	12/2011	Taylor	N/A	N/A
8323188	12/2011	Tran	N/A	N/A
8323189	12/2011	Tran et al.	N/A	N/A
8323204	12/2011	Stahmann et al.	N/A	N/A
8326418	12/2011	Sommer et al.	N/A	N/A
8326420	12/2011	Skelton et al.	N/A	N/A
8326433	12/2011	Blum et al.	N/A	N/A
8328718	12/2011	Tran	N/A	N/A
8332017	12/2011	Tarassenko et al.	N/A	N/A
8332024	12/2011	Rapoport et al.	N/A	N/A
8332038	12/2011	Heruth et al.	N/A	N/A
8332041	12/2011	Skelton et al.	N/A	N/A
8332191	12/2011	Rosthal et al.	N/A	N/A
8334690	12/2011	Kitching et al.	N/A	N/A
8335561	12/2011	Modarres	N/A	N/A
8335664	12/2011	Eberle	N/A	N/A
8335715	12/2011	Pradeep et al.	N/A	N/A
8335716	12/2011	Pradeep et al.	N/A	N/A
8337404	12/2011	Osorio	N/A	N/A
8340752	12/2011	Cox et al.	N/A	N/A
8340753	12/2011	Hardt	N/A	N/A
8340771	12/2011	Thimineur et al.	N/A	N/A
8343026	12/2012	Gardiner et al.	N/A	N/A
8343027	12/2012	DiMino et al.	N/A	N/A
8343066	12/2012	Eagleman et al.	N/A	N/A
8346331	12/2012	Bunce et al.	N/A	N/A
8346342	12/2012	Kalafut	N/A	N/A
8346349	12/2012	Gutttag et al.	N/A	N/A
8346354	12/2012	Hyde et al.	N/A	N/A
8346365	12/2012	Lozano	N/A	N/A
8350804	12/2012	Moll	N/A	N/A
8352023	12/2012	John et al.	N/A	N/A
8352031	12/2012	Rousso et al.	N/A	N/A
8353837	12/2012	John et al.	N/A	N/A
8354438	12/2012	Chez	N/A	N/A
8354881	12/2012	Denison	N/A	N/A
8355768	12/2012	Masmanidis et al.	N/A	N/A
8356004	12/2012	Jung et al.	N/A	N/A
8356594	12/2012	Ujhazy et al.	N/A	N/A
8358818	12/2012	Miga et al.	N/A	N/A
8359080	12/2012	Diab et al.	N/A	N/A
8362780	12/2012	Rosthal et al.	N/A	N/A
8364226	12/2012	Diab et al.	N/A	N/A
8364254	12/2012	Jacquín et al.	N/A	N/A
8364255	12/2012	Isenhardt et al.	N/A	N/A
8364271	12/2012	De Ridder	N/A	N/A
8364272	12/2012	Goetz	N/A	N/A
8369940	12/2012	Sun et al.	N/A	N/A
8374411	12/2012	Ernst et al.	N/A	N/A
8374412	12/2012	Kimura	N/A	N/A
8374690	12/2012	Ma	N/A	N/A

8374696	12/2012	Sanchez et al.	N/A	N/A
8374701	12/2012	Hyde et al.	N/A	N/A
8374703	12/2012	Imran	N/A	N/A
8376965	12/2012	Schuette et al.	N/A	N/A
8379947	12/2012	Garg et al.	N/A	N/A
8379952	12/2012	McIntyre et al.	N/A	N/A
8380289	12/2012	Zellers et al.	N/A	N/A
8380290	12/2012	Scarantino et al.	N/A	N/A
8380296	12/2012	Lee et al.	N/A	N/A
8380314	12/2012	Panken et al.	N/A	N/A
8380316	12/2012	Hagedorn et al.	N/A	N/A
8380658	12/2012	Jung et al.	N/A	N/A
8382667	12/2012	Osorio	N/A	N/A
8386188	12/2012	Taylor et al.	N/A	N/A
8386244	12/2012	Ricci et al.	N/A	N/A
8386312	12/2012	Pradeep et al.	N/A	N/A
8386313	12/2012	Pradeep et al.	N/A	N/A
RE44097	12/2012	Wilber et al.	N/A	N/A
8388529	12/2012	Fueyo et al.	N/A	N/A
8388530	12/2012	Shusterman	N/A	N/A
8388555	12/2012	Panken et al.	N/A	N/A
8391942	12/2012	Benni	N/A	N/A
8391956	12/2012	Zellers et al.	N/A	N/A
8391966	12/2012	Luo et al.	N/A	N/A
8392250	12/2012	Pradeep et al.	N/A	N/A
8392251	12/2012	Pradeep et al.	N/A	N/A
8392253	12/2012	Pradeep et al.	N/A	N/A
8392254	12/2012	Pradeep et al.	N/A	N/A
8392255	12/2012	Pradeep et al.	N/A	N/A
8396542	12/2012	Johnson et al.	N/A	N/A
8396545	12/2012	Berridge et al.	N/A	N/A
8396546	12/2012	Hirata et al.	N/A	N/A
8396557	12/2012	DiLorenzo	N/A	N/A
8396565	12/2012	Singhal et al.	N/A	N/A
8396744	12/2012	Pradeep et al.	N/A	N/A
8398692	12/2012	Deisseroth et al.	N/A	N/A
8401624	12/2012	Govari	N/A	N/A
8401626	12/2012	Mietus et al.	N/A	N/A
8401634	12/2012	Whitehurst et al.	N/A	N/A
8401654	12/2012	Foster et al.	N/A	N/A
8401655	12/2012	De Ridder	N/A	N/A
8401666	12/2012	Skelton et al.	N/A	N/A
8403848	12/2012	Mietus et al.	N/A	N/A
8406838	12/2012	Kato	N/A	N/A
8406841	12/2012	Lin et al.	N/A	N/A
8406848	12/2012	Wu et al.	N/A	N/A
8406862	12/2012	Hopenfeld	N/A	N/A
8406890	12/2012	Goetz	N/A	N/A
8412334	12/2012	Whitehurst et al.	N/A	N/A
8412335	12/2012	Gliner et al.	N/A	N/A
8412337	12/2012	Lozano	N/A	N/A
8412338	12/2012	Faltys	N/A	N/A
8412655	12/2012	Colman et al.	N/A	N/A
8415123	12/2012	Pilla et al.	N/A	N/A
8417344	12/2012	Colborn et al.	N/A	N/A
8423118	12/2012	Wenzel et al.	N/A	N/A
8423125	12/2012	Rousso et al.	N/A	N/A
8423144	12/2012	Tass et al.	N/A	N/A
8423155	12/2012	Jaax et al.	N/A	N/A
8423297	12/2012	Wilber	N/A	N/A
8425415	12/2012	Tran	N/A	N/A
8425583	12/2012	Nofzinger	N/A	N/A
8428696	12/2012	Foo	N/A	N/A
8428703	12/2012	Hopenfeld	N/A	N/A
8428704	12/2012	Johnson et al.	N/A	N/A
8428726	12/2012	Ignagni et al.	N/A	N/A
8429225	12/2012	Jung et al.	N/A	N/A
8430805	12/2012	Burnett et al.	N/A	N/A
8430816	12/2012	Avinash et al.	N/A	N/A
8431537	12/2012	Gong et al.	N/A	N/A
8433388	12/2012	Blunt et al.	N/A	N/A
8433410	12/2012	Stevenson et al.	N/A	N/A
8433414	12/2012	Gliner et al.	N/A	N/A
8433418	12/2012	DeRidder	N/A	N/A
8435166	12/2012	Burnett et al.	N/A	N/A

8437843	12/2012	Kayyali et al.	N/A	N/A
8437844	12/2012	Syed Momen et al.	N/A	N/A
8437861	12/2012	Skelton et al.	N/A	N/A
8439845	12/2012	Folkerts et al.	N/A	N/A
8442626	12/2012	Zavoronkovs et al.	N/A	N/A
8444571	12/2012	Folkerts et al.	N/A	N/A
8445021	12/2012	Akhtari et al.	N/A	N/A
8445851	12/2012	Rousso et al.	N/A	N/A
8447392	12/2012	Llinas	N/A	N/A
8447407	12/2012	Talathi et al.	N/A	N/A
8447411	12/2012	Skelton et al.	N/A	N/A
8449471	12/2012	Tran	N/A	N/A
8452387	12/2012	Osorio et al.	N/A	N/A
8452544	12/2012	Hymel	N/A	N/A
8454555	12/2012	Struijk et al.	N/A	N/A
8456164	12/2012	Subbarao	N/A	N/A
8456166	12/2012	DePavia et al.	N/A	N/A
8456309	12/2012	Sachanandani et al.	N/A	N/A
8457730	12/2012	Makinen	N/A	N/A
8457746	12/2012	Libbus	N/A	N/A
8457747	12/2012	Terry, Jr.	N/A	N/A
8461988	12/2012	Tran	N/A	N/A
8463006	12/2012	Prokoski	N/A	N/A
8463007	12/2012	Steinberg et al.	N/A	N/A
8463349	12/2012	Diab et al.	N/A	N/A
8463370	12/2012	Korhonen et al.	N/A	N/A
8463374	12/2012	Hudson et al.	N/A	N/A
8463378	12/2012	Tass	N/A	N/A
8463386	12/2012	Tass	N/A	N/A
8463387	12/2012	De Ridder	N/A	N/A
8464288	12/2012	Pradeep et al.	N/A	N/A
8465408	12/2012	Phillips et al.	N/A	N/A
8467877	12/2012	Imran	N/A	N/A
8467878	12/2012	Lozano et al.	N/A	N/A
8473024	12/2012	Causevic et al.	N/A	N/A
8473044	12/2012	Lee et al.	N/A	N/A
8473306	12/2012	Seely	N/A	N/A
8473345	12/2012	Pradeep et al.	N/A	N/A
8475354	12/2012	Phillips et al.	N/A	N/A
8475368	12/2012	Tran et al.	N/A	N/A
8475371	12/2012	Derchak et al.	N/A	N/A
8475387	12/2012	Derchak et al.	N/A	N/A
8475506	12/2012	Bendett et al.	N/A	N/A
8478389	12/2012	Brockway et al.	N/A	N/A
8478394	12/2012	Prichep et al.	N/A	N/A
8478402	12/2012	Wahlstrand et al.	N/A	N/A
8478417	12/2012	Drew et al.	N/A	N/A
8478428	12/2012	Cowley	N/A	N/A
8480554	12/2012	Phillips et al.	N/A	N/A
8483795	12/2012	Okada	N/A	N/A
8483815	12/2012	Liley	N/A	N/A
8483816	12/2012	Payton et al.	N/A	N/A
8484081	12/2012	Pradeep et al.	N/A	N/A
8484270	12/2012	Kurtz et al.	N/A	N/A
8485979	12/2012	Giftakis et al.	N/A	N/A
8487760	12/2012	Kangas et al.	N/A	N/A
8489185	12/2012	Kilgard et al.	N/A	N/A
8492336	12/2012	Masure	N/A	N/A
8494610	12/2012	Pradeep et al.	N/A	N/A
8494829	12/2012	Teixeira	N/A	N/A
8494857	12/2012	Pakhomov	N/A	N/A
8494905	12/2012	Pradeep et al.	N/A	N/A
8496594	12/2012	Taylor et al.	N/A	N/A
8498697	12/2012	Yong et al.	N/A	N/A
8498699	12/2012	Wells et al.	N/A	N/A
8498708	12/2012	Bentwich	N/A	N/A
RE44408	12/2012	Lindsay	N/A	N/A
8500282	12/2012	Bolger et al.	N/A	N/A
8500636	12/2012	Tran	N/A	N/A
8504150	12/2012	Skelton	N/A	N/A
8506469	12/2012	Dietrich et al.	N/A	N/A
8509879	12/2012	Durkin et al.	N/A	N/A
8509881	12/2012	Thiagarajan et al.	N/A	N/A
8509885	12/2012	Snyder et al.	N/A	N/A
8509904	12/2012	Rickert et al.	N/A	N/A

8512219	12/2012	Ferren et al.	N/A	N/A
8512221	12/2012	Kaplan et al.	N/A	N/A
8512240	12/2012	Zuckerman-Stark et al.	N/A	N/A
8515535	12/2012	Hopper et al.	N/A	N/A
8515538	12/2012	Osorio et al.	N/A	N/A
8515541	12/2012	Jaax et al.	N/A	N/A
8515549	12/2012	Panken et al.	N/A	N/A
8515550	12/2012	Skelton et al.	N/A	N/A
8517909	12/2012	Honeycutt et al.	N/A	N/A
8517912	12/2012	Clare	N/A	N/A
8519705	12/2012	Savukov et al.	N/A	N/A
8519853	12/2012	Eskandarian et al.	N/A	N/A
8520974	12/2012	Fujita et al.	N/A	N/A
8521284	12/2012	Kim et al.	N/A	N/A
8523779	12/2012	Taylor et al.	N/A	N/A
8525673	12/2012	Tran	N/A	N/A
8525687	12/2012	Tran	N/A	N/A
8527029	12/2012	Okada	N/A	N/A
8527035	12/2012	Diamond	N/A	N/A
8527435	12/2012	Han et al.	N/A	N/A
8529463	12/2012	Della Santina et al.	N/A	N/A
8531291	12/2012	Tran	N/A	N/A
8532756	12/2012	Schalk et al.	N/A	N/A
8532757	12/2012	Molnar et al.	N/A	N/A
8533042	12/2012	Pradeep et al.	N/A	N/A
8536667	12/2012	de Graff et al.	N/A	N/A
8538108	12/2012	Shekhar et al.	N/A	N/A
8538512	12/2012	Bibian et al.	N/A	N/A
8538513	12/2012	Molnar et al.	N/A	N/A
8538514	12/2012	Sun et al.	N/A	N/A
8538523	12/2012	Sommer et al.	N/A	N/A
8538536	12/2012	Rezai et al.	N/A	N/A
8538543	12/2012	McIntyre et al.	N/A	N/A
8538700	12/2012	Badri et al.	N/A	N/A
8538705	12/2012	Greenwald	N/A	N/A
8542900	12/2012	Tolkowsky et al.	N/A	N/A
8542916	12/2012	Tognoli et al.	N/A	N/A
8543189	12/2012	Paitel et al.	N/A	N/A
8543199	12/2012	Snyder et al.	N/A	N/A
8543214	12/2012	Osorio et al.	N/A	N/A
8543219	12/2012	Tass	N/A	N/A
8545378	12/2012	Peterchev	N/A	N/A
8545416	12/2012	Kayyali et al.	N/A	N/A
8545420	12/2012	Einav et al.	N/A	N/A
8545436	12/2012	Robertson et al.	N/A	N/A
8548583	12/2012	Rouso et al.	N/A	N/A
8548594	12/2012	Thimineur et al.	N/A	N/A
8548604	12/2012	Whitehurst et al.	N/A	N/A
8548786	12/2012	Plenz	N/A	N/A
8548852	12/2012	Pradeep et al.	N/A	N/A
8553956	12/2012	Wu et al.	N/A	N/A
8554311	12/2012	Warner et al.	N/A	N/A
8554325	12/2012	Molnar et al.	N/A	N/A
8559645	12/2012	Corona-Strauss et al.	N/A	N/A
8560034	12/2012	Diab et al.	N/A	N/A
8560041	12/2012	Flaherty et al.	N/A	N/A
8560073	12/2012	Osorio	N/A	N/A
8562525	12/2012	Nakashima et al.	N/A	N/A
8562526	12/2012	Heneghan et al.	N/A	N/A
8562527	12/2012	Braun et al.	N/A	N/A
8562536	12/2012	Osorio et al.	N/A	N/A
8562540	12/2012	Goodall et al.	N/A	N/A
8562548	12/2012	Shimada et al.	N/A	N/A
8562660	12/2012	Peyman	N/A	N/A
8562951	12/2012	Suffin et al.	N/A	N/A
8565606	12/2012	Kim et al.	N/A	N/A
8565864	12/2012	Drew et al.	N/A	N/A
8565867	12/2012	Armstrong et al.	N/A	N/A
8565883	12/2012	Lozano	N/A	N/A
8565886	12/2012	Nelson et al.	N/A	N/A
8568231	12/2012	Solanki et al.	N/A	N/A
8568329	12/2012	Lee et al.	N/A	N/A
8571293	12/2012	Ernst et al.	N/A	N/A
8571629	12/2012	Faro et al.	N/A	N/A
8571642	12/2012	Gill et al.	N/A	N/A

8571643	12/2012	Osorio et al.	N/A	N/A
8571653	12/2012	Ben-David et al.	N/A	N/A
8574164	12/2012	Mashiach	N/A	N/A
8574279	12/2012	Schiffer	N/A	N/A
8577103	12/2012	Vija et al.	N/A	N/A
8577464	12/2012	Mashiach	N/A	N/A
8577465	12/2012	Mashiach	N/A	N/A
8577466	12/2012	Mashiach	N/A	N/A
8577467	12/2012	Mashiach et al.	N/A	N/A
8577468	12/2012	Mashiach et al.	N/A	N/A
8577472	12/2012	Mashiach et al.	N/A	N/A
8577478	12/2012	Mashiach et al.	N/A	N/A
8579786	12/2012	Osorio et al.	N/A	N/A
8579793	12/2012	Honeycutt et al.	N/A	N/A
8579795	12/2012	Martel	N/A	N/A
8579834	12/2012	Davis et al.	N/A	N/A
8583238	12/2012	Heldman et al.	N/A	N/A
8583252	12/2012	Skelton et al.	N/A	N/A
8585568	12/2012	Phillips et al.	N/A	N/A
8586019	12/2012	Satchi-Fainaro et al.	N/A	N/A
8586932	12/2012	Rousso et al.	N/A	N/A
8587304	12/2012	Budker et al.	N/A	N/A
8588486	12/2012	Virtue et al.	N/A	N/A
8588552	12/2012	Garg et al.	N/A	N/A
8588899	12/2012	Schiff	N/A	N/A
8588929	12/2012	Skelton et al.	N/A	N/A
8588933	12/2012	Floyd et al.	N/A	N/A
8588941	12/2012	Mashiach	N/A	N/A
8589316	12/2012	Lujan et al.	N/A	N/A
8591419	12/2012	Tyler	N/A	N/A
8591498	12/2012	John	N/A	N/A
8593141	12/2012	Radparvar et al.	N/A	N/A
8593154	12/2012	Ross	N/A	N/A
8594798	12/2012	Osorio et al.	N/A	N/A
8594800	12/2012	Butson et al.	N/A	N/A
8594950	12/2012	Taylor	N/A	N/A
8597171	12/2012	Altman et al.	N/A	N/A
8597193	12/2012	Grunwald et al.	N/A	N/A
8600493	12/2012	Tanner et al.	N/A	N/A
8600502	12/2012	Lovett et al.	N/A	N/A
8600513	12/2012	Aur et al.	N/A	N/A
8600521	12/2012	Armstrong et al.	N/A	N/A
8600696	12/2012	Zafiris	N/A	N/A
8603790	12/2012	Deisseroth et al.	N/A	N/A
8606349	12/2012	Rousso et al.	N/A	N/A
8606351	12/2012	Wheeler	N/A	N/A
8606356	12/2012	Lee et al.	N/A	N/A
8606360	12/2012	Butson et al.	N/A	N/A
8606361	12/2012	Gliner et al.	N/A	N/A
8606530	12/2012	Taylor	N/A	N/A
8606592	12/2012	Hyde et al.	N/A	N/A
8612005	12/2012	Rezai et al.	N/A	N/A
8613695	12/2012	Von Ohlsen et al.	N/A	N/A
8613905	12/2012	El-Agnaf	N/A	N/A
8614254	12/2012	Llinas et al.	N/A	N/A
8614873	12/2012	Beran	N/A	N/A
8615293	12/2012	Jacobson et al.	N/A	N/A
8615309	12/2012	Craig	N/A	N/A
8615479	12/2012	Jung et al.	N/A	N/A
8615664	12/2012	Jung et al.	N/A	N/A
8618799	12/2012	Radparvar et al.	N/A	N/A
8620206	12/2012	Brown et al.	N/A	N/A
8620419	12/2012	Rotenberg et al.	N/A	N/A
8626264	12/2013	Beran	N/A	N/A
8626301	12/2013	Libbus	N/A	N/A
8628328	12/2013	Palacios	N/A	N/A
8628480	12/2013	Derchak	N/A	N/A
8630699	12/2013	Baker et al.	N/A	N/A
8630705	12/2013	Mann et al.	N/A	N/A
8630812	12/2013	Taylor	N/A	N/A
8632465	12/2013	Brockway	N/A	N/A
8632750	12/2013	Suffin et al.	N/A	N/A
8634616	12/2013	Den Harder et al.	N/A	N/A
8634922	12/2013	Osorio et al.	N/A	N/A
8635105	12/2013	Pradeep et al.	N/A	N/A

8636640	12/2013	Chang	N/A	N/A
8638950	12/2013	Anderson et al.	N/A	N/A
8641632	12/2013	Quintin et al.	N/A	N/A
8641646	12/2013	Colborn	N/A	N/A
8644754	12/2013	Brown	N/A	N/A
8644910	12/2013	Rouso et al.	N/A	N/A
8644914	12/2013	Hunt	N/A	N/A
8644921	12/2013	Wilson	N/A	N/A
8644945	12/2013	Skelton et al.	N/A	N/A
8644946	12/2013	Butson et al.	N/A	N/A
8644954	12/2013	Jaax et al.	N/A	N/A
8644957	12/2013	Mashiach	N/A	N/A
8647278	12/2013	Ji et al.	N/A	N/A
8648017	12/2013	Umansky et al.	N/A	N/A
8649845	12/2013	McIntyre et al.	N/A	N/A
8649866	12/2013	Brooke	N/A	N/A
8649871	12/2013	Frei et al.	N/A	N/A
8652038	12/2013	Tran et al.	N/A	N/A
8652187	12/2013	Wells et al.	N/A	N/A
8652189	12/2013	Gafni et al.	N/A	N/A
8655428	12/2013	Pradeep et al.	N/A	N/A
8655437	12/2013	Pradeep et al.	N/A	N/A
8655817	12/2013	Hasey et al.	N/A	N/A
8657732	12/2013	Vasishta	N/A	N/A
8657756	12/2013	Stahmann et al.	N/A	N/A
8658149	12/2013	Satchi-Fainaro et al.	N/A	N/A
8660642	12/2013	Ferren et al.	N/A	N/A
8660649	12/2013	Ruffini et al.	N/A	N/A
8660666	12/2013	Craig	N/A	N/A
8660799	12/2013	Watson et al.	N/A	N/A
8664258	12/2013	Teegarden et al.	N/A	N/A
8666099	12/2013	Nielsen et al.	N/A	N/A
8666467	12/2013	Lynn et al.	N/A	N/A
8666478	12/2013	LaViolette et al.	N/A	N/A
8666501	12/2013	Kilgard et al.	N/A	N/A
8668496	12/2013	Nolen	N/A	N/A
8670603	12/2013	Tolkowsky et al.	N/A	N/A
8672852	12/2013	Gavish	N/A	N/A
8675936	12/2013	Vija et al.	N/A	N/A
8675945	12/2013	Barnhorst et al.	N/A	N/A
8675983	12/2013	Yahil	N/A	N/A
8676324	12/2013	Simon et al.	N/A	N/A
8676325	12/2013	Lindenthaler et al.	N/A	N/A
8676330	12/2013	Simon et al.	N/A	N/A
8679009	12/2013	Osorio	N/A	N/A
8680119	12/2013	Teegarden et al.	N/A	N/A
8680991	12/2013	Tran	N/A	N/A
8682422	12/2013	Hopenfeld	N/A	N/A
8682441	12/2013	De Ridder	N/A	N/A
8682449	12/2013	Simon	N/A	N/A
8682687	12/2013	Hyde et al.	N/A	N/A
8684742	12/2013	Siefert	N/A	N/A
8684900	12/2013	Tran	N/A	N/A
8684921	12/2013	Osorio	N/A	N/A
8684922	12/2013	Tran	N/A	N/A
8684926	12/2013	Arndt	N/A	N/A
8688209	12/2013	Verbitskiy	N/A	N/A
8690748	12/2013	Fu	N/A	N/A
8693756	12/2013	Tolkowsky et al.	N/A	N/A
8693765	12/2013	Mercier et al.	N/A	N/A
8694087	12/2013	Schiff	N/A	N/A
8694089	12/2013	Arad	N/A	N/A
8694092	12/2013	Ferren et al.	N/A	N/A
8694107	12/2013	Falci	N/A	N/A
8694118	12/2013	Armstrong	N/A	N/A
8694157	12/2013	Wenderow et al.	N/A	N/A
8696722	12/2013	Deisseroth et al.	N/A	N/A
8696724	12/2013	Rogers	N/A	N/A
8698639	12/2013	Fung et al.	N/A	N/A
8700137	12/2013	Albert	N/A	N/A
8700141	12/2013	Causevic	N/A	N/A
8700142	12/2013	John et al.	N/A	N/A
8700163	12/2013	Terry, Jr. et al.	N/A	N/A
8700167	12/2013	Sabel	N/A	N/A
8700174	12/2013	Skelton et al.	N/A	N/A

8700183	12/2013	Mashiach	N/A	N/A
8703114	12/2013	Satchi-Fainaro et al.	N/A	N/A
8706183	12/2013	Cui et al.	N/A	N/A
8706205	12/2013	Shahaf et al.	N/A	N/A
8706206	12/2013	Kanai et al.	N/A	N/A
8706207	12/2013	Flint	N/A	N/A
8706237	12/2013	Giftakis et al.	N/A	N/A
8706241	12/2013	Firlik et al.	N/A	N/A
8706518	12/2013	Hyde et al.	N/A	N/A
8708903	12/2013	Tran	N/A	N/A
8708934	12/2013	Skelton et al.	N/A	N/A
8711655	12/2013	Gzara et al.	N/A	N/A
8712507	12/2013	Cazares et al.	N/A	N/A
8712512	12/2013	Doidge et al.	N/A	N/A
8712513	12/2013	Modarres	N/A	N/A
8712547	12/2013	Whitehurst et al.	N/A	N/A
8716447	12/2013	Deisseroth et al.	N/A	N/A
8717430	12/2013	Simon et al.	N/A	N/A
8718747	12/2013	Bjornerud et al.	N/A	N/A
8718776	12/2013	Mashiach et al.	N/A	N/A
8718777	12/2013	Lowry et al.	N/A	N/A
8718779	12/2013	Whitehurst et al.	N/A	N/A
8721695	12/2013	Tass et al.	N/A	N/A
8724871	12/2013	Biagiotti et al.	N/A	N/A
8725238	12/2013	Liu et al.	N/A	N/A
8725243	12/2013	Dilorenzo et al.	N/A	N/A
8725311	12/2013	Breed	N/A	N/A
8725668	12/2013	Georgopoulos	N/A	N/A
8725669	12/2013	Fu	N/A	N/A
8725796	12/2013	Serena	N/A	N/A
8727978	12/2013	Tran et al.	N/A	N/A
8728001	12/2013	Lynn	N/A	N/A
8729040	12/2013	Deisseroth et al.	N/A	N/A
8731650	12/2013	Sajda et al.	N/A	N/A
8731656	12/2013	Bourget et al.	N/A	N/A
8731987	12/2013	Chen et al.	N/A	N/A
8733290	12/2013	Gerashchenko	N/A	N/A
8734356	12/2013	Taylor	N/A	N/A
8734357	12/2013	Taylor	N/A	N/A
8734498	12/2013	DiMauro et al.	N/A	N/A
8738121	12/2013	Virag et al.	N/A	N/A
8738126	12/2013	Craig	N/A	N/A
8738136	12/2013	Frei et al.	N/A	N/A
8738140	12/2013	De Ridder	N/A	N/A
8738395	12/2013	Hyde et al.	N/A	N/A
8744562	12/2013	Giftakis et al.	N/A	N/A
8744563	12/2013	Yoshida	N/A	N/A
8747313	12/2013	Tran et al.	N/A	N/A
8747336	12/2013	Tran	N/A	N/A
8747382	12/2013	D'Souza et al.	N/A	N/A
8750971	12/2013	Tran	N/A	N/A
8750974	12/2013	Baker et al.	N/A	N/A
8750992	12/2013	Hopper et al.	N/A	N/A
8751008	12/2013	Carlton et al.	N/A	N/A
8751011	12/2013	Skelton et al.	N/A	N/A
8753296	12/2013	Einav et al.	N/A	N/A
8754238	12/2013	Teegarden et al.	N/A	N/A
8755854	12/2013	Addison et al.	N/A	N/A
8755856	12/2013	Diab et al.	N/A	N/A
8755868	12/2013	Yazicioglu	N/A	N/A
8755869	12/2013	Zhang et al.	N/A	N/A
8755871	12/2013	Weng et al.	N/A	N/A
8755877	12/2013	Zoica	N/A	N/A
8755901	12/2013	Skelton et al.	N/A	N/A
8756017	12/2013	Hu et al.	N/A	N/A
8758274	12/2013	Sahasrabudhe et al.	N/A	N/A
8761438	12/2013	Lee et al.	N/A	N/A
8761866	12/2013	Chance	N/A	N/A
8761868	12/2013	Giftakis et al.	N/A	N/A
8761869	12/2013	Leuthardt et al.	N/A	N/A
8761889	12/2013	Wingeier et al.	N/A	N/A
8762065	12/2013	DiLorenzo	N/A	N/A
8762202	12/2013	Pradeep et al.	N/A	N/A
8764651	12/2013	Tran	N/A	N/A
8764652	12/2013	Lee et al.	N/A	N/A

8764653	12/2013	Kaminska et al.	N/A	N/A
8764673	12/2013	McCraty et al.	N/A	N/A
8768022	12/2013	Miga et al.	N/A	N/A
8768427	12/2013	Sjaaheim et al.	N/A	N/A
8768431	12/2013	Ross et al.	N/A	N/A
8768446	12/2013	Drew et al.	N/A	N/A
8768447	12/2013	Ermes et al.	N/A	N/A
8768449	12/2013	Pesaran et al.	N/A	N/A
8768471	12/2013	Colborn et al.	N/A	N/A
8768477	12/2013	Spitzer et al.	N/A	N/A
8768718	12/2013	Cazares et al.	N/A	N/A
8771194	12/2013	John et al.	N/A	N/A
8774923	12/2013	Rom	N/A	N/A
8775340	12/2013	Waxman et al.	N/A	N/A
8781193	12/2013	Steinberg et al.	N/A	N/A
8781197	12/2013	Wang et al.	N/A	N/A
8781557	12/2013	Dean et al.	N/A	N/A
8781563	12/2013	Foo	N/A	N/A
8781595	12/2013	Grevious et al.	N/A	N/A
8781597	12/2013	DiLorenzo	N/A	N/A
8781796	12/2013	Mott et al.	N/A	N/A
8784109	12/2013	Gottfried	N/A	N/A
8784322	12/2013	Kim et al.	N/A	N/A
8785441	12/2013	Teegarden et al.	N/A	N/A
8786624	12/2013	Echaz et al.	N/A	N/A
8787637	12/2013	Duchesnay et al.	N/A	N/A
8788030	12/2013	Payton et al.	N/A	N/A
8788033	12/2013	Rossi	N/A	N/A
8788044	12/2013	John	N/A	N/A
8788055	12/2013	Gerber et al.	N/A	N/A
8788057	12/2013	Stevenson et al.	N/A	N/A
8790255	12/2013	Behar	N/A	N/A
8790272	12/2013	Sackner et al.	N/A	N/A
8790297	12/2013	Bromander et al.	N/A	N/A
8792972	12/2013	Zaidel et al.	N/A	N/A
8792974	12/2013	Rothman	N/A	N/A
8792991	12/2013	Gerber et al.	N/A	N/A
8795175	12/2013	Funane et al.	N/A	N/A
8798717	12/2013	Roscher	N/A	N/A
8798728	12/2013	Drew et al.	N/A	N/A
8798735	12/2013	Bibian et al.	N/A	N/A
8798736	12/2013	Sullivan et al.	N/A	N/A
8798773	12/2013	Mashiach	N/A	N/A
8801620	12/2013	Melker et al.	N/A	N/A
8805516	12/2013	Bentwich	N/A	N/A
8805518	12/2013	King et al.	N/A	N/A
8812126	12/2013	Butson et al.	N/A	N/A
8812237	12/2013	Wilt et al.	N/A	N/A
8812245	12/2013	Taylor	N/A	N/A
8812246	12/2013	Taylor	N/A	N/A
8814923	12/2013	Nissila et al.	N/A	N/A
8815582	12/2013	Deisseroth et al.	N/A	N/A
8821376	12/2013	Tolkowsky	N/A	N/A
8821408	12/2013	Hu et al.	N/A	N/A
8821559	12/2013	DiMauro et al.	N/A	N/A
8825149	12/2013	Kraus et al.	N/A	N/A
8825166	12/2013	John	N/A	N/A
8825167	12/2013	Tass et al.	N/A	N/A
8825428	12/2013	Addison et al.	N/A	N/A
8827912	12/2013	Bukhman	N/A	N/A
8827917	12/2013	Watson et al.	N/A	N/A
8829908	12/2013	Roshtal et al.	N/A	N/A
8831705	12/2013	Dobak	N/A	N/A
8831731	12/2013	Blum et al.	N/A	N/A
8831732	12/2013	Frei et al.	N/A	N/A
8834392	12/2013	Panken et al.	N/A	N/A
8834546	12/2013	Deisseroth et al.	N/A	N/A
8838201	12/2013	Mori et al.	N/A	N/A
8838225	12/2013	Ahonen et al.	N/A	N/A
8838226	12/2013	Bibian et al.	N/A	N/A
8838227	12/2013	Causevic et al.	N/A	N/A
8838247	12/2013	Hagedorn et al.	N/A	N/A
8843199	12/2013	Kim et al.	N/A	N/A
8843201	12/2013	Heldman et al.	N/A	N/A
8843210	12/2013	Simon et al.	N/A	N/A

8845545	12/2013	Folkerts et al.	N/A	N/A
8849390	12/2013	Echauz et al.	N/A	N/A
8849392	12/2013	Lozano	N/A	N/A
8849407	12/2013	Danilov et al.	N/A	N/A
8849409	12/2013	Colborn et al.	N/A	N/A
8849632	12/2013	Sparks et al.	N/A	N/A
8849681	12/2013	Hargrove et al.	N/A	N/A
8852073	12/2013	Genereux et al.	N/A	N/A
8852100	12/2013	Osorio	N/A	N/A
8852103	12/2013	Rothberg et al.	N/A	N/A
8855758	12/2013	Rodriquez-Villegas et al.	N/A	N/A
8855773	12/2013	Kokones et al.	N/A	N/A
8855775	12/2013	Leyde	N/A	N/A
8858440	12/2013	Tyler	N/A	N/A
8858449	12/2013	Inan et al.	N/A	N/A
8861819	12/2013	Lee et al.	N/A	N/A
8862196	12/2013	Lynn	N/A	N/A
8862210	12/2013	Yazicioglu et al.	N/A	N/A
8862236	12/2013	Wolpaw et al.	N/A	N/A
8862581	12/2013	Zhang et al.	N/A	N/A
8864310	12/2013	Gross et al.	N/A	N/A
8864806	12/2013	Wells et al.	N/A	N/A
8868148	12/2013	Engelbrecht et al.	N/A	N/A
8868163	12/2013	Guttag et al.	N/A	N/A
8868172	12/2013	Leyde et al.	N/A	N/A
8868173	12/2013	Nelson et al.	N/A	N/A
8868174	12/2013	Sato et al.	N/A	N/A
8868175	12/2013	Arad	N/A	N/A
8868177	12/2013	Simon et al.	N/A	N/A
8868189	12/2013	Stevenson et al.	N/A	N/A
8868201	12/2013	Roberts et al.	N/A	N/A
8870737	12/2013	Phillips et al.	N/A	N/A
8871797	12/2013	Teegarden et al.	N/A	N/A
8872640	12/2013	Horseman	N/A	N/A
8874205	12/2013	Simon et al.	N/A	N/A
8874218	12/2013	Terry, Jr.	N/A	N/A
8874227	12/2013	Simon et al.	N/A	N/A
8874439	12/2013	Kim et al.	N/A	N/A
8880207	12/2013	Abeyratne et al.	N/A	N/A
8880576	12/2013	Ochs et al.	N/A	N/A
8886299	12/2013	Yazicioglu et al.	N/A	N/A
8886302	12/2013	Skelton et al.	N/A	N/A
8888672	12/2013	Phillips et al.	N/A	N/A
8888673	12/2013	Phillips et al.	N/A	N/A
8888702	12/2013	Osorio	N/A	N/A
8888708	12/2013	Diab et al.	N/A	N/A
8888723	12/2013	Einav	N/A	N/A
8892207	12/2013	Nelson et al.	N/A	N/A
8893120	12/2013	Pinsky et al.	N/A	N/A
8898037	12/2013	Watson et al.	N/A	N/A
8900284	12/2013	DiMauro et al.	N/A	N/A
8902070	12/2013	Kobetski et al.	N/A	N/A
8903479	12/2013	Zoicas	N/A	N/A
8903483	12/2013	Sun et al.	N/A	N/A
8903486	12/2013	Bourget et al.	N/A	N/A
8903494	12/2013	Goldwasser et al.	N/A	N/A
8906360	12/2013	Deisseroth et al.	N/A	N/A
8907668	12/2013	Okada	N/A	N/A
8909345	12/2013	Danilov et al.	N/A	N/A
8910638	12/2013	Boyden et al.	N/A	N/A
8913810	12/2013	Panin et al.	N/A	N/A
8914100	12/2013	Adachi et al.	N/A	N/A
8914115	12/2013	Giftakis et al.	N/A	N/A
8914119	12/2013	Wu et al.	N/A	N/A
8914122	12/2013	Simon et al.	N/A	N/A
8915741	12/2013	Hatlestad et al.	N/A	N/A
8915871	12/2013	Einav	N/A	N/A
8918162	12/2013	Prokoski	N/A	N/A
8918176	12/2013	Nelson et al.	N/A	N/A
8918178	12/2013	Simon et al.	N/A	N/A
8918183	12/2013	Carlton et al.	N/A	N/A
8921320	12/2013	Paul et al.	N/A	N/A
8922376	12/2013	Kangas et al.	N/A	N/A
8922788	12/2013	Addison et al.	N/A	N/A
8923958	12/2013	Gupta et al.	N/A	N/A

8924235	12/2013	Seely	N/A	N/A
RE45336	12/2014	Teegarden et al.	N/A	N/A
RE45337	12/2014	Teegarden et al.	N/A	N/A
8926959	12/2014	Deisseroth et al.	N/A	N/A
8929991	12/2014	Fowler et al.	N/A	N/A
8929999	12/2014	Maschiach	N/A	N/A
8932218	12/2014	Thompson	N/A	N/A
8932227	12/2014	Lynn	N/A	N/A
8932562	12/2014	Deisseroth et al.	N/A	N/A
8933696	12/2014	Nishikawa	N/A	N/A
8934685	12/2014	Avinash et al.	N/A	N/A
8934965	12/2014	Rogers et al.	N/A	N/A
8934967	12/2014	Kilgard et al.	N/A	N/A
8934979	12/2014	Moffitt	N/A	N/A
8934986	12/2014	Goetz	N/A	N/A
8936629	12/2014	Boyden et al.	N/A	N/A
8936630	12/2014	Denison et al.	N/A	N/A
8938102	12/2014	Carroll	N/A	N/A
8938289	12/2014	Einav et al.	N/A	N/A
8938290	12/2014	Wingeier et al.	N/A	N/A
8938301	12/2014	Hagedorn	N/A	N/A
8939903	12/2014	Roberts et al.	N/A	N/A
8942777	12/2014	Diab et al.	N/A	N/A
8942813	12/2014	Hagedorn et al.	N/A	N/A
8942817	12/2014	Hyde et al.	N/A	N/A
8945006	12/2014	Osorio	N/A	N/A
8948834	12/2014	Diab et al.	N/A	N/A
8948849	12/2014	Diamond et al.	N/A	N/A
8948855	12/2014	Osorio et al.	N/A	N/A
8948860	12/2014	Causevic	N/A	N/A
8951189	12/2014	Osorio	N/A	N/A
8951190	12/2014	Chmiel et al.	N/A	N/A
8951192	12/2014	Osorio	N/A	N/A
8951203	12/2014	Patangay et al.	N/A	N/A
8954139	12/2014	Hopenfeld et al.	N/A	N/A
8954146	12/2014	Hopper et al.	N/A	N/A
8954293	12/2014	Klinkenbusch	N/A	N/A
8955010	12/2014	Pradeep et al.	N/A	N/A
8955974	12/2014	Gross et al.	N/A	N/A
8956277	12/2014	Mishelevich	N/A	N/A
8956363	12/2014	Schneider et al.	N/A	N/A
8958868	12/2014	Ghovanloo et al.	N/A	N/A
8958870	12/2014	Gerber et al.	N/A	N/A
8958882	12/2014	Hagedorn	N/A	N/A
8961187	12/2014	Boers et al.	N/A	N/A
8961385	12/2014	Pilla et al.	N/A	N/A
8961386	12/2014	Phillips et al.	N/A	N/A
8962042	12/2014	Geng	N/A	N/A
8962589	12/2014	Deisseroth et al.	N/A	N/A
8964298	12/2014	Haddick et al.	N/A	N/A
8965492	12/2014	Baker et al.	N/A	N/A
8965513	12/2014	Wingeier et al.	N/A	N/A
8965514	12/2014	Bikson et al.	N/A	N/A
8968172	12/2014	Wang et al.	N/A	N/A
8968176	12/2014	Altman et al.	N/A	N/A
8968195	12/2014	Tran	N/A	N/A
8968376	12/2014	Wells et al.	N/A	N/A
8971936	12/2014	Derchak	N/A	N/A
8972004	12/2014	Simon et al.	N/A	N/A
8972013	12/2014	Maschino	N/A	N/A
8974365	12/2014	Best	N/A	N/A
8977024	12/2014	Rex et al.	N/A	N/A
8977110	12/2014	Pradeep et al.	N/A	N/A
8977362	12/2014	Saab	N/A	N/A
8980891	12/2014	Stirn et al.	N/A	N/A
8983155	12/2014	McIntyre et al.	N/A	N/A
8983591	12/2014	Leininger et al.	N/A	N/A
8983620	12/2014	Cinbis	N/A	N/A
8983628	12/2014	Simon et al.	N/A	N/A
8983629	12/2014	Simon et al.	N/A	N/A
8985119	12/2014	Webb et al.	N/A	N/A
8986207	12/2014	Li et al.	N/A	N/A
8989835	12/2014	Badower et al.	N/A	N/A
8989836	12/2014	Machon et al.	N/A	N/A
8989863	12/2014	Osorio	N/A	N/A

8989867	12/2014	Chow et al.	N/A	N/A
8989868	12/2014	Mashiach et al.	N/A	N/A
8989871	12/2014	Ollivier	N/A	N/A
8992230	12/2014	Tuchschmid et al.	N/A	N/A
8993623	12/2014	Goodenowe	N/A	N/A
8996112	12/2014	Brooke	N/A	N/A
8996120	12/2014	Calle et al.	N/A	N/A
8998828	12/2014	Reichow et al.	N/A	N/A
9002458	12/2014	Pal et al.	N/A	N/A
9002471	12/2014	Stevenson et al.	N/A	N/A
9002477	12/2014	Burnett	N/A	N/A
9004687	12/2014	Stack	N/A	N/A
9005102	12/2014	Burnett et al.	N/A	N/A
9005126	12/2014	Beach et al.	N/A	N/A
9005649	12/2014	Ho et al.	N/A	N/A
9008367	12/2014	Tolkowsky et al.	N/A	N/A
9008754	12/2014	Steinberg et al.	N/A	N/A
9008771	12/2014	Dong et al.	N/A	N/A
9008780	12/2014	Nudo et al.	N/A	N/A
9008970	12/2014	Donderici et al.	N/A	N/A
9011329	12/2014	Ferren et al.	N/A	N/A
9014216	12/2014	Lazar et al.	N/A	N/A
9014453	12/2014	Steinberg et al.	N/A	N/A
9014804	12/2014	Giftakis et al.	N/A	N/A
9014811	12/2014	Pal et al.	N/A	N/A
9014819	12/2014	Lee et al.	N/A	N/A
9014823	12/2014	Simon et al.	N/A	N/A
9015057	12/2014	Phillips et al.	N/A	N/A
9015087	12/2014	Li et al.	N/A	N/A
9020576	12/2014	Nagatani	N/A	N/A
9020582	12/2014	Osorio et al.	N/A	N/A
9020585	12/2014	John et al.	N/A	N/A
9020586	12/2014	Yamada et al.	N/A	N/A
9020598	12/2014	Simon et al.	N/A	N/A
9020612	12/2014	Danilov et al.	N/A	N/A
9020789	12/2014	Butson et al.	N/A	N/A
9022930	12/2014	Sachanandani et al.	N/A	N/A
9022936	12/2014	Rothberg et al.	N/A	N/A
9025845	12/2014	Carroll	N/A	N/A
9026194	12/2014	Okada	N/A	N/A
9026202	12/2014	Albert	N/A	N/A
9026217	12/2014	Kokones et al.	N/A	N/A
9026218	12/2014	Lozano et al.	N/A	N/A
9026372	12/2014	O'Donnell, Jr. et al.	N/A	N/A
9028405	12/2014	Tran	N/A	N/A
9028412	12/2014	Rothberg et al.	N/A	N/A
9031644	12/2014	Johnson et al.	N/A	N/A
9031653	12/2014	Mashiach	N/A	N/A
9031655	12/2014	Osorio et al.	N/A	N/A
9031658	12/2014	Chiao et al.	N/A	N/A
9033884	12/2014	Rothberg et al.	N/A	N/A
9034055	12/2014	Vinjamuri et al.	N/A	N/A
9034911	12/2014	Selvey et al.	N/A	N/A
9034923	12/2014	Goodenowe	N/A	N/A
9035657	12/2014	Zhang et al.	N/A	N/A
9036844	12/2014	Suhami et al.	N/A	N/A
9037224	12/2014	Fu	N/A	N/A
9037225	12/2014	Saliga et al.	N/A	N/A
9037254	12/2014	John	N/A	N/A
9037256	12/2014	Bokil	N/A	N/A
9037530	12/2014	Tan et al.	N/A	N/A
9042074	12/2014	Beran	N/A	N/A
9042201	12/2014	Tyler et al.	N/A	N/A
9042952	12/2014	Lynn et al.	N/A	N/A
9042958	12/2014	Karmarkar et al.	N/A	N/A
9042988	12/2014	DiLorenzo	N/A	N/A
9043001	12/2014	Simon et al.	N/A	N/A
9044188	12/2014	DiLorenzo et al.	N/A	N/A
9044612	12/2014	Mashiach et al.	N/A	N/A
9050469	12/2014	Osorio et al.	N/A	N/A
9050470	12/2014	Carlton et al.	N/A	N/A
9050471	12/2014	Skelton et al.	N/A	N/A
9053516	12/2014	Stempora	N/A	N/A
9053534	12/2014	Ross et al.	N/A	N/A
9055871	12/2014	Inan et al.	N/A	N/A

9055974	12/2014	Goetz	N/A	N/A
9056195	12/2014	Sabesan	N/A	N/A
9058473	12/2014	Navratil et al.	N/A	N/A
9060671	12/2014	Badower et al.	N/A	N/A
9060683	12/2014	Tran	N/A	N/A
9060695	12/2014	Peters	N/A	N/A
9060722	12/2014	Teixeira	N/A	N/A
9060746	12/2014	Weng et al.	N/A	N/A
9061132	12/2014	Zweber et al.	N/A	N/A
9061133	12/2014	Wurster et al.	N/A	N/A
9061151	12/2014	Mashiach et al.	N/A	N/A
9061153	12/2014	Lebovitz et al.	N/A	N/A
9063183	12/2014	Toda et al.	N/A	N/A
9063643	12/2014	Sparks et al.	N/A	N/A
9064036	12/2014	Hyde et al.	N/A	N/A
9067052	12/2014	Moses et al.	N/A	N/A
9067054	12/2014	Simon et al.	N/A	N/A
9067070	12/2014	Connor	N/A	N/A
9069031	12/2014	Guedes et al.	N/A	N/A
9069097	12/2014	Zhang et al.	N/A	N/A
9070492	12/2014	Yarmush et al.	N/A	N/A
9072449	12/2014	Semenov	N/A	N/A
9072482	12/2014	Sarkela et al.	N/A	N/A
9072832	12/2014	Frei et al.	N/A	N/A
9072870	12/2014	Wu et al.	N/A	N/A
9072905	12/2014	Kokones et al.	N/A	N/A
9074976	12/2014	Adolphi et al.	N/A	N/A
9076212	12/2014	Ernst et al.	N/A	N/A
9078564	12/2014	Taylor	N/A	N/A
9078577	12/2014	He et al.	N/A	N/A
9078584	12/2014	Jorge et al.	N/A	N/A
9079039	12/2014	Carlson et al.	N/A	N/A
9079940	12/2014	Deisseroth et al.	N/A	N/A
9081488	12/2014	Soederstroem	N/A	N/A
9081882	12/2014	Taylor	N/A	N/A
9081890	12/2014	An et al.	N/A	N/A
9082169	12/2014	Thomson et al.	N/A	N/A
9084584	12/2014	Weiland et al.	N/A	N/A
9084885	12/2014	Deisseroth et al.	N/A	N/A
9084896	12/2014	Kokones et al.	N/A	N/A
9084900	12/2014	Hershey et al.	N/A	N/A
9087147	12/2014	Fonte	N/A	N/A
9089310	12/2014	Isenhardt et al.	N/A	N/A
9089400	12/2014	Nofzinger	N/A	N/A
9089683	12/2014	Mishelevich	N/A	N/A
9089707	12/2014	Kilgard et al.	N/A	N/A
9089713	12/2014	John	N/A	N/A
9089719	12/2014	Simon et al.	N/A	N/A
9091785	12/2014	Donderici et al.	N/A	N/A
9092556	12/2014	Amble et al.	N/A	N/A
9092895	12/2014	Ross et al.	N/A	N/A
9095266	12/2014	Fu	N/A	N/A
9095268	12/2014	Kurtz et al.	N/A	N/A
9095295	12/2014	Eagleman et al.	N/A	N/A
9095303	12/2014	Osorio	N/A	N/A
9095314	12/2014	Osorio et al.	N/A	N/A
9095618	12/2014	Satchi-Fainaro et al.	N/A	N/A
9095713	12/2014	Foster et al.	N/A	N/A
9100758	12/2014	Adachi et al.	N/A	N/A
9101263	12/2014	Jung et al.	N/A	N/A
9101276	12/2014	Georgopoulos	N/A	N/A
9101279	12/2014	Ritchey et al.	N/A	N/A
9101690	12/2014	Deisseroth et al.	N/A	N/A
9101759	12/2014	Delp et al.	N/A	N/A
9101766	12/2014	Nekhendzy	N/A	N/A
9102717	12/2014	Huang et al.	N/A	N/A
9107586	12/2014	Tran	N/A	N/A
9107595	12/2014	Smyth	N/A	N/A
9108041	12/2014	Craig	N/A	N/A
9113777	12/2014	Mittal	N/A	N/A
9113801	12/2014	DiLorenzo	N/A	N/A
9113803	12/2014	Zhang	N/A	N/A
9113830	12/2014	Galen et al.	N/A	N/A
9116201	12/2014	Shah et al.	N/A	N/A
9116835	12/2014	Smyth	N/A	N/A

9118775	12/2014	Lim et al.	N/A	N/A
9119533	12/2014	Ghaffari	N/A	N/A
9119551	12/2014	Qi et al.	N/A	N/A
9119583	12/2014	Tass	N/A	N/A
9119597	12/2014	Dripps et al.	N/A	N/A
9119598	12/2014	Engelbrecht et al.	N/A	N/A
9121964	12/2014	Lewis et al.	N/A	N/A
9125574	12/2014	Zia et al.	N/A	N/A
9125581	12/2014	Wu et al.	N/A	N/A
9125788	12/2014	Tee et al.	N/A	N/A
9126050	12/2014	Simon et al.	N/A	N/A
9131864	12/2014	Korenberg	N/A	N/A
9133024	12/2014	Phan et al.	N/A	N/A
9133709	12/2014	Huh et al.	N/A	N/A
9135221	12/2014	Shahaf et al.	N/A	N/A
9135400	12/2014	McIntyre et al.	N/A	N/A
9138156	12/2014	Wu et al.	N/A	N/A
9138175	12/2014	Ernst et al.	N/A	N/A
9138183	12/2014	McKenna et al.	N/A	N/A
9138579	12/2014	Wolpaw et al.	N/A	N/A
9138580	12/2014	Ignagni et al.	N/A	N/A
9142145	12/2014	Tuchschmid et al.	N/A	N/A
9142185	12/2014	Fateh	N/A	N/A
9144392	12/2014	Santosh et al.	N/A	N/A
RE45766	12/2014	Lindsay	N/A	N/A
9149195	12/2014	Hadley	N/A	N/A
9149197	12/2014	Taylor	N/A	N/A
9149210	12/2014	Sahasrabudhe et al.	N/A	N/A
9149214	12/2014	Adachi et al.	N/A	N/A
9149226	12/2014	Jadidi	N/A	N/A
9149255	12/2014	Rothberg et al.	N/A	N/A
9149577	12/2014	Robertson et al.	N/A	N/A
9149599	12/2014	Walter et al.	N/A	N/A
9149719	12/2014	Guan et al.	N/A	N/A
9152757	12/2014	Taylor	N/A	N/A
9155373	12/2014	Allen et al.	N/A	N/A
9155484	12/2014	Baker et al.	N/A	N/A
9155487	12/2014	Linderman et al.	N/A	N/A
9155521	12/2014	Rothberg et al.	N/A	N/A
9161715	12/2014	Jung et al.	N/A	N/A
9162051	12/2014	Morrell	N/A	N/A
9162052	12/2014	Morrell	N/A	N/A
9165472	12/2014	Hagedorn et al.	N/A	N/A
9167970	12/2014	Gratton et al.	N/A	N/A
9167974	12/2014	Taylor	N/A	N/A
9167976	12/2014	Wingeier et al.	N/A	N/A
9167977	12/2014	Wingeier et al.	N/A	N/A
9167978	12/2014	Wingeier et al.	N/A	N/A
9167979	12/2014	Skidmore et al.	N/A	N/A
9171353	12/2014	Vija et al.	N/A	N/A
9171366	12/2014	Declerck et al.	N/A	N/A
9173582	12/2014	Popovic et al.	N/A	N/A
9173609	12/2014	Nelson	N/A	N/A
9173610	12/2014	Navakatikyan	N/A	N/A
9174045	12/2014	Simon et al.	N/A	N/A
9174055	12/2014	Davis et al.	N/A	N/A
9174066	12/2014	Simon et al.	N/A	N/A
9175095	12/2014	Deisseroth et al.	N/A	N/A
9177379	12/2014	Biagiotti et al.	N/A	N/A
9177416	12/2014	Sharp	N/A	N/A
9179850	12/2014	Wingeier et al.	N/A	N/A
9179854	12/2014	Doidge et al.	N/A	N/A
9179855	12/2014	Burdea et al.	N/A	N/A
9179858	12/2014	Hasson et al.	N/A	N/A
9179875	12/2014	Hua	N/A	N/A
9179876	12/2014	Ochs et al.	N/A	N/A
9183351	12/2014	Shusterman	N/A	N/A
9186060	12/2014	De Graff et al.	N/A	N/A
9186106	12/2014	Osorio	N/A	N/A
9186503	12/2014	Lindenthaler et al.	N/A	N/A
9186510	12/2014	Gliner et al.	N/A	N/A
9187745	12/2014	Deisseroth et al.	N/A	N/A
9192300	12/2014	Jung et al.	N/A	N/A
9192309	12/2014	Hopenfeld et al.	N/A	N/A
9198563	12/2014	Ferren et al.	N/A	N/A

9198612	12/2014	Fueyo et al.	N/A	N/A
9198621	12/2014	Fernstrom et al.	N/A	N/A
9198624	12/2014	Funane et al.	N/A	N/A
9198637	12/2014	Rothberg et al.	N/A	N/A
9198707	12/2014	McKay et al.	N/A	N/A
9198733	12/2014	Neal, II et al.	N/A	N/A
9204796	12/2014	Tran	N/A	N/A
9204835	12/2014	Parsey et al.	N/A	N/A
9204838	12/2014	Osorio	N/A	N/A
9204998	12/2014	Kilgard et al.	N/A	N/A
9208430	12/2014	Solari	N/A	N/A
9208557	12/2014	Pautot	N/A	N/A
9208558	12/2014	Dean et al.	N/A	N/A
9211076	12/2014	Kim	N/A	N/A
9211077	12/2014	Jung et al.	N/A	N/A
9211212	12/2014	Nofzinger et al.	N/A	N/A
9211411	12/2014	Wu et al.	N/A	N/A
9211417	12/2014	Heldman et al.	N/A	N/A
9213074	12/2014	van der Kouwe et al.	N/A	N/A
9213076	12/2014	Liu	N/A	N/A
9215298	12/2014	Schiff	N/A	N/A
9215978	12/2014	Knight et al.	N/A	N/A
9220910	12/2014	Colborn	N/A	N/A
9220917	12/2014	Boyden et al.	N/A	N/A
9221755	12/2014	Teegarden et al.	N/A	N/A
9226672	12/2015	Taylor	N/A	N/A
9227056	12/2015	Heldman et al.	N/A	N/A
9229080	12/2015	Lin	N/A	N/A
9230065	12/2015	Hasegawa et al.	N/A	N/A
9230539	12/2015	Pakhomov	N/A	N/A
9232910	12/2015	Alshaer et al.	N/A	N/A
9232984	12/2015	Guthart et al.	N/A	N/A
9233244	12/2015	Pal et al.	N/A	N/A
9233245	12/2015	Lamensdorf et al.	N/A	N/A
9233246	12/2015	Simon et al.	N/A	N/A
9233258	12/2015	Simon et al.	N/A	N/A
9235679	12/2015	Taylor	N/A	N/A
9235685	12/2015	McIntyre et al.	N/A	N/A
9238142	12/2015	Heldman et al.	N/A	N/A
9238150	12/2015	Deisseroth et al.	N/A	N/A
9241647	12/2015	Osorio et al.	N/A	N/A
9241665	12/2015	deCharms	N/A	N/A
9242067	12/2015	Shore et al.	N/A	N/A
9242092	12/2015	Simon et al.	N/A	N/A
9247890	12/2015	Turnbull et al.	N/A	N/A
9247911	12/2015	Galloway et al.	N/A	N/A
9247924	12/2015	Rothberg et al.	N/A	N/A
9248003	12/2015	Wright et al.	N/A	N/A
9248280	12/2015	Moffitt et al.	N/A	N/A
9248286	12/2015	Simon et al.	N/A	N/A
9248288	12/2015	Panken et al.	N/A	N/A
9248290	12/2015	Mashiach	N/A	N/A
9248291	12/2015	Mashiach	N/A	N/A
9248296	12/2015	Carcieri et al.	N/A	N/A
9249200	12/2015	Deisseroth et al.	N/A	N/A
9249234	12/2015	Deisseroth et al.	N/A	N/A
9251566	12/2015	Bajic	N/A	N/A
9254097	12/2015	Espy et al.	N/A	N/A
9254099	12/2015	Connor	N/A	N/A
9254383	12/2015	Simon et al.	N/A	N/A
9254387	12/2015	Blum et al.	N/A	N/A
9256982	12/2015	Sharp et al.	N/A	N/A
9259177	12/2015	Drew et al.	N/A	N/A
9259482	12/2015	Satchi-Fainaro et al.	N/A	N/A
9259591	12/2015	Brown et al.	N/A	N/A
9261573	12/2015	Radparvar et al.	N/A	N/A
9265458	12/2015	Stack	N/A	N/A
9265660	12/2015	Kilgard et al.	N/A	N/A
9265661	12/2015	Kilgard et al.	N/A	N/A
9265662	12/2015	Kilgard et al.	N/A	N/A
9265663	12/2015	Kilgard et al.	N/A	N/A
9265931	12/2015	Morrell	N/A	N/A
9265943	12/2015	Yun et al.	N/A	N/A
9265946	12/2015	Morrell	N/A	N/A
9265965	12/2015	Fox et al.	N/A	N/A

9265974	12/2015	You et al.	N/A	N/A
9268014	12/2015	Rothberg et al.	N/A	N/A
9268015	12/2015	Rothberg et al.	N/A	N/A
9268902	12/2015	Taylor et al.	N/A	N/A
9271651	12/2015	Avinash et al.	N/A	N/A
9271657	12/2015	Taylor	N/A	N/A
9271660	12/2015	Luo et al.	N/A	N/A
9271674	12/2015	Deisseroth et al.	N/A	N/A
9271679	12/2015	Cho et al.	N/A	N/A
9272091	12/2015	Skelton et al.	N/A	N/A
9272139	12/2015	Hamilton et al.	N/A	N/A
9272145	12/2015	Kilgard et al.	N/A	N/A
9272153	12/2015	Blum et al.	N/A	N/A
9273035	12/2015	Teegarden et al.	N/A	N/A
9275191	12/2015	Dean et al.	N/A	N/A
9275451	12/2015	Ben-Haim et al.	N/A	N/A
9277871	12/2015	Keenan et al.	N/A	N/A
9277873	12/2015	Sarma et al.	N/A	N/A
9278159	12/2015	Deisseroth et al.	N/A	N/A
9278231	12/2015	Vasishta	N/A	N/A
9280784	12/2015	Barnett et al.	N/A	N/A
9282927	12/2015	Hyde et al.	N/A	N/A
9282930	12/2015	Machon et al.	N/A	N/A
9282934	12/2015	Liley et al.	N/A	N/A
9283279	12/2015	Satchi-Fainaro et al.	N/A	N/A
9283394	12/2015	Whitehurst et al.	N/A	N/A
9284353	12/2015	Deisseroth et al.	N/A	N/A
9285249	12/2015	Schober et al.	N/A	N/A
9289143	12/2015	Wingeier et al.	N/A	N/A
9289595	12/2015	Floyd et al.	N/A	N/A
9289599	12/2015	Craig	N/A	N/A
9289603	12/2015	Giuffrida et al.	N/A	N/A
9289609	12/2015	Moffitt	N/A	N/A
9292471	12/2015	Fung et al.	N/A	N/A
9292858	12/2015	Marci et al.	N/A	N/A
9292920	12/2015	Dean et al.	N/A	N/A
9295838	12/2015	Starr et al.	N/A	N/A
9296382	12/2015	Fung et al.	N/A	N/A
9302069	12/2015	Tass et al.	N/A	N/A
9302093	12/2015	Mashiach	N/A	N/A
9302103	12/2015	Nirenberg	N/A	N/A
9302109	12/2015	Sabesan	N/A	N/A
9302110	12/2015	Kokones et al.	N/A	N/A
9302114	12/2015	Rossi	N/A	N/A
9302116	12/2015	Vo-Dinh et al.	N/A	N/A
9305376	12/2015	Lee et al.	N/A	N/A
9307925	12/2015	Russell et al.	N/A	N/A
9307944	12/2015	Colman et al.	N/A	N/A
9308372	12/2015	Sparks et al.	N/A	N/A
9308392	12/2015	Deisseroth et al.	N/A	N/A
9309296	12/2015	Deisseroth et al.	N/A	N/A
9310985	12/2015	Blum et al.	N/A	N/A
9311335	12/2015	Simon	N/A	N/A
9314190	12/2015	Giuffrida et al.	N/A	N/A
9314613	12/2015	Mashiach	N/A	N/A
9314633	12/2015	Osorio et al.	N/A	N/A
9314635	12/2015	Libbus	N/A	N/A
9320449	12/2015	Gu	N/A	N/A
9320450	12/2015	Badower	N/A	N/A
9320451	12/2015	Feldkamp et al.	N/A	N/A
9320900	12/2015	DiLorenzo	N/A	N/A
9320913	12/2015	Dimino et al.	N/A	N/A
9320914	12/2015	Toselli et al.	N/A	N/A
9322895	12/2015	Santosh et al.	N/A	N/A
9326705	12/2015	Derchak	N/A	N/A
9326720	12/2015	McLaughlin	N/A	N/A
9326742	12/2015	Hirschman et al.	N/A	N/A
9327069	12/2015	Foster et al.	N/A	N/A
9327070	12/2015	Skelton et al.	N/A	N/A
9328107	12/2015	Teegarden et al.	N/A	N/A
9329758	12/2015	Guzak et al.	N/A	N/A
9330206	12/2015	Dean et al.	N/A	N/A
9330523	12/2015	Sutton et al.	N/A	N/A
9331841	12/2015	Kim et al.	N/A	N/A
9332939	12/2015	Osorio	N/A	N/A

9333334	12/2015	Jeffery et al.	N/A	N/A
9333347	12/2015	Simon et al.	N/A	N/A
9333350	12/2015	Rise et al.	N/A	N/A
9336302	12/2015	Swamy	N/A	N/A
9336535	12/2015	Pradeep et al.	N/A	N/A
9336611	12/2015	Bilgic et al.	N/A	N/A
9339200	12/2015	Fonte	N/A	N/A
9339227	12/2015	D'arcy et al.	N/A	N/A
9339641	12/2015	Rajguru et al.	N/A	N/A
9339654	12/2015	Kilgard et al.	N/A	N/A
9340589	12/2015	Deisseroth et al.	N/A	N/A
9345412	12/2015	Horne	N/A	N/A
9345609	12/2015	Hyde et al.	N/A	N/A
9345886	12/2015	Kilgard et al.	N/A	N/A
9345901	12/2015	Peterchev	N/A	N/A
9348974	12/2015	Goetz	N/A	N/A
9349178	12/2015	Itu et al.	N/A	N/A
9351640	12/2015	Tran	N/A	N/A
9351651	12/2015	Nagasaka	N/A	N/A
9352145	12/2015	Whitehurst et al.	N/A	N/A
9352152	12/2015	Lindenthaler et al.	N/A	N/A
9352156	12/2015	Lane et al.	N/A	N/A
9357240	12/2015	Pradeep et al.	N/A	N/A
9357298	12/2015	Hiroe	N/A	N/A
9357941	12/2015	Simon	N/A	N/A
9357949	12/2015	Drew	N/A	N/A
9357970	12/2015	Clark et al.	N/A	N/A
9358361	12/2015	Hyde et al.	N/A	N/A
9358381	12/2015	Simon et al.	N/A	N/A
9358392	12/2015	Mashiach	N/A	N/A
9358393	12/2015	Lozano	N/A	N/A
9358398	12/2015	Moffitt et al.	N/A	N/A
9359449	12/2015	Deisseroth et al.	N/A	N/A
9360472	12/2015	Deisseroth et al.	N/A	N/A
9364462	12/2015	Simpson, Jr.	N/A	N/A
9364665	12/2015	Bokil et al.	N/A	N/A
9364674	12/2015	Cook et al.	N/A	N/A
9364679	12/2015	John	N/A	N/A
9365628	12/2015	Deisseroth et al.	N/A	N/A
9367131	12/2015	Klappert et al.	N/A	N/A
9367738	12/2015	Harumatsu et al.	N/A	N/A
9368018	12/2015	Kangas et al.	N/A	N/A
9368265	12/2015	Park et al.	N/A	N/A
9370309	12/2015	Ko et al.	N/A	N/A
9370667	12/2015	Schmidt	N/A	N/A
9375145	12/2015	Chin et al.	N/A	N/A
9375151	12/2015	Hopenfeld et al.	N/A	N/A
9375171	12/2015	Teixeira	N/A	N/A
9375564	12/2015	Wingeier et al.	N/A	N/A
9375571	12/2015	Errico et al.	N/A	N/A
9375573	12/2015	Dilorenzo	N/A	N/A
9377348	12/2015	Kataoka	N/A	N/A
9377515	12/2015	Kim et al.	N/A	N/A
9380976	12/2015	Stack	N/A	N/A
9381346	12/2015	Lee et al.	N/A	N/A
9381352	12/2015	Yun et al.	N/A	N/A
9383208	12/2015	Mohanty	N/A	N/A
9387320	12/2015	Wingeier et al.	N/A	N/A
9387338	12/2015	Burnett	N/A	N/A
9390233	12/2015	Fueyo et al.	N/A	N/A
9392955	12/2015	Folkerts et al.	N/A	N/A
9393406	12/2015	Ollivier	N/A	N/A
9393418	12/2015	Giuffrida et al.	N/A	N/A
9394347	12/2015	Deisseroth et al.	N/A	N/A
9395425	12/2015	Diamond et al.	N/A	N/A
9396533	12/2015	Skidmore	N/A	N/A
9396669	12/2015	Karkanias et al.	N/A	N/A
9398873	12/2015	Van Dooren et al.	N/A	N/A
9399126	12/2015	Pal et al.	N/A	N/A
9399133	12/2015	Besio	N/A	N/A
9399134	12/2015	Simon et al.	N/A	N/A
9399144	12/2015	Howard	N/A	N/A
9401021	12/2015	Biagiotti et al.	N/A	N/A
9401033	12/2015	Bajic	N/A	N/A
9402558	12/2015	John et al.	N/A	N/A

9402994	12/2015	Chow et al.	N/A	N/A
9403000	12/2015	Lyons et al.	N/A	N/A
9403001	12/2015	Simon et al.	N/A	N/A
9403009	12/2015	Mashiach	N/A	N/A
9403010	12/2015	Fried et al.	N/A	N/A
9403038	12/2015	Tyler	N/A	N/A
9405366	12/2015	Segal	N/A	N/A
9408530	12/2015	Ferren et al.	N/A	N/A
9409013	12/2015	Mashiach et al.	N/A	N/A
9409022	12/2015	Jaax et al.	N/A	N/A
9409028	12/2015	Whitehurst et al.	N/A	N/A
9410885	12/2015	Schober et al.	N/A	N/A
9411033	12/2015	He et al.	N/A	N/A
9411935	12/2015	Moffitt et al.	N/A	N/A
9412076	12/2015	Sapiro et al.	N/A	N/A
9412233	12/2015	Bagherzadeh et al.	N/A	N/A
9414029	12/2015	Miyazaki et al.	N/A	N/A
9414749	12/2015	Semenov	N/A	N/A
9414763	12/2015	Semenov	N/A	N/A
9414764	12/2015	Semenov	N/A	N/A
9414776	12/2015	Sillay et al.	N/A	N/A
9414780	12/2015	Rhoads	N/A	N/A
9414907	12/2015	Wortz et al.	N/A	N/A
9415215	12/2015	Mashiach	N/A	N/A
9415216	12/2015	Mashiach	N/A	N/A
9415219	12/2015	Simon et al.	N/A	N/A
9415222	12/2015	DiLorenzo	N/A	N/A
9415233	12/2015	Pilla et al.	N/A	N/A
9418368	12/2015	Jung et al.	N/A	N/A
9420970	12/2015	Dagum	N/A	N/A
9421258	12/2015	Deisseroth et al.	N/A	N/A
9421372	12/2015	Mashiach et al.	N/A	N/A
9421373	12/2015	DiLorenzo	N/A	N/A
9421379	12/2015	Zhu	N/A	N/A
9424761	12/2015	Tuchschrnid et al.	N/A	N/A
9427474	12/2015	Satchi-Fainaro et al.	N/A	N/A
9427581	12/2015	Simon et al.	N/A	N/A
9427585	12/2015	Gliner	N/A	N/A
9427598	12/2015	Pilla et al.	N/A	N/A
9430615	12/2015	Michaelis et al.	N/A	N/A
9432777	12/2015	Lunner et al.	N/A	N/A
9433797	12/2015	Pilla et al.	N/A	N/A
9434692	12/2015	Xiong et al.	N/A	N/A
9436989	12/2015	Uber, III	N/A	N/A
9438650	12/2015	Serena	N/A	N/A
9439150	12/2015	Carlson et al.	N/A	N/A
9440063	12/2015	Ho et al.	N/A	N/A
9440064	12/2015	Wingeier et al.	N/A	N/A
9440070	12/2015	Goldwasser et al.	N/A	N/A
9440084	12/2015	Davis et al.	N/A	N/A
9440089	12/2015	Pilla et al.	N/A	N/A
9440646	12/2015	Fung et al.	N/A	N/A
9442088	12/2015	Feldkamp et al.	N/A	N/A
9442525	12/2015	Choi et al.	N/A	N/A
9443141	12/2015	Mirowski et al.	N/A	N/A
9444998	12/2015	Kim et al.	N/A	N/A
9445713	12/2015	Douglas et al.	N/A	N/A
9445730	12/2015	Snyder et al.	N/A	N/A
9445739	12/2015	Payton et al.	N/A	N/A
9445763	12/2015	Davis et al.	N/A	N/A
9446238	12/2015	Lozano	N/A	N/A
9448289	12/2015	Wang et al.	N/A	N/A
9449147	12/2015	Taylor	N/A	N/A
9451303	12/2015	Kothuri et al.	N/A	N/A
9451734	12/2015	Onuma et al.	N/A	N/A
9451883	12/2015	Gallant et al.	N/A	N/A
9451886	12/2015	Teixeira	N/A	N/A
9451899	12/2015	Ritchey et al.	N/A	N/A
9452287	12/2015	Rosenbluth et al.	N/A	N/A
9453215	12/2015	Deisseroth et al.	N/A	N/A
9454646	12/2015	Siefert	N/A	N/A
9458208	12/2015	Deisseroth et al.	N/A	N/A
9459597	12/2015	Kahn et al.	N/A	N/A
9460400	12/2015	De Bruin et al.	N/A	N/A
9462733	12/2015	Hokari	N/A	N/A

9462956	12/2015	Pandia et al.	N/A	N/A
9462975	12/2015	Sackner et al.	N/A	N/A
9462977	12/2015	Horseman	N/A	N/A
9463327	12/2015	Lempka et al.	N/A	N/A
9468541	12/2015	Contreras-Vidal et al.	N/A	N/A
9468761	12/2015	Frei et al.	N/A	N/A
9470728	12/2015	George et al.	N/A	N/A
9471978	12/2015	Chen et al.	N/A	N/A
9472000	12/2015	Dempsey et al.	N/A	N/A
9474481	12/2015	Dagum	N/A	N/A
9474852	12/2015	Lozano et al.	N/A	N/A
9474903	12/2015	Chen et al.	N/A	N/A
9475502	12/2015	Fung et al.	N/A	N/A
RE46189	12/2015	Prichet et al.	N/A	N/A
RE46209	12/2015	Gong et al.	N/A	N/A
9480402	12/2015	Leuthardt et al.	N/A	N/A
9480425	12/2015	Culver et al.	N/A	N/A
9480812	12/2015	Thompson	N/A	N/A
9480841	12/2015	Hershey et al.	N/A	N/A
9480845	12/2015	Harris et al.	N/A	N/A
9480854	12/2015	Von Ohlsen et al.	N/A	N/A
9483117	12/2015	Karkkainen et al.	N/A	N/A
9483613	12/2015	Fueyo et al.	N/A	N/A
9486168	12/2015	Bonmassar et al.	N/A	N/A
9486381	12/2015	Juto et al.	N/A	N/A
9486389	12/2015	Tass	N/A	N/A
9486618	12/2015	Wingeier et al.	N/A	N/A
9486632	12/2015	Saab	N/A	N/A
9489854	12/2015	Haruta et al.	N/A	N/A
9492084	12/2015	Behar et al.	N/A	N/A
9492114	12/2015	Reiman	N/A	N/A
9492120	12/2015	Horseman	N/A	N/A
9492313	12/2015	Nofzinger	N/A	N/A
9492656	12/2015	Chow et al.	N/A	N/A
9492678	12/2015	Chow	N/A	N/A
9495684	12/2015	Jung et al.	N/A	N/A
9497017	12/2015	Kim et al.	N/A	N/A
9498134	12/2015	Trobaugh et al.	N/A	N/A
9498628	12/2015	Kaemmerer et al.	N/A	N/A
9498634	12/2015	De Ridder	N/A	N/A
9500722	12/2015	Takahashi	N/A	N/A
9501829	12/2015	Carlton et al.	N/A	N/A
9504390	12/2015	Osorio	N/A	N/A
9504410	12/2015	Gal	N/A	N/A
9504420	12/2015	Davis et al.	N/A	N/A
9504788	12/2015	Hyde et al.	N/A	N/A
9505402	12/2015	Fung et al.	N/A	N/A
9505817	12/2015	Deisseroth et al.	N/A	N/A
9510790	12/2015	Kang et al.	N/A	N/A
9513398	12/2015	Wilson et al.	N/A	N/A
9517020	12/2015	Shacham-Diamand et al.	N/A	N/A
9517031	12/2015	Jung	N/A	N/A
9517222	12/2015	Goodenowe	N/A	N/A
9519981	12/2015	Sudarsky et al.	N/A	N/A
9521958	12/2015	Nagasaka et al.	N/A	N/A
9522085	12/2015	Kilgard et al.	N/A	N/A
9522278	12/2015	Heldman et al.	N/A	N/A
9522282	12/2015	Chow et al.	N/A	N/A
9522288	12/2015	Deisseroth et al.	N/A	N/A
9526419	12/2015	Derchak et al.	N/A	N/A
9526902	12/2015	Blum et al.	N/A	N/A
9526906	12/2015	Mashiach	N/A	N/A
9526913	12/2015	Vo-Dinh et al.	N/A	N/A
9526914	12/2015	Vo-Dinh et al.	N/A	N/A
9533113	12/2016	Lain et al.	N/A	N/A
9533144	12/2016	Bahmer	N/A	N/A
9533147	12/2016	Osorio	N/A	N/A
9533148	12/2016	Carcieri	N/A	N/A
9533150	12/2016	Nudo et al.	N/A	N/A
9533151	12/2016	Craig	N/A	N/A
9534044	12/2016	El-Agnaf	N/A	N/A
9538635	12/2016	Beran	N/A	N/A
9538948	12/2016	Dagum	N/A	N/A
9538951	12/2016	Osorio	N/A	N/A
9539118	12/2016	Leuthardt et al.	N/A	N/A

9541383	12/2016	Abovitz et al.	N/A	N/A
9545221	12/2016	Adhikari et al.	N/A	N/A
9545222	12/2016	Derchak et al.	N/A	N/A
9545225	12/2016	Cavuoto et al.	N/A	N/A
9545226	12/2016	Osorio	N/A	N/A
9545285	12/2016	Ghaffari et al.	N/A	N/A
9545510	12/2016	Kokones et al.	N/A	N/A
9545515	12/2016	Wolpaw et al.	N/A	N/A
9549691	12/2016	Tran	N/A	N/A
9550064	12/2016	Mashiach	N/A	N/A
9556149	12/2016	Krishnan et al.	N/A	N/A
9556487	12/2016	Umansky et al.	N/A	N/A
9557439	12/2016	Wilson et al.	N/A	N/A
9558558	12/2016	Stehle et al.	N/A	N/A
9560458	12/2016	Lunner et al.	N/A	N/A
9560967	12/2016	Hyde et al.	N/A	N/A
9560984	12/2016	Pradeep et al.	N/A	N/A
9560986	12/2016	Varcoe	N/A	N/A
9561380	12/2016	Carcieri et al.	N/A	N/A
9562988	12/2016	Wilson et al.	N/A	N/A
9563273	12/2016	Mann	N/A	N/A
9563740	12/2016	Abdelghani et al.	N/A	N/A
9563950	12/2016	Raj	N/A	N/A
9566426	12/2016	Simon et al.	N/A	N/A
9567327	12/2016	Xiong et al.	N/A	N/A
9568564	12/2016	Ma et al.	N/A	N/A
9568635	12/2016	Suhami	N/A	N/A
9572996	12/2016	Tass et al.	N/A	N/A
9577992	12/2016	Zizi et al.	N/A	N/A
9578425	12/2016	Hakansson	N/A	N/A
9579035	12/2016	Sarkela	N/A	N/A
9579048	12/2016	Rayner et al.	N/A	N/A
9579247	12/2016	Juto et al.	N/A	N/A
9579457	12/2016	Osorio	N/A	N/A
9579506	12/2016	Osorio	N/A	N/A
9582072	12/2016	Connor	N/A	N/A
9582152	12/2016	Gulaka et al.	N/A	N/A
9582925	12/2016	Durand et al.	N/A	N/A
9584928	12/2016	Laudanski et al.	N/A	N/A
9585581	12/2016	Mullins et al.	N/A	N/A
9585723	12/2016	Taylor	N/A	N/A
9586047	12/2016	Osorio et al.	N/A	N/A
9586053	12/2016	Moffitt et al.	N/A	N/A
9588203	12/2016	Zhu et al.	N/A	N/A
9588490	12/2016	Tsang	N/A	N/A
9590986	12/2016	Zizi et al.	N/A	N/A
9592003	12/2016	Osorio et al.	N/A	N/A
9592004	12/2016	DiLorenzo et al.	N/A	N/A
9592384	12/2016	Tass	N/A	N/A
9592387	12/2016	Skelton et al.	N/A	N/A
9592389	12/2016	Moffitt	N/A	N/A
9592409	12/2016	Yoo et al.	N/A	N/A
9596224	12/2016	Woods et al.	N/A	N/A
9597493	12/2016	Wingeier et al.	N/A	N/A
9597494	12/2016	Wingeier et al.	N/A	N/A
9597501	12/2016	Danilov et al.	N/A	N/A
9597504	12/2016	Danilov et al.	N/A	N/A
9600138	12/2016	Thomas et al.	N/A	N/A
9600778	12/2016	Sapiro et al.	N/A	N/A
9604056	12/2016	Starr et al.	N/A	N/A
9604067	12/2016	Kothandaraman et al.	N/A	N/A
9604073	12/2016	Deisseroth et al.	N/A	N/A
9607023	12/2016	Swamy	N/A	N/A
9607377	12/2016	Lovberg et al.	N/A	N/A
9609453	12/2016	Jabri	N/A	N/A
9610442	12/2016	Yoo et al.	N/A	N/A
9610456	12/2016	Linke et al.	N/A	N/A
9610459	12/2016	Burnett et al.	N/A	N/A
9612295	12/2016	Toda et al.	N/A	N/A
9613184	12/2016	Giftakis et al.	N/A	N/A
9613186	12/2016	Fonte	N/A	N/A
9615746	12/2016	Horseman	N/A	N/A
9615749	12/2016	Clifton et al.	N/A	N/A
9615789	12/2016	Deisseroth et al.	N/A	N/A
9616166	12/2016	Kalafut et al.	N/A	N/A

9616227	12/2016	Lindenthaler et al.	N/A	N/A
9618591	12/2016	Radparvar et al.	N/A	N/A
9622660	12/2016	Le et al.	N/A	N/A
9622672	12/2016	Yoshida et al.	N/A	N/A
9622675	12/2016	Leyde et al.	N/A	N/A
9622676	12/2016	Masmanidis et al.	N/A	N/A
9622700	12/2016	Sahasrabudhe et al.	N/A	N/A
9622702	12/2016	Badower et al.	N/A	N/A
9622703	12/2016	Badower et al.	N/A	N/A
9623240	12/2016	Simon et al.	N/A	N/A
9623241	12/2016	Wagner et al.	N/A	N/A
9626756	12/2016	Dean et al.	N/A	N/A
9629548	12/2016	Sachanandani et al.	N/A	N/A
9629568	12/2016	Hagedorn et al.	N/A	N/A
9629976	12/2016	Acton	N/A	N/A
9630004	12/2016	Rajguru et al.	N/A	N/A
9630008	12/2016	McLaughlin et al.	N/A	N/A
9630011	12/2016	Lipani	N/A	N/A
9630029	12/2016	Wurster et al.	N/A	N/A
9636019	12/2016	Hendler et al.	N/A	N/A
9636185	12/2016	Quaid et al.	N/A	N/A
9640167	12/2016	DeFranks et al.	N/A	N/A
9641665	12/2016	Lee et al.	N/A	N/A
9642552	12/2016	Hua	N/A	N/A
9642553	12/2016	Hokari	N/A	N/A
9642554	12/2016	Simola et al.	N/A	N/A
9642699	12/2016	Wortz et al.	N/A	N/A
9643015	12/2016	Moffitt et al.	N/A	N/A
9643017	12/2016	Carcieri et al.	N/A	N/A
9643019	12/2016	Higgins et al.	N/A	N/A
9646248	12/2016	Benvenuto et al.	N/A	N/A
9649030	12/2016	Gross et al.	N/A	N/A
9649036	12/2016	Teixeira	N/A	N/A
9649439	12/2016	John	N/A	N/A
9649493	12/2016	Mashiach	N/A	N/A
9649494	12/2016	Gerber et al.	N/A	N/A
9649501	12/2016	Best	N/A	N/A
9651368	12/2016	Abovitz et al.	N/A	N/A
9651706	12/2016	Mandviwala et al.	N/A	N/A
9652626	12/2016	Son et al.	N/A	N/A
9652871	12/2016	Han et al.	N/A	N/A
9655573	12/2016	Majewski et al.	N/A	N/A
9655669	12/2016	Palti et al.	N/A	N/A
9656069	12/2016	Danilov et al.	N/A	N/A
9656075	12/2016	Osorio	N/A	N/A
9656078	12/2016	Danilov et al.	N/A	N/A
9656096	12/2016	Pilla	N/A	N/A
9659186	12/2016	Pinsky et al.	N/A	N/A
9659229	12/2016	Clifton et al.	N/A	N/A
9662049	12/2016	Scarantino et al.	N/A	N/A
9662069	12/2016	De Graff et al.	N/A	N/A
9662083	12/2016	Sakaue	N/A	N/A
9662490	12/2016	Tracey et al.	N/A	N/A
9662492	12/2016	Tucker et al.	N/A	N/A
9662502	12/2016	Giuffrida et al.	N/A	N/A
9664856	12/2016	Nagasaka	N/A	N/A
9665824	12/2016	Chang et al.	N/A	N/A
9665987	12/2016	Fateh	N/A	N/A
9668694	12/2016	Badower	N/A	N/A
9669185	12/2016	Nofzinger	N/A	N/A
9669239	12/2016	Carpentier	N/A	N/A
9672302	12/2016	Dean et al.	N/A	N/A
9672617	12/2016	Dean et al.	N/A	N/A
9674621	12/2016	Bahmer	N/A	N/A
9675254	12/2016	Semenov	N/A	N/A
9675255	12/2016	Semenov	N/A	N/A
9675292	12/2016	Fadem	N/A	N/A
9675794	12/2016	Miller	N/A	N/A
9675809	12/2016	Chow	N/A	N/A
9681814	12/2016	Galloway et al.	N/A	N/A
9681820	12/2016	Wagner	N/A	N/A
9682232	12/2016	Shore et al.	N/A	N/A
9682241	12/2016	Hyde et al.	N/A	N/A
9684051	12/2016	Nieminen et al.	N/A	N/A
9684335	12/2016	Kim et al.	N/A	N/A

9685600	12/2016	Washington, II et al.	N/A	N/A
9687187	12/2016	Dagum	N/A	N/A
9687562	12/2016	Satchi-Fainaro et al.	N/A	N/A
9693684	12/2016	Lopez et al.	N/A	N/A
9693724	12/2016	Dagum	N/A	N/A
9693725	12/2016	Soza	N/A	N/A
9693734	12/2016	Horseman	N/A	N/A
9694155	12/2016	Panova et al.	N/A	N/A
9694178	12/2016	Ruffini et al.	N/A	N/A
9694197	12/2016	Segal	N/A	N/A
9697330	12/2016	Taylor	N/A	N/A
9697336	12/2016	Hyde et al.	N/A	N/A
9700256	12/2016	Osorio et al.	N/A	N/A
9700716	12/2016	Faltys et al.	N/A	N/A
9700723	12/2016	Sabesan	N/A	N/A
9704205	12/2016	Akutagawa et al.	N/A	N/A
9706910	12/2016	Blaha et al.	N/A	N/A
9706925	12/2016	Taylor	N/A	N/A
9706957	12/2016	Wu et al.	N/A	N/A
9706963	12/2016	Gupta et al.	N/A	N/A
9707372	12/2016	Smith	N/A	N/A
9707390	12/2016	Ahmed	N/A	N/A
9707391	12/2016	Ahmed	N/A	N/A
9707396	12/2016	Su et al.	N/A	N/A
9710788	12/2016	Horseman	N/A	N/A
9712736	12/2016	Kearns et al.	N/A	N/A
9713428	12/2016	Chon et al.	N/A	N/A
9713433	12/2016	Gadot et al.	N/A	N/A
9713444	12/2016	Severson	N/A	N/A
9713712	12/2016	Wingeier et al.	N/A	N/A
9715032	12/2016	Song et al.	N/A	N/A
9717461	12/2016	Yu et al.	N/A	N/A
9717904	12/2016	Simon et al.	N/A	N/A
9717920	12/2016	Heldman et al.	N/A	N/A
9724517	12/2016	Giftakis et al.	N/A	N/A
9729252	12/2016	Tyler et al.	N/A	N/A
9732039	12/2016	Xiong et al.	N/A	N/A
9734589	12/2016	Yu et al.	N/A	N/A
9734601	12/2016	Bresler et al.	N/A	N/A
9734632	12/2016	Thomas et al.	N/A	N/A
9737230	12/2016	Sarma et al.	N/A	N/A
9740710	12/2016	Han et al.	N/A	N/A
9740946	12/2016	Varkuti et al.	N/A	N/A
9741114	12/2016	Varkuti	N/A	N/A
9743197	12/2016	Petersen et al.	N/A	N/A
9743835	12/2016	Taylor	N/A	N/A
9744358	12/2016	Hehrmann et al.	N/A	N/A
9763592	12/2016	Le et al.	N/A	N/A
2001/0003799	12/2000	Boveja	N/A	N/A
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2004/0015205	12/2003	Whitehurst et al.	N/A	N/A
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2005/0159670	12/2004	Sneddon	N/A	N/A
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2005/0167588	12/2004	Donnangelo	N/A	N/A
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2005/0182287	12/2004	Becker	N/A	N/A
2005/0182288	12/2004	Zabara	N/A	N/A
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2005/0283053	12/2004	deCharms	N/A	N/A
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2006/0217781	12/2005	John	N/A	N/A
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2006/0239482	12/2005	Hatoum	N/A	N/A
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2007/0255135	12/2006	Kalafut et al.	N/A	N/A
2007/0255155	12/2006	Drew et al.	N/A	N/A
2007/0255320	12/2006	Inman et al.	N/A	N/A
2007/0255379	12/2006	Williams et al.	N/A	N/A
2007/0255531	12/2006	Drew	N/A	N/A
2007/0259323	12/2006	Brown et al.	N/A	N/A
2007/0260151	12/2006	Clifford	N/A	N/A
2007/0265508	12/2006	Sheikhzadeh-Nadjar et al.	N/A	N/A
2007/0265533	12/2006	Tran	N/A	N/A
2007/0273504	12/2006	Tran	N/A	N/A
2007/0273611	12/2006	Torch	N/A	N/A
2007/0276270	12/2006	Tran	N/A	N/A
2007/0276278	12/2006	Coyle et al.	N/A	N/A
2007/0276279	12/2006	Echaz et al.	N/A	N/A
2007/0276441	12/2006	Goetz	N/A	N/A
2007/0276609	12/2006	Greenwald	N/A	N/A
2007/0280508	12/2006	Ernst et al.	N/A	N/A
2007/0282228	12/2006	Einav et al.	N/A	N/A
2007/0287896	12/2006	Derchak et al.	N/A	N/A
2007/0291832	12/2006	Diab et al.	N/A	N/A
2007/0293760	12/2006	Schaafsma	N/A	N/A
2007/0299370	12/2006	Bystritsky	N/A	N/A
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2008/0001600	12/2007	deCharms	N/A	N/A
2008/0001735	12/2007	Tran	N/A	N/A
2008/0004514	12/2007	Diab et al.	N/A	N/A
2008/0004550	12/2007	Einav et al.	N/A	N/A
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2008/0009685	12/2007	Kim et al.	N/A	N/A
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2008/0013747	12/2007	Tran	N/A	N/A
2008/0015458	12/2007	Buarque de Macedo et al.	N/A	N/A
2008/0015459	12/2007	Llinas	N/A	N/A
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2008/0021336	12/2007	Dobak	N/A	N/A
2008/0021340	12/2007	Sarkela	N/A	N/A
2008/0021341	12/2007	Harris et al.	N/A	N/A
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2009/0209831	12/2008	Kucharczyk et al.	N/A	N/A
2009/0209835	12/2008	Diab et al.	N/A	N/A
2009/0209845	12/2008	Christen et al.	N/A	N/A
2009/0210018	12/2008	Lozano	N/A	N/A
2009/0216091	12/2008	Arndt	N/A	N/A
2009/0216146	12/2008	Teicher et al.	N/A	N/A
2009/0216288	12/2008	Schiff et al.	N/A	N/A
2009/0220425	12/2008	Moxon et al.	N/A	N/A
2009/0220429	12/2008	Johnsen et al.	N/A	N/A
2009/0221904	12/2008	Shealy et al.	N/A	N/A
2009/0221928	12/2008	Einav et al.	N/A	N/A
2009/0221930	12/2008	Laken	N/A	N/A
2009/0227876	12/2008	Tran	N/A	N/A
2009/0227877	12/2008	Tran	N/A	N/A
2009/0227882	12/2008	Foo	N/A	N/A
2009/0227889	12/2008	John et al.	N/A	N/A
2009/0234419	12/2008	Maschino et al.	N/A	N/A
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2009/0243756	12/2008	Stevenson et al.	N/A	N/A
2009/0246138	12/2008	Santosh et al.	N/A	N/A
2009/0247893	12/2008	Lapinlampi et al.	N/A	N/A
2009/0247894	12/2008	Causevic	N/A	N/A
2009/0259277	12/2008	Cornejo Cruz et al.	N/A	N/A
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2009/0264952	12/2008	Jassemidis et al.	N/A	N/A
2009/0264954	12/2008	Rise et al.	N/A	N/A

2009/0264955	12/2008	Giftakis et al.	N/A	N/A
2009/0264956	12/2008	Rise et al.	N/A	N/A
2009/0264957	12/2008	Giftakis et al.	N/A	N/A
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2009/0270694	12/2008	Hyde et al.	N/A	N/A
2009/0270754	12/2008	Moridaira	N/A	N/A
2009/0270758	12/2008	Eagleman et al.	N/A	N/A
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2009/0287108	12/2008	Levy	N/A	N/A
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2009/0287274	12/2008	De Ridder	N/A	N/A
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2009/0299169	12/2008	deCharms	N/A	N/A
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2010/0056276	12/2009	Silberstein	N/A	N/A
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2010/0063563	12/2009	Craig	N/A	N/A
2010/0068751	12/2009	Eberle	N/A	N/A
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2010/0069739	12/2009	deCharms	N/A	N/A

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2010/0069775	12/2009	Milgramm et al.	N/A	N/A
2010/0069777	12/2009	Marks	N/A	N/A
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2010/0076274	12/2009	Severson	N/A	N/A
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2010/0076334	12/2009	Rothblatt	N/A	N/A
2010/0076338	12/2009	Kwak	N/A	N/A
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2010/0087900	12/2009	Flint	N/A	N/A
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2010/0286549	12/2009	John et al.	N/A	N/A

2010/0286747	12/2009	Sabesan et al.	N/A	N/A
2010/0292602	12/2009	Worrell et al.	N/A	N/A
2010/0292752	12/2009	Bardakjian et al.	N/A	N/A
2010/0293002	12/2009	Firminger et al.	N/A	N/A
2010/0293115	12/2009	Seyed Momen	N/A	N/A
2010/0298624	12/2009	Becker	N/A	N/A
2010/0298735	12/2009	Suffin	N/A	N/A
2010/0303101	12/2009	Lazar et al.	N/A	N/A
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2010/0312188	12/2009	Robertson et al.	N/A	N/A
2010/0312579	12/2009	Firminger et al.	N/A	N/A
2010/0318025	12/2009	John	N/A	N/A
2010/0318160	12/2009	Stevenson et al.	N/A	N/A
2010/0322488	12/2009	Virtue et al.	N/A	N/A
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2010/0324441	12/2009	Hargrove et al.	N/A	N/A
2010/0331649	12/2009	Chou	N/A	N/A
2010/0331715	12/2009	Addison et al.	N/A	N/A
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2011/0015536	12/2010	Milgramm et al.	N/A	N/A
2011/0015539	12/2010	deCharms	N/A	N/A
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2011/0021970	12/2010	Vo-Dinh et al.	N/A	N/A
2011/0022981	12/2010	Mahajan et al.	N/A	N/A
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2011/0087125	12/2010	Causevic	N/A	N/A
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2011/0112394	12/2010	Mishelevich	N/A	N/A
2011/0112426	12/2010	Causevic	N/A	N/A
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2011/0160608	12/2010	Hargrove	N/A	N/A
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2011/0172564	12/2010	Drew	N/A	N/A
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2011/0208094	12/2010	Mishelevich	N/A	N/A
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2011/0208539	12/2010	Lynn	N/A	N/A
2011/0213200	12/2010	Mishelevich	N/A	N/A
2011/0213222	12/2010	Leyde et al.	N/A	N/A
2011/0217240	12/2010	Ferris	N/A	N/A
2011/0218405	12/2010	Avinash et al.	N/A	N/A
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2011/0224569	12/2010	Isenhardt et al.	N/A	N/A
2011/0224570	12/2010	Causevic	N/A	N/A
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2011/0230738	12/2010	Chance	N/A	N/A
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2011/0238136	12/2010	Bourget et al.	N/A	N/A
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2012/0059246	12/2011	Taylor	N/A	N/A
2012/0059273	12/2011	Meggiolaro et al.	N/A	N/A
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2012/0060851	12/2011	Amberg	N/A	N/A
2012/0065536	12/2011	Causevic et al.	N/A	N/A
2012/0070044	12/2011	Avinash et al.	N/A	N/A
2012/0071771	12/2011	Behar	N/A	N/A
2012/0078115	12/2011	Lonky	N/A	N/A
2012/0078323	12/2011	Osorio	N/A	N/A
2012/0078327	12/2011	Sloan et al.	N/A	N/A
2012/0080305	12/2011	Koruga	N/A	N/A
2012/0083668	12/2011	Pradeep et al.	N/A	N/A
2012/0083690	12/2011	Semenov	N/A	N/A
2012/0083700	12/2011	Osorio	N/A	N/A
2012/0083701	12/2011	Osorio	N/A	N/A
2012/0083708	12/2011	Rajdev et al.	N/A	N/A
2012/0088987	12/2011	Braun et al.	N/A	N/A
2012/0088992	12/2011	Armitstead	N/A	N/A
2012/0089004	12/2011	Hsu et al.	N/A	N/A
2012/0089205	12/2011	Boyden et al.	N/A	N/A
2012/0092156	12/2011	Tran	N/A	N/A
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2012/0095352	12/2011	Tran	N/A	N/A
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2012/0100514	12/2011	Desain et al.	N/A	N/A
2012/0101326	12/2011	Simon et al.	N/A	N/A
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2012/0101401	12/2011	Faul et al.	N/A	N/A
2012/0101402	12/2011	Nguyen	N/A	N/A
2012/0101430	12/2011	Robertson et al.	N/A	N/A
2012/0101544	12/2011	Hoberman et al.	N/A	N/A
2012/0108909	12/2011	Slobounov et al.	N/A	N/A
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2012/0109020	12/2011	Wagner et al.	N/A	N/A
2012/0116149	12/2011	Pilla et al.	N/A	N/A
2012/0116179	12/2011	Drew et al.	N/A	N/A
2012/0116235	12/2011	Trumble et al.	N/A	N/A
2012/0116244	12/2011	McIntyre et al.	N/A	N/A
2012/0116475	12/2011	Nelson et al.	N/A	N/A
2012/0116741	12/2011	Choi et al.	N/A	N/A
2012/0123232	12/2011	Najarian et al.	N/A	N/A
2012/0123290	12/2011	Kidmose et al.	N/A	N/A

2012/0125337	12/2011	Asanoi	N/A	N/A
2012/0128683	12/2011	Shantha	N/A	N/A
2012/0130204	12/2011	Basta et al.	N/A	N/A
2012/0130228	12/2011	Zellers et al.	N/A	N/A
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2012/0130300	12/2011	Stavchansky et al.	N/A	N/A
2012/0130641	12/2011	Morrison et al.	N/A	N/A
2012/0136242	12/2011	Qi et al.	N/A	N/A
2012/0136274	12/2011	Burdea et al.	N/A	N/A
2012/0136605	12/2011	Addison et al.	N/A	N/A
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2012/0143074	12/2011	Shin et al.	N/A	N/A
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2012/0143104	12/2011	Tee et al.	N/A	N/A
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2012/0149997	12/2011	Diab et al.	N/A	N/A
2012/0150255	12/2011	Lindenthaler et al.	N/A	N/A
2012/0150257	12/2011	Aur et al.	N/A	N/A
2012/0150262	12/2011	Gliner et al.	N/A	N/A
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2012/0157804	12/2011	Rogers et al.	N/A	N/A
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2012/0158092	12/2011	Thimineur et al.	N/A	N/A
2012/0159656	12/2011	Gerber et al.	N/A	N/A
2012/0162002	12/2011	Semenov	N/A	N/A
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2012/0165631	12/2011	Diab et al.	N/A	N/A
2012/0165696	12/2011	Arns	N/A	N/A
2012/0165898	12/2011	Moffitt	N/A	N/A
2012/0165899	12/2011	Gliner	N/A	N/A
2012/0165904	12/2011	Lee et al.	N/A	N/A
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2012/0179071	12/2011	Skelton	N/A	N/A
2012/0179228	12/2011	DeCharms	N/A	N/A
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2012/0191158	12/2011	Craig	N/A	N/A
2012/0191542	12/2011	Nurmi	N/A	N/A
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2012/0197163	12/2011	Mishelevich	N/A	N/A
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2012/0203079	12/2011	McLaughlin	N/A	N/A
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2012/0203130	12/2011	Bernhard	N/A	N/A
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2012/0302845	12/2011	Lynn et al.	N/A	N/A
2012/0302856	12/2011	Chang et al.	N/A	N/A
2012/0302867	12/2011	Ichimura	N/A	N/A
2012/0302894	12/2011	Diab et al.	N/A	N/A
2012/0302912	12/2011	Moffitt et al.	N/A	N/A
2012/0303080	12/2011	Ben-David et al.	N/A	N/A
2012/0303087	12/2011	Moffitt et al.	N/A	N/A
2012/0310050	12/2011	Osorio	N/A	N/A
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2012/0310105	12/2011	Feingold et al.	N/A	N/A
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2012/0316793	12/2011	Jung et al.	N/A	N/A
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2012/0321759	12/2011	Marinkovich et al.	N/A	N/A
2012/0323108	12/2011	Carroll	N/A	N/A
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2013/0006124	12/2012	Eyal et al.	N/A	N/A
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2013/0054214	12/2012	Taylor	N/A	N/A
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2013/0144353	12/2012	Lozano	N/A	N/A
2013/0144537	12/2012	Schalk et al.	N/A	N/A
2013/0150650	12/2012	Phillips et al.	N/A	N/A
2013/0150651	12/2012	Phillips et al.	N/A	N/A
2013/0150659	12/2012	Shaw et al.	N/A	N/A
2013/0150702	12/2012	Hokari	N/A	N/A
2013/0150921	12/2012	Singhal et al.	N/A	N/A
2013/0151163	12/2012	Taylor et al.	N/A	N/A
2013/0158883	12/2012	Hasegawa et al.	N/A	N/A
2013/0159041	12/2012	Jayaraman et al.	N/A	N/A
2013/0165766	12/2012	Nishikawa et al.	N/A	N/A
2013/0165804	12/2012	Johnson et al.	N/A	N/A
2013/0165812	12/2012	Aksenova et al.	N/A	N/A
2013/0165846	12/2012	Peyman	N/A	N/A
2013/0165996	12/2012	Meadows et al.	N/A	N/A
2013/0167360	12/2012	Masmanidis et al.	N/A	N/A

2013/0172663	12/2012	Leonard	N/A	N/A
2013/0172686	12/2012	Addison et al.	N/A	N/A
2013/0172691	12/2012	Tran	N/A	N/A
2013/0172716	12/2012	Lozano et al.	N/A	N/A
2013/0172763	12/2012	Wheeler	N/A	N/A
2013/0172767	12/2012	Dripps et al.	N/A	N/A
2013/0172772	12/2012	Alshaer et al.	N/A	N/A
2013/0172774	12/2012	Crowder et al.	N/A	N/A
2013/0178693	12/2012	Neuvonen et al.	N/A	N/A
2013/0178718	12/2012	Tran et al.	N/A	N/A
2013/0178733	12/2012	Langleben	N/A	N/A
2013/0178913	12/2012	Lozano	N/A	N/A
2013/0182860	12/2012	Adachi et al.	N/A	N/A
2013/0184218	12/2012	Paul et al.	N/A	N/A
2013/0184516	12/2012	Genereux et al.	N/A	N/A
2013/0184552	12/2012	Westermann et al.	N/A	N/A
2013/0184558	12/2012	Gallant et al.	N/A	N/A
2013/0184597	12/2012	Hopenfeld	N/A	N/A
2013/0184603	12/2012	Rothman	N/A	N/A
2013/0184639	12/2012	Whitehurst et al.	N/A	N/A
2013/0184728	12/2012	Mishelevich	N/A	N/A
2013/0184781	12/2012	Eskandar et al.	N/A	N/A
2013/0184786	12/2012	Goetz	N/A	N/A
2013/0184792	12/2012	Simon et al.	N/A	N/A
2013/0184997	12/2012	Mott	N/A	N/A
2013/0185144	12/2012	Pradeep et al.	N/A	N/A
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2013/0188830	12/2012	Ernst et al.	N/A	N/A
2013/0188854	12/2012	Bilgic et al.	N/A	N/A
2013/0189663	12/2012	Tuchschmid et al.	N/A	N/A
2013/0190577	12/2012	Brunner et al.	N/A	N/A
2013/0190642	12/2012	Muesch et al.	N/A	N/A
2013/0197321	12/2012	Wilson	N/A	N/A
2013/0197322	12/2012	Tran	N/A	N/A
2013/0197328	12/2012	Diab et al.	N/A	N/A
2013/0197339	12/2012	Bardakjian et al.	N/A	N/A
2013/0197401	12/2012	Sato et al.	N/A	N/A
2013/0197944	12/2012	Drew et al.	N/A	N/A
2013/0203019	12/2012	Nolen	N/A	N/A
2013/0204085	12/2012	Alexander et al.	N/A	N/A
2013/0204122	12/2012	Hendler et al.	N/A	N/A
2013/0204144	12/2012	Colborn et al.	N/A	N/A
2013/0204150	12/2012	Similowski et al.	N/A	N/A
2013/0211183	12/2012	Schiffer	N/A	N/A
2013/0211224	12/2012	Isenhardt et al.	N/A	N/A
2013/0211238	12/2012	DeCharms	N/A	N/A
2013/0211276	12/2012	Luo et al.	N/A	N/A
2013/0211291	12/2012	Tran	N/A	N/A
2013/0211728	12/2012	Taylor et al.	N/A	N/A
2013/0217982	12/2012	Behzadi	N/A	N/A
2013/0218043	12/2012	Yoshida	N/A	N/A
2013/0218053	12/2012	Kaiser et al.	N/A	N/A
2013/0218232	12/2012	Giftakis et al.	N/A	N/A
2013/0218233	12/2012	Warschewske et al.	N/A	N/A
2013/0218819	12/2012	Lujan et al.	N/A	N/A
2013/0221961	12/2012	Liu	N/A	N/A
2013/0223709	12/2012	Wagner	N/A	N/A
2013/0225940	12/2012	Fujita et al.	N/A	N/A
2013/0225953	12/2012	Oliviero et al.	N/A	N/A
2013/0225992	12/2012	Osorio	N/A	N/A
2013/0226261	12/2012	Sparks et al.	N/A	N/A
2013/0226408	12/2012	Fung et al.	N/A	N/A
2013/0226464	12/2012	Marci et al.	N/A	N/A
2013/0231574	12/2012	Tran	N/A	N/A
2013/0231580	12/2012	Chen et al.	N/A	N/A
2013/0231709	12/2012	Lozano	N/A	N/A
2013/0231716	12/2012	Skelton et al.	N/A	N/A
2013/0231721	12/2012	DeCharms	N/A	N/A
2013/0231947	12/2012	Shusterman	N/A	N/A
2013/0234823	12/2012	Kahn et al.	N/A	N/A
2013/0235550	12/2012	Stevenson et al.	N/A	N/A
2013/0237541	12/2012	Teegarden et al.	N/A	N/A
2013/0237874	12/2012	Zoicas	N/A	N/A
2013/0238049	12/2012	Simon et al.	N/A	N/A
2013/0238050	12/2012	Simon et al.	N/A	N/A

2013/0238053	12/2012	Ignagni et al.	N/A	N/A
2013/0238063	12/2012	Nofzinger	N/A	N/A
2013/0242262	12/2012	Lewis	N/A	N/A
2013/0243287	12/2012	Thomson et al.	N/A	N/A
2013/0244323	12/2012	Deisseroth et al.	N/A	N/A
2013/0245416	12/2012	Yarmush et al.	N/A	N/A
2013/0245422	12/2012	D'arcy et al.	N/A	N/A
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2013/0245886	12/2012	Fung et al.	N/A	N/A
2013/0251641	12/2012	Akhtari et al.	N/A	N/A
2013/0253363	12/2012	Juffali et al.	N/A	N/A
2013/0253612	12/2012	Chow	N/A	N/A
2013/0255586	12/2012	Gerashchenko	N/A	N/A
2013/0261490	12/2012	Truccolo et al.	N/A	N/A
2013/0261506	12/2012	Mishelevich	N/A	N/A
2013/0261703	12/2012	Chow et al.	N/A	N/A
2013/0266163	12/2012	Morikawa et al.	N/A	N/A
2013/0267760	12/2012	Jin	N/A	N/A
2013/0267866	12/2012	Nakashima et al.	N/A	N/A
2013/0267928	12/2012	Imran et al.	N/A	N/A
2013/0274562	12/2012	Ghaffari et al.	N/A	N/A
2013/0274580	12/2012	Madsen et al.	N/A	N/A
2013/0274586	12/2012	Miyazaki et al.	N/A	N/A
2013/0274625	12/2012	Sarma et al.	N/A	N/A
2013/0275159	12/2012	Seely	N/A	N/A
2013/0281758	12/2012	Solvason et al.	N/A	N/A
2013/0281759	12/2012	Hagedorn et al.	N/A	N/A
2013/0281811	12/2012	Imran	N/A	N/A
2013/0281879	12/2012	Raniere	N/A	N/A
2013/0281890	12/2012	Mishelevich	N/A	N/A
2013/0282075	12/2012	De Ridder	N/A	N/A
2013/0282339	12/2012	Ricci et al.	N/A	N/A
2013/0289360	12/2012	Hyde et al.	N/A	N/A
2013/0289364	12/2012	Colman et al.	N/A	N/A
2013/0289385	12/2012	Lozano et al.	N/A	N/A
2013/0289386	12/2012	Deisseroth et al.	N/A	N/A
2013/0289401	12/2012	Colbaugh et al.	N/A	N/A
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2013/0289433	12/2012	Jin et al.	N/A	N/A
2013/0289653	12/2012	Kilgard et al.	N/A	N/A
2013/0289669	12/2012	Deisseroth et al.	N/A	N/A
2013/0293844	12/2012	Gross et al.	N/A	N/A
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2013/0296406	12/2012	Deisseroth et al.	N/A	N/A
2013/0296637	12/2012	Kilgard et al.	N/A	N/A
2013/0300573	12/2012	Brown et al.	N/A	N/A
2013/0303828	12/2012	Hargrove	N/A	N/A
2013/0303934	12/2012	Collura	N/A	N/A
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2013/0304159	12/2012	Simon et al.	N/A	N/A
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2013/0308099	12/2012	Stack	N/A	N/A
2013/0309278	12/2012	Peyman	N/A	N/A
2013/0310422	12/2012	Brown et al.	N/A	N/A
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2013/0310909	12/2012	Simon et al.	N/A	N/A
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2013/0317384	12/2012	Le	N/A	N/A
2013/0317474	12/2012	Rezai et al.	N/A	N/A
2013/0317568	12/2012	Skelton	N/A	N/A
2013/0317580	12/2012	Simon et al.	N/A	N/A
2013/0318546	12/2012	Kothuri et al.	N/A	N/A
2013/0324880	12/2012	Adachi et al.	N/A	N/A
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2013/0338459	12/2012	Lynn et al.	N/A	N/A
2013/0338518	12/2012	Zoica	N/A	N/A
2013/0338526	12/2012	Howard	N/A	N/A
2013/0338738	12/2012	Garcia Molina et al.	N/A	N/A
2013/0338803	12/2012	Maoz et al.	N/A	N/A
2013/0339043	12/2012	Bakar et al.	N/A	N/A
2013/0344465	12/2012	Dickinson et al.	N/A	N/A
2013/0345522	12/2012	Sun et al.	N/A	N/A
2013/0345523	12/2012	Diab et al.	N/A	N/A
2014/0000630	12/2013	Ford	N/A	N/A
2014/0003696	12/2013	Taghva	N/A	N/A
2014/0005518	12/2013	Ko et al.	N/A	N/A
2014/0005743	12/2013	Giuffrida et al.	N/A	N/A
2014/0005744	12/2013	Hershey et al.	N/A	N/A
2014/0005988	12/2013	Brockway	N/A	N/A
2014/0012061	12/2013	Song et al.	N/A	N/A
2014/0012110	12/2013	Watson et al.	N/A	N/A
2014/0012133	12/2013	Sverdlik et al.	N/A	N/A
2014/0012153	12/2013	Greenwald	N/A	N/A
2014/0015852	12/2013	Kantartzis et al.	N/A	N/A
2014/0018649	12/2013	Jespersen et al.	N/A	N/A
2014/0018792	12/2013	Gang et al.	N/A	N/A
2014/0019165	12/2013	Horseman	N/A	N/A
2014/0023999	12/2013	Greder	N/A	N/A
2014/0025133	12/2013	Lozano	N/A	N/A
2014/0025396	12/2013	Horseman	N/A	N/A
2014/0025397	12/2013	Horseman	N/A	N/A
2014/0029830	12/2013	Vija et al.	N/A	N/A
2014/0031703	12/2013	Rayner et al.	N/A	N/A
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2014/0031903	12/2013	Mashiach	N/A	N/A
2014/0032512	12/2013	Drew et al.	N/A	N/A
2014/0038147	12/2013	Morrow	N/A	N/A
2014/0039279	12/2013	Jarvik et al.	N/A	N/A
2014/0039290	12/2013	De Graff et al.	N/A	N/A
2014/0039336	12/2013	Osorio et al.	N/A	N/A
2014/0039571	12/2013	Wolpaw et al.	N/A	N/A
2014/0039577	12/2013	Kothandaraman et al.	N/A	N/A
2014/0039578	12/2013	Whitehurst et al.	N/A	N/A
2014/0039975	12/2013	Hill	N/A	N/A
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2014/0046208	12/2013	Sejdic et al.	N/A	N/A
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2014/0051044	12/2013	Badower et al.	N/A	N/A
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2014/0051961	12/2013	Badower et al.	N/A	N/A
2014/0052213	12/2013	Osorio	N/A	N/A
2014/0055284	12/2013	Tran et al.	N/A	N/A
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2014/0066763	12/2013	Rothberg et al.	N/A	N/A
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2014/0077946	12/2013	Tran	N/A	N/A
2014/0081071	12/2013	Simon et al.	N/A	N/A
2014/0081114	12/2013	Shachar et al.	N/A	N/A
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2014/0081353	12/2013	Cook et al.	N/A	N/A
2014/0088341	12/2013	Altman et al.	N/A	N/A
2014/0088377	12/2013	Manzke et al.	N/A	N/A
2014/0089399	12/2013	Chun	709/204	G06Q 50/01
2014/0094710	12/2013	Sarma et al.	N/A	N/A
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2014/0100467	12/2013	Baker et al.	N/A	N/A
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2014/0101084	12/2013	Li et al.	N/A	N/A
2014/0104059	12/2013	Tran	N/A	N/A
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2014/0114165	12/2013	Walker et al.	N/A	N/A
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2014/0142448	12/2013	Bae et al.	N/A	N/A
2014/0142653	12/2013	Osorio	N/A	N/A
2014/0142654	12/2013	Simon et al.	N/A	N/A
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2014/0143064	12/2013	Tran	N/A	N/A
2014/0148479	12/2013	Chesworth et al.	N/A	N/A
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2014/0148693	12/2013	Taylor	N/A	N/A
2014/0148716	12/2013	Hopenfeld et al.	N/A	N/A
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2014/0151563	12/2013	Rouso et al.	N/A	N/A
2014/0152673	12/2013	Lynn et al.	N/A	N/A
2014/0154647	12/2013	Nolen	N/A	N/A
2014/0154650	12/2013	Stack	N/A	N/A
2014/0155430	12/2013	Chesworth et al.	N/A	N/A
2014/0155706	12/2013	Kochs et al.	N/A	N/A
2014/0155714	12/2013	Gavish	N/A	N/A
2014/0155730	12/2013	Bansal et al.	N/A	N/A
2014/0155740	12/2013	Semenov	N/A	N/A
2014/0155770	12/2013	Taylor	N/A	N/A
2014/0155772	12/2013	Frei et al.	N/A	N/A
2014/0155952	12/2013	Lozano et al.	N/A	N/A
2014/0156000	12/2013	Campin et al.	N/A	N/A
2014/0159862	12/2013	Yang et al.	N/A	N/A
2014/0161352	12/2013	Buyens et al.	N/A	N/A
2014/0163328	12/2013	Geva et al.	N/A	N/A
2014/0163330	12/2013	Horseman	N/A	N/A
2014/0163331	12/2013	Horseman	N/A	N/A
2014/0163332	12/2013	Horseman	N/A	N/A
2014/0163333	12/2013	Horseman	N/A	N/A
2014/0163335	12/2013	Horseman	N/A	N/A
2014/0163336	12/2013	Horseman	N/A	N/A
2014/0163337	12/2013	Horseman	N/A	N/A
2014/0163368	12/2013	Rouso et al.	N/A	N/A
2014/0163385	12/2013	Kelleher et al.	N/A	N/A
2014/0163409	12/2013	Arndt	N/A	N/A
2014/0163425	12/2013	Tran	N/A	N/A
2014/0163627	12/2013	Starr et al.	N/A	N/A
2014/0163643	12/2013	Craig	N/A	N/A
2014/0163893	12/2013	Harumatsu et al.	N/A	N/A
2014/0163897	12/2013	Lynn et al.	N/A	N/A
2014/0171749	12/2013	Chin et al.	N/A	N/A
2014/0171757	12/2013	Kawato et al.	N/A	N/A
2014/0171819	12/2013	Patterson	N/A	N/A
2014/0171820	12/2013	Causevic	N/A	N/A
2014/0174277	12/2013	Mann	N/A	N/A
2014/0175261	12/2013	Addison et al.	N/A	N/A
2014/0176944	12/2013	Addison et al.	N/A	N/A
2014/0179980	12/2013	Phillips et al.	N/A	N/A
2014/0180088	12/2013	Rothberg et al.	N/A	N/A
2014/0180092	12/2013	Rothberg et al.	N/A	N/A
2014/0180093	12/2013	Rothberg et al.	N/A	N/A
2014/0180094	12/2013	Rothberg et al.	N/A	N/A
2014/0180095	12/2013	Rothberg et al.	N/A	N/A
2014/0180096	12/2013	Rothberg et al.	N/A	N/A
2014/0180097	12/2013	Rothberg et al.	N/A	N/A
2014/0180099	12/2013	Rothberg et al.	N/A	N/A
2014/0180100	12/2013	Rothberg et al.	N/A	N/A
2014/0180112	12/2013	Rothberg et al.	N/A	N/A
2014/0180113	12/2013	Rothberg et al.	N/A	N/A
2014/0180145	12/2013	Kanai et al.	N/A	N/A
2014/0180153	12/2013	Zia et al.	N/A	N/A
2014/0180160	12/2013	Brown et al.	N/A	N/A
2014/0180161	12/2013	Bolger et al.	N/A	N/A
2014/0180176	12/2013	Rothberg et al.	N/A	N/A
2014/0180177	12/2013	Rothberg et al.	N/A	N/A
2014/0180194	12/2013	Lozano	N/A	N/A
2014/0180358	12/2013	Giftakis et al.	N/A	N/A
2014/0180597	12/2013	Brown et al.	N/A	N/A
2014/0184550	12/2013	Hennessey et al.	N/A	N/A
2014/0187901	12/2013	Cui et al.	N/A	N/A
2014/0187994	12/2013	Thornton	N/A	N/A

2014/0188006	12/2013	Alshaer et al.	N/A	N/A
2014/0188770	12/2013	Agrafioti et al.	N/A	N/A
2014/0193336	12/2013	Rousso et al.	N/A	N/A
2014/0194702	12/2013	Tran	N/A	N/A
2014/0194720	12/2013	Hua	N/A	N/A
2014/0194726	12/2013	Mishelevich et al.	N/A	N/A
2014/0194758	12/2013	Korenberg	N/A	N/A
2014/0194759	12/2013	Weiland et al.	N/A	N/A
2014/0194768	12/2013	Nierenberg et al.	N/A	N/A
2014/0194769	12/2013	Nierenberg et al.	N/A	N/A
2014/0194780	12/2013	Alshaer et al.	N/A	N/A
2014/0194793	12/2013	Nakata et al.	N/A	N/A
2014/0200414	12/2013	Osorio	N/A	N/A
2014/0200432	12/2013	Banerji et al.	N/A	N/A
2014/0200623	12/2013	Lindenthaler et al.	N/A	N/A
2014/0203797	12/2013	Stivoric et al.	N/A	N/A
2014/0206981	12/2013	Nagasaka	N/A	N/A
2014/0207224	12/2013	Simon	N/A	N/A
2014/0207432	12/2013	Taylor	N/A	N/A
2014/0211593	12/2013	Tyler et al.	N/A	N/A
2014/0213842	12/2013	Simon et al.	N/A	N/A
2014/0213843	12/2013	Pilla et al.	N/A	N/A
2014/0213844	12/2013	Pilla et al.	N/A	N/A
2014/0213937	12/2013	Bianchi et al.	N/A	N/A
2014/0213961	12/2013	Whitehurst et al.	N/A	N/A
2014/0214135	12/2013	Ben-David et al.	N/A	N/A
2014/0214330	12/2013	Iyer et al.	N/A	N/A
2014/0214335	12/2013	Siefert	N/A	N/A
2014/0221726	12/2013	Pilla et al.	N/A	N/A
2014/0221866	12/2013	Quy	N/A	N/A
2014/0222113	12/2013	Gliner et al.	N/A	N/A
2014/0222406	12/2013	Taylor	N/A	N/A
2014/0226131	12/2013	Lopez et al.	N/A	N/A
2014/0226888	12/2013	Skidmore	N/A	N/A
2014/0228620	12/2013	Vasishta	N/A	N/A
2014/0228649	12/2013	Rayner et al.	N/A	N/A
2014/0228651	12/2013	Causevic et al.	N/A	N/A
2014/0228653	12/2013	Kiraly	N/A	N/A
2014/0228702	12/2013	Shahaf et al.	N/A	N/A
2014/0232516	12/2013	Stivoric et al.	N/A	N/A
2014/0235826	12/2013	Deisseroth et al.	N/A	N/A
2014/0235965	12/2013	Tran	N/A	N/A
2014/0236039	12/2013	Strokova Aksenova et al.	N/A	N/A
2014/0236077	12/2013	Robertson et al.	N/A	N/A
2014/0236272	12/2013	Simon et al.	N/A	N/A
2014/0236492	12/2013	Taylor	N/A	N/A
2014/0237073	12/2013	Schiff	N/A	N/A
2014/0243608	12/2013	Hunt	N/A	N/A
2014/0243613	12/2013	Osorio	N/A	N/A
2014/0243614	12/2013	Rothberg et al.	N/A	N/A
2014/0243621	12/2013	Weng et al.	N/A	N/A
2014/0243628	12/2013	Ochs et al.	N/A	N/A
2014/0243647	12/2013	Clark et al.	N/A	N/A
2014/0243652	12/2013	Pashko	N/A	N/A
2014/0243663	12/2013	Taylor	N/A	N/A
2014/0243694	12/2013	Baker et al.	N/A	N/A
2014/0243714	12/2013	Ward et al.	N/A	N/A
2014/0243926	12/2013	Carcieri	N/A	N/A
2014/0243934	12/2013	Vo-Dinh et al.	N/A	N/A
2014/0245191	12/2013	Serena	N/A	N/A
2014/0247970	12/2013	Taylor	N/A	N/A
2014/0249360	12/2013	Jaeger et al.	N/A	N/A
2014/0249396	12/2013	Shacham-Diamand et al.	N/A	N/A
2014/0249429	12/2013	Tran	N/A	N/A
2014/0249445	12/2013	Deadwyler et al.	N/A	N/A
2014/0249447	12/2013	Sereno et al.	N/A	N/A
2014/0249454	12/2013	Carpentier	N/A	N/A
2014/0249608	12/2013	Rogers	N/A	N/A
2014/0249791	12/2013	Taylor	N/A	N/A
2014/0249792	12/2013	Taylor	N/A	N/A
2014/0257047	12/2013	Sillay et al.	N/A	N/A
2014/0257073	12/2013	Machon et al.	N/A	N/A
2014/0257118	12/2013	DiLorenzo et al.	N/A	N/A
2014/0257128	12/2013	Moxon et al.	N/A	N/A
2014/0257132	12/2013	Kilgard et al.	N/A	N/A

2014/0257147	12/2013	John et al.	N/A	N/A
2014/0257430	12/2013	Kilgard et al.	N/A	N/A
2014/0257437	12/2013	Simon et al.	N/A	N/A
2014/0257438	12/2013	Simon et al.	N/A	N/A
2014/0266696	12/2013	Addison et al.	N/A	N/A
2014/0266787	12/2013	Tran	N/A	N/A
2014/0270438	12/2013	Declerck et al.	N/A	N/A
2014/0271483	12/2013	Satchi-Fainaro et al.	N/A	N/A
2014/0275716	12/2013	Connor	N/A	N/A
2014/0275741	12/2013	Vandenbelt et al.	N/A	N/A
2014/0275807	12/2013	Redei	N/A	N/A
2014/0275847	12/2013	Perryman et al.	N/A	N/A
2014/0275851	12/2013	Amble et al.	N/A	N/A
2014/0275886	12/2013	Teixeira	N/A	N/A
2014/0275889	12/2013	Addison et al.	N/A	N/A
2014/0275891	12/2013	Muehlemann et al.	N/A	N/A
2014/0275944	12/2013	Semenov	N/A	N/A
2014/0276012	12/2013	Semenov	N/A	N/A
2014/0276013	12/2013	Muehlemann et al.	N/A	N/A
2014/0276014	12/2013	Khanicheh et al.	N/A	N/A
2014/0276090	12/2013	Breed	N/A	N/A
2014/0276123	12/2013	Yang	N/A	N/A
2014/0276130	12/2013	Mirelman et al.	N/A	N/A
2014/0276181	12/2013	Sun et al.	N/A	N/A
2014/0276183	12/2013	Badower	N/A	N/A
2014/0276185	12/2013	Carlson et al.	N/A	N/A
2014/0276187	12/2013	Iasemidis et al.	N/A	N/A
2014/0276194	12/2013	Osorio	N/A	N/A
2014/0276549	12/2013	Osorio	N/A	N/A
2014/0276702	12/2013	McKay et al.	N/A	N/A
2014/0276944	12/2013	Farritor et al.	N/A	N/A
2014/0277255	12/2013	Sabesan	N/A	N/A
2014/0277256	12/2013	Osorio	N/A	N/A
2014/0277282	12/2013	Jaax	N/A	N/A
2014/0277286	12/2013	Cinbis	N/A	N/A
2014/0277582	12/2013	Leuthardt et al.	N/A	N/A
2014/0279341	12/2013	Bhardwaj et al.	N/A	N/A
2014/0279746	12/2013	De Bruin et al.	N/A	N/A
2014/0288381	12/2013	Faarbaek et al.	N/A	N/A
2014/0288614	12/2013	Hagedorn et al.	N/A	N/A
2014/0288620	12/2013	DiLorenzo	N/A	N/A
2014/0288953	12/2013	Lynn et al.	N/A	N/A
2014/0289172	12/2013	Rothman et al.	N/A	N/A
2014/0296646	12/2013	Wingeier et al.	N/A	N/A
2014/0296655	12/2013	Akhbardeh et al.	N/A	N/A
2014/0296724	12/2013	Guttag et al.	N/A	N/A
2014/0296733	12/2013	Omurtag et al.	N/A	N/A
2014/0296750	12/2013	Einav et al.	N/A	N/A
2014/0297397	12/2013	Bakalash et al.	N/A	N/A
2014/0300532	12/2013	Karkkainen et al.	N/A	N/A
2014/0303424	12/2013	Glass	N/A	N/A
2014/0303425	12/2013	Pilla et al.	N/A	N/A
2014/0303452	12/2013	Ghaffari	N/A	N/A
2014/0303453	12/2013	Seely et al.	N/A	N/A
2014/0303454	12/2013	Clifton et al.	N/A	N/A
2014/0303486	12/2013	Baumgartner et al.	N/A	N/A
2014/0303508	12/2013	Plotnik-Peleg et al.	N/A	N/A
2014/0303511	12/2013	Sajda et al.	N/A	N/A
2014/0304773	12/2013	Woods et al.	N/A	N/A
2014/0309484	12/2013	Chang	N/A	N/A
2014/0309614	12/2013	Frei et al.	N/A	N/A
2014/0309881	12/2013	Fung et al.	N/A	N/A
2014/0309943	12/2013	Grundlehner et al.	N/A	N/A
2014/0313303	12/2013	Davis et al.	N/A	N/A
2014/0315169	12/2013	Bohbot	N/A	N/A
2014/0316191	12/2013	de Zambotti et al.	N/A	N/A
2014/0316192	12/2013	de Zambotti et al.	N/A	N/A
2014/0316217	12/2013	Purdon et al.	N/A	N/A
2014/0316221	12/2013	Rothman	N/A	N/A
2014/0316230	12/2013	Denison et al.	N/A	N/A
2014/0316235	12/2013	Davis et al.	N/A	N/A
2014/0316243	12/2013	Niedermeyer	N/A	N/A
2014/0316248	12/2013	deCharms	N/A	N/A
2014/0316278	12/2013	Addison et al.	N/A	N/A
2014/0323849	12/2013	Deisseroth et al.	N/A	N/A

2014/0323899	12/2013	Silberstein	N/A	N/A
2014/0323900	12/2013	Bibian et al.	N/A	N/A
2014/0323924	12/2013	Mishelevich	N/A	N/A
2014/0323946	12/2013	Bourke, Jr. et al.	N/A	N/A
2014/0324118	12/2013	Simon et al.	N/A	N/A
2014/0324138	12/2013	Wentz et al.	N/A	N/A
2014/0328487	12/2013	Hiroe	N/A	N/A
2014/0330093	12/2013	Pedro	N/A	N/A
2014/0330102	12/2013	Zbrzeski et al.	N/A	N/A
2014/0330157	12/2013	Snook	N/A	N/A
2014/0330159	12/2013	Costa et al.	N/A	N/A
2014/0330268	12/2013	Palti et al.	N/A	N/A
2014/0330334	12/2013	Errico et al.	N/A	N/A
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2014/0330336	12/2013	Errico et al.	N/A	N/A
2014/0330337	12/2013	Linke et al.	N/A	N/A
2014/0330345	12/2013	John	N/A	N/A
2014/0330357	12/2013	Stevenson et al.	N/A	N/A
2014/0330394	12/2013	Leuthardt et al.	N/A	N/A
2014/0330404	12/2013	Abdelghani et al.	N/A	N/A
2014/0330580	12/2013	Grima et al.	N/A	N/A
2014/0335489	12/2013	DeCharms	N/A	N/A
2014/0336473	12/2013	Greco	N/A	N/A
2014/0336489	12/2013	Angotzi et al.	N/A	N/A
2014/0336514	12/2013	Peyman	N/A	N/A
2014/0336547	12/2013	Tass et al.	N/A	N/A
2014/0336730	12/2013	Simon et al.	N/A	N/A
2014/0340084	12/2013	Alon	N/A	N/A
2014/0343397	12/2013	Kim et al.	N/A	N/A
2014/0343399	12/2013	Posse	N/A	N/A
2014/0343408	12/2013	Tolkowsky	N/A	N/A
2014/0343463	12/2013	Mishelevich	N/A	N/A
2014/0343882	12/2013	Taulu et al.	N/A	N/A
2014/0347265	12/2013	Aimone et al.	N/A	N/A
2014/0347491	12/2013	Connor	N/A	N/A
2014/0348183	12/2013	Kim et al.	N/A	N/A
2014/0348412	12/2013	Taylor	N/A	N/A
2014/0350353	12/2013	Connor	N/A	N/A
2014/0350369	12/2013	Budiman et al.	N/A	N/A
2014/0350380	12/2013	Eidelberg	N/A	N/A
2014/0350431	12/2013	Hagedorn	N/A	N/A
2014/0350436	12/2013	Nathan et al.	N/A	N/A
2014/0350634	12/2013	Grill et al.	N/A	N/A
2014/0350636	12/2013	King et al.	N/A	N/A
2014/0350864	12/2013	Fang et al.	N/A	N/A
2014/0354278	12/2013	Subbarao	N/A	N/A
2014/0355859	12/2013	Taylor et al.	N/A	N/A
2014/0357507	12/2013	Umansky et al.	N/A	N/A
2014/0357932	12/2013	Lozano	N/A	N/A
2014/0357935	12/2013	Ilmoniemä et al.	N/A	N/A
2014/0357936	12/2013	Simon et al.	N/A	N/A
2014/0357962	12/2013	Harrington et al.	N/A	N/A
2014/0358024	12/2013	Nelson et al.	N/A	N/A
2014/0358025	12/2013	Parhi et al.	N/A	N/A
2014/0358067	12/2013	Deisseroth et al.	N/A	N/A
2014/0358193	12/2013	Lyons et al.	N/A	N/A
2014/0358199	12/2013	Lim	N/A	N/A
2014/0364721	12/2013	Lee et al.	N/A	N/A
2014/0364746	12/2013	Addison et al.	N/A	N/A
2014/0369537	12/2013	Pontoppidan et al.	N/A	N/A
2014/0370479	12/2013	Gazzaley	N/A	N/A
2014/0371515	12/2013	John	N/A	N/A
2014/0371516	12/2013	Tsai et al.	N/A	N/A
2014/0371544	12/2013	Wu et al.	N/A	N/A
2014/0371573	12/2013	Komoto et al.	N/A	N/A
2014/0371599	12/2013	Wu et al.	N/A	N/A
2014/0371611	12/2013	Kim	N/A	N/A
2014/0371984	12/2013	Fung et al.	N/A	N/A
2014/0378809	12/2013	Weitnauer et al.	N/A	N/A
2014/0378810	12/2013	Davis et al.	N/A	N/A
2014/0378815	12/2013	Huang et al.	N/A	N/A
2014/0378830	12/2013	Li	N/A	N/A
2014/0378851	12/2013	Frei et al.	N/A	N/A
2014/0378941	12/2013	Su et al.	N/A	N/A
2014/0379620	12/2013	Sarrafzadeh et al.	N/A	N/A

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2015/0003698	12/2014	Davis et al.	N/A	N/A
2015/0003699	12/2014	Davis et al.	N/A	N/A
2015/0005592	12/2014	Osorio	N/A	N/A
2015/0005594	12/2014	Chamoun et al.	N/A	N/A
2015/0005640	12/2014	Davis et al.	N/A	N/A
2015/0005644	12/2014	Rhoads	N/A	N/A
2015/0005646	12/2014	Balakrishnan et al.	N/A	N/A
2015/0005660	12/2014	Kraus et al.	N/A	N/A
2015/0005680	12/2014	Lipani	N/A	N/A
2015/0005839	12/2014	Sabesan et al.	N/A	N/A
2015/0005840	12/2014	Pal et al.	N/A	N/A
2015/0005841	12/2014	Pal et al.	N/A	N/A
2015/0006186	12/2014	Davis et al.	N/A	N/A
2015/0008916	12/2014	Le Prado et al.	N/A	N/A
2015/0010223	12/2014	Sapiro et al.	N/A	N/A
2015/0011866	12/2014	Baumgartner	N/A	N/A
2015/0011877	12/2014	Baumgartner	N/A	N/A
2015/0011907	12/2014	Purdon et al.	N/A	N/A
2015/0012054	12/2014	Kilgard et al.	N/A	N/A
2015/0012057	12/2014	Carlson et al.	N/A	N/A
2015/0012111	12/2014	Contreras-Vidal et al.	N/A	N/A
2015/0012466	12/2014	Sapiro et al.	N/A	N/A
2015/0016618	12/2014	Adachi et al.	N/A	N/A
2015/0017115	12/2014	Satchi-Fainaro et al.	N/A	N/A
2015/0018665	12/2014	Jasanoff et al.	N/A	N/A
2015/0018699	12/2014	Zeng et al.	N/A	N/A
2015/0018702	12/2014	Galloway et al.	N/A	N/A
2015/0018705	12/2014	Barlow et al.	N/A	N/A
2015/0018706	12/2014	Segal	N/A	N/A
2015/0018758	12/2014	John	N/A	N/A
2015/0018893	12/2014	Kilgard et al.	N/A	N/A
2015/0018905	12/2014	Nofzinger et al.	N/A	N/A
2015/0019241	12/2014	Bennett et al.	N/A	N/A
2015/0019266	12/2014	Stempora	N/A	N/A
2015/0024356	12/2014	Hillyer et al.	N/A	N/A
2015/0025351	12/2014	Govari	N/A	N/A
2015/0025408	12/2014	Wingeier et al.	N/A	N/A
2015/0025410	12/2014	Wolpaw et al.	N/A	N/A
2015/0025421	12/2014	Wagner et al.	N/A	N/A
2015/0025422	12/2014	Tyler	N/A	N/A
2015/0025610	12/2014	Wingeier et al.	N/A	N/A
2015/0025917	12/2014	Stempora	N/A	N/A
2015/0026446	12/2014	Kim et al.	N/A	N/A
2015/0029087	12/2014	Klappert et al.	N/A	N/A
2015/0030220	12/2014	Cho et al.	N/A	N/A
2015/0032017	12/2014	Babaeizadeh et al.	N/A	N/A
2015/0032044	12/2014	Peyman	N/A	N/A
2015/0032178	12/2014	Simon et al.	N/A	N/A
2015/0033245	12/2014	Klappert et al.	N/A	N/A
2015/0033258	12/2014	Klappert et al.	N/A	N/A
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2015/0033262	12/2014	Klappert et al.	N/A	N/A
2015/0033266	12/2014	Klappert et al.	N/A	N/A
2015/0033363	12/2014	Pinsky et al.	N/A	N/A
2015/0035959	12/2014	Amble et al.	N/A	N/A
2015/0038804	12/2014	Younes	N/A	N/A
2015/0038812	12/2014	Ayaz et al.	N/A	N/A
2015/0038822	12/2014	Wingeier et al.	N/A	N/A
2015/0038869	12/2014	Simon et al.	N/A	N/A
2015/0039066	12/2014	Wingeier et al.	N/A	N/A
2015/0039110	12/2014	Abeyratne et al.	N/A	N/A
2015/0042477	12/2014	Kobetski et al.	N/A	N/A
2015/0044138	12/2014	Lansbergen et al.	N/A	N/A
2015/0045606	12/2014	Hagedorn et al.	N/A	N/A
2015/0045607	12/2014	Hakansson	N/A	N/A
2015/0045686	12/2014	Lynn	N/A	N/A
2015/0051655	12/2014	Kilgard et al.	N/A	N/A
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2015/0051658	12/2014	Kilgard et al.	N/A	N/A
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2015/0051663	12/2014	Hagedorn	N/A	N/A
2015/0051668	12/2014	Bahmer	N/A	N/A
2015/0057512	12/2014	Kapoor	N/A	N/A

2015/0057715	12/2014	Kilgard et al.	N/A	N/A
2015/0065803	12/2014	Douglas et al.	N/A	N/A
2015/0065831	12/2014	Popovic et al.	N/A	N/A
2015/0065838	12/2014	Wingeier et al.	N/A	N/A
2015/0065839	12/2014	Farah et al.	N/A	N/A
2015/0065845	12/2014	Takiguchi	N/A	N/A
2015/0066124	12/2014	Stevenson et al.	N/A	N/A
2015/0068069	12/2014	Tran et al.	N/A	N/A
2015/0069846	12/2014	Hokari	N/A	N/A
2015/0071907	12/2014	Crombez et al.	N/A	N/A
2015/0072394	12/2014	Deisseroth et al.	N/A	N/A
2015/0073141	12/2014	Teegarden et al.	N/A	N/A
2015/0073237	12/2014	Osorio	N/A	N/A
2015/0073249	12/2014	Musha	N/A	N/A
2015/0073294	12/2014	Zhang et al.	N/A	N/A
2015/0073306	12/2014	Abeyratne et al.	N/A	N/A
2015/0073505	12/2014	Errico et al.	N/A	N/A
2015/0073722	12/2014	Taylor et al.	N/A	N/A
2015/0080327	12/2014	Paul et al.	N/A	N/A
2015/0080671	12/2014	Christensen et al.	N/A	N/A
2015/0080674	12/2014	Drew et al.	N/A	N/A
2015/0080695	12/2014	Rogers et al.	N/A	N/A
2015/0080703	12/2014	Reiman	N/A	N/A
2015/0080746	12/2014	Bleich et al.	N/A	N/A
2015/0080753	12/2014	Miyazaki et al.	N/A	N/A
2015/0080985	12/2014	Yun et al.	N/A	N/A
2015/0081226	12/2014	Baki	N/A	N/A
2015/0081299	12/2014	Jasinschi et al.	N/A	N/A
2015/0087931	12/2014	Banerjee et al.	N/A	N/A
2015/0088015	12/2014	Taylor	N/A	N/A
2015/0088024	12/2014	Sackellares et al.	N/A	N/A
2015/0088093	12/2014	Goetz	N/A	N/A
2015/0088120	12/2014	Garcia et al.	N/A	N/A
2015/0088224	12/2014	Goldwasser et al.	N/A	N/A
2015/0088228	12/2014	Moffitt	N/A	N/A
2015/0088478	12/2014	Taylor	N/A	N/A
2015/0091730	12/2014	Kangas et al.	N/A	N/A
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2015/0092949	12/2014	Adachi et al.	N/A	N/A
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2015/0099941	12/2014	Tran	N/A	N/A
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2015/0099959	12/2014	Bonmassar et al.	N/A	N/A
2015/0099962	12/2014	Weiss et al.	N/A	N/A
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2015/0105641	12/2014	Austin et al.	N/A	N/A
2015/0105701	12/2014	Mayer et al.	N/A	N/A
2015/0105837	12/2014	Aguilar Domingo	N/A	N/A
2015/0105844	12/2014	Tass et al.	N/A	N/A
2015/0112222	12/2014	Sun et al.	N/A	N/A
2015/0112403	12/2014	Ruffini et al.	N/A	N/A
2015/0112409	12/2014	Hagedorn	N/A	N/A
2015/0112899	12/2014	Dagum	N/A	N/A
2015/0119652	12/2014	Hyde et al.	N/A	N/A
2015/0119658	12/2014	Osorio	N/A	N/A
2015/0119689	12/2014	Pascual-Leone et al.	N/A	N/A
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2015/0119743	12/2014	Maksym et al.	N/A	N/A
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2015/0119794	12/2014	Peyman	N/A	N/A
2015/0119898	12/2014	Desalles et al.	N/A	N/A
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2015/0133830	12/2014	Dirks et al.	N/A	N/A
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2015/0134264	12/2014	Tansey	N/A	N/A
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2015/0141529	12/2014	Hargrove	N/A	N/A
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2015/0150475	12/2014	Varcoe	N/A	N/A
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2015/0150753	12/2014	Racette	N/A	N/A
2015/0151142	12/2014	Tyler et al.	N/A	N/A
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2015/0182417	12/2014	Nagatani	N/A	N/A
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2015/0190094	12/2014	Lee et al.	N/A	N/A
2015/0190636	12/2014	Simon et al.	N/A	N/A
2015/0190637	12/2014	Simon et al.	N/A	N/A
2015/0192532	12/2014	Clevenson et al.	N/A	N/A
2015/0192776	12/2014	Lee et al.	N/A	N/A
2015/0196213	12/2014	Pandia et al.	N/A	N/A
2015/0196246	12/2014	Osorio	N/A	N/A
2015/0196249	12/2014	Brown et al.	N/A	N/A
2015/0196800	12/2014	Macri et al.	N/A	N/A
2015/0199010	12/2014	Coleman	345/156	H04L 67/01
2015/0199121	12/2014	Gulaka et al.	N/A	N/A
2015/0200046	12/2014	Park et al.	N/A	N/A
2015/0201849	12/2014	Taylor	N/A	N/A

2015/0201879	12/2014	Hargrove	N/A	N/A
2015/0202330	12/2014	Yang et al.	N/A	N/A
2015/0202428	12/2014	Miller	N/A	N/A
2015/0202447	12/2014	Afshar et al.	N/A	N/A
2015/0203822	12/2014	Tremolada et al.	N/A	N/A
2015/0206051	12/2014	McIntosh et al.	N/A	N/A
2015/0206174	12/2014	Barnett et al.	N/A	N/A
2015/0208940	12/2014	Addison et al.	N/A	N/A
2015/0208975	12/2014	Ghajar	N/A	N/A
2015/0208978	12/2014	Osorio et al.	N/A	N/A
2015/0208982	12/2014	Ho et al.	N/A	N/A
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2015/0223721	12/2014	De Ridder	N/A	N/A
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2015/0231405	12/2014	Okada	N/A	N/A
2015/0231408	12/2014	Williams et al.	N/A	N/A
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2015/0238112	12/2014	Park et al.	N/A	N/A
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2015/0242608	12/2014	Kim et al.	N/A	N/A
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2015/0250393	12/2014	Tran	N/A	N/A
2015/0250401	12/2014	Tveit	N/A	N/A
2015/0250415	12/2014	Leininger et al.	N/A	N/A
2015/0251016	12/2014	Vo-Dinh et al.	N/A	N/A
2015/0253391	12/2014	Toda et al.	N/A	N/A
2015/0253410	12/2014	Warfield et al.	N/A	N/A
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2015/0257645	12/2014	Bae et al.	N/A	N/A
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2015/0262016	12/2014	Rothblatt	N/A	N/A
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2015/0272448	12/2014	Fonte et al.	N/A	N/A
2015/0272461	12/2014	Morimoto et al.	N/A	N/A
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2015/0272510	12/2014	Chin	N/A	N/A
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2015/0289813	12/2014	Lipov	N/A	N/A
2015/0289929	12/2014	Toth et al.	N/A	N/A
2015/0290419	12/2014	Kare et al.	N/A	N/A
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2015/0304048	12/2014	Kim et al.	N/A	N/A

2015/0304101	12/2014	Gupta et al.	N/A	N/A
2015/0305685	12/2014	Shahaf et al.	N/A	N/A
2015/0305686	12/2014	Coleman et al.	N/A	N/A
2015/0305689	12/2014	Gourmelon et al.	N/A	N/A
2015/0305799	12/2014	Trieu	N/A	N/A
2015/0305800	12/2014	Trieu	N/A	N/A
2015/0305801	12/2014	Trieu	N/A	N/A
2015/0306057	12/2014	Goodenowe	N/A	N/A
2015/0306340	12/2014	Giap et al.	N/A	N/A
2015/0306390	12/2014	Zalay et al.	N/A	N/A
2015/0306391	12/2014	Wu et al.	N/A	N/A
2015/0306392	12/2014	Sabesan	N/A	N/A
2015/0309563	12/2014	Connor	N/A	N/A
2015/0309582	12/2014	Gupta	N/A	N/A
2015/0310862	12/2014	Dauphin et al.	N/A	N/A
2015/0313496	12/2014	Connor	N/A	N/A
2015/0313498	12/2014	Coleman et al.	N/A	N/A
2015/0313535	12/2014	Alshaer et al.	N/A	N/A
2015/0313539	12/2014	Connor	N/A	N/A
2015/0313540	12/2014	Deuchar et al.	N/A	N/A
2015/0313949	12/2014	Cutillo	N/A	N/A
2015/0313971	12/2014	Haslett et al.	N/A	N/A
2015/0315554	12/2014	Shekdar et al.	N/A	N/A
2015/0317447	12/2014	Helleputte et al.	N/A	N/A
2015/0317796	12/2014	Schett et al.	N/A	N/A
2015/0320591	12/2014	Smith et al.	N/A	N/A
2015/0321000	12/2014	Rosenbluth et al.	N/A	N/A
2015/0324544	12/2014	Maslowski et al.	N/A	N/A
2015/0324545	12/2014	Fonte	N/A	N/A
2015/0324692	12/2014	Ritchey et al.	N/A	N/A
2015/0325151	12/2014	Tuchschrnid et al.	N/A	N/A
2015/0327813	12/2014	Fu	N/A	N/A
2015/0327837	12/2014	Qi et al.	N/A	N/A
2015/0328330	12/2014	Satchi-Fainaro et al.	N/A	N/A
2015/0328455	12/2014	Meadows et al.	N/A	N/A
2015/0331929	12/2014	El-Saban et al.	N/A	N/A
2015/0332015	12/2014	Taylor	N/A	N/A
2015/0335281	12/2014	Scroggins	N/A	N/A
2015/0335288	12/2014	Toth et al.	N/A	N/A
2015/0335292	12/2014	Mittal	N/A	N/A
2015/0335294	12/2014	Witcher et al.	N/A	N/A
2015/0335295	12/2014	Park et al.	N/A	N/A
2015/0335303	12/2014	Chandelier et al.	N/A	N/A
2015/0335876	12/2014	Jeffery et al.	N/A	N/A
2015/0335877	12/2014	Jeffery et al.	N/A	N/A
2015/0338915	12/2014	Publicover et al.	N/A	N/A
2015/0339363	12/2014	Moldoveanu et al.	N/A	N/A
2015/0339459	12/2014	Taylor	N/A	N/A
2015/0342472	12/2014	Semenov	N/A	N/A
2015/0342478	12/2014	Galen et al.	N/A	N/A
2015/0342493	12/2014	Hardt	N/A	N/A
2015/0343215	12/2014	De Ridder	N/A	N/A
2015/0343222	12/2014	Kilgard et al.	N/A	N/A
2015/0343242	12/2014	Tyler et al.	N/A	N/A
2015/0351655	12/2014	Coleman	N/A	N/A
2015/0351690	12/2014	Toth et al.	N/A	N/A
2015/0351701	12/2014	Moxon et al.	N/A	N/A
2015/0352362	12/2014	Craig	N/A	N/A
2015/0352363	12/2014	McIntyre et al.	N/A	N/A
2015/0359431	12/2014	Bakalash et al.	N/A	N/A
2015/0359441	12/2014	Giovangrandi et al.	N/A	N/A
2015/0359450	12/2014	Lee et al.	N/A	N/A
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2015/0359467	12/2014	Tran	N/A	N/A
2015/0359486	12/2014	Kovacs et al.	N/A	N/A
2015/0359492	12/2014	Giovangrandi et al.	N/A	N/A
2015/0360026	12/2014	Wagner	N/A	N/A
2015/0360030	12/2014	Cartledge et al.	N/A	N/A
2015/0360039	12/2014	Lempka et al.	N/A	N/A
2015/0363941	12/2014	Taylor	N/A	N/A
2015/0366482	12/2014	Lee	N/A	N/A
2015/0366497	12/2014	Cavuoto et al.	N/A	N/A
2015/0366503	12/2014	Sjaaheim et al.	N/A	N/A
2015/0366504	12/2014	Connor	N/A	N/A
2015/0366516	12/2014	Dripps et al.	N/A	N/A

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2015/0366656	12/2014	Wortz et al.	N/A	N/A
2015/0366659	12/2014	Wortz et al.	N/A	N/A
2015/0369864	12/2014	Marlow et al.	N/A	N/A
2015/0370320	12/2014	Connor	N/A	N/A
2015/0370325	12/2014	Jarosiewicz et al.	N/A	N/A
2015/0374250	12/2014	Hatano et al.	N/A	N/A
2015/0374285	12/2014	Chang et al.	N/A	N/A
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2015/0374300	12/2014	Najarian et al.	N/A	N/A
2015/0374973	12/2014	Morrell	N/A	N/A
2015/0374983	12/2014	Simon et al.	N/A	N/A
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2015/0375006	12/2014	Denison et al.	N/A	N/A
2015/0379230	12/2014	Taylor	N/A	N/A
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2015/0379878	12/2014	Walter et al.	N/A	N/A
2015/0380009	12/2014	Chang et al.	N/A	N/A
2016/0000348	12/2015	Kitajo et al.	N/A	N/A
2016/0000354	12/2015	Hagedorn et al.	N/A	N/A
2016/0000383	12/2015	Lee et al.	N/A	N/A
2016/0001065	12/2015	Wingeier et al.	N/A	N/A
2016/0001096	12/2015	Mishelevich	N/A	N/A
2016/0001098	12/2015	Wingeier et al.	N/A	N/A
2016/0002523	12/2015	Huh et al.	N/A	N/A
2016/0004298	12/2015	Mazed et al.	N/A	N/A
2016/0004396	12/2015	Gulaka et al.	N/A	N/A
2016/0004821	12/2015	Fueyo et al.	N/A	N/A
2016/0004957	12/2015	Solari	N/A	N/A
2016/0005235	12/2015	Fateh	N/A	N/A
2016/0005320	12/2015	deCharms et al.	N/A	N/A
2016/0007899	12/2015	Durkee et al.	N/A	N/A
2016/0007904	12/2015	Vardy	N/A	N/A
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2016/0008489	12/2015	Korzus	N/A	N/A
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2016/0008620	12/2015	Stubbeman	N/A	N/A
2016/0008632	12/2015	Wetmore et al.	N/A	N/A
2016/0012011	12/2015	Llinas et al.	N/A	N/A
2016/0012583	12/2015	Cales et al.	N/A	N/A
2016/0012749	12/2015	Connor	N/A	N/A
2016/0015281	12/2015	McKenna et al.	N/A	N/A
2016/0015289	12/2015	Simon et al.	N/A	N/A
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2016/0015673	12/2015	Goodenowe	N/A	N/A
2016/0016014	12/2015	Wagner et al.	N/A	N/A
2016/0019434	12/2015	Caldwell	N/A	N/A
2016/0019693	12/2015	Silbersweig et al.	N/A	N/A
2016/0022141	12/2015	Mittal et al.	N/A	N/A
2016/0022156	12/2015	Kovacs et al.	N/A	N/A
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2016/0023016	12/2015	Bonmassar et al.	N/A	N/A
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2016/0027178	12/2015	Yu et al.	N/A	N/A
2016/0027293	12/2015	Esteller et al.	N/A	N/A
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2016/0029896	12/2015	Lee et al.	N/A	N/A
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2016/0029946	12/2015	Simon et al.	N/A	N/A
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2016/0029958	12/2015	Le et al.	N/A	N/A

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2016/0030666	12/2015	Lozano et al.	N/A	N/A
2016/0030702	12/2015	Yang	N/A	N/A
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2016/0031479	12/2015	Fung et al.	N/A	N/A
2016/0035093	12/2015	Kateb et al.	N/A	N/A
2016/0038037	12/2015	Kovacs	N/A	N/A
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2016/0038042	12/2015	Mulligan et al.	N/A	N/A
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2016/0038559	12/2015	Palmer et al.	N/A	N/A
2016/0038770	12/2015	Tyler et al.	N/A	N/A
2016/0040514	12/2015	Rahmani et al.	N/A	N/A
2016/0044841	12/2015	Chamberlain	N/A	N/A
2016/0045128	12/2015	Sitt et al.	N/A	N/A
2016/0045150	12/2015	Leininger et al.	N/A	N/A
2016/0045162	12/2015	De Graff et al.	N/A	N/A
2016/0045731	12/2015	Simon et al.	N/A	N/A
2016/0045756	12/2015	Phillips et al.	N/A	N/A
2016/0048659	12/2015	Pereira et al.	N/A	N/A
2016/0048948	12/2015	Bajic	N/A	N/A
2016/0048965	12/2015	Stehle et al.	N/A	N/A
2016/0051161	12/2015	Labyt et al.	N/A	N/A
2016/0051162	12/2015	Durand et al.	N/A	N/A
2016/0051187	12/2015	Damadian	N/A	N/A
2016/0051195	12/2015	Pang et al.	N/A	N/A
2016/0051793	12/2015	Gibson-Horn	N/A	N/A
2016/0051812	12/2015	Montgomery, Jr. et al.	N/A	N/A
2016/0051818	12/2015	Simon et al.	N/A	N/A
2016/0055236	12/2015	Frank et al.	N/A	N/A
2016/0055304	12/2015	Russell et al.	N/A	N/A
2016/0055415	12/2015	Baxi	N/A	N/A
2016/0055842	12/2015	DeFranks et al.	N/A	N/A
2016/0058301	12/2015	Shusterman	N/A	N/A
2016/0058304	12/2015	Emblem et al.	N/A	N/A
2016/0058322	12/2015	Brister et al.	N/A	N/A
2016/0058354	12/2015	Phan et al.	N/A	N/A
2016/0058359	12/2015	Osorio	N/A	N/A
2016/0058366	12/2015	Choi et al.	N/A	N/A
2016/0058376	12/2015	Baek et al.	N/A	N/A
2016/0058392	12/2015	Hasson et al.	N/A	N/A
2016/0058673	12/2015	Francis	N/A	N/A
2016/0060926	12/2015	Kim et al.	N/A	N/A
2016/0062459	12/2015	Publicover et al.	N/A	N/A
2016/0063207	12/2015	Schmidt	N/A	N/A
2016/0063883	12/2015	Jeyanandarajan	N/A	N/A
2016/0065724	12/2015	Lee et al.	N/A	N/A
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2016/0066788	12/2015	Tran et al.	N/A	N/A
2016/0066789	12/2015	Rogers et al.	N/A	N/A
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2016/0066838	12/2015	DeCharms	N/A	N/A
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2016/0067492	12/2015	Wolpaw et al.	N/A	N/A
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2016/0073886	12/2015	Connor	N/A	N/A
2016/0073916	12/2015	Aksenova et al.	N/A	N/A
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2016/0073991	12/2015	Taylor	N/A	N/A
2016/0074657	12/2015	Kwan et al.	N/A	N/A
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2016/0074661	12/2015	Lipani	N/A	N/A
2016/0077547	12/2015	Aimone et al.	N/A	N/A
2016/0078780	12/2015	Alexander et al.	N/A	N/A
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2016/0081610	12/2015	Osorio et al.	N/A	N/A
2016/0081613	12/2015	Braun et al.	N/A	N/A
2016/0081616	12/2015	Li	N/A	N/A
2016/0081625	12/2015	Kim et al.	N/A	N/A
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2016/0082180	12/2015	Toth et al.	N/A	N/A
2016/0082319	12/2015	Macri et al.	N/A	N/A
2016/0084925	12/2015	Le Prado et al.	N/A	N/A
2016/0085302	12/2015	Publicover et al.	N/A	N/A
2016/0086622	12/2015	Yamamoto	N/A	N/A
2016/0087603	12/2015	Ricci et al.	N/A	N/A
2016/0089031	12/2015	Hu	N/A	N/A
2016/0091448	12/2015	Soleimani	N/A	N/A
2016/0095546	12/2015	Sahasrabudhe et al.	N/A	N/A
2016/0095838	12/2015	Satchi-Fainaro et al.	N/A	N/A
2016/0096025	12/2015	Moffitt et al.	N/A	N/A
2016/0097824	12/2015	Fujii et al.	N/A	N/A
2016/0100769	12/2015	Kim et al.	N/A	N/A
2016/0101260	12/2015	Austin et al.	N/A	N/A
2016/0102500	12/2015	Donderici et al.	N/A	N/A
2016/0103487	12/2015	Crawford et al.	N/A	N/A
2016/0103963	12/2015	Mishra	N/A	N/A
2016/0104006	12/2015	Son et al.	N/A	N/A
2016/0106331	12/2015	Zorick et al.	N/A	N/A
2016/0106344	12/2015	Nazari	N/A	N/A
2016/0106513	12/2015	De Stavola et al.	N/A	N/A
2016/0106950	12/2015	Vasapollo	N/A	N/A
2016/0106997	12/2015	Arendash et al.	N/A	N/A
2016/0107309	12/2015	Walsh et al.	N/A	N/A
2016/0107653	12/2015	Fung et al.	N/A	N/A
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2016/0109959	12/2015	Heo	N/A	N/A
2016/0110517	12/2015	Taylor	N/A	N/A
2016/0110866	12/2015	Taylor	N/A	N/A
2016/0110867	12/2015	Taylor	N/A	N/A
2016/0112022	12/2015	Butts	N/A	N/A
2016/0112684	12/2015	Connor	N/A	N/A
2016/0113517	12/2015	Lee et al.	N/A	N/A
2016/0113528	12/2015	Taylor	N/A	N/A
2016/0113539	12/2015	Sinharay et al.	N/A	N/A
2016/0113545	12/2015	Kim et al.	N/A	N/A
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2016/0114165	12/2015	Levine et al.	N/A	N/A
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2016/0116553	12/2015	Kim et al.	N/A	N/A
2016/0117815	12/2015	Taylor	N/A	N/A
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2016/0117819	12/2015	Taylor	N/A	N/A
2016/0119726	12/2015	Pontoppidan et al.	N/A	N/A
2016/0120048	12/2015	Seo et al.	N/A	N/A
2016/0120428	12/2015	Yoshida et al.	N/A	N/A
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2016/0120433	12/2015	Hughes et al.	N/A	N/A
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2016/0121114	12/2015	Simon et al.	N/A	N/A
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2016/0125228	12/2015	Son et al.	N/A	N/A
2016/0125572	12/2015	Yoo et al.	N/A	N/A
2016/0128589	12/2015	Tabib-Azar	N/A	N/A
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2016/0133015	12/2015	Taylor	N/A	N/A
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2016/0136443	12/2015	Grandhe et al.	N/A	N/A
2016/0139215	12/2015	Fujii	N/A	N/A
2016/0140306	12/2015	Hua et al.	N/A	N/A
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2016/0143582	12/2015	Connor	N/A	N/A
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2016/0144175	12/2015	Simon et al.	N/A	N/A
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2016/0152233	12/2015	Fung et al.	N/A	N/A
2016/0155005	12/2015	Varkuti et al.	N/A	N/A
2016/0157742	12/2015	Huang et al.	N/A	N/A
2016/0157773	12/2015	Baek et al.	N/A	N/A
2016/0157777	12/2015	Attal et al.	N/A	N/A
2016/0157828	12/2015	Sumi et al.	N/A	N/A
2016/0158553	12/2015	Panken et al.	N/A	N/A
2016/0158554	12/2015	Graig	N/A	N/A
2016/0162652	12/2015	Siekmeier	N/A	N/A
2016/0164813	12/2015	Anderson et al.	N/A	N/A
2016/0165852	12/2015	Goldfain	N/A	N/A
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2016/0166169	12/2015	Badower et al.	N/A	N/A
2016/0166197	12/2015	Venkatraman et al.	N/A	N/A
2016/0166199	12/2015	Sun et al.	N/A	N/A
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2016/0166208	12/2015	Girouard et al.	N/A	N/A
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2016/0167672	12/2015	Krueger	N/A	N/A
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2016/0170996	12/2015	Frank et al.	N/A	N/A
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2016/0174099	12/2015	Goldfain	N/A	N/A
2016/0174862	12/2015	Yu et al.	N/A	N/A
2016/0174863	12/2015	Foerster et al.	N/A	N/A
2016/0174867	12/2015	Hatano et al.	N/A	N/A
2016/0174907	12/2015	Colman et al.	N/A	N/A
2016/0175557	12/2015	Tass	N/A	N/A
2016/0175607	12/2015	Deisseroth et al.	N/A	N/A
2016/0176053	12/2015	Rognini et al.	N/A	N/A
2016/0178392	12/2015	Goldfain	N/A	N/A
2016/0180042	12/2015	Menon et al.	N/A	N/A
2016/0180054	12/2015	Luo et al.	N/A	N/A
2016/0180055	12/2015	Fonte	N/A	N/A
2016/0183812	12/2015	Zhang et al.	N/A	N/A
2016/0183828	12/2015	Ouyang et al.	N/A	N/A
2016/0183861	12/2015	Hayes et al.	N/A	N/A

2016/0183881	12/2015	Keenan et al.	N/A	N/A
2016/0184029	12/2015	Peng et al.	N/A	N/A
2016/0184596	12/2015	Fried et al.	N/A	N/A
2016/0184599	12/2015	Segal	N/A	N/A
2016/0187524	12/2015	Suhami	N/A	N/A
2016/0191517	12/2015	Bae et al.	N/A	N/A
2016/0192841	12/2015	Inagaki et al.	N/A	N/A
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2016/0192879	12/2015	Yamashita	N/A	N/A
2016/0193499	12/2015	Kim et al.	N/A	N/A
2016/0196185	12/2015	Gu et al.	N/A	N/A
2016/0196393	12/2015	Avinash et al.	N/A	N/A
2016/0196635	12/2015	Cho et al.	N/A	N/A
2016/0196758	12/2015	Causevic et al.	N/A	N/A
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2016/0199577	12/2015	Hyde et al.	N/A	N/A
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2016/0203597	12/2015	Chang et al.	N/A	N/A
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2016/0204937	12/2015	Edwards et al.	N/A	N/A
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2016/0213947	12/2015	Han et al.	N/A	N/A
2016/0216760	12/2015	Trutna et al.	N/A	N/A
2016/0217586	12/2015	Dickrell et al.	N/A	N/A
2016/0217595	12/2015	Han et al.	N/A	N/A
2016/0219345	12/2015	Knight et al.	N/A	N/A
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2016/0223622	12/2015	Yu et al.	N/A	N/A
2016/0223627	12/2015	Shah et al.	N/A	N/A
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2016/0235323	12/2015	Tadi et al.	N/A	N/A
2016/0235324	12/2015	Mershin et al.	N/A	N/A
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2016/0235359	12/2015	Cho et al.	N/A	N/A
2016/0235980	12/2015	Berman et al.	N/A	N/A
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2016/0238673	12/2015	Honkura	N/A	N/A
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2016/0239966	12/2015	Parsey et al.	N/A	N/A
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2016/0240765	12/2015	Washington et al.	N/A	N/A
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2016/0256109	12/2015	Semenov	N/A	N/A
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2016/0259085	12/2015	Wilson et al.	N/A	N/A
2016/0259905	12/2015	Park et al.	N/A	N/A
2016/0260216	12/2015	Wu et al.	N/A	N/A
2016/0261962	12/2015	Petersen et al.	N/A	N/A
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2016/0262664	12/2015	Linderman	N/A	N/A
2016/0262680	12/2015	Martucci et al.	N/A	N/A
2016/0262685	12/2015	Wagner et al.	N/A	N/A
2016/0262695	12/2015	Zhang et al.	N/A	N/A
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2016/0263376	12/2015	Yoo et al.	N/A	N/A
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2016/0263393	12/2015	Vo-Dinh et al.	N/A	N/A
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2016/0270656	12/2015	Samec et al.	N/A	N/A
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2016/0274660	12/2015	Publicover et al.	N/A	N/A
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2016/0278651	12/2015	Lu et al.	N/A	N/A
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2016/0278687	12/2015	Xia	N/A	N/A
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2016/0278713	12/2015	Shoaran et al.	N/A	N/A
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2016/0278870	12/2015	Quaid et al.	N/A	N/A
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2016/0287308	12/2015	Grant et al.	N/A	N/A
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2016/0287869	12/2015	Errico et al.	N/A	N/A
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2016/0296746	12/2015	Wingeier et al.	N/A	N/A
2016/0298449	12/2015	Orban	N/A	N/A
2016/0299568	12/2015	Segal	N/A	N/A
2016/0300252	12/2015	Frank et al.	N/A	N/A
2016/0300352	12/2015	Raj	N/A	N/A
2016/0302683	12/2015	Lawrence et al.	N/A	N/A
2016/0302704	12/2015	Lynn et al.	N/A	N/A
2016/0302709	12/2015	Mossbridge	N/A	N/A
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2016/0303322	12/2015	John	N/A	N/A
2016/0303396	12/2015	Deisseroth et al.	N/A	N/A
2016/0303397	12/2015	Hirschman et al.	N/A	N/A
2016/0303402	12/2015	Tyler	N/A	N/A
2016/0306844	12/2015	Frank et al.	N/A	N/A
2016/0306942	12/2015	Rapaka et al.	N/A	N/A
2016/0310031	12/2015	Sarkar	N/A	N/A
2016/0310070	12/2015	Sabesan	N/A	N/A
2016/0310071	12/2015	Kim	N/A	N/A
2016/0313408	12/2015	Hatano et al.	N/A	N/A
2016/0313417	12/2015	Kawabata et al.	N/A	N/A
2016/0313418	12/2015	Fujii et al.	N/A	N/A
2016/0313798	12/2015	Connor	N/A	N/A
2016/0317056	12/2015	Moon et al.	N/A	N/A
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2016/0317077	12/2015	Sillay	N/A	N/A
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2016/0321742	12/2015	Phillips et al.	N/A	N/A
2016/0324445	12/2015	Kim et al.	N/A	N/A
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2016/0331952	12/2015	Faltys et al.	N/A	N/A
2016/0331970	12/2015	Lozano	N/A	N/A
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2016/0338634	12/2015	Neu et al.	N/A	N/A
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2017/0007122	12/2016	Samec et al.	N/A	N/A
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2017/0007165	12/2016	Jain et al.	N/A	N/A
2017/0007173	12/2016	Adamczyk et al.	N/A	N/A
2017/0007182	12/2016	Samec et al.	N/A	N/A
2017/0007450	12/2016	Samec et al.	N/A	N/A
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2017/0014083	12/2016	Diab et al.	N/A	N/A
2017/0014625	12/2016	Rosenbluth et al.	N/A	N/A
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2017/0079573	12/2016	Osorio	N/A	N/A
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2017/0112427	12/2016	Simon et al.	N/A	N/A
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2017/0188916	12/2016	Wang et al.	N/A	N/A
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2017/0188932	12/2016	Singer et al.	N/A	N/A
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Background/Summary

CROSS REFERENCE TO RELATED APPLICATIONS (1) The present application is a Continuation of U.S. patent application Ser. No. 17/693,395, filed Mar. 12, 2022, now pending, which is a Continuation of U.S. patent application Ser. No. 16/237,483, filed Dec. 31, 2018, now U.S. Pat. No. 11,273,283, issued Mar. 15, 2022, and is a Continuation of Ser. No. 16/237,471, filed Dec. 31, 2018, now U.S. Pat. No. 11,478,603, issued Oct. 25, 2022, and is a Continuation of U.S. patent application Ser. No. 16/237,497, filed Dec. 31, 2018, now U.S. Pat. No. 11,318,277, issued May 3, 2022, which are each a Non-provisional of, and claims benefit of priority from, U.S. Provisional Patent Applications No. 62/612,565, filed Dec. 31, 2017, and No. 62/660,839 filed on Apr. 20, 2018, each of which are expressly incorporated herein by reference in their entirety. This application is related to PCT/US18/68220, filed Dec. 18, 2018.

FIELD OF THE INVENTION

(1) The present invention generally relates to the fields of neuroenhancement, neuromodulation, neurostimulation, and brain entrainment, and, more specifically, to devices, systems, and methods for selectively inducing brainwave activity patterns in humans or animals that correspond to, or enhance, an emotion or emotional response.

BACKGROUND OF THE INVENTION

(2) People often substitute an authentic experience by a replica thereof. Those who cannot visit the Louvre Museum, can look at the Mona Lisa on a reproduction. Anybody who has seen the real Mona Lisa in the Louvre can testify that the emotional experience is completely different from just looking at a reproduction. Yet, people often substitute reproductions for authentic works of art, when the latter are not readily accessible. The emotional response to viewing a reproduction pales in comparison to the emotional response to viewing an authentic piece of art in a museum. Looking at a photograph of the Grand Canyon is incomparable with experiencing the real thing—visiting the Grand Canyon, which is a breathtaking experience. Yet, people unable to travel, often replace the authentic experience of traveling and visiting new places with watching videos on the Travel Channel or on the Internet. Needless to say, watching TV or a video on the Internet is a poor substitute for the real experience of traveling and does not elicit the strong emotions, a person experiences when visiting new places.

(3) Because of lack of excitement in their daily lives people seek excitement in the movies. Movies tend to be more immersive experiences and can produce strong emotional responses. Many movie-goers cry while watching movies. A sentimental, emotionally-charged movie is referred to as a tear-jerker due to its ability to elicit a strong emotional response, resulting in tears. However, the emotions experience of watching a movie cannot be compared with the broad range of emotions experienced in real life.

(4) Recent advancements in 3D viewing technology and the emergence of Virtual Reality (VR) devices produce more realistic representation of reality they depict. However, even VR devices are incapable of producing emotional responses comparable to the emotions experienced in real life.

(5) A viewer may benefit from enhanced emotional responses associated with viewing art reproductions, watching TV, movies, Internet videos, or Virtual Reality.

(6) Some people lack certain emotions. For example, sociopathic personalities are incapable of experiencing emotions of empathy and compassion. A number of neurologic, psychiatric and psychological pathologies may affect the ability to experience certain emotions. Patients suffering from advanced stages of Parkinson and Alzheimer's diseases often exhibit subdued emotional response. Patients affected by paranoid schizophrenia, brain injury, or dementia sometimes experience Capgras delusion. They see a familiar face of a spouse or another family member but do not experience emotional response they expect to experience when seeing a face of a close family member, which leads them to believe that they live with an imposter that only "looks like" their family member, they complaint about a doppelganger living with them. It may be beneficial to artificially enhance the emotional response of a patient, bringing it to the normal level expected of a healthy person.

(7) It is well known that memory retention is affected by the emotional state of the person. Emotionally-charged experiences are etched in the memory, whereas experiences not associated with high emotions are easily forgotten. Artificially raising emotional levels during study may significantly increase the retention of the information and ease its subsequent recall.

(8) It has been observed in neuroscience that various emotions correlate with different frequency and location of the brainwaves. Accordingly, inducing in a subject the brainwaves of particular frequency in a particular location may induce and/or enhance the desired emotional response.

(9) Emotions are viewed as discrete and dimensional. The discrete framework classifies emotional states as physiological and behavioral manifestations of discrete emotions such as anger, happiness, etc. The dimensional perspective organizes emotional states by two factors, valence (positive/negative) and arousal (calm/exciting).

(10) Emotions are thought to be associated with different parts of the brain:

(11) Frontal Lobe (movement of the body; personality; concentration, planning, problem solving; meaning of words; emotional reactions; speech; smell); Parietal Lobe (touch and pressure; taste; body awareness); Temporal Lobe (hearing; recognizing faces; emotion; long-term memory); Occipital Lobe (sight); Cerebellum (Latin for little brain, fine motor (muscle) control; balance and coordination (avoid objects and keep from falling)); Limbic Lobe (controls emotions like happiness, sadness, and love).

(12) Each reference and document cited herein is expressly incorporated herein by reference in its entirety, for all purposes.

(13) Time in a biological matter Almost everything in biology is subject to change over time. These changes occur on many different time scales, which vary greatly. For example, there are evolutionary changes that affect entire populations over time rather than a single organism. Evolutionary changes are often slower than a human time scale that spans many years (usually, a human lifetime). Faster variations of the timing and duration of biological activity in living organisms occur, for example, in many essential biological processes in everyday life: in humans and animals, these variations occur, for example, in eating, sleeping, mating, hibernating, migration, cellular regeneration, etc. Other fast changes may include the transmission of a neural signal, for example, through a synapse, such as the Calyx of Held, a particularly large synapse in the auditory central nervous system of mammals that can reach transmission frequencies of up to 50 Hz. With recruitment modulation, the effective frequencies can be higher. A single nerve impulse can reach a speed as high as one hundred meters (0.06 mile) per second (Kraus, David. Concepts in Modern Biology. New York: Globe Book Company, 1969: 170.). Myelination of axons can increase the speed of transmission by segmenting the membrane depolarization process.

(14) Many of these changes over time are repetitive or rhythmic and are described as some frequency or oscillation. The field of chronobiology, examines such periodic (cyclic) phenomena in living organisms and their adaptation, for example, to solar and lunar-related rhythms (DeCoursey, et al. (2003).) These cycles are also known as biological rhythms. The related terms "chronomics" and "chronome" have been used in some cases to describe either the molecular mechanisms invoked in chronobiological phenomena or the more quantitative aspects of chronobiology, particularly where comparison of cycles between organisms is required. Chronobiological studies include, but are not limited to, comparative anatomy, physiology, genetics, molecular biology and behavior of organisms within biological rhythms' mechanics (DeCoursey et al. (2003).). Other aspects include epigenetics, development, reproduction, ecology, and evolution.

(15) The most important rhythms in chronobiology are the circadian rhythms, roughly 24-hour cycles shown by physiological processes in all

organisms. They are regulated by circadian clocks. The circadian rhythms can be further broken down into routine cycles during the 24-hour day (Nelson R. J. 2005. *An Introduction to Behavioral Endocrinology*. Sinauer Associates, Inc.: Massachusetts. Pg. 587.) All animals can be classified according to their activity cycles: Diurnal, which describes organisms active during daytime; Nocturnal, which describes organisms active in the night and Crepuscular, which describes animals primarily active during the dawn and dusk hours (e.g., white-tailed deer, some bats).

(16) While circadian rhythms are defined as regulated by endogenous processes, other biological cycles may be regulated by exogenous signals. In some cases, multi-trophic systems may exhibit rhythms driven by the circadian clock of one of the members (which may also be influenced or reset by external factors).

(17) Many other important cycles are also studied, including: Infradian rhythms, which are cycles longer than a day. Examples include circannual or annual cycles that govern migration or reproduction cycles in many plants and animals, or the human menstrual cycle; ultradian rhythms, which are cycles shorter than 24 hours, such as the 90-minute REM cycle, the 4-hour nasal cycle, or the 3-hour cycle of growth hormone production; tidal rhythms, commonly observed in marine life, which follow the roughly 12.4-hour transition from high to low tide and back; lunar rhythms, which follow the lunar month (29.5 days). They are relevant, for example, to marine life, as the level of the tides is modulated across the lunar cycle; and gene oscillations—some genes are expressed more during certain hours of the day than during other hours.

(18) Within each cycle, the time period during which the process is more active is called the acrophase (Refinetti, Roberto (2006). *Circadian Physiology*. CRC Press/Taylor & Francis Group. ISBN 0-8493-2233-2. Lay summary). When the process is less active, the cycle is in its bathyphase, or trough phase. The particular moment of highest activity is the peak or maximum; the lowest point is the nadir. How high (or low) the process gets is measured by the amplitude.

(19) Neural Correlates A neural correlate of an emotional or mental state is an electro-neuro-biological state or the state assumed by some biophysical subsystem of the brain, whose presence necessarily and regularly correlates with such specific emotional or mental states. All properties credited to the mind, including consciousness, emotion, and desires are thought to have direct neural correlates. For our purposes, neural correlates of an emotional or mental state can be defined as the minimal set of neuronal oscillations that correspond to the given emotional or mental state. Neuroscience uses empirical approaches to discover neural correlates of emotional or mental state.

(20) Mental State A mental state is a state of mind that a subject is in. Some mental states are pure and unambiguous, while humans are capable of complex states that are a combination of mental representations, which may have in their pure state contradictory characteristics. There are several paradigmatic states of mind that a subject has: love, hate, pleasure, fear, and pain. Mental states can also include a waking state, a sleeping state, a flow (or being in the “zone”), a will (desire) for something, and a mood (a mental state). A mental state is a hypothetical state that corresponds to thinking and feeling, and consists of a conglomeration of mental representations. A mental state is related to an emotion, though it can also relate to cognitive processes. Because the mental state itself is complex and potentially possesses inconsistent attributes, clear interpretation of mental state through external analysis (other than self-reporting) is difficult or impossible. However, some studies report that certain attributes of mental state or thought processes may, in fact, be determined through passive monitoring, such as EEG, or fMRI with some degree of statistical reliability. In most studies, the characterization of mental state was an endpoint, and the raw signals, after statistical classification or semantic labeling, are superseded. The remaining signal energy treated as noise.

(21) A number of studies report that certain attributes of mental state or thought processes may in fact be determined through passive monitoring, such as EEG, with some degree of statistical reliability. In most studies, the characterization of mental state was an endpoint, and the raw signals, after statistically classification or semantic labelling, are superseded and the remaining signal energy treated as noise.

(22) Brain The brain is a key part of the central nervous system, enclosed in the skull. In humans, and mammals more generally, the brain controls both autonomic processes, as well as cognitive processes. The brain (and to a lesser extent, the spinal cord) controls all volitional functions of the body and interprets information from the outside world. Intelligence, memory, emotions, speech, thoughts, movements and creativity are controlled by the brain. The central nervous system also controls autonomic functions and many homeostatic and reflex actions, such as breathing, heart rate, etc. The human brain consists of the cerebrum, cerebellum, and brainstem. The brainstem includes the midbrain, the pons, and the medulla oblongata. Sometimes the diencephalon, the caudal part of the forebrain, is included.

(23) The brain is composed of neurons, neuroglia (a.k.a., glia), and other cell types in connected networks that integrate sensory inputs, control movements, facilitate learning and memory, activate and express emotions, and control all other behavioral and cognitive functions. Neurons communicate primarily through electrochemical pulses that transmit signals between connected cells within and between brain areas. Thus, the desire to noninvasively capture and replicate neural activity associated with cognitive states has been a subject of interest to behavioral and cognitive neuroscientists.

(24) Technological advances now allow for non-invasive recording of large quantities of information from the brain at multiple spatial and temporal scales. Examples include electroencephalogram (“EEG”) data using multi-channel electrode arrays placed on the scalp or inside the brain, magnetoencephalography (“MEG”), magnetic resonance imaging (“MRI”), functional data using functional magnetic resonance imaging (“fMRI”), positron emission tomography (“PET”), near-infrared spectroscopy (“NIRS”), single-photon emission computed tomography (“SPECT”), and others.

(25) Noninvasive neuromodulation technologies have also been developed that can modulate the pattern of neural activity, and thereby cause altered behavior, cognitive states, perception, and motor output Integration of noninvasive measurement and neuromodulation techniques for identifying and transplanting brain states from neural activity would be very valuable for clinical therapies, such as brain stimulation and related technologies often attempting to treat disorders of cognition.

(26) The brainstem provides the main motor and sensory innervation to the face and neck via the cranial nerves. Of the twelve pairs of cranial nerves, ten pairs come from the brainstem. This is an extremely important part of the brain, as the nerve connections of the motor and sensory systems from the main part of the brain to the rest of the body pass through the brainstem. This includes the corticospinal tract (motor), the posterior column-medial lemniscus pathway (fine touch, vibration sensation, and proprioception), and the spinothalamic tract (pain, temperature, itch, and crude touch). The brainstem also plays an important role in the regulation of cardiac and respiratory function. It also regulates the central nervous system and is pivotal in maintaining consciousness and regulating the sleep cycle. The brainstem has many basic functions including controlling heart rate, breathing, sleeping, and eating.

(27) The function of the skull is to protect delicate brain tissue from injury. The skull consists of eight fused bones: the frontal, two parietal, two temporal, sphenoid, occipital and ethmoid. The face is formed by 14 paired bones including the maxilla, zygoma, nasal, palatine, lacrimal, inferior nasal conchae, mandible, and vomer. The bony skull is separated from the brain by the dura, a membranous organ, which in turn contains cerebrospinal fluid. The cortical surface of the brain typically is not subject to localized pressure from the skull. The skull, therefore, imposes a barrier to electrical access to the brain functions, and in a healthy human, breaching the dura to access the brain is highly disfavored. The result is that electrical readings of brain activity are filtered by the dura, the cerebrospinal fluid, the skull, the scalp, hair, resulting in a loss of potential spatial resolution and amplitude of signals emanating from the brain. While magnetic fields resulting from brain electrical activity are accessible, the spatial resolution using feasible sensors is also limited.

(28) The cerebrum is the largest part of the brain and is composed of right and left hemispheres. It performs higher functions, such as interpreting inputs from the senses, as well as speech, reasoning, emotions, learning, and fine control of movement. The surface of the cerebrum has a folded appearance called the cortex. The human cortex contains about 70% of the nerve cells (neurons) and gives an appearance of gray color (grey matter). Beneath the cortex are long connecting fibers between neurons, called axons, which make up the white matter.

(29) The cerebellum is located behind the cerebrum and brainstem. It coordinates muscle movements, helps to maintain balance and posture. The cerebellum may also be involved in some cognitive functions such as attention and language, as well as in regulating fear and pleasure responses.

There is considerable evidence that the cerebellum plays an essential role in some types of motor learning. The tasks where the cerebellum most clearly comes into play are those in which it is necessary to make fine adjustments to the way an action is performed. There is a dispute about whether learning takes place within the cerebellum itself, or whether it merely serves to provide signals that promote learning in other brain structures. Cerebellum also plays an important role in sleep and long-term memory formation.

(30) The brain communicates with the body through the spinal cord and twelve pairs of cranial nerves. Ten of the twelve pairs of cranial nerves that control hearing, eye movement, facial sensations, taste, swallowing and movement of the face, neck, shoulder and tongue muscles originate in the brainstem. The cranial nerves for smell and vision originate in the cerebrum.

(31) The right and left hemispheres of the brain are joined by a structure consisting of fibers called the corpus callosum. Each hemisphere controls the opposite side of the body. The right eye sends visual signals to the left hemisphere and vice versa. However, the right ear sends signals to the right hemisphere, and the left ear sends signals to the left hemisphere. Not all functions of the hemispheres are shared. For example, speech is processed exclusively in the left hemisphere.

(32) The cerebral hemispheres have distinct structures, which divide the brain into lobes. Each hemisphere has four lobes: frontal, temporal, parietal, and occipital. There are very complex relationships between the lobes of the brain and between the right and left hemispheres:

(33) Frontal lobes control judgment, planning, problem-solving, behavior, emotions, personality, speech, self-awareness, concentration, intelligence, body movements.

(34) Temporal lobes control understanding of language, memory, organization, and hearing.

(35) Parietal lobes control the interpretation of language; input from vision, hearing, sensory, and motor, temperature, pain, tactile signals, memory, spatial and visual perception.

(36) Occipital lobes interpret visual input (movement, light, color).

(37) A neuron is a fundamental unit of the nervous system, which comprises the autonomic nervous system and the central nervous system.

(38) Brain structures and particular areas within brain structures include but are not limited to hindbrain structures (e.g., myelencephalon structures (e.g., medulla oblongata, medullary pyramids, olivary body, inferior olivary nucleus, respiratory center, cuneate nucleus, gracile nucleus, intercalated nucleus, medullary cranial nerve nuclei, inferior salivatory nucleus, nucleus ambiguus, dorsal nucleus of the vagus nerve, hypoglossal nucleus, solitary nucleus, etc.), metencephalon structures (e.g., pons, pontine cranial nerve nuclei, chief or pontine nucleus of the trigeminal nerve sensory nucleus (V), motor nucleus for the trigeminal nerve (V), abducens nucleus (VI), facial nerve nucleus (VII), vestibulocochlear nuclei (vestibular nuclei and cochlear nuclei) (VIII), superior salivatory nucleus, pontine tegmentum, respiratory centers, pneumotaxic center, apneustic center, pontine micturition center (Barrington's nucleus), locus coeruleus, pedunculopontine nucleus, laterodorsal tegmental nucleus, tegmental pontine reticular nucleus, superior olivary complex, paramedian pontine reticular formation, cerebella peduncles, superior cerebella peduncle, middle cerebella peduncle, inferior cerebella peduncle, fourth ventricle, cerebellum, cerebella vermis, cerebella hemispheres, anterior lobe, posterior lobe, flocculonodular lobe, cerebella nuclei, fastigial nucleus, interposed nucleus, globose nucleus, emboliform nucleus, dentate nucleus, etc.)), midbrain structures (e.g., tectum, corpora quadrigemina, inferior colliculi, superior colliculi, pretectum, tegmentum, periaqueductal gray, parabrachial area, medial parabrachial nucleus, lateral parabrachial nucleus, subparabrachial nucleus (Kolliker-Fuse nucleus), rostral interstitial nucleus of medial longitudinal fasciculus, midbrain reticular formation, dorsal raphe nucleus, red nucleus, ventral tegmental area, substantia nigra, pars compacta, pars reticulata, interpeduncular nucleus, cerebral peduncle, cms cerebri, mesencephalic cranial nerve nuclei, oculomotor nucleus (III), trochlear nucleus (IV), mesencephalic duct (cerebral aqueduct, aqueduct of Sylvius), etc.), forebrain structures (e.g., diencephalon, epithalamus structures (e.g., pineal body, habenular nuclei, stria medullares, tения thalami, etc.), third ventricle, thalamus structures (e.g., anterior nuclear group, anteroventral nucleus (a.k.a. ventral anterior nucleus), anterodorsal nucleus, anteromedial nucleus, medial nuclear group, medial dorsal nucleus, midline nuclear group, paratenial nucleus, reuniens nucleus, rhomboidal nucleus, intralaminar nuclear group, centromedial nucleus, parafascicular nucleus, paracentral nucleus, central lateral nucleus, central medial nucleus, lateral nuclear group, lateral dorsal nucleus, lateral posterior nucleus, pulvinar, ventral nuclear group, ventral anterior nucleus, ventral lateral nucleus, ventral posterior nucleus, ventral posterior lateral nucleus, ventral posterior medial nucleus, metathalamus, medial geniculate body, lateral geniculate body, thalamic reticular nucleus, etc.), hypothalamus structures (e.g., anterior, medial area, parts of preoptic area, medial preoptic nucleus, suprachiasmatic nucleus, paraventricular nucleus, supraoptic nucleus (mainly), anterior hypothalamic nucleus, lateral area, parts of preoptic area, lateral preoptic nucleus, anterior part of lateral nucleus, part of supraoptic nucleus, other nuclei of preoptic area, median preoptic nucleus, periventricular preoptic nucleus, tuberal, medial area, dorsomedial hypothalamic nucleus, ventromedial nucleus, arcuate nucleus, lateral area, tuberal part of lateral nucleus, lateral tuberal nuclei, posterior, medial area, mammillary nuclei (part of mammillary bodies), posterior nucleus, lateral area, posterior part of lateral nucleus, optic chiasm, subfornical organ, periventricular nucleus, pituitary stalk, tuber cinereum, tuberal nucleus, tuberomammillary nucleus, tuberal region, mammillary bodies, mammillary nucleus, etc.), subthalamus structures (e.g., thalamic nucleus, zona incerta, etc.), pituitary gland structures (e.g., neurohypophysis, pars intermedia (intermediate lobe), adenohypophysis, etc.), telencephalon structures, white matter structures (e.g., corona radiata, internal capsule, external capsule, extreme capsule, arcuate fasciculus, uncinate fasciculus, perforant path, etc.), subcortical structures (e.g., hippocampus (medial temporal lobe), dentate gyrus, cornu ammonis (CA fields), cornu ammonis area 1, cornu ammonis area 2, cornu ammonis area 3, cornu ammonis area 4, amygdala (limbic system) (limbic lobe), central nucleus (autonomic nervous system), medial nucleus (accessory olfactory system), cortical and basomedial nuclei (main olfactory system), lateral) and basolateral nuclei (frontotemporal cortical system), claustrum, basal ganglia, stratum, dorsal stratum (a.k.a. neostriatum), putamen, caudate nucleus, ventral stratum, nucleus accumbens, olfactory tubercle, globus pallidus (forms nucleus lentiformis with putamen), subthalamus structures, basal forebrain, anterior perforated substance, substantia innominata, nucleus basalis, diagonal band of Broca, medial septal nuclei, etc.), rhinencephalon structures (e.g., olfactory bulb, piriform cortex, anterior olfactory nucleus, olfactory tract, anterior commissure, uncus, etc.), cerebral cortex structures (e.g., frontal lobe, cortex, primary motor cortex (precentral gyrus, M1), supplementary motor cortex, premotor cortex, prefrontal cortex, gyri, superior frontal gyrus, middle frontal gyrus, inferior frontal gyrus, Brodmann areas: 4, 6, 8, 9, 10, 11, 12, 24, 25, 32, 33, 44, 45, 46, 47, parietal lobe, cortex, primary somatosensory cortex (S1), secondary somatosensory cortex (S2), posterior parietal cortex, gyri, postcentral gyrus (primary somesthetic area), precuneus, Brodmann areas 1, 2, 3 (primary somesthetic area); 5, 7, 23, 26, 29, 31, 39, 40, occipital lobe, cortex, primary visual cortex (V1), V2, V3, V4, V5/MT, lateral occipital gyrus, cuneus, Brodmann areas 17 (V1, primary visual cortex); 18, 19, temporal lobe, primary auditory cortex (A1), secondary auditory cortex (A2), inferior temporal cortex, posterior inferior temporal cortex, superior temporal gyms, middle temporal gyrus, inferior temporal gyrus, entorhinal cortex, perirhinal cortex, parahippocampal gyrus, fusiform gyrus, Brodmann areas: 9, 20, 21, 22, 27, 34, 35, 36, 37, 38, 41, 42, medial superior temporal area (MST), insular cortex, cingulate cortex, anterior cingulate, Posterior cingulate, Retrosplenial cortex, Indusium griseum, Subgenual area 25, Brodmann areas 23, 24; 26, 29, 30 (retrosplenial areas); 31, 32, etc.).

(39) The brain is the largest sex organ controlling the biological urge, mediating all thoughts, experiences and physiological responses to sex. The euphoric and pleasurable experience of sex stems primarily from the limbic system including the amygdala, hippocampus and limbic lobe (dentate and cingulate gyrus).

(40) Neurons Neurons are electrically excitable cells that receive, process, and transmit information, and based on that information sends a signal to other neurons, muscles, or glands through electrical and chemical signals. These signals between neurons occur via specialized connections called synapses. Neurons can connect to each other to form neural networks. The basic purpose of a neuron is to receive incoming information and, based upon that information send a signal to other neurons, muscles, or glands. Neurons are designed to rapidly send signals across physiologically long distances. They do this using electrical signals called nerve impulses or action potentials. When a nerve impulse reaches the end of a neuron, it triggers the release of a chemical, or neurotransmitter. The neurotransmitter travels rapidly across the short gap between cells (the synapse) and acts

to signal the adjacent cell. See [www.biologyonline.com/Mod-Nu/Neuron.html#ixzz5AVxCuM5a](#).

(41) Neurons can receive thousands of inputs from other neurons through synapses. Synaptic integration is a mechanism whereby neurons integrate these inputs before the generation of a nerve impulse, or action potential. The ability of synaptic inputs to effect neuronal output is determined by a number of factors: Size, shape and relative timing of electrical potentials generated by synaptic inputs; the geometric structure of the target neuron; the physical location of synaptic inputs within that structure; and the expression of voltage-gated channels in different regions of the neuronal membrane.

(42) Neurons within a neural network receive information from, and send information to, many other cells, at specialized junctions called synapses. Synaptic integration is the computational process by which an individual neuron processes its synaptic inputs and converts them into an output signal. Synaptic potentials occur when neurotransmitter binds to and opens ligand-operated channels in the dendritic membrane, allowing ions to move into or out of the cell according to their electrochemical gradient. Synaptic potentials can be either excitatory or inhibitory depending on the direction and charge of ion movement. Action potentials occur if the summed synaptic inputs to a neuron reach a threshold level of depolarization and trigger regenerative opening of voltage-gated ion channels. Synaptic potentials are often brief and of small amplitude, therefore summation of inputs in time (temporal summation) or from multiple synaptic inputs (spatial summation) is usually required to reach action potential firing threshold.

(43) There are two types of synapses: electrical synapses and chemical synapses. Electrical synapses are a direct electrical coupling between two cells mediated by gap junctions, which are pores constructed of connexin proteins—essentially result in the passing of a gradient potential (may be depolarizing or hyperpolarizing) between two cells. Electrical synapses are very rapid (no synaptic delay). It is a passive process where signal can degrade with distance and may not produce a large enough depolarization to initiate an action potential in the postsynaptic cell. Electrical synapses are bidirectional, i.e., postsynaptic cell can actually send messages to the “presynaptic cell.”

(44) Chemical synapses are a coupling between two cells through neuro-transmitters, ligand or voltage gated channels, receptors. They are influenced by the concentration and types of ions on either side of the membrane. Among the neurotransmitters, Glutamate, sodium, potassium, and calcium are positively charged. GABA and chloride are negatively charged. Neurotransmitter junctions provide an opportunity for pharmacological intervention, and many different drugs, including illicit drugs, act at synapses.

(45) An excitatory postsynaptic potential (EPSP) is a postsynaptic potential that makes the postsynaptic neuron more likely to fire an action potential. An electrical charge (hyperpolarization) in the membrane of a postsynaptic neuron is caused by the binding of an inhibitory neurotransmitter from a presynaptic cell to a postsynaptic receptor. It makes it more difficult for a postsynaptic neuron to generate an action potential. An electrical change (depolarization) in the membrane of a postsynaptic neuron caused by the binding of an excitatory neurotransmitter from a presynaptic cell to a postsynaptic receptor. It makes it more likely for a postsynaptic neuron to generate an action potential. In a neuronal synapse that uses glutamate as receptor, for example, receptors open ion channels that are non-selectively permeable to cations. When these glutamate receptors are activated, both Na^+ and K^+ flow across the postsynaptic membrane. The reversal potential (Erev) for the post-synaptic current is approximately 0 mV. The resting potential of neurons is approximately -60 mV. The resulting EPSP will depolarize the post synaptic membrane potential, bringing it toward 0 mV.

(46) An inhibitory postsynaptic potential (IPSP) is a kind of synaptic potential that makes a postsynaptic neuron less likely to generate an action potential. An example of inhibitory post synaptic action is a neuronal synapse that uses γ -Aminobutyric acid (GABA) as its transmitter. At such synapses, the GABA receptors typically open channels that are selectively permeable to Cl^- . When these channels open, negatively charged chloride ions can flow across the membrane. The postsynaptic neuron has a resting potential of -60 mV and an action potential threshold of -40 mV. Transmitter release at this synapse will inhibit the postsynaptic cell. Since ECI is more negative than the action potential threshold, e.g., -70 mV, it reduces the probability that the postsynaptic cell will fire an action potential.

(47) Some types of neurotransmitters, such as glutamate, consistently result in EPSPs. Others, such as GABA, consistently result in IPSPs. The action potential lasts about one millisecond (1 msec). In contrast, the EPSPs and IPSPs can last as long as 5 to 10 msec. This allows the effect of one postsynaptic potential to build upon the next and so on.

(48) Membrane leakage, and to a lesser extent, potentials per se, can be influenced by external electrical and magnetic fields. These fields may be generated focally, such as through implanted electrodes, or less specifically, such as through transcranial stimulation. Transcranial stimulation may be subthreshold or superthreshold. In the former case, the external stimulation acts to modulate resting membrane potential, making nerves more or less excitable. Such stimulation may be direct current or alternating current. In the latter case, this will tend to synchronize neuron depolarization with the signals. Superthreshold stimulation can be painful (at least because the stimulus directly excites pain neurons) and must be pulsed. Since this has correspondence to electroconvulsive therapy, superthreshold transcranial stimulation is sparingly used.

(49) A number of neurotransmitters are known, as are pharmaceutical interventions and therapies that influence these compounds. Typically, the major neurotransmitters are small monoamine molecules, such as dopamine, epinephrine, norepinephrine, serotonin, GABA, histamine, etc., as well as acetylcholine. In addition, neurotransmitters also include amino acids, gas molecules such as nitric oxide, carbon monoxide, carbon dioxide, and hydrogen sulfide, as well as peptides. The presence, metabolism, and modulation of these molecules may influence learning and memory. Supply of neurotransmitter precursors, control of oxidative and mental stress conditions, and other influences on learning and memory-related brain chemistry, may be employed to facilitate memory, learning, and learning adaption transfer.

(50) The neuropeptides, as well as their respective receptors, are widely distributed throughout the mammalian central nervous system. During learning and memory processes, besides structural synaptic remodeling, changes are observed at molecular and metabolic levels with the alterations in neurotransmitter and neuropeptide synthesis and release. While there is a consensus that brain cholinergic neurotransmission plays a critical role in the processes related to learning and memory, it is also well known that these functions are influenced by a tremendous number of neuropeptides and non-peptide molecules. Arginine vasopressin (AVP), oxytocin, angiotensin II, insulin, growth factors, serotonin (5-HT), melanin-concentrating hormone, histamine, bombesin and gastrin-releasing peptide (GRP), glucagon-like peptide-1 (GLP-1), cholecystokinin (CCK), dopamine, corticotropin-releasing factor (CRF) have modulatory effects on learning and memory. Among these peptides, CCK, 5-HT, and CRF play strategic roles in the modulation of memory processes under stressful conditions. CRF is accepted as the main neuropeptide involved in both physical and emotional stress, with a protective role during stress, possibly through the activation of the hypothalamo-pituitary (HPA) axis. The peptide CCK has been proposed to facilitate memory processing, and CCK-like immunoreactivity in the hypothalamus was observed upon stress exposure, suggesting that CCK may participate in the central control of stress response and stress-induced memory dysfunction. On the other hand, 5-14T appears to play a role in behaviors that involve a high cognitive demand and stress exposure activates serotonergic systems in a variety of brain regions. See: Mehmetali Gülpinar, Berrak C Yeğen, “The Physiology of Learning and Memory: Role of Peptides and Stress”, *Current Protein and Peptide Science*, 2004(5) www.researchgate.net/publication/8147320_The_Physiology_of_Learning_and_Memory_Role_of_Peptides_and_Stress. Deep brain stimulation is described in NIH Research Matters, “A noninvasive deep brain stimulation technique”, (2017), Brainworks, “QEEG Brain Mapping”. Carmon, A., Mor, J., & Goldberg, J. (1976). Evoked cerebral responses to noxious thermal stimuli in humans. *Experimental Brain Research*, 25(1), 103-107.

(51) Brainwaves At the root of all our thoughts, emotions and behaviors is the communication between neurons within our brains, a rhythmic or repetitive neural activity in the central nervous system. The oscillation can be produced by a single neuron or by synchronized electrical pulses from ensembles of neurons communicating with each other. The interaction between neurons can give rise to oscillations at a different frequency than the firing frequency of individual neurons. The synchronized activity of large numbers of neurons produces macroscopic oscillations, which can be observed in an electroencephalogram. They are divided into bandwidths to describe their purported functions or functional relationships. Oscillatory activity in the brain is widely observed at different levels of organization and is thought to play a key role in processing neural information. Numerous experimental studies support a functional role of neural oscillations. A unified interpretation, however, is still not determined. Neural

oscillations and synchronization have been linked to many cognitive functions such as perception, motor control and memory. Electroencephalographic (EEG) signals are relatively easy and safe to acquire, have a long history of analysis, and can have high dimensionality, e.g., up to 128 or 256 separate recording electrodes. While the information represented in each electrode is not independent of the others, and the noise in the signals high, there is much information available through such signals that has not been fully characterized to date.

(52) Brainwaves have been widely studied in neural activity generated by large groups of neurons, mostly by EEG. In general, EEG signals reveal oscillatory activity (groups of neurons periodically firing in synchrony), in specific frequency bands: alpha (7.5-12.5 Hz) that can be detected from the occipital lobe during relaxed wakefulness and which increases when the eyes are closed; delta (1-4 Hz), theta (4-8 Hz), beta (13-30 Hz), low gamma (30-70 Hz), and high gamma (70-150 Hz) frequency bands, where faster rhythms such as gamma activity have been linked to cognitive processing. Higher frequencies imply multiple groups of neurons firing in coordination, either in parallel or in series, or both, since individual neurons do not fire at rates of 100 Hz. Neural oscillations of specific characteristics have been linked to cognitive states, such as awareness and consciousness and different sleep stages.

(53) Nyquist Theorem states that the highest frequency that can be accurately represented is one-half of the sampling rate. Practically, the sampling rate should be ten times higher than the highest frequency of the signal. (See, www.slideshare.net/ertvk/eeg-examples). While EEG signals are largely band limited, the superimposed noise may not be. Further, the EEG signals themselves represent components from a large number of neurons, which fire independently. Therefore, large bandwidth signal acquisition may have utility.

(54) It is a useful analogy to think of brainwaves as music. In orchestral music, where various instrument groups (string groups, such as violins, violas, cellos and double basses, brass, woodwind, and percussion instruments) produce particular sounds based on their respective characteristic frequencies of vibrations that all come together in a musical composition. Similarly, in the brain, groups of neurons oscillate in unison producing specific frequencies that combine in brainwaves. Like in a symphony, the higher and lower frequencies link and cohere with each other through harmonics, especially when one considers that neurons may be coordinated not only based on transitions, but also on phase delay. Oscillatory activity is observed throughout the central nervous system at all levels of organization. Each respective mental state is associated with the dominant neuro oscillation frequency. Moreover, the nuances of each mental state may be associated with secondary and tertiary harmonics or, using musical analogy, the “overtones.” Some hypothesize that very slow brainwaves serve to synchronize various lobes and neuronal groups in the brain (similarly to low-frequency instruments, such as drums and double basses, serve to provide overall rhythm to the orchestra).

(55) The functions of brainwaves are wide-ranging and vary for different types of oscillatory activity. Neural oscillations also play an important role in many neurological disorders.

(56) Delta wave is the frequency range from 0.5 Hz to 4 Hz. It tends to be the highest in amplitude and the slowest waves (except for very-slow waves that have frequency less than 0.5 Hz). It is normally seen in adults in NREM (en.wikipedia.org/wiki/NREM). It is also seen normally in babies. It may occur focally with subcortical lesions and in general distribution with diffuse lesions, metabolic encephalopathy hydrocephalus or deep midline lesions. It is usually most prominent frontally in adults (e.g., FIRDA-frontal intermittent rhythmic delta) and posteriorly in children (e.g., OIRDA-occipital intermittent rhythmic delta).

(57) Theta is the frequency range from 4 Hz to 7 Hz. Theta is normally seen in young children. It may be seen in drowsiness or arousal in older children and adults; it can also be seen in meditation. Excess theta for age represents abnormal activity. It can be seen as a focal disturbance in focal subcortical lesions; it can be seen in generalized distribution in diffuse disorder or metabolic encephalopathy or deep midline disorders or some instances of hydrocephalus. On the other hand, this range has been associated with reports of relaxed, meditative, and creative states.

(58) Alpha is the frequency range from 7.5 Hz to 12.5 Hz. This is the “posterior basic rhythm” (also called the “posterior dominant rhythm,” the “posterior alpha rhythm” or the Berger’s wave), arising from the synchronous and coherent electrical activity in the thalamic pacemaker cells and seen in the posterior regions of the head on both sides, higher in amplitude on the dominant side. They predominantly originate from the occipital lobe during wakeful relaxation with closed eyes. Alpha wave emerges with the closing of the eyes and with relaxation and attenuates with eye opening or mental exertion. The posterior basic rhythm is actually slower than 8 Hz in young children (therefore technically in the theta range). In addition to the posterior basic rhythm, there are other normal alpha rhythms such as the sensorimotor, or mu rhythm (alpha activity in the contralateral sensory and motor cortical areas) that emerges when the hands and arms are idle; and the “third rhythm” (alpha activity in the temporal or frontal lobes). Alpha can be abnormal; for example, an EEG that has diffuse alpha occurring in coma and is not responsive to external stimuli is referred to as “alpha coma.”

(59) Beta is the frequency range from 15 Hz to about 30 Hz. It is usually seen on both sides in symmetrical distribution and is most evident frontally. Beta activity is closely linked to motor behavior and is generally attenuated during active movements. Low-amplitude beta with multiple and varying frequencies is often associated with active, busy or anxious thinking and active concentration. Rhythmic beta with a dominant set of frequencies is associated with various pathologies, such as Dup15q syndrome, and drug effects, especially benzodiazepines. It may be absent or reduced in areas of cortical damage. It is the dominant rhythm in patients who are alert or anxious or who have their eyes open.

(60) Gamma is the frequency range approximately 250-100 Hz. Gamma rhythms are thought to represent binding of different populations of neurons together into a network to carry out a certain cognitive or motor function. Low gamma (25-70 Hz), and high gamma (70-150 Hz) frequency bands are also recognized with higher frequencies being associated with cognitive processing.

(61) Mu range is 8-13 Hz and partly overlaps with other frequencies, but is generally considered one of the two types of alpha wave (the second type being the third rhythm). It reflects the synchronous firing of motor neurons in a rest state. Mu suppression is thought to reflect motor mirror neuron systems, because when an action is observed, the pattern extinguishes, possibly because of the normal neuronal system and the mirror neuron system “go out of sync” and interfere with each other. See: Abeles M, Local Cortical Circuits (1982) New York: Springer-Verlag. Braitenberg V and Schuz A (1991) Anatomy of the Cortex. Statistics and Geometry. New York: Springer-Verlag. Ebersole J S (1997) Defining epileptogenic foci: past, present, future. J. Clin. Neurophysiology 14: 470-483. Edelman G M and Tononi G (2000) A Universe of Consciousness, New York: Basic Books. Freeman W J (1975) Mass Action in the Nervous System, New York Academic Press. Gevins A S and Cuttillo B A (1995) Neuroelectric measures of mind. In: P L Nunez (Au), Neocortical Dynamics and Human EEG Rhythms. N Y: Oxford U. Press, pp. 304-338. Gevins A S, Le J, Martin N, Brickett P, Desmond J, and Reuter B (1994) High resolution EEG: 124-channel recording, spatial enhancement, and MRI integration methods.

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(62) TABLE-US-00001 TABLE 1 Comparison of EEG bands Freq. Band (Hz) Location Normally Pathologically Delta <4 frontally in adults, adult slow-wave sleep subcortical lesions posteriorly in babies diffuse lesions children; high- Has been found during metabolic encephalopathy hydrocephalus amplitude waves some continuous- deep midline lesions attention tasks Theta 4-7 Found in locations higher in young children focal subcortical lesions not related to task drowsiness in adults metabolic encephalopathy at hand and teens deep midline disorders idling some instances of hydrocephalus Associated with inhibition of elicited responses (has been found to spike in situations where a person is actively trying to repress a response or action). Alpha 7.5-12.5 posterior regions relaxed/reflecting Coma of head, both closing the eyes sides, higher in Also associated with amplitude on inhibition control, dominant side. seemingly with the Central sites (c3-c4) purpose of timing at rest inhibitory activity in different locations across the brain. Beta 12.5-30 both sides, range span: active calm Benzodiazepines (en.wikipedia.org/wiki/Benzodiazepines) symmetrical .fwdarw. intense .fwdarw. stressed Dup15q syndrome distribution, most .fwdarw. mild obsessive evident frontally; active thinking, focus, low-amplitude high alert, anxious waves Gamma 25-100 Somatosensory Displays during cross- A decrease in gamma-band activity may be associated with cortex modal sensory cognitive decline, especially when related to the theta band; processing (perception however, this has not been proven for use as a clinical that combines two diagnostic measurement different senses, such as sound and sight) Also is shown during short-term memory matching of recognized objects, sounds, or tactile sensations Mu 8-12 Sensorimotor Shows rest-state motor Mu suppression could indicate that motor mirror neurons are cortex neurons. working. Deficits in Mu suppression, and thus in mirror neurons, might play a role in autism.

(63) EEG AND qEEG An EEG electrode will mainly detect the neuronal activity in the brain region just beneath it. However, the electrodes receive the activity from thousands of neurons. One square millimeter of cortex surface, for example, has more than 100,000 neurons. It is only when the input to a region is synchronized with electrical activity occurring at the same time that simple periodic waveforms in the EEG become distinguishable. The temporal pattern associated with specific brainwaves can be digitized and encoded a non-transient memory, and embodied in or referenced by, computer software.

(64) EEG (electroencephalography) and MEG (magnetoencephalography) are available technologies to monitor brain electrical activity. Each generally has sufficient temporal resolution to follow dynamic changes in brain electrical activity. Electroencephalography (EEG) and quantitative electroencephalography (qEEG) are electrophysiological monitoring methods that analyze the electrical activity of the brain to measure and display patterns that correspond to cognitive states and/or diagnostic information. It is typically noninvasive, with the electrodes placed on the scalp, although invasive electrodes are also used in some cases. EEG signals may be captured and analyzed by a mobile device, often referred as "brain wearables". There are a variety of "brain wearables" readily available on the market today. EEGs can be obtained with a non-invasive method where the aggregate oscillations of brain electric potentials are recorded with numerous electrodes attached to the scalp of a person. Most EEG signals originate in the brain's outer layer (the cerebral cortex), believed largely responsible for our thoughts, emotions, and behavior. Cortical synaptic action generates electrical signals that change in the 10 to 100-millisecond range. Transcutaneous EEG signals are limited by the relatively insulating nature of the skull surrounding the brain, the conductivity of the cerebrospinal fluid and brain tissue, relatively low amplitude of individual cellular electrical activity, and distances between the cellular current flows and the electrodes. EEG is characterized by: (1) Voltage; (2) Frequency; (3) Spatial location; (4) Inter-hemispheric symmetries; (5) Reactivity (reaction to state change); (6) Character of waveform occurrence (random, serial, continuous); and (7) Morphology of transient events. EEGs can be separated into two main categories. Spontaneous EEG which occur in the absence of specific sensory stimuli and evoked potentials (EPs) which are associated with sensory stimuli like repeated light flashes, auditory tones, finger pressure or mild electric shocks. The latter is recorded for example by time averaging to remove effects of spontaneous EEG. Non-sensory triggered potentials are also known. EPs typically are time synchronized with the bigger, and thus have an organization principle. Event-related potentials (ERPs) provide evidence of a direct link between cognitive events and brain electrical activity in a wide range of cognitive paradigms. It has generally been held that an ERP is the result of a set of discrete stimulus-evoked brain events. Event-related potentials (ERPs) are recorded in the same way as EPs, but occur at longer latencies from the stimuli and are more associated with an endogenous brain state.

(65) In standard EEG recording practice, 19 recording electrodes are placed uniformly on the scalp (the International 10-20 System). In addition, one or two reference electrodes (often placed on earlobes) and a ground electrode (often placed on the nose to provide amplifiers with reference voltages) are required. However, additional electrodes may add minimal useful information unless supplemented by computer algorithms to reduce raw EEG data to a manageable form. When large numbers of electrodes are employed, the potential at each location may be measured with respect to the average of all potentials (the common average reference), which often provides a good estimate of potential at infinity. The common average reference is not appropriate when electrode coverage is sparse (perhaps less than 64 electrodes). (See, Paul L. Nunez and Ramesh Srinivasan (2007) *Electroencephalogram*. Scholarpedia, 2(2):1348, scholarpedia.org/article/Electroencephalogram. Dipole localization algorithms may be useful to determine spatial emission patterns in EEG.)

(66) Scalp potential may be expressed as a volume integral of dipole moment per unit volume over the entire brain provided $P(r,t)$ defined generally rather than in columnar terms. For the important case of dominant cortical sources, scalp potential may be approximated by the following integral

over the cortical volume Θ , $VS(r,t) = \iiint_{\Theta} \mathbf{d} \cdot \nabla G(r,r') \mathbf{d}\Theta(r')$. If the volume element $\mathbf{d}\Theta(r')$ is defined in terms of cortical columns, the volume integral may be reduced to an integral over the folded cortical surface. The time-dependence of scalp potential is the weighted sum of all dipole time variations in the brain, although deep dipole volumes typically make negligible contributions. The vector Green's function $G(r,r')$ contains all geometric and conductive information about the head volume conductor and weights the integral accordingly. Thus, each scalar component of the Green's function is essentially an inverse electrical distance between each source component and scalp location. For the idealized case of sources in an infinite medium of constant conductivity, the electrical distance equals the geometric distance. The Green's function accounts for the tissue's finite spatial extent and its inhomogeneity and anisotropy. The forward problem in EEG consists of choosing a head model to provide $G(r,r')$ and carrying out the integral for some assumed source distribution. The inverse problem consists of using the recorded scalp potential distribution $VS(r,t)$ plus some constraints (usual assumptions) on $P(r,t)$ to find the best fit source distribution $P(r,t)$. Since the inverse problem has no unique solution, any inverse solution depends critically on the chosen constraints, for example, only one or two isolated sources, distributed sources confined to the cortex, or spatial and temporal smoothness criteria. High-resolution EEG uses the experimental scalp potential $VS(r,t)$ to predict the potential on the dura surface (the unfolded membrane surrounding the cerebral cortex) $VD(r,t)$. This may be accomplished using a head model Green's function $G(r,r')$ or by estimating the surface Laplacian with either spherical or 3D splines. These two approaches typically provide very similar dura potentials $VD(r,t)$; the estimates of dura potential distribution are unique subject to head model, electrode density, and noise issues.

(67) In an EEG recording system, each electrode is connected to one input of a differential amplifier (one amplifier per pair of electrodes); a common system reference electrode (or synthesized reference) is connected to the other input of each differential amplifier. These amplifiers amplify the voltage between the active electrode and the reference (typically 1,000-100,000 times, or 60-100 dB of voltage gain). The amplified signal is digitized via an analog-to-digital converter, after being passed through an anti-aliasing filter. Analog-to-digital sampling typically occurs at 256-512 Hz in clinical scalp EEG; sampling rates of up to 20 kHz are used in some research applications. The EEG signals can be captured with open source hardware such as OpenBCI, and the signal can be processed by freely available EEG software such as EEGLAB or the Neurophysiological Biomarker Toolbox. A typical adult human EEG signal is about 10 μV to 100 μV in amplitude when measured from the scalp and is about 10-20 mV when measured from subdural electrodes.

(68) Typically, a magnetic sensor with sufficient sensitivity to individual cell depolarization or small groups is a superconducting quantum interference device (SQUID), which requires cryogenic temperature operation, either at liquid nitrogen temperatures (high temperature superconductors, HTS) or at liquid helium temperatures (low temperature superconductors, LTS). However, current research shows possible feasibility of room temperature superconductors (20 C). Magnetic sensing has an advantage, due to the dipole nature of sources, of having better potential volumetric localization; however, due to this added information, complexity of signal analysis is increased.

(69) In general, the electromagnetic signals detected represent action potentials, an automatic response of a nerve cell to depolarization beyond a threshold, which briefly opens conduction channels. The cells have ion pumps which seek to maintain a depolarized state. Once triggered, the action potential propagates along the membrane in two-dimensions, causing a brief high level of depolarizing ion flow. There is a quiescent period after depolarization that generally prevents oscillation within a single cell. Since the axon extends from the body of the neuron, the action potential will typically proceed along the length of the axon, which terminates in a synapse with another cell. While direct electrical connections between cells occur, often the axon releases a neurotransmitter compound into the synapse, which causes a depolarization or hyperpolarization of the target cell. Indeed, the result may also be release of a hormone or peptide, which may have a local or more distant effect.

(70) The electrical fields detectable externally tend to not include signals which low frequency signals, such as static levels of polarization, or cumulative depolarizing or hyperpolarizing effects between action potentials. In myelinated tracts, the current flows at the segments tend to be small, and therefore the signals from individual cells are small. Therefore, the largest signal components are from the synapses and cell bodies. In the cerebrum and cerebellum, these structures are mainly in the cortex, which is largely near the skull, making electroencephalography useful, since it provides spatial discrimination based on electrode location. However, deep signals are attenuated, and poorly localized. Magnetoencephalography detects dipoles, which derive from current flow, rather than voltage changes. In the case of a radially or spherically symmetric current flow within a short distance, the dipoles will tend to cancel, while net current flows long axons will reinforce. Therefore, an electroencephalogram reads a different signal than a magnetoencephalogram.

(71) EEG-based studies of emotional specificity at the single-electrode level demonstrated that asymmetric activity at the frontal site, especially in the alpha (8-12 Hz) band, is associated with emotion. Voluntary facial expressions of smiles of enjoyment produce higher left frontal activation. Decreased left frontal activity is observed during the voluntary facial expressions of fear. In addition to alpha band activity, theta band power at the frontal midline (Fm) has also been found to relate to emotional states. Pleasant (as opposed to unpleasant) emotions are associated with an increase in frontal midline theta power. Many studies have sought to utilize pattern classification, such as neural networks, statistical classifiers, clustering algorithms, etc., to differentiate between various emotional states reflected in EEG.

(72) EEG-based studies of emotional specificity at the single-electrode level demonstrated that asymmetric activity at the frontal site, especially in the alpha (8-12 Hz) band, is associated with emotion. Ekman and Davidson found that voluntary facial expressions of smiles of enjoyment produced higher left frontal activation (Ekman P, Davidson R J (1993) Voluntary Smiling Changes Regional Brain Activity. *Psychol Sci* 4: 342-345). Another study by Coan et al. found decreased left frontal activity during the voluntary facial expressions of fear (Coan J A, Allen J J, Harmon-Jones E (2001) Voluntary facial expression and hemispheric asymmetry over the frontal cortex. *Psychophysiology* 38: 912-925). In addition to alpha band activity, theta band power at the frontal midline (Fm) has also been found to relate to emotional states. Sammler and colleagues, for example, showed that pleasant (as opposed to unpleasant) emotion is associated with an increase in frontal midline theta power (Sammler D, Grigutsch M, Fritz T, Koelsch S (2007) Music and emotion: Electrophysiological correlates of the processing of pleasant and unpleasant music. *Psychophysiology* 44: 293-304). To further demonstrate whether these emotion-specific EEG characteristics are strong enough to differentiate between various emotional states, some studies have utilized a pattern classification analysis approach. See, for example: Dan N, Xiao-Wei W, Li-Chen S, Bao-Liang L. EEG-based emotion recognition during watching movies; 2011 Apr. 27 2011-May 1. 2011: 667-670; Lin Y P, Wang C H, Jung T P, Wu T L, Jeng S K, et al. (2010) EEG-Based Emotion Recognition in Music Listening. *Ieee T Bio Med Eng* 57: 1798-1806; Murugappan M, Nagarajan R, Yaacob S (2010) Classification of human emotion from EEG using discrete wavelet transform. *J Biomed Sci Eng* 3: 390-396; Murugappan M, Nagarajan R, Yaacob S (2011) Combining Spatial Filtering and Wavelet Transform for Classifying Human Emotions Using EEG Signals. *J Med. Bio. Eng.* 31: 45-51.

(73) Detecting different emotional states by EEG may be more appropriate using EEG-based functional connectivity. There are various ways to estimate EEG-based functional brain connectivity: correlation, coherence and phase synchronization indices between each pair of EEG electrodes had been used. The assumption is that a higher correlation map indicates a stronger relationship between two signals. (Brazier M A, Casby J U (1952) Cross-correlation and autocorrelation studies of electroencephalographic potentials. *Electroen clin neuro* 4: 201-211). Coherence gives information similar to correlation, but also includes the covariation between two signals as a function of frequency. (Cantero J L, Atienza M, Salas R M, Gomez C M (1999) Alpha EEG coherence in different brain states: an electrophysiological index of the arousal level in human subjects. *Neurosci lett* 271: 167-70.) The assumption is that higher coherence indicates a stronger relationship between two signals. (Guevara M A, Corsi-Cabrera M (1996) EEG coherence or EEG correlation? *Int J Psychophysiology* 23: 145-153; Centeno J L, Atienza M, Salas R M, Gomez C M (1999) Alpha EEG coherence in different brain states: an electrophysiological index of the arousal level in human subjects. *Neurosci lett* 271: 167-70; Adler G, Brassen S, Jajcevic A (2003) EEG coherence in Alzheimer's dementia. *J Neural Transm* 110: 1051-1058; Deeny S P, Hillman C H, Janelle C M, Hatfield B D (2003) Cortico-cortical communication and superior performance in skilled marksmen: An EEG coherence analysis. *J Sport Exercise Psy* 25: 188-204.) Phase synchronization among the neuronal groups estimated based on the phase difference between two signals is another way to estimate the EEG-based functional connectivity among brain areas. It is. (Fraszczuk P J, Bergey G K (1999) An autoregressive method for the

(74) A number of groups have examined emotional specificity using EEG-based functional brain connectivity. For example, Shin and Park showed that, when emotional states become more negative at high room temperatures, correlation coefficients between the channels in temporal and occipital sites increase (Shin J-H, Park D-H. (2011) Analysis for Characteristics of Electroencephalogram (EEG) and Influence of Environmental Factors According to Emotional Changes. In Lee G, Howard D, Ślęzak D, editors. *Convergence and Hybrid Information Technology*. Springer Berlin Heidelberg, 488-500.) Hinrichs and Machleidt demonstrated that coherence decreases in the alpha band during sadness, compared to happiness (Hinrichs H, Machleidt W (1992) Basic emotions reflected in EEG-coherences. *Int J Psychophysiol* 13: 225-232). Miskovic and Schmidt found that EEG coherence between the prefrontal cortex and the posterior cortex increased while viewing highly emotionally arousing (i.e., threatening) images, compared to viewing neutral images (Miskovic V, Schmidt L A (2010) Cross-regional cortical synchronization during affective image viewing. *Brain Res* 1362: 102-111). Costa and colleagues applied the synchronization index to detect interaction in different brain sites under different emotional states (Costa T, Rognoni E, Galati D (2006) EEG phase synchronization during emotional response to positive and negative film stimuli. *Neurosci Lett* 406: 159-164). Costa's results showed an overall increase in the synchronization index among frontal channels during emotional stimulation, particularly during negative emotion (i.e., sadness). Furthermore, phase synchronization patterns were found to differ between positive and negative emotions. Costa also found that sadness was more synchronized than happiness at each frequency band and was associated with a wider synchronization both between the right and left frontal sites and within the left hemisphere. In contrast, happiness was associated with a wider synchronization between the frontal and occipital sites.

(75) Different connectivity indices are sensitive to different characteristics of EEG signals. Correlation is sensitive to phase and polarity, but is independent of amplitudes. Changes in both amplitude and phase lead to a change in coherence (Guevara M A, Corsi-Cabrera M (1996) EEG coherence or EEG correlation? *Int J Psychophysiol* 23: 145-153). The phase synchronization index is only sensitive to a change in phase (Lachaux J P, Rodriguez E, Martinerie J, Varela F J (1999) Measuring phase synchrony in brain signals. *Hum Brain Mapp* 8: 194-208).

(76) A number of studies have tied to classify emotional states by means of recording and statistically analyzing EEG signals from the central nervous systems. See for example: Lin Y P, Wang C H, Jung T P, Wu T L, Jeng S K, et al. (2010) EEG-Based Emotion Recognition in Music Listening. *IEEE T Bio Med Eng* 57: 1798-1806 Murugappan M, Nagarajan R, Yaacob S (2010) Classification of human emotion from EEG using discrete wavelet transform. *J Biomed Sci Eng* 3: 390-396. Murugappan M, Nagarajan R, Yaacob S (2011) Combining Spatial Filtering and Wavelet Transform for Classifying Human Emotions Using EEG Signals. *J Med. Bio. Eng.* 31: 45-51. Berkman E, Wong D K, Guimaraes M P, Uy E T, Gross J J, et al. (2004) Brain wave recognition of emotions in EEG. *Psychophysiology* 41: S71-S71. Chanel G, Kronegg J, Grandjean D, Pun T (2006) Emotion assessment Arousal evaluation using EEG's and peripheral physiological signals. *Multimedia Content Representation, Classification and Security* 4105: 530-537. Hagiwara K (2003) A Feeling Estimation System Using a Simple Electroencephalograph. *IEEE International Conference on Systems, Man and Cybernetics*. 4204-4209. You-Yun Lee and Shulan Hsieh studied different emotional states by means of EEG-based functional connectivity patterns. They used emotional film clips to elicit three different emotional states.

(77) The dimensional theory of emotion, which asserts that there are neutral, positive, and negative emotional states, may be used to classify emotional states, because numerous studies have suggested that the responses of the central nervous system correlate with emotional valence and arousal. As suggested by Mauss and Robins (2009), "measures of emotional responding appear to be structured along dimensions (e.g., valence, arousal) rather than discrete emotional states (e.g., sadness, fear, anger)". See for example: Davidson R J (1993) Cerebral Asymmetry and Emotion—Conceptual and Methodological Conundrums. *Cognition Emotion* 7:115-138; Jones N A, Fox N A (1992) Electroencephalogram asymmetry during emotionally evocative films and its relation to positive and negative affectivity. *Brain Cogn* 20: 280-299; Schmidt L A, Trainor L J (2001) Frontal brain electrical activity (EEG) distinguishes valence and intensity of musical emotions. *Cognition Emotion* 15: 487-500; Tomarken A J, Davidson R J, Henriques J B (1990) Resting frontal brain asymmetry predicts affective responses to films. *J Pers Soc Psychol* 59: 791-801.)

(78) EEG-based functional connectivity change was found to be significantly different among emotional states of neutral, positive, or negative. Lee Y-Y, Hsieh S (2014) Classifying Different Emotional States by Means of EEG-Based Functional Connectivity Patterns. *PLoS ONE* 9(4): e95415. doi.org/10.1371/journal.pone.0095415. A connectivity pattern may be detected by pattern classification analysis using Quadratic Discriminant Analysis. The results indicated that the classification rate was better than chance. The authors found the following correlations:

(79) Theta band. Compared to neutral emotions, a significantly lower correlation at the frontal site and higher correlations at the temporal and occipital sites were found when watching negative films. No differences between a negative state and a positive state were found in the theta band. A significantly lower correlation was found in a positive state than in a neutral state at the frontal and parietal sites. A positive state showed higher correlations than a neutral state mainly at the temporal, parietal and occipital sites.

(80) Alpha band. A significantly higher correlation was found in a neutral state only in the case of F7-P7 activity. A negative state showed a significantly higher correlation than a positive state, especially at the parietal and occipital sites. A neutral state showed a lower correlation than a positive state mainly at the right temporal site.

(81) Beta band. No significant difference in correlation was observed among emotional states in the beta band.

(82) Gamma band. No significant difference in correlation was observed among emotional states in the gamma band.

(83) They concluded that estimating EEG-based functional connectivity provides a useful tool for studying the relationship between brain activity and emotional states.

(84) Emotions affect learning. Intelligent Tutoring Systems (ITS) learner model initially composed of a cognitive module was extended to include a psychological module and an emotional module. Alicia Heraz et al. introduced an emotional agent. It interacts with an ITS to communicate the emotional state of the learner based upon his mental state. The mental state was obtained from the learner's brainwaves. The agent learns to predict the learner's emotions by using ML techniques. (Alicia Heraz, Ryad Razaki; Claude Frasson, "Using machine learning to predict learner emotional state from brainwaves" *Advanced Learning Technologies*, 2007. *ICALT 2007*. Seventh IEEE International Conference on Advanced Learning Technologies (ICALT 2007)) See also: Ella T. Mampusti, Jose S. Ng, Jarren James I. Quinto, Grizelda L. Teng, Merlin Teodosia C. Suarez, Rhia S. Trogo, "Measuring Academic Affective States of Students via Brainwave Signals", *Knowledge and Systems Engineering (KSE) 2011 Third International Conference on*, pp. 226-231, 2011 Judith J. Azcarraga, John Francis Ibanez Jr., Ianne Robert Lim, Nestor Lumanas Jr., "Use of Personality Profile in Predicting Academic Emotion Based on Brainwaves Signals and Mouse Behavior", *Knowledge and Systems Engineering (KSE) 2011 Third International Conference on*, pp. 239-244, 2011. Yi-Hung Liu, Chien-Te Wu, Yung-Hwa Kao, Ya-Ting Chen, "Single-trial EEG-based emotion recognition using kernel Eigen-emotion pattern and adaptive support vector machine", *Engineering in Medicine and Biology Society (EMBC) 2013 35th Annual International Conference of the IEEE*, pp. 4306-4309, 2013, ISSN 1557-170X. Thong Tri Vo, Nam Phuong Nguyen, Toi Vo Van, *IFMBE Proceedings*, vol. 63, pp. 621, 2018, ISSN 1680-0737, ISBN 978-981-10-4360-4. Adrian Rodriguez Aguiñaga, Miguel Angel Lopez Ramirez, *Lecture Notes in Computer Science*, vol. 9456, pp. 177, 2015, ISSN 0302-9743, ISBN 978-3-319-26507-0. Judith Azcarraga, Merlin Teodosia Suarez, "Recognizing Student Emotions using Brainwaves and Mouse Behavior Data", *International Journal of Distance Education Technologies*, vol. 11, pp. 1, 2013, ISSN 1539-3100. Tri Thong Vo, Phuong Nam Nguyen, Van Toi Vo, *IFMBE Proceedings*, vol. 61, pp. 67, 2017, ISSN 1680-0737, ISBN 978-981-10-4219-5. Alicia Heraz, Claude Frasson, *Lecture Notes in Computer Science*, vol. 5535, pp. 367, 2009, ISSN 0302-9743, ISBN 978-3-642-02246-3. Hamwira Yaacob, Wahab Abdul, Norhaslinda Kamaruddin, "Classification of EEG signals using MLP based on categorical and dimensional perceptions of emotions", *Information and Communication Technology for the Muslim World (ICT4M) 2013 5th International Conference on*, pp. 1-6, 2013. Yuan-Pin Lin, Chi-Hong Wang, Tzyy-Ping Jung, Tien-Lin Wu, Shyh-Kang Jeng, Jeng-Ren Duann, Jyh-Hong Chen, "EEG-Based Emotion Recognition in Music Listening", *Biomedical Engineering IEEE Transactions on*, vol. 57, pp. 1798-1806, 2010, ISSN 0018-9294. Yi-Hung Liu, Wei-Teng Cheng, Yu-Tsung Hsiao, Chien-Te Wu, Mu-Der Jeng, "EEG-based emotion recognition based on kernel

(85) Using EEG to assess the emotional state has numerous practical applications. One of the first such applications was the development of a travel guide based on emotions by measuring brainwaves by the Singapore tourism group. "By studying the brainwaves of a family on vacation, the researchers drew up the Singapore Emotion Travel Guide, which advises future visitors of the emotions they can expect to experience at different attractions." (www.lonelyplanet.com/news/2017/04/12/singapore-emotion-travel-guide) Joel Pearson at University of New South Wales and his group developed the protocol of measuring brainwaves of travelers using EEG and decoding specific emotional states.

(86) Another recently released application pertains to virtual reality (VR) technology. On Sep. 18, 2017 Looxid Labs launched a technology that harnesses EEG from a subject wearing a VR headset Looxid Labs intention is to factor in brainwaves into VR applications in order to accurately infer emotions. Other products such as MindMaze and even Samsung have tied creating similar applications through facial muscles recognition. (scottamymx.com/2017/10/13/looxid-labs-vr-brain-waves-human-emotions/). According to its website (looxidlabs.com/device-2/), the Looxid Labs Development Kit provides a VR headset embedded with miniaturized eye and brain sensors. It uses 6 EEG channels: Fp1, Fp2, AF7, AFB, AF3, AF4 in international 10-20 system.

(87) To assess a user's state of mind, a computer may be used to analyze the EEG signals produced by the brain of the user. However, the emotional states of a brain are complex, and the brainwaves associated with specific emotions seem to change over time. Wei-Long Zheng at Shanghai Jiao Tong University used machine learning (ML) to identify the emotional brain states and to repeat it reliably. The ML algorithm found a set of patterns that clearly distinguished positive, negative, and neutral emotions that worked for different subjects and for the same subjects over time with an accuracy of about 80 percent (See Wei-Long Zheng, Jia-Yi Zhu, Bao-Liang Lu, Identifying Stable Patterns over Time for Emotion Recognition from EEG, arxiv.org/abs/1601.02197; see also How One Intelligent Machine Learned to Recognize Human Emotions, MIT Technology Review, Jan. 23, 2016.)

(88) MEG Magnetoencephalography (MEG) is a functional neuroimaging technique for mapping brain activity by recording magnetic fields produced by electrical currents occurring naturally in the brain, using very sensitive magnetometers. Arrays of SQUIDS (superconducting quantum interference devices) are currently the most common magnetometer, while the SERF (spin exchange relaxation-free) magnetometer is being investigated (Hämäläinen, Matti; Hari, Riitta; Ilmoniemi, Risto J; Knuutila, Jukka; Lounasmaa, Olli V. (1993). "Magnetoencephalography-theory, instrumentation, and applications to noninvasive studies of the working human brain". *Reviews of Modern Physics*. 65 (2): 413-497. ISSN 0034-6861. doi:10.1103/RevModPhys.65.413.) It is known that "neuronal activity causes local changes in cerebral blood flow, blood volume, and blood oxygenation" (Dynamic magnetic resonance imaging of human brain activity during primary sensory stimulation. K. K. Kwong, J. W. Belliveau, D. A. Chesler, I. E. Goldberg, R. M. Weisskoff, B. P. Poncelet D. N. Kennedy, B. E. Hoppel, M. S. Cohen, and R. Turner). Using "a 122-channel D.C. SQUID magnetometer with a helmet-shaped detector array covering the subject's head" it has been shown that the "system allows simultaneous recording of magnetic activity all over the head." (122-channel squid instrument for investigating the magnetic signals from the human brain.) A. I. Ahonen, M. S. Hämäläinen, M. J. Kajola, J. E. T. Knuutila, P. P. Laine, O. V. Lounasmaa, L. T. Parkkonen, J. T. Simola, and C. D. Tesche *Physica Scripta*, Volume 1993, T49A).

(89) In some cases, magnetic fields cancel, and thus the detectable electrical activity may fundamentally differ from the detectable electrical activity obtained via EEG. However, the main types of brain rhythms are detectable by both methods.

(90) See: U.S. Pat. Nos. 5,059,814; 5,118,606; 5,136,687; 5,224,203; 5,303,705; 5,325,862; 5,461,699; 5,522,863; 5,640,493; 5,715,821; 5,719,561; 5,722,418; 5,730,146; 5,736,543; 5,737,485; 5,747,492; 5,791,342; 5,816,247; 6,497,658; 6,510,340; 6,654,729; 6,893,407; 6,950,697; 8,135,957; 8,620,206; 8,644,754; 9,118,775; 9,179,875; 9,642,552; 20030018278; 20030171689; 20060293578; 20070156457; 20070259323; 20080015458; 20080154148; 20080229408; 20100010365; 20100076334; 20100090835; 20120046531; 20120052905; 20130041281; 20150081299; 20150262016. See EP1304073A2; EP1304073A3; WO2000025668A1; and WO2001087153A1.

(91) MEGs seek to detect the magnetic dipole emission from an electrical discharge in cells, e.g., neural action potentials. Typical sensors for MEGs are superconducting quantum interference devices (SQUIDS). These currently require cooling to liquid nitrogen or liquid helium temperatures. However, the development of room temperature, or near room temperature superconductors, and miniature cryocoolers, may permit field deployments and portable or mobile detectors. Because MEGs are less influenced by medium conductivity and dielectric properties, and because they inherently detect the magnetic field vector, MEG technology permits volumetric mapping of brain activity and distinction of complementary activity that might suppress detectable EEG signals. MEG technology also supports vector mapping of fields, since magnetic emitters are inherently dipoles, and therefore a larger amount of information is inherently available.

(92) See, U.S. Pat. Nos. 4,862,359; 5,027,817; 5,198,977; 5,230,346; 5,269,315; 5,309,923; 5,325,862; 5,331,970; 5,546,943; 5,568,816; 5,662,109; 5,724,987; 5,797,853; 5,840,040; 5,845,639; 6,042,548; 6,080,164; 6,088,611; 6,097,980; 6,144,872; 6,161,031; 6,171,239; 6,240,308; 6,241,686; 6,280,393; 6,309,361; 6,319,205; 6,322,515; 6,356,781; 6,370,414; 6,377,833; 6,385,479; 6,390,979; 6,402,689; 6,419,629; 6,466,816; 6,490,472; 6,526,297; 6,527,715; 6,530,884; 6,547,746; 6,551,243; 6,553,252; 6,622,036; 6,644,976; 6,648,880; 6,663,571; 6,684,098; 6,697,660; 6,728,564; 6,740,032; 6,743,167; 6,773,400; 6,907,280; 6,947,790; 6,950,698; 6,963,770; 6,963,771; 6,996,261; 7,010,340; 7,011,814; 7,022,083; 7,092,748; 7,104,947; 7,105,824; 7,120,486; 7,130,673; 7,171,252; 7,177,675; 7,231,245; 7,254,500; 7,283,861; 7,286,871; 7,338,455; 7,346,395; 7,378,056; 7,461,045; 7,489,964; 7,490,085; 7,499,745; 7,510,699; 7,539,528; 7,547,284; 7,565,193; 7,567,693; 7,577,472; 7,613,502; 7,627,370; 7,647,098; 7,653,433; 7,697,979; 7,729,755; 7,754,190; 7,756,568; 7,766,827; 7,769,431; 7,778,692; 7,787,937; 7,787,946; 7,787,946; 7,791,403; 7,831,305; 7,840,250; 7,856,264; 7,860,552; 7,899,524; 7,904,139; 7,904,144; 7,933,645; 7,962,204; 7,983,740; 7,986,991; 8,000,773; 8,000,793; 8,002,553; 8,014,847; 8,036,434; 8,065,360; 8,069,125; 8,086,296; 8,121,694; 8,190,248; 8,190,264; 8,197,437; 8,224,433; 8,233,682; 8,233,965; 8,236,038; 8,262,714; 8,280,514; 8,295,914; 8,306,607; 8,306,610; 8,313,441; 8,326,433; 8,337,404; 8,346,331; 8,346,342; 8,356,004; 8,358,818; 8,364,271; 8,380,289; 8,380,290; 8,380,314; 8,391,942; 8,391,956; 8,423,125; 8,425,583; 8,429,225; 8,445,851; 8,457,746; 8,467,878; 8,473,024; 8,498,708; 8,509,879; 8,527,035; 8,532,756; 8,538,513; 8,543,189; 8,554,325; 8,562,951; 8,571,629; 8,586,932; 8,591,419; 8,606,349; 8,606,356; 8,615,479; 8,626,264; 8,626,301; 8,632,750; 8,644,910; 8,655,817; 8,657,756; 8,666,478; 8,679,009; 8,684,926; 8,690,748; 8,696,722; 8,706,205; 8,706,241; 8,706,518; 8,712,512; 8,717,430; 8,725,669; 8,738,395; 8,761,869; 8,761,889; 8,768,022; 8,805,516; 8,814,923; 8,831,731; 8,834,546; 8,838,227; 8,849,392; 8,849,632; 8,852,103; 8,855,773; 8,858,440; 8,868,174; 8,888,702; 8,915,741; 8,918,162; 8,938,289; 8,938,290; 8,951,189; 8,951,192; 8,956,277; 8,965,513; 8,977,362; 8,989,836; 8,998,828; 9,005,126; 9,020,576; 9,022,936; 9,026,217; 9,026,218; 9,028,412; 9,033,884; 9,037,224; 9,042,201; 9,050,470; 9,067,052; 9,072,905; 9,084,896; 9,089,400; 9,089,683; 9,092,556; 9,095,266; 9,101,276; 9,107,595; 9,116,835; 9,133,024; 9,144,392; 9,149,255; 9,155,521; 9,167,970; 9,167,976; 9,167,977; 9,167,978; 9,171,366; 9,173,609; 9,179,850; 9,179,854; 9,179,858; 9,179,875; 9,192,300; 9,198,637; 9,198,707; 9,204,835; 9,211,077; 9,211,212; 9,213,074; 9,242,067; 9,247,890; 9,247,924; 9,248,288; 9,254,097; 9,254,383; 9,268,014; 9,268,015; 9,271,651; 9,271,674; 9,282,930; 9,289,143; 9,302,110; 9,308,372; 9,320,449; 9,322,895; 9,326,742; 9,332,939; 9,336,611; 9,339,227; 9,357,941; 9,367,131; 9,370,309; 9,375,145; 9,375,564; 9,387,320; 9,395,425; 9,402,558; 9,403,038; 9,414,029; 9,436,989; 9,440,064; 9,463,327; 9,470,728; 9,471,978; 9,474,852; 9,486,632; 9,492,313; 9,560,967; 9,579,048; 9,592,409; 9,597,493; 9,597,494; 9,615,789; 9,616,166; 9,655,573; 9,655,669; 9,662,049; 9,662,492; 9,669,185; 9,675,292; 9,682,232; 9,687,187; 9,707,396; 9,713,433; 9,713,444; 20010020127; 20010021800; 20010051774; 20020005784; 20020016552; 20020017994; 20020042563; 20020058867; 20020099273; 20020099295; 20020103428; 20020103429; 20020128638; 20030001098; 20030009096; 20030013981; 20030032870; 20030040660; 20030068605; 20030074032; 20030093004; 20030093005; 20030120140; 20030128801; 20030135128; 20030153818; 20030163027; 20030163028; 20030181821; 20030187359; 20030204135; 20030225335; 20030236458; 20040030585; 20040059241; 20040072133; 20040077960; 20040092809; 20040096395; 20040097802; 20040116798; 20040122787; 20040122790; 20040144925; 20040204656; 20050004489; 20050007091;

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(93) EEGs and MEGs can monitor the state of consciousness. For example, states of deep sleep are associated with slower EEG oscillations of larger amplitude. Various signal analysis methods allow for robust identifications of distinct sleep stages, depth of anesthesia, epileptic seizures and connections to detailed cognitive events.

(94) Positron Emission Tomography (PET) Scan A PET scan is an imaging test that helps reveal how tissues and organs are functioning (Bailey, D. L.; D. W. Townsend; P. E. Valk; M. N. Maisey (2005). *Positron Emission Tomography: Basic Sciences*. Secaucus, NJ: Springer-Verlag. ISBN 1-85233-798-2.). A PET scan uses a radioactive drug (positron-emitting tracer) to show this activity. It uses this radiation to produce 3-D, images colored for the different activity of the brain. See, e.g.: Jarden, Jens O., Vijay Dhawan, Alexander Poltorak, Jerome B. Posner, and David A. Rottenberg. "Positron emission tomographic measurement of blood-to-brain and blood-to-tumor transport of 82Rb: The effect of dexamethasone and whole-brain radiation therapy." *Annals of neurology* 18, no. 6 (1985): 636-646. Dhawan, V. I. J. A. Y., A. Poltorak, J. R. Moeller, J. O. Jarden, S. C. Strother, H. Thaler, and D. A. Rottenberg. "Positron emission tomographic measurement of blood-to-brain and blood-to-tumour transport of 82Rb. I: Error analysis and computer simulations." *Physics in medicine and biology* 34, no. 12 (1989):1773.

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(96) fMRI Functional magnetic resonance imaging or functional MRI (fMRI) is a functional neuroimaging procedure using MRI technology that measures brain activity by detecting changes associated with blood flow ("Magnetic Resonance, a critical peer-reviewed introduction; functional MRI". European Magnetic Resonance Forum. Retrieved 17 Nov. 2014; Huettel, Song & McCarthy (2009)).

(97) Yukiyasu Kamitani et al., *Neuron* (DOI: 10.1016/j.neuron.2008.11.004) used an image of brain activity taken in a functional MRI scanner to recreate a black-and-white image from scratch. See also 'Mind-reading' software could record your dreams" By Celeste Biever. *New Scientist* 12 Dec. 2008. (www.newscientist.com/article/dn16267-mind-reading-software-could-record-your-dreams/)

(98) See, U.S. Pat. Nos. 6,622,036; 7,120,486; 7,177,675; 7,209,788; 7,489,964; 7,697,979; 7,754,190; 7,856,264; 7,873,411; 7,962,204; 8,060,181; 8,224,433; 8,315,962; 8,320,649; 8,326,433; 8,356,004; 8,380,314; 8,386,312; 8,392,253; 8,532,756; 8,562,951; 8,626,264; 8,632,750; 8,655,817;

8,679,009; 8,684,742; 8,684,926; 8,698,639; 8,706,241; 8,725,669; 8,831,731; 8,849,632; 8,855,773; 8,868,174; 8,915,871; 8,918,162; 8,939,903; 8,951,189; 8,951,192; 9,026,217; 9,037,224; 9,042,201; 9,050,470; 9,072,905; 9,084,896; 9,095,266; 9,101,276; 9,101,279; 9,135,221; 9,161,715; 9,192,300; 9,230,065; 9,248,286; 9,248,288; 9,265,458; 9,265,974; 9,292,471; 9,296,382; 9,302,110; 9,308,372; 9,345,412; 9,367,131; 9,420,970; 9,440,646; 9,451,899; 9,454,646; 9,463,327; 9,468,541; 9,474,481; 9,475,502; 9,489,854; 9,505,402; 9,538,948; 9,579,247; 9,579,457; 9,615,746; 9,693,724; 9,693,734; 9,694,155; 9,713,433; 9,713,444; 20030093129; 20030135128; 20040059241; 20050131311; 20050240253; 20060015034; 20060074822; 20060129324; 20060161218; 20060167564; 20060189899; 20060241718; 20070179534; 20070244387; 20080009772; 20080091118; 20080125669; 20080228239; 20090006001; 20090009284; 20090030930; 20090062676; 20090062679; 20090082829; 20090132275; 20090137923; 20090157662; 20090164132; 20090209845; 20090216091; 20090220429; 20090270754; 20090287271; 20090287272; 20090287273; 20090287467; 20090290767; 20090292713; 20090297000; 20090312808; 20090312817; 20090312998; 20090318773; 20090326604; 20090327068; 20100036233; 20100049276; 20100076274; 20100094154; 20100143256; 20100145215; 20100191124; 20100298735; 20110004412; 20110028827; 20110034821; 20110092882; 20110106750; 20110119212; 20110256520; 20110306845; 20110306846; 20110313268; 20110313487; 20120035428; 20120035765; 20120052469; 20120060851; 20120083668; 20120108909; 20120165696; 20120203725; 20120212353; 20120226185; 20120253219; 20120265267; 20120271376; 20120296569; 20130031038; 20130063550; 20130080127; 20130085678; 20130130799; 20130131755; 20130185145; 20130185145; 20130185145; 20130185145; 20130185145; 20130185145; 20130226408; 20130245886; 20130253363; 20130338803; 20140058528; 20140114889; 20140135642; 20140142654; 20140154650; 20140163328; 20140163409; 20140171757; 20140200414; 20140200432; 20140211593; 20140214335; 20140243652; 20140276549; 20140279746; 20140309881; 20140315169; 20140347265; 20140371984; 20150024356; 20150029087; 20150033245; 20150033258; 20150033259; 20150033262; 20150033266; 20150038812; 20150080753; 20150094962; 20150112899; 20150119658; 20150164431; 20150174362; 20150174418; 20150196800; 20150227702; 20150248470; 20150257700; 20150290453; 20150290454; 20150297893; 20150305685; 20150324692; 20150327813; 20150339363; 20150343242; 20150351655; 20150359431; 20150360039; 20150366482; 20160015307; 20160027342; 20160031479; 20160038049; 20160048659; 20160051161; 20160051162; 20160055304; 20160107653; 20160120437; 20160144175; 20160152233; 20160158553; 20160206380; 20160213276; 20160226280; 20160226318; 20160302711; 20160306942; 20160324457; 20160357256; 20160366462; 20170027812; 20170031440; 20170032098; 20170042474; 20170043160; 20170043167; 20170061034; 20170065349; 20170085547; 20170086727; 20170087302; 20170091418; 20170113046; 20170188876; 20170196501; 20170202476; 20170202518; and 20170206913.

(99) Functional Near Infrared Spectroscopy (fNIRS) fNIR is a non-invasive imaging method involving the quantification of chromophore concentration resolved from the measurement of near infrared (NIR) light attenuation or temporal or phasic changes. NIR spectrum light takes advantage of the optical window in which skin, tissue, and bone are mostly transparent to NIR light in the spectrum of 700-900 nm, while hemoglobin (Hb) and deoxygenated-hemoglobin (deoxy-Hb) are stronger absorbers of light. Differences in the absorption spectra of deoxy-Hb and oxy-Hb allow the measurement of relative changes in hemoglobin concentration through the use of light attenuation at multiple wavelengths. Two or more wavelengths are selected, with one wavelength above and one below the isosbestic point of 810 nm at which deoxy-Hb and oxy-Hb have identical absorption coefficients. Using the modified Beer-Lambert law (mBLL), relative concentration can be calculated as a function of total photon path length. Typically, the light emitter and detector are placed ipsilaterally on the subject's skull so recorded measurements are due to back-scattered (reflected) light following elliptical pathways. The use of fNIR as a functional imaging method relies on the principle of neuro-vascular coupling also known as the hemodynamic response or blood-oxygen-level dependent (BOLD) response. This principle also forms the core of fMRI techniques. Through neuro-vascular coupling, neuronal activity is linked to related changes in localized cerebral blood flow. fNIR and fMRI are sensitive to similar physiologic changes and are often comparative methods. Studies relating fMRI and fNIR show highly correlated results in cognitive tasks. fNIR has several advantages in cost and portability over fMRI, but cannot be used to measure cortical activity more than 4 cm deep due to limitations in light emitter power and has more limited spatial resolution. fNIR includes the use of diffuse optical tomography (DOT/NIRDOT) for functional purposes. Multiplexing fNIRS channels can allow 2D topographic functional maps of brain activity (e.g. with Hitachi ETG-4000 or Minis Oxymon) while using multiple emitter spacings may be used to build 3D tomographic maps.

(100) Beste Yuksel and Robert Jacob, Brain Automated Chorales (BACH), ACM CHI 2016, DOI: 10.1145/2858036.2858388, provides a system that helps beginners learn to play Bach chorales on piano by measuring how hard their brains are working. This is accomplished by estimating the brain's workload using functional Near-Infrared Spectroscopy (fNIRS), a technique that measures oxygen levels in the brain—in this case in the prefrontal cortex. A brain that's working hard pulls in more oxygen. Sensors strapped to the player's forehead talk to a computer, which delivers the new music, one line at a time. See also "Mind-reading tech helps beginners quickly learn to play Bach." By Anna Nowogrodzki, New Scientist 9 Feb. 2016 available online at www.newscientist.com/article/2076899-mind-reading-tech-helps-beginners-quickly-learn-to-play-bach/.

(101) LORETA Low-resolution brain electromagnetic tomography often referred as LORETA is a functional imaging technology usually using a linearly constrained minimum variance vector beamformer in the time-frequency domain as described in Gross et al., "Dynamic imaging of coherent sources: Studying neural interactions in the human brain," PNAS 98, 694-699, 2001. It allows to the image (mostly 3D) evoked and induced oscillatory activity in a variable time-frequency range, where time is taken relative to a triggered event. There are three categories of imaging related to the technique used for LORETA. See, wiki.besa.de/index.php?title=Source_Analysis_3D_Imaging#Multiple_Source_Beamformer_.28MSBF.29. The Multiple Source Beamformer (MSBF) is a tool for imaging brain activity. It is applied in the time-frequency domain and based on single-trial data. Therefore, it can image not only evoked, but also induced activity, which is not visible in time-domain averages of the data. Dynamic Imaging of Coherent Sources (Dios) can find coherence between any two pairs of voxels in the brain or between an external source and brain voxels. DICS requires time-frequency-transformed data and can find coherence for evoked and induced activity. The following imaging methods provides an image of brain activity based on a distributed multiple source model: CLARA is an iterative application of LORETA images, focusing the obtained 3D image in each iteration step. LAURA uses a spatial weighting function that has the form of a local autoregressive function. LORETA has the 3D Laplacian operator implemented as spatial weighting prior. sLORETA is an unweighted minimum norm that is standardized by the resolution matrix. swLORETA is equivalent to sLORETA, except for an additional depth weighting. SSLOFO is an iterative application of standardized minimum norm images with consecutive shrinkage of the source space. A User-defined volume image allows experimenting with the different imaging techniques. It is possible to specify user-defined parameters for the family of distributed source images to create a new imaging technique. If no individual MRI is available, the minimum norm image is displayed on a standard brain surface and computed for standard source locations. If available, an individual brain surface is used to construct the distributed source model and to image the brain activity. Unlike classical LORETA, cortical LORETA is not computed in a 3D volume, but on the cortical surface. Unlike classical CLARA, cortical CLARA is not computed in a 3D volume, but on the cortical surface. The Multiple Source Probe Scan (MSPS) is a tool for the validation of a discrete multiple source model. The Source Sensitivity image displays the sensitivity of a selected source in the current discrete source model and is, therefore, data independent.

(102) See U.S. Pat. Nos. 4,562,540; 4,594,662; 5,650,726; 5,859,533; 6,026,173; 6,182,013; 6,294,917; 6,332,087; 6,393,363; 6,534,986; 6,703,838; 6,791,331; 6,856,830; 6,863,127; 7,030,617; 7,092,748; 7,119,553; 7,170,294; 7,239,731; 7,276,916; 7,286,871; 7,295,019; 7,353,065; 7,363,164; 7,454,243; 7,499,894; 7,648,498; 7,804,441; 7,809,434; 7,841,986; 7,852,087; 7,937,222; 8,000,795; 8,046,076; 8,131,526; 8,174,430; 8,188,749; 8,244,341; 8,263,574; 8,332,191; 8,346,365; 8,362,780; 8,456,166; 8,538,700; 8,565,883; 8,593,154; 8,600,513; 8,706,205; 8,711,655; 8,731,987; 8,756,017; 8,761,438; 8,812,237; 8,829,908; 8,958,882; 9,008,970; 9,035,657; 9,069,097; 9,072,449; 9,091,785; 9,092,895; 9,121,964; 9,133,709; 9,165,472; 9,179,854; 9,320,451; 9,367,738; 9,414,749; 9,414,763; 9,414,764; 9,442,088; 9,468,541; 9,513,398; 9,545,225; 9,557,439; 9,562,988; 9,568,635; 9,651,706; 9,675,254; 9,675,255; 9,675,292; 9,713,433; 9,715,032; 20020000808; 20020017905; 20030018277; 20030093004; 20040097802; 20040116798; 20040131998; 20040140811; 20040145370; 20050156602; 20060058856; 20060069059; 20060136135;

20060149160; 20060152227; 20060170424; 20060176062; 20060184058; 20070060974; 20070159185; 20070191727; 20080033513; 20080097235; 20080125830; 20080125831; 20080183072; 20080242976; 20080255816; 20080281667; 20090039889; 20090054801; 20090082688; 20090099783; 20090216146; 20090261832; 20090306534; 20090312663; 20100010366; 20100030097; 20100042011; 20100056276; 20100092934; 20100132448; 20100134113; 20100168053; 20100198519; 20100231221; 20100238763; 20110004115; 20110050232; 20110160607; 20110308789; 20120010493; 20120011927; 20120016430; 20120083690; 20120130641; 20120150257; 20120162002; 20120215448; 20120245474; 20120268272; 20120269385; 20120296569; 20130091941; 20130096408; 20130141103; 20130231709; 20130289385; 20130303934; 20140015852; 20140025133; 20140058528; 20140066739; 20140107519; 20140128763; 20140155740; 20140161352; 20140163328; 20140163893; 20140228702; 20140243714; 20140275944; 20140276012; 20140323899; 20150051663; 20150112409; 20150119689; 20150137817; 20150145519; 20150157235; 20150167459; 20150177413; 20150248615; 20150257648; 20150257649; 20150301218; 20150342472; 20160002523; 20160038049; 20160040514; 20160051161; 20160051162; 20160091448; 20160102500; 20160120436; 20160136427; 20160187524; 20160213276; 20160220821; 20160223703; 20160235983; 20160245952; 20160256109; 20160259085; 20160262623; 20160298449; 20160334534; 20160345856; 20160356911; 20160367812; 20170001016; 20170067323; 20170138132; and 20170151436.

(103) Neurofeedback Neurofeedback (NFB), also called neurotherapy or neurobiofeedback, is a type of biofeedback that uses real-time displays of brain activity-most commonly electroencephalography (EEG), to teach self-regulation of brain function. Typically, sensors are placed on the scalp to measure activity, with measurements displayed using video displays or sound. The feedback may be in various other forms as well. Typically, the feedback is sought to be presented through primary sensory inputs, but this is not a limitation on the technique.

(104) The applications of neurofeedback to enhance performance extend to the arts in fields such as music, dance, and acting. A study with conservatoire musicians found that alpha-theta training benefitted the three music domains of musicality, communication, and technique. Historically, alpha-theta training, a form of neurofeedback, was created to assist creativity by inducing hypnagogia, a “borderline waking state associated with creative insights”, through facilitation of neural connectivity. Alpha-theta training has also been shown to improve novice singing in children. Alpha-theta neurofeedback, in conjunction with heart rate variability training, a form of biofeedback, has also produced benefits in dance by enhancing performance in competitive ballroom dancing and increasing cognitive creativity in contemporary dancers. Additionally, neurofeedback has also been shown to instill a superior flow state in actors, possibly due to greater immersion while performing.

(105) Several studies of brain wave activity in experts while performing a task related to their respective area of expertise revealed certain characteristic telltale signs of so-called “flow” associated with top-flight performance. Mihaly Csikszentmihalyi (University of Chicago) found that the most skilled chess players showed less EEG activity in the prefrontal cortex, which is typically associated with higher cognitive processes such as working memory and verbalization, during a game.

(106) Chris Berka et al., *Advanced Brain Monitoring*, Carlsbad, California, The International J. Sport and Society, vol 1, p 87, looked at the brainwaves of Olympic archers and professional golfers. A few seconds before the archers fired off an arrow or the golfers hit the ball, the team spotted a small increase in alpha band patterns. This may correspond to the contingent negative variation observed in evoked potential studies, and the Bereitschaftspotential or BP (from German, “readiness potential”), also called the pre-motor potential or readiness potential (RP), a measure of activity in the motor cortex and supplementary motor area of the brain leading up to voluntary muscle movement. Berka also trained novice marksmen using neurofeedback. Each person was hooked up to electrodes that tease out and display specific brainwaves, along with a monitor that measured their heartbeat. By controlling their breathing and learning to deliberately manipulate the waveforms on the screen in front of them, the novices managed to produce the alpha waves characteristic of the flow state. This, in turn, helped them improve their accuracy at hitting the targets.

(107) Low Energy Neurofeedback System (LENS) The LENS, or Low Energy Neurofeedback System, uses a very low power electromagnetic field, to carry feedback to the person receiving it. The feedback travels down the same wires carrying the brainwaves to the amplifier and computer. Although the feedback signal is weak, it produces a measurable change in the brainwaves without conscious effort from the individual receiving the feedback. The system is software controlled, to receive input from EEG electrodes, to control the stimulation. Through the scalp. Neurofeedback uses a feedback frequency that is different from, but correlates with, the dominant brainwave frequency. When exposed to this feedback frequency, the EEG amplitude distribution changes in power. Most of the time the brainwaves reduce in power, but at times they also increase in power. In either case the result is a changed brainwave state, and much greater ability for the brain to regulate itself.

(108) Content-Based Brainwave Analysis Memories are not unique. Janice Chen, *Nature Neuroscience*, DOI: 10.1038/nn.4450, showed that when people describe the episode from Sherlock Holmes drama, their brain activity patterns were almost exactly the same as each other's, for each scene. Moreover, there's also evidence that, when a person tells someone else about it, they implant that same activity into their brain as well. Moreover, research in which people who have not seen a movie listen to someone else's description of it. Chen et al. have found that the listener's brain activity looks much like that of the person who has seen it. See also “Our brains record and remember things in exactly the same way” by Andy Coghlan, *New Scientist* Dec. 5, 2016 (www.newscientist.com/article/2115093-our-brains-record-and-remember-things-in-exactly-the-same-way/)

(109) Brian Pasley, *Frontiers in Neuroengineering*, doi.org/whb, developed a technique for reading thoughts. The team hypothesized that hearing speech and thinking to oneself might spark some of the same neural signatures in the brain. They supposed that an algorithm trained to identify speech heard out loud might also be able to identify words that are thought. In the experiment, the decoder trained on speech was able to reconstruct which words several of the volunteers were thinking, using neural activity alone. See also “Hearing our inner voice” by Helen Thomson, *New Scientist* Oct. 29, 2014 (www.newscientist.com/article/mg22429934-000-brain-decoder-can-eavesdrop-on-your-inner-voice/)

(110) Jack Gallant et al. were able to detect which of a set of images someone was looking at from a brain scan, using software that compared the subjects' brain activity while looking at an image with that captured while they were looking at “training” photographs. The program then picked the most likely match from a set of previously unseen pictures.

(111) Ann Graybiel and Mark Howe used electrodes to analyze brainwaves in the ventromedial striatum of rats while they were taught to navigate a maze. As rats were learning the task, their brain activity showed bursts of fast gamma waves. Once the rats mastered the task, their brainwaves slowed to almost a quarter of their initial frequency, becoming beta waves. Graybiel's team posited that this transition reflects when learning becomes a habit.

(112) Bernard Balleine, *Proceedings of the National Academy of Sciences*, DOI: 10.1073/pnas.1113158108. See also “Habits form when brainwaves slow down” by Wendy Zukerman, *New Scientist* Sep. 26, 2011 (www.newscientist.com/article/dn20964-habits-form-when-brainwaves-slow-down/) posits that the slower brainwaves may be the brain weeding out excess activity to refine behavior. He suggests it might be possible to boost the rate at which they learn a skill by enhancing such beta-wave activity.

(113) U.S. Pat. No. 9,763,592 provides a system for instructing a user behavior change comprising: collecting and analyzing bioelectrical signal datasets; and providing a behavior change suggestion based upon the analysis. A stimulus may be provided to prompt an action by the user, which may be visual, auditory, or haptic. See also U.S. Pat. No. 9,622,660, 20170041699; 20130317384; 20130317382; 20130314243; 20070173733; and 20070066914.

(114) The chess game is a good example of a cognitive task which needs a lot of training and experience. A number of EEG studies have been done on chess players. Pawel Stepień, Włodzimierz Klonowski and Nikolay Suvorov, *Nonlinear analysis of EEG in chess players*, *EPJ Nonlinear Biomedical Physics* 20153:1, showed better applicability of Higuchi Fractal Dimension method for analysis of EEG signals related to chess tasks than that of Sliding Window Empirical Mode Decomposition. The paper shows that the EEG signal during the game is more complex, non-linear, and non-stationary even when there are no significant differences between the game and relaxed state in the contribution of different EEG bands to total power of the signal. There is the need of gathering more data from more chess experts and of comparing them with data from novice chess

players. See also Junior, L. R. S., Cesar, F. H. G., Rocha, F. T., and Thomaz, C. E. EEG and Eye Movement Maps of Chess Players. *Proceedings of the Sixth International Conference on Pattern Recognition Applications and Methods. (ICPRAM 2017)* pp. 343-441. (fei.edu.br/~cet/icpram17_LaercioJunior.pdf).

(115) Estimating EEG-based functional connectivity provides a useful tool for studying the relationship between brain activity and emotional states. See You-Yun Lee, Shulan Hsieh. *Classifying Different Emotional States by Means of EEG-Based Functional Connectivity Patterns*. Apr. 17, 2014, (doi.org/10.1371/journal.pone.0095415), which aimed to classify different emotional states by means of EEG-based functional connectivity patterns, and showed that the EEG-based functional connectivity change was significantly different among emotional states. Furthermore, the connectivity pattern was detected by pattern classification analysis using Quadratic Discriminant Analysis. The results indicated that the classification rate was better than chance. Estimating EEG-based functional connectivity provides a useful tool for studying the relationship between brain activity and emotional states.

(116) Neuromodulation/Neuroenhancement Neuromodulation is the alteration of nerve activity through targeted delivery of a stimulus, such as electrical stimulation or chemical agents, to specific neurological sites in the body. It is carried out to normalize—or modulate—nervous tissue function. Neuromodulation is an evolving therapy that can involve a range of electromagnetic stimuli such as a magnetic field (TMS, rTMS), an electric current (TES, e.g., tDCS, HD-tDCS, tACS, osc-tDCS, electrosleep), or a drug instilled directly in the subdural space (intrathecal drug delivery). Emerging applications involve targeted introduction of genes or gene regulators and light (optogenetics). The most clinical experience has been with electrical stimulation. Neuromodulation, whether electrical or magnetic, employs the body's natural biological response by stimulating nerve cell activity that can influence populations of nerves by releasing transmitters, such as dopamine, or other chemical messengers such as the peptide Substance P, that can modulate the excitability and firing patterns of neural circuits. There may also be more direct electrophysiological effects on neural membranes. According to some applications, the end effect is a “normalization” of a neural network function from its perturbed state. Presumed mechanisms of action for neurostimulation include depolarizing blockade, stochastic normalization of neural firing, axonal blockade, reduction of neural firing keratosis, and suppression of neural network oscillations. Although the exact mechanisms of neurostimulation are not entirely clear, the empirical effectiveness has led to considerable application clinically.

(117) Neuroenhancement refers to the targeted enhancement and extension of cognitive, affective, and motor abilities based on an understanding of their underlying neurobiology in healthy persons who do not have any mental illness. As such, it can be thought of as an umbrella term that encompasses pharmacological and non-pharmacological methods of improving cognitive, affective, and motor functionality. Critically, for any agent to qualify as a neuroenhancer, it must reliably engender substantial cognitive, affective, or motor benefits beyond normal functioning in healthy individuals (or in select groups of individuals having pathology), while causing few side effects: at most at the level of commonly used comparable legal substances or activities, such as caffeine, alcohol, and sleep-deprivation. Pharmacological neuroenhancement agents include the well-validated nootropics, such as racetam, vinpocetine, and phosphatidylserine, as well as other drugs used for treating patients suffering from neurological disorders. Non-pharmacological measures include non-invasive brain stimulation, which has been employed to improve various cognitive and affective functions, and brain-machine interfaces, which hold much potential to extend the repertoire of motor and cognitive actions available to humans.

(118) Brain Stimulation The entrainment hypothesis, suggests the possibility of inducing a particular oscillation frequency in the brain using an external oscillatory force (e.g., rTMS, tDCS, tACS, binaural beats, isochronic tones, light stimulation). The physiological basis of oscillatory cortical activity lies in the timing of the interacting neurons; when groups of neurons synchronize their firing activities, brain rhythms emerge, network oscillations are generated, and the basis for interactions between brain areas may develop. Synchronization of spatially separated lobes of the brain may also play a role. Because of the variety of experimental protocols for brain stimulation, limits on descriptions of the actual protocols employed, and limited controls, consistency of reported studies is lacking, and extrapolability is limited. Thus, while there is some consensus in various aspects of the effects of extra cranial brain stimulation, the results achieved have a degree of uncertainty dependent on details of implementation. On the other hand, within a specific experimental protocol, it is possible to obtain statistically significant and repeatable results. This implies that feedback control might be effective to control implementation of the stimulation for a given purpose; however, prior studies that employ feedback control are lacking.

(119) Changes in the neuronal threshold result from changes in membrane permeability (Liebetanz et al., 2002), which influence the response of the task-related network. The same mechanism of action may be responsible for both TES methods and TMS, i.e., the induction of noise in the system. However, the neural activity induced by TES will be highly influenced by the state of the system because it is a neuromodulatory method (Paulus, 2011), and its effect will depend on the activity of the stimulated area. Therefore, the final result will depend strongly on the task characteristics, the system state and the way in which TES will interact with such a state.

(120) In TMS, the magnetic pulse causes a rapid increase in current flow, which can in some cases cause and above-threshold depolarization of cell membranes affected by the current, triggering an action potential, and leading to the trans-synaptic depolarization or hyperpolarization of connected cortical neurons, depending on their natural response to the firing of the stimulated neuron(s). Therefore, TMS activates a neural population that, depending on several factors, can be congruent (facilitate) or incongruent (inhibit) with task execution. TES induces a polarization of cortical neurons at a subthreshold level that is too weak to evoke an action potential. However, by inducing a polarity shift in the intrinsic neuronal excitability, TES can alter the spontaneous firing rate of neurons and modulate the response to afferent signals. In this sense, TES-induced effects are even more bound to the state of the stimulated area that is determined by the conditions. In short NIBS leads to a stimulation-induced modulation of the state that can be substantially defined as noise induction. Induced noise will not be just random activity, but will depend on the interaction of many parameters, from the characteristics of the stimulation to the state.

(121) The noise induced by NIBS will be influenced by the state of the neural population of the stimulated area. Although the types and number of neurons “triggered” by NIBS are theoretically random, the induced change in neuronal activity is likely to be correlated with ongoing activity, yet even if we are referring to a non-deterministic process, the noise introduced will not be a totally random element. Because it will be partially determined by the experimental variables, the level of noise that will be introduced by the stimulation and by the context can be estimated, as well as the interaction between the two levels of noise (stimulation and context). Although, HD-tDCS made a significantly more focused spatial application of TES possible, generally, known transcranial stimulation does not permit stimulation with a focused and highly targeted signal to a clearly defined area of the brain to establish a unique brain-behavior relationship; therefore, the known introduced stimulus activity in the brain stimulation is ‘noise.’

(122) Cosmetic neuroscience has emerged as a new field of research. Roy Hamilton, Samuel Messing, and Anjan Chatterjee, “Rethinking the thinking cap—Ethics of neural enhancement using noninvasive brain stimulation.” *Neurology*, Jan. 11, 2011, vol. 76 no. 2 187-193.

(www.neurology.org/content/76/2/187.) discuss the use noninvasive brain stimulation techniques such as transcranial magnetic stimulation and transcranial direct current stimulation to enhance neurologic function: cognitive skills, mood, and social cognition.

(123) Electrical brain stimulation (EBS), also known as, focal brain stimulation (FBS), is a form of clinical neurobiology electrotherapy used to stimulate a neuron or neural network in the brain through the direct or indirect excitation of cell membranes using an electric current. See: en.wikipedia.org/wiki/Electrical_brain_stimulation; U.S. Pat. Nos. 7,753,836; 7,946,73; 8,545,378; 9,345,901; 9,610,456; 9,694,178; 20140330337; 20150112403; and 20150119689.

(124) Motor skills can be affected by CNS stimulation.

(125) See, U.S. Pat. Nos. 5,343,871; 5,742,748; 6,057,846; 6,390,979; 6,644,976; 6,656,137; 7,063,535; 7,558,622; 7,618,381; 7,733,224; 7,829,562; 7,863,272; 8,016,597; 8,065,240; 8,069,125; 8,108,036; 8,126,542; 8,150,796; 8,195,593; 8,356,004; 8,449,471; 8,461,988; 8,525,673; 8,525,687;

8,591,291; 8,591,419; 8,606,592; 8,615,479; 8,680,991; 8,682,449; 8,706,518; 8,747,336; 8,750,971; 8,764,651; 8,784,109; 8,858,440; 8,862,236; 8,938,289; 8,962,042; 9,005,649; 9,064,036; 9,107,586; 9,125,788; 9,138,579; 9,149,599; 9,173,582; 9,204,796; 9,211,077; 9,265,458; 9,351,640; 9,358,361; 9,380,976; 9,403,038; 9,418,368; 9,468,541; 9,495,684; 9,545,515; 9,549,691; 9,560,967; 9,577,992; 9,590,986; 20030068605; 20040072133; 20050020483; 20050032827; 20050059689; 20050153268; 20060014753; 20060052386; 20060106326; 20060191543; 20060229164; 20070031798; 20070138886; 20070276270; 20080001735; 20080004904; 20080243005; 20080287821; 20080294019; 20090005654; 20090018407; 20090024050; 20090118593; 20090119154; 20090132275; 20090156907; 20090156955; 20090157323; 20090157481; 20090157482; 20090157625; 20090157660; 20090157751; 20090157813; 20090163777; 20090164131; 20090164132; 20090164302; 20090164401; 20090164403; 20090164458; 20090164503; 20090164549; 20090171164; 20090172540; 20090221928; 20090267758; 20090271011; 20090271120; 20090271347; 20090312595; 20090312668; 20090318773; 20090318779; 20090319002; 20100004762; 20100015583; 20100017001; 20100022820; 20100041958; 20100042578; 20100063368; 20100069724; 20100076249; 20100081860; 20100081861; 20100100036; 20100125561; 20100130811; 20100145219; 20100163027; 20100163035; 20100168525; 20100168602; 20100280332; 20110015209; 20110015469; 20110082154; 20110105859; 20110115624; 20110152284; 20110178441; 20110181422; 20110288119; 20120092156; 20120092157; 20120130300; 20120143104; 20120164613; 20120177716; 20120316793; 20120330109; 20130009783; 20130018592; 20130034837; 20130053656; 20130054215; 20130085678; 20130121984; 20130132029; 20130137717; 20130144537; 20130184728; 20130184997; 20130211291; 20130231574; 20130281890; 20130289385; 20130330428; 20140039571; 20140058528; 20140077946; 20140094720; 20140104059; 20140148479; 20140155430; 20140163425; 20140207224; 20140235965; 20140249429; 20150025410; 20150025422; 20150068069; 20150071907; 20150141773; 20150208982; 20150265583; 20150290419; 20150294067; 20150294085; 20150294086; 20150359467; 20150379878; 20160001096; 20160007904; 20160007915; 20160030749; 20160030750; 20160067492; 20160074657; 20160120437; 20160140834; 20160198968; 20160206671; 20160220821; 20160303402; 20160351069; 20160360965; 20170046971; 20170065638; 20170080320; 20170084187; 20170086672; 20170112947; 20170127727; 20170131293; 20170143966; 20170151436; 20170157343; and 20170193831. 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(126) Michael A. Nitsche, and Armin Kibele. "Noninvasive brain stimulation and neural entrainment enhance athletic performance—a review." *J. Cognitive Enhancement* 1.1 (2017): 73-79, discusses that non-invasive brain stimulation (NIBS) bypasses the correlative approaches of other imaging techniques, making it possible to establish a causal relationship between cognitive processes and the functioning of specific brain areas. NIBS can provide information about where a particular process occurs. NIBS offers the opportunity to study brain mechanisms beyond process localization, providing information about when activity in a given brain region is involved in a cognitive process, and even how it is involved. When using NIBS to explore cognitive processes, it is important to understand not only how NIBS functions but also the functioning of the neural structures themselves. Non-invasive brain stimulation (NIBS) methods, which include transcranial magnetic stimulation (TMS) and transcranial electric stimulation (tES), are used in cognitive neuroscience to induce transient changes in brain activity and thereby alter the behavior of the subject. The application of NIBS aims at establishing the role of a given cortical area in an ongoing specific motor, perceptual or cognitive process (Hallett, 2000; Walsh and Cowey, 2000). Physically, NIBS techniques affect neuronal states through different mechanisms. In TMS, a solenoid (coil) is used to deliver a strong and transient magnetic field, or "pulse," to induce a transitory electric current at the cortical surface beneath the coil. (US 2004078056) The pulse causes the rapid and above-threshold depolarization of cell membranes affected by the current (Barker et al., 1985, 1987), followed by the transsynaptic depolarization or hyperpolarization of interconnected neurons. Therefore, TMS induces a current that elicits action potentials in neurons. A complex set of coils can deliver a complex 3D excitation field. By contrast, in tES techniques, the stimulation involves the application of weak electrical currents directly to the scalp through a pair of electrodes (Nitsche and Paulus, 2000; Priori et al., 1998). As a result, tES induces a subthreshold polarization of cortical neurons that is too weak to generate an action potential. However, by changing the intrinsic neuronal excitability, tES can induce changes in the resting membrane potential and the postsynaptic activity of cortical neurons. This, in turn, can alter the spontaneous firing rate of neurons and modulate their response to afferent signals (Bindman et al., 1962, 1964, 1979; Creutzfeldt et al., 1962), leading to changes in synaptic efficacy. The typical application of NIBS involves different types of protocols: TMS can be delivered as a single pulse (spTMS) at a precise time, as pairs of pulses separated by a variable interval, or as a series of stimuli in conventional or patterned protocols of repetitive TMS (rTMS) (for a complete classification see Rossi et al., 2009). In tES, different protocols are established by the electrical current used and by its polarity, which can be direct (anodal or cathodal transcranial direct current stimulation: tDCS), high-definition transcranial direct current stimulation (HD-tDCS), oscillating transcranial direct current stimulation (osc-tDCS), alternating at a fix frequency (transcranial alternating current stimulation: tACS) transcranial pulsed current stimulation (tPCS) (electrosleep), or at random frequencies (transcranial random noise stimulation: tRNS) (Nitsche et al., 2008; Paulus, 2011).

(127) NIBS also includes brain entrainment using light stimulation and sound stimulation. The latter can be binaural beats (BB) or isochronic tones.

(128) In general, the final effects of NIBS on the central nervous system depend on a lengthy list of parameters (e.g., frequency, temporal characteristics, intensity, geometric configuration of the coil/electrode, i.e., "montage," current direction), when it is delivered before (off-line) or during (on-line) the task as part of the experimental procedure (e.g., Jacobson et al., 2011; Nitsche and Paulus, 2011; Sandrini et al., 2011). In addition, these factors interact with several variables related to the brain anatomy and morphology (e.g., brain size, properties of the brain tissue and its location, Radman et al., 2007), as well as physiological (e.g., gender and age, Landi and Rossini, 2010; Lang et al., 2011; Ridding and Ziemann, 2010) and cognitive (e.g., Miniussi et al., 2010; Silvanto et al., 2008; Walsh et al., 1998) states of the stimulated area/subject. Transcranial Direct Current Stimulation (tDCS) Cranial electrotherapy stimulation (CES) is a form of non-invasive brain stimulation that applies a small, pulsed electric current across a person's head to treat a variety of conditions such as anxiety, depression and insomnia. See, en.wikipedia.org/wiki/Cranial_electrotherapy_stimulation. Transcranial direct current stimulation (tDCS, HD-tDCS, osc-tDCS, tPCS) is a form of neurostimulation that uses constant, low current delivered to the brain area of interest via electrodes on the scalp. It was originally developed to help patients with brain injuries or psychiatric conditions like major depressive disorder. tDCS appears to have some potential for treating depression. See, en.wikipedia.org/wiki/Transcranial_direct-current_stimulation.

(129) The hypotheses concerning the application of tDCS in cognition are very similar to those of TMS, with the exception that tDCS was never considered a virtual lesion method. tDCS can increase or decrease cortical excitability in the stimulated brain regions and facilitate or inhibit behavior accordingly. TES does not induce action potentials but instead modulates the neuronal response threshold so that it can be defined as subthreshold stimulation.

(130) tDCS is being studied for acceleration of learning. The mild electrical shock (usually, a 2-milliamp current) is used to depolarize the neuronal membranes, making the cells more excitable and responsive to inputs. Weisend, *Experimental Brain Research*, vol 213, p 9 (DARPA) showed that tDCS accelerates the formation of new neural pathways during the time that someone practices a skill. tDCS appears to bring about the flow state. The movements of the subjects become more automatic; they report calm, focused concentration, and their performance improves immediately. (See Adey, Sally, "Zap your brain into the zone: Fast track to pure focus", *New Scientist* No. 2850, Feb. 1, 2012, www.newscientist.com/article/mg21328501-600-zap-your-brain-into-the-zone-fast-track-to-pure-focus/).

(131) U.S. Pat. Nos. 7,856,264; 8,706,241; 8,725,669; 9,037,224; 9,042,201; 9,095,266; 9,248,286; 9,349,178; 9,629,568; 9,693,725; 9,713,433; 20040195512; 20070179534; 20110092882; 20110311021; 20120165696; 20140142654; 20140200432; 20140211593; 20140316243; 20140347265; 20150099946; 20150174418; 20150257700; 20150327813; 20150343242; 20150351655; 20160000354; 20160038049; 20160113569; 20160144175; 20160148371; 20160180042; 20160213276; 20160228702; and 20160235323.

(132) Reinhart, Robert M G. "Disruption and rescue of interareal theta phase coupling and adaptive behavior." *Proceedings of the National Academy of Sciences* (2017): provide evidence for a causal relation between interareal theta phase synchronization in frontal cortex and multiple components of adaptive human behavior. Reinhart's results support the idea that the precise timing of rhythmic population activity spatially distributed in frontal cortex conveys information to direct behavior. Given prior work showing that phase synchronization can change spike time-dependent plasticity, together with Reinhart's findings showing stimulation effects on neural activity and behavior can outlast a 20-min period of electrical stimulation, it is reasonable to suppose that the externally modulated interareal coupling changed behavior by causing neuroplastic modifications in functional connectivity. Reinhart suggests that we may be able to noninvasively intervene in the temporal coupling of distant rhythmic activity in the human brain to optimize (or impede) the postsynaptic effect of spikes from one area on the other, improving (or impairing) the cross-area communication necessary for cognitive action control and learning. Moreover, these neuroplastic alterations in functional connectivity were induced with a 0° phase, suggesting that inducing synchronization does not require a meticulous accounting of the communication delay between regions such as medial frontal cortex (MFC) and lateral prefrontal cortex (IPFC) to effectively modify behavior and learning. This conforms to work showing that despite long axonal conduction delays between distant brain areas, theta phase synchronizations at 0° phase lag can occur between these regions and underlie meaningful functions of cognition and action. It is also possible that a third subcortical or posterior region with a nonzero time lag interacted with

these two frontal areas to drive changes in goal-directed behavior. Alexander W H & Brown J W (2011) Medial prefrontal cortex as an action-outcome predictor. *Nature Neuroscience* 14(10):1338-1344. Alexander W H & Brown J W (2015) Hierarchical error representation: A computational model of anterior cingulate and dorsolateral prefrontal cortex. *Neural Computation* 27:2354-2410. Anguera J A, et al. (2013) Video game training enhances cognitive control in older adults. *Nature* 501:97-101. Aron A R, Fletcher P C, Bullmore E T, Sahakian B J, Robbins T W (2003) Stop-signal inhibition disrupted by damage to right inferior frontal gyms in humans. *Nat Neurosci* 6:115-116. Au J, et al. (2015) Improving fluid intelligence with training on working memory: a meta-analysis. *Psychonomic Bulletin & Review* 22:366-377. Bellman R, Kalaba R (1959) A mathematical theory of adaptive control processes. *Proc Natl Acad Sci USA* 45:1288-1290. 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(133) High-Definition-tDCS High-Definition transcranial Direct Current Stimulation (HD-tDCS) was invented by Dr. Marom Bikson's group at The City College of New York with the introduction of the 4x1 HD-tDCS montage. The 4x1 HD-tDCS montage allows precise targeting of cortical structures. The region of current flow is circumscribed by the area of the 4x ring, such that decreasing ring radius increases focality. 4x1 HD-tDCS allows for unifocal stimulation, meaning the polarity of the center 1x electrode will determine the direction of neuromodulation under the ring. This is in contrast to conventional tDCS where the need for one anode and one cathode always produces bidirectional modulation (even when an extra-cephalic electrode is used). 4x1 HD-tDCS thus provides the ability not only to select a cortical brain region to target, but to modulate the excitability of that brain region with a designed polarity without having to consider return counter-electrode flow.

(134) Osc-tDCS Oscillating transcranial direct current stimulation (osc-tDCS) is a tDCS wherein the amplitude of the current is modulated with a sinusoid waveform of a certain frequency. Osc-tDCS modulates the spontaneous brain activity in a frequency-specific manner. Osc-tDCS mainly affects brain oscillatory activity. Anodal oscillatory stimulation at 0.75 Hz (slow osc-tDCS) in frontal areas during sleep stage 2 of a diurnal nap or during nocturnal sleep can induce a frequency-specific enhancement of Slow-Wave Activity (SWA, 0.5-4 Hz) during sleep. The enhancement in normal subjects of SWA induced by osc-tDCS at 0.75 Hz during sleep significantly improves performance in a memory task after sleep. See Bergmann T O, Groppa S, Seeger M, Male M, Marshall L, Siebner H R. "Acute changes in motor cortical excitability during slow oscillatory and constant anodal transcranial direct current stimulation." *J Neurophysiol.* 2009 October; 102(4):2303-11. Marshall L, Helgadottir H, Mölle M, Born J. "Boosting slow oscillations during sleep potentiates memory." *Nature.* 2006 Nov. 30; 444(7119):610-3. Marshall L, Kirov R, Brade J, Mölle M, Born J

(135) "Transcranial electrical currents to probe EEG brain rhythms and memory consolidation during sleep in humans." *PLoS One.* 2011 Feb. 14; 6(2):e16905.

(136) Transcranial Alternative Current Stimulation (tACS) Transcranial alternating current stimulation (tACS) is a noninvasive means by which alternating electrical current applied through the skin and skull entrains in a frequency-specific fashion the neural oscillations of the underlying brain. See, en.wikipedia.org/wiki/Transcranial_alternating_current_stimulation

(137) U.S. Pub. App. No. 20170197081 discloses transdermal electrical stimulation of nerves to modify or induce a cognitive state using transdermal electrical stimulation (TES).

(138) U.S. Pat. Nos. 6,804,558; 7,149,773; 7,181,505; 7,278,966; 9,042,201; 9,629,568; 9,713,433; 20010051787; 20020013613; 20020052539; 20020082665; 20050171410; 20140211593; 20140316243; 20150174418; 20150343242; 20160000354; 20160038049; 20160106513; 20160213276; 20160228702; 20160232330; 20160235323; and 20170113056.

(139) Transcranial Random Noise Stimulation (tRNS) Transcranial random noise stimulation (tRNS) is a non-invasive brain stimulation technique and a form of transcranial electrical stimulation (tES). See, en.wikipedia.org/wiki/Transcranial_random_noise_stimulation; U.S. Pat. Nos. 9,198,733; 9,713,433; 20140316243; 20160038049; and 20160213276.

(140) Transcranial pulsed current stimulation (tPCS) The stimulus may comprise transcranial pulsed current stimulation (tPCS). See: Shapour Jaberzadeh, Andisheh Bastani, Maryam Zoghi, "Anodal transcranial pulsed current stimulation: A novel technique to enhance corticospinal excitability," *Clin. Neurophysiology*, Volume 125, Issue 2, February 2014, Pages 344-351, doi.org/10.1016/j.clinph.2013.08.025; earthpulse.net/Apcs-transcranial-pulsed-current-stimulation/; help.foc.us/article/16-tpcs-transcranial-pulsed-current-stimulation.

(141) Transcranial Magnetic Stimulation Transcranial magnetic stimulation (TMS) is a method in which a changing magnetic field is used to cause electric current to flow in a small region of the brain via electromagnetic induction. During a TMS procedure, a magnetic field generator, or "coil", is placed near the head of the person receiving the treatment. The coil is connected to a pulse generator, or stimulator, that delivers a changing electric current to the coil. TMS is used diagnostically to measure the connection between the central nervous system and skeletal muscle to evaluate damage in a wide variety of disease states, including stroke, multiple sclerosis, amyotrophic lateral sclerosis, movement disorders, and motor neuron diseases. Evidence is available suggesting that TMS is useful in treating neuropathic pain, major depressive disorder, and other conditions.

(142) See, en.wikipedia.org/wiki/Transcranial_magnetic_stimulation,

(143) See U.S. Pat. Nos. 4,296,756; 4,367,527; 5,069,218; 5,088,497; 5,359,363; 5,384,588; 5,459,536; 5,711,305; 5,877,801; 5,891,131; 5,954,662; 5,971,923; 6,188,924; 6,259,399; 6,487,441; 6,603,502; 7,714,936; 7,844,324; 7,856,264; 8,221,330; 8,655,817; 8,706,241; 8,725,669; 8,914,115; 9,037,224; 9,042,201; 9,095,266; 9,149,195; 9,248,286; 9,265,458; 9,414,776; 9,445,713; 9,713,433; 20020097332; 20040088732; 20070179534; 20070249949; 20080194981; 20090006001; 20110004412; 20110007129; 20110087127; 20110092882; 20110119212; 20110137371; 20120165696; 20120296569; 20130339043; 20140142654; 20140163328; 20140200432; 20140211593; 20140257047; 20140279746; 20140316243; 20140350369; 20150065803; 20150099946; 20150148617; 20150174418; 20150257700; 20150327813; 20150343242; 20150351655; 20160038049; 20160140306; 20160144175; 20160213276; 20160235323; 20160284082; 20160306942; 20160317077; 20170084175; and 20170113056.

(144) Pulsed electromagnetic field (PEMF) Pulsed electromagnetic field (PEMF) when applied to the brain is referred to as Transcranial magnetic stimulation, and has been FDA approved since 2008 for use in people who failed to respond to antidepressants. Weak magnetic stimulation of the brain is often called transcranial pulsed electromagnetic field (tPEMF) therapy. See, en.wikipedia.org/wiki/Pulsed_electromagnetic_field_therapy,

(145) See, U.S. Pat. Nos. 7,280,861; 8,343,027; 8,415,123; 8,430,805; 8,435,166; 8,571,642; 8,657,732; 8,775,340; 8,961,385; 8,968,172; 9,002,477; 9,005,102; 9,278,231; 9,320,913; 9,339,641; 9,387,338; 9,415,233; 9,427,598; 9,433,797; 9,440,089; 9,610,459; 9,630,004; 9,656,096; 20030181791; 20060129022; 20100057655; 20100197993; 20120101544; 20120116149; 20120143285; 20120253101; 20130013339;

20140131843; 20140213844; 20140221726; 20140228620; 20140303425; 20160235983; 20170087367; and 20170165496.

(146) Deep Brain Stimulation (DBS) Deep brain stimulation (DBS) is a neurosurgical procedure involving the implantation of a medical device called a neurostimulator (sometimes referred to as a ‘brain pacemaker’), which sends electrical impulses, through implanted electrodes, to specific targets in the brain (brain nuclei) for the treatment of movement and neuropsychiatric disorders. See, en.wikipedia.org/wiki/Deep_brain_stimulation.

(147) Transcranial Pulse Ultrasound (TPU) Transcranial pulsed ultrasound (TPU) uses low intensity, low frequency ultrasound (LILFU) as a method to stimulate the brain. See, en.wikipedia.org/wiki/Transcranial_pulsed_ultrasound;

(148) U.S. Pat. Nos. 8,591,419; 8,858,440; 8,903,494; 8,921,320; 9,002,458; 9,014,811; 9,036,844; 9,042,201; 9,061,133; 9,233,244; 9,333,334; 9,399,126; 9,403,038; 9,440,070; 9,630,029; 9,669,239; 20120259249; 20120283502; 20120289869; 20130079621; 20130144192; 20130184218; 20140058219; 20140211593; 20140228653; 20140249454; 20140316243; 20150080327; 20150133716; 20150343242; 20160143541; 20160176053; and 20160220850.

(149) Sensory Stimulation Light, sound or electromagnetic fields may be used to remotely convey a temporal pattern of brainwaves. See:

(150) U.S. Pat. Nos. 5,293,187; 5,422,689; 5,447,166; 5,491,492; 5,546,943; 5,622,168; 5,649,061; 5,720,619; 5,740,812; 5,983,129; 6,050,962; 6,092,058; 6,149,586; 6,325,475; 6,377,833; 6,394,963; 6,428,490; 6,482,165; 6,503,085; 6,520,921; 6,522,906; 6,527,730; 6,556,695; 6,565,518; 6,652,458; 6,652,470; 6,701,173; 6,726,624; 6,743,182; 6,746,409; 6,758,813; 6,843,774; 6,896,655; 6,996,261; 7,037,260; 7,070,571; 7,107,090; 7,120,486; 7,212,851; 7,215,994; 7,260,430; 7,269,455; 7,280,870; 7,392,079; 7,407,485; 7,463,142; 7,478,108; 7,488,294; 7,515,054; 7,567,693; 7,647,097; 7,740,592; 7,751,877; 7,831,305; 7,856,264; 7,881,780; 7,970,734; 7,972,278; 7,974,787; 7,991,461; 8,012,107; 8,032,486; 8,033,996; 8,060,194; 8,095,209; 8,209,224; 8,239,030; 8,262,714; 8,320,649; 8,358,818; 8,376,965; 8,380,316; 8,386,312; 8,386,313; 8,392,250; 8,392,253; 8,392,254; 8,392,255; 8,437,844; 8,464,288; 8,475,371; 8,483,816; 8,494,905; 8,517,912; 8,533,042; 8,545,420; 8,560,041; 8,655,428; 8,672,852; 8,682,687; 8,684,742; 8,694,157; 8,706,241; 8,706,518; 8,738,395; 8,753,296; 8,762,202; 8,764,673; 8,768,022; 8,788,030; 8,790,255; 8,790,297; 8,821,376; 8,838,247; 8,864,310; 8,872,640; 8,888,723; 8,915,871; 8,938,289; 8,938,301; 8,942,813; 8,955,010; 8,955,974; 8,958,882; 8,964,298; 8,971,936; 8,989,835; 8,992,230; 8,998,828; 9,004,687; 9,060,671; 9,101,279; 9,135,221; 9,142,145; 9,165,472; 9,173,582; 9,179,855; 9,208,558; 9,215,978; 9,232,984; 9,241,665; 9,242,067; 9,254,099; 9,271,660; 9,275,191; 9,282,927; 9,292,858; 9,292,920; 9,320,450; 9,326,705; 9,330,206; 9,357,941; 9,396,669; 9,398,873; 9,414,780; 9,414,907; 9,424,761; 9,445,739; 9,445,763; 9,451,303; 9,451,899; 9,454,646; 9,462,977; 9,468,541; 9,483,117; 9,492,120; 9,504,420; 9,504,788; 9,526,419; 9,541,383; 9,545,221; 9,545,222; 9,545,225; 9,560,967; 9,560,984; 9,563,740; 9,582,072; 9,596,224; 9,615,746; 9,622,702; 9,622,703; 9,626,756; 9,629,568; 9,642,699; 9,649,030; 9,651,368; 9,655,573; 9,668,694; 9,672,302; 9,672,617; 9,682,232; 9,693,734; 9,694,155; 9,704,205; 9,706,910; 9,710,788; RE44408; RE45766; 20020024450; 20020103428; 20020103429; 20020112732; 20020128540; 20030028081; 20030028121; 20030070685; 20030083596; 20030100844; 20030120172; 20030149351; 20030158496; 20030158497; 20030171658; 20040019257; 20040024287; 20040068172; 20040092809; 20040101146; 20040116784; 20040143170; 20040267152; 20050010091; 20050019734; 20050025704; 20050038354; 20050113713; 20050124851; 20050148828; 20050228785; 20050240253; 20050245796; 20050267343; 20050267344; 20050283053; 20060020184; 20060061544; 20060078183; 20060087746; 20060102171; 20060129277; 20060161218; 20060189866; 20060200013; 20060241718; 20060252978; 20060252979; 20070050715; 20070179534; 20070191704; 20070238934; 20070273611; 20070282228; 20070299371; 20080004550; 20080009772; 20080058668; 20080081963; 20080119763; 20080123927; 20080132383; 20080228239; 20080234113; 20080234601; 20080242521; 20080255949; 20090018419; 20090058660; 20090062698; 20090076406; 20090099474; 20090112523; 20090221928; 20090267758; 20090270687; 20090270688; 20090270692; 20090270693; 20090270694; 20090270786; 20090281400; 20090287108; 20090297000; 20090299169; 20090311655; 20090312808; 20090312817; 20090318794; 20090326604; 20100004977; 20100010289; 20100010366; 20100041949; 20100069739; 20100069780; 20100163027; 20100163028; 20100163035; 20100165593; 20100168525; 20100168529; 20100168602; 20100268055; 20100293115; 20110004412; 20110009777; 20110015515; 20110015539; 20110043759; 20110054272; 20110077548; 20110092882; 20110105859; 20110130643; 20110172500; 20110218456; 20110256520; 20110270074; 20110301488; 20110307079; 20120004579; 20120021394; 20120036004; 20120071771; 20120108909; 20120108995; 20120136274; 20120150545; 20120203130; 20120262558; 20120271377; 20120310106; 20130012804; 20130046715; 20130063434; 20130063550; 20130080127; 20130120246; 20130127980; 20130185144; 20130189663; 20130204085; 20130211238; 20130226484; 20130242262; 20130245424; 20130281759; 20130289360; 20130293844; 20130308099; 20130318546; 20140058528; 20140155714; 20140171757; 20140200432; 20140214335; 20140221866; 20140243608; 20140243614; 20140243652; 20140276130; 20140276944; 20140288614; 20140296750; 20140300532; 20140303508; 20140304773; 20140313303; 20140315169; 20140316191; 20140316192; 20140316235; 20140316248; 20140323899; 20140335489; 20140343408; 20140347491; 20140350353; 20140350431; 20140364721; 20140378810; 20150002815; 20150003698; 20150003699; 20150005640; 20150005644; 20150006186; 20150012111; 20150038869; 20150045606; 20150051663; 20150099946; 20150112409; 20150120007; 20150124220; 20150126845; 20150126873; 20150133812; 20150141773; 20150145676; 20150154889; 20150174362; 20150196800; 20150213191; 20150223731; 20150234477; 20150235088; 20150235370; 20150235441; 20150235447; 20150241705; 20150241959; 20150242575; 20150242943; 20150243100; 20150243105; 20150243106; 20150247723; 20150247975; 20150247976; 20150248169; 20150248170; 20150248787; 20150248788; 20150248789; 20150248791; 20150248792; 20150248793; 20150290453; 20150290454; 20150305685; 20150306340; 20150309563; 20150313496; 20150313539; 20150324692; 20150325151; 20150335288; 20150339363; 20150351690; 20150366497; 20150366504; 20150366656; 20150366659; 20150369864; 20150370320; 20160000354; 20160004298; 20160005320; 20160007915; 20160008620; 20160012749; 20160015289; 20160022167; 20160022206; 20160029946; 20160029965; 20160038069; 20160051187; 20160051793; 20160066838; 20160073886; 20160077547; 20160078780; 20160106950; 20160112684; 20160120436; 20160143582; 20160166219; 20160167672; 20160176053; 20160180054; 20160198950; 20160199577; 20160202755; 20160216760; 20160220439; 20160228640; 20160232625; 20160232811; 20160235323; 20160239084; 20160248994; 20160249826; 20160256108; 20160267809; 20160270656; 20160287157; 20160302711; 20160306942; 20160313798; 20160317060; 20160317383; 20160324478; 20160324580; 20160334866; 20160338644; 20160338825; 20160339300; 20160345901; 20160357256; 20160360970; 20160363483; 20170000324; 20170000325; 20170000326; 20170000329; 20170000330; 20170000331; 20170000332; 20170000333; 20170000334; 20170000335; 20170000337; 20170000340; 20170000341; 20170000342; 20170000343; 20170000345; 20170000454; 20170000683; 20170001032; 20170006931; 20170007111; 20170007115; 20170007116; 20170007122; 20170007123; 20170007165; 20170007182; 20170007450; 20170007799; 20170007843; 20170010469; 20170010470; 20170017083; 20170020447; 20170020454; 20170020627; 20170027467; 20170027651; 20170027812; 20170031440; 20170032098; 20170035344; 20170043160; 20170055900; 20170060298; 20170061034; 20170071523; 20170071537; 20170071546; 20170071551; 20170080320; 20170086729; 20170095157; 20170099479; 20170100540; 20170103440; 20170112427; 20170112671; 20170113046; 20170113056; 20170119994; 20170135597; 20170135633; 20170136264; 20170136265; 20170143249; 20170143442; 20170148340; 20170156662; 20170162072; 20170164876; 20170164878; 20170168568; 20170173262; 20170173326; 20170177023; 20170188947; 20170202633; 20170209043; 20170209094; and 20170209737.

(151) Light Stimulation The functional relevance of brain oscillations in the alpha frequency range (7.5-12 Hz) has been repeatedly investigated through the use of rhythmic visual stimulation. There are two hypotheses on the origin of steady-state visual evoked potential (SSVEP) measured in EEG during rhythmic stimulation: entrainment of brain oscillations and superposition of event-related responses (ERPs). The entrainment but not the superposition hypothesis justifies rhythmic visual stimulation as a means to manipulate brain oscillations, because superposition assumes a linear summation of single responses, independent from ongoing brain oscillations. Participants stimulated with rhythmic flickering light of different frequencies and intensities, and entrainment was measured by comparing the phase coupling of brain oscillations stimulated by rhythmic visual flicker with the oscillations induced by arrhythmic jittered stimulation, varying the time, stimulation frequency, and intensity conditions. Phase

coupling was found to be more pronounced with increasing stimulation intensity as well as at stimulation frequencies closer to each participant's intrinsic frequency. Even in a single sequence of an SSVEP, non-linear features (intermittency of phase locking) was found that contradict the linear summation of single responses, as assumed by the superposition hypothesis. Thus, evidence suggests that visual rhythmic stimulation entrains brain oscillations, validating the approach of rhythmic stimulation as a manipulation of brain oscillations. See, Notbohm A, Kurths J, Herrmann C S, Modification of Brain Oscillations via Rhythmic Light Stimulation Provides Evidence for Entrainment but Not for Superposition of Event-Related Responses, *Front Hum Neurosci*. 2016 Feb. 3; 10:10. doi: 10.3389/fnhum.2016.00010. eCollection 2016.

(152) It is also known that periodic visual stimulation can trigger epileptic seizures.

(153) Cochlear Implant A cochlea implant is a surgically implanted electronic device that provides a sense of sound to a person who is profoundly deaf or severely hard of hearing in both ears. See, en.wikipedia.org/wiki/Cochlear_implant;

(154) See, U.S. Pat. Nos. 5,999,856; 6,354,299; 6,427,086; 6,430,443; 6,665,562; 6,873,872; 7,359,837; 7,440,806; 7,493,171; 7,610,083; 7,610,100; 7,702,387; 7,747,318; 7,765,088; 7,853,321; 7,890,176; 7,917,199; 7,920,916; 7,957,806; 8,014,870; 8,024,029; 8,065,017; 8,108,033; 8,108,042; 8,140,152; 8,165,687; 8,175,700; 8,195,295; 8,209,018; 8,224,431; 8,315,704; 8,332,024; 8,401,654; 8,433,410; 8,478,417; 8,515,541; 8,538,543; 8,560,041; 8,565,864; 8,574,164; 8,577,464; 8,577,465; 8,577,466; 8,577,467; 8,577,468; 8,577,472; 8,577,478; 8,588,941; 8,594,800; 8,644,946; 8,644,957; 8,652,187; 8,676,325; 8,696,724; 8,700,183; 8,718,776; 8,768,446; 8,768,477; 8,788,057; 8,798,728; 8,798,773; 8,812,126; 8,864,806; 8,868,189; 8,929,999; 8,968,376; 8,989,868; 8,996,120; 9,002,471; 9,044,612; 9,061,132; 9,061,151; 9,095,713; 9,135,400; 9,186,503; 9,235,685; 9,242,067; 9,248,290; 9,248,291; 9,259,177; 9,302,093; 9,314,613; 9,327,069; 9,352,145; 9,352,152; 9,358,392; 9,358,393; 9,403,009; 9,409,013; 9,415,215; 9,415,216; 9,421,372; 9,432,777; 9,501,829; 9,526,902; 9,533,144; 9,545,510; 9,550,064; 9,561,380; 9,578,425; 9,592,389; 9,604,067; 9,616,227; 9,643,017; 9,649,493; 9,674,621; 9,682,232; 9,743,197; 9,744,358; 20010014818; 20010029391; 20020099412; 20030114886; 20040073273; 20050149157; 20050182389; 20050182450; 20050182467; 20050182468; 20050182469; 20050187600; 20050192647; 20050209664; 20050209665; 20050209666; 20050228451; 20050240229; 20060064140; 20060094970; 20060094971; 20060094972; 20060095091; 20060095092; 20060161217; 20060173259; 20060178709; 20060195039; 20060206165; 20060235484; 20060235489; 20060247728; 20060282123; 20060287691; 20070038264; 20070049988; 20070156180; 20070198063; 20070213785; 20070244407; 20070255155; 20070255531; 20080049376; 20080140149; 20080161886; 20080208280; 20080235469; 20080249589; 20090163980; 20090163981; 20090243756; 20090259277; 20090270944; 20090280153; 20100030287; 20100100164; 20100198282; 20100217341; 20100231327; 20100241195; 20100268055; 20100268288; 20100318160; 20110004283; 20110060382; 20110166471; 20110295344; 20110295345; 20110295346; 20110295347; 20120035698; 20120116179; 20120116741; 20120150255; 20120245655; 20120262250; 20120265270; 20130165996; 20130197944; 20130235550; 20140032512; 20140098981; 20140200623; 20140249608; 20140275847; 20140330357; 20140350634; 20150018699; 20150045607; 20150051668; 20150065831; 20150066124; 20150080674; 20150328455; 20150374986; 20150374987; 20160067485; 20160243362; 20160261962; 20170056655; 20170087354; 20170087355; 20170087356; 20170113046; 20170117866; 20170135633; and 20170182312.

(155) Vagus Nerve Stimulation Vagus nerve stimulation (VNS) is a medical treatment that involves delivering electrical impulses to the vagus nerve. It is used as an adjunctive treatment for certain types of intractable epilepsy and treatment-resistant depression. See, en.wikipedia.org/wiki/Vagus_nerve_stimulation;

(156) See, U.S. Pat. Nos. 5,215,086; 5,231,988; 5,299,569; 5,335,657; 5,571,150; 5,928,272; 5,995,868; 6,104,956; 6,167,311; 6,205,359; 6,208,902; 6,248,126; 6,269,270; 6,339,725; 6,341,236; 6,356,788; 6,366,814; 6,418,344; 6,497,699; 6,549,804; 6,556,868; 6,560,486; 6,587,727; 6,591,137; 6,597,954; 6,609,030; 6,622,047; 6,665,562; 6,671,556; 6,684,105; 6,708,064; 6,735,475; 6,782,292; 6,788,975; 6,873,872; 6,879,859; 6,882,881; 6,920,357; 6,961,618; 7,003,352; 7,151,961; 7,155,279; 7,167,751; 7,177,678; 7,203,548; 7,209,787; 7,228,167; 7,231,254; 7,242,984; 7,277,758; 7,292,890; 7,313,442; 7,324,485; 7,346,395; 7,366,571; 7,386,347; 7,389,144; 7,403,820; 7,418,290; 7,422,555; 7,44,184; 7,454,245; 7,457,665; 7,463,927; 7,486,986; 7,493,172; 7,499,752; 7,561,918; 7,620,455; 7,623,927; 7,623,928; 7,630,757; 7,634,317; 7,643,881; 7,653,433; 7,657,316; 7,676,263; 7,680,526; 7,684,858; 7,706,871; 7,711,432; 7,734,355; 7,736,382; 7,747,325; 7,747,326; 7,769,461; 7,783,362; 7,801,601; 7,805,203; 7,840,280; 7,848,803; 7,853,321; 7,853,329; 7,860,548; 7,860,570; 7,865,244; 7,869,867; 7,869,884; 7,869,885; 7,890,185; 7,894,903; 7,899,539; 7,904,134; 7,904,151; 7,904,175; 7,908,008; 7,920,915; 7,925,353; 7,945,316; 7,957,796; 7,962,214; 7,962,219; 7,962,220; 7,974,688; 7,974,693; 7,974,697; 7,974,701; 7,996,079; 8,000,788; 8,027,730; 8,036,745; 8,041,418; 8,041,419; 8,046,076; 8,064,994; 8,068,911; 8,097,926; 8,108,038; 8,112,148; 8,112,153; 8,116,883; 8,150,508; 8,150,524; 8,160,696; 8,172,759; 8,180,601; 8,190,251; 8,190,264; 8,204,603; 8,209,009; 8,209,019; 8,214,035; 8,219,188; 8,224,444; 8,224,451; 8,229,559; 8,239,028; 8,260,426; 8,280,505; 8,306,627; 8,315,703; 8,315,704; 8,326,418; 8,337,404; 8,340,771; 8,346,354; 8,352,031; 8,374,696; 8,374,701; 8,379,952; 8,382,667; 8,401,634; 8,412,334; 8,412,338; 8,417,344; 8,423,155; 8,428,726; 8,452,387; 8,454,555; 8,457,747; 8,467,878; 8,478,428; 8,485,979; 8,489,185; 8,498,699; 8,515,538; 8,536,667; 8,538,523; 8,538,543; 8,548,583; 8,548,594; 8,548,604; 8,560,073; 8,562,536; 8,562,660; 8,565,867; 8,571,643; 8,571,653; 8,588,933; 8,591,419; 8,600,521; 8,603,790; 8,606,360; 8,615,309; 8,630,705; 8,634,922; 8,641,646; 8,644,954; 8,649,871; 8,652,187; 8,660,666; 8,666,501; 8,676,324; 8,676,330; 8,684,921; 8,694,118; 8,700,163; 8,712,547; 8,716,447; 8,718,779; 8,725,243; 8,738,126; 8,744,562; 8,761,868; 8,762,065; 8,768,471; 8,781,597; 8,815,582; 8,827,912; 8,831,732; 8,843,210; 8,849,409; 8,852,100; 8,855,775; 8,858,440; 8,864,806; 8,868,172; 8,868,177; 8,874,205; 8,874,218; 8,874,227; 8,888,702; 8,914,122; 8,918,178; 8,934,967; 8,942,817; 8,945,006; 8,948,855; 8,965,514; 8,968,376; 8,972,004; 8,972,013; 8,983,155; 8,983,628; 8,983,629; 8,985,119; 8,989,863; 8,989,867; 9,014,804; 9,014,823; 9,020,582; 9,020,598; 9,020,789; 9,026,218; 9,031,655; 9,042,201; 9,042,988; 9,043,001; 9,044,188; 9,050,469; 9,056,195; 9,067,054; 9,067,070; 9,079,940; 9,089,707; 9,089,719; 9,095,303; 9,095,314; 9,108,041; 9,113,801; 9,119,533; 9,135,400; 9,138,580; 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(157) Brain-To-Brain Interface A brain-brain interface is a direct communication pathway between the brain of one animal and the brain of another animal. Brain to brain interfaces have been used to help rats collaborate with each other. When a second rat was unable to choose the correct lever, the first rat noticed (not getting a second reward), and produced a round of task-related neuron firing that made the second rat more likely to choose the correct lever. Human studies have also been conducted.

(158) In 2013, researcher from the University of Washington were able to use electrical brain recordings and a form of magnetic stimulation to send a brain signal to a recipient, which caused the recipient to hit the fire button on a computer game. In 2015, researchers linked up multiple brains, of both monkeys and rats, to form an “organic computer.” It is hypothesized that by using brain-to-brain interfaces (BTBIs) a biological computer, or brain-net, could be constructed using animal brains as its computational units. Initial exploratory work demonstrated collaboration between rats in distant cages linked by signals from cortical microelectrode arrays implanted in their brains. The rats were rewarded when actions were performed by the “decoding rat” which conformed to incoming signals and when signals were transmitted by the “encoding rat” which resulted in the desired action. In the initial experiment the rewarded action was pushing a lever in the remote location corresponding to the position of a lever near a lighted LED at the home location. About a month was required for the rats to acclimate themselves to incoming “brainwaves.” When a decoding rat was unable to choose the correct lever, the encoding rat noticed (not getting an expected reward), and produced a round of task-related neuron firing that made the second rat more likely to choose the correct lever.

(159) In another study, electrical brain readings were used to trigger a form of magnetic stimulation, to send a brain signal based on brain activity on a subject to a recipient, which caused the recipient to hit the fire button on a computer game.

(160) Brain-To-Computer Interface A brain-computer interface (BCI), sometimes called a neural-control interface (NCI), mind-machine interface (MMI), direct neural interface (DNI), or brain-machine interface (BMI), is a direct communication pathway between an enhanced or wired brain and an external device. BCI differs from neuromodulation in that it allows for bidirectional information flow. BCIs are often directed at researching, mapping, assisting, augmenting, or repairing human cognitive or sensory-motor functions.

(161) Synthetic telepathy, also known as techlepathy or psychotronics (geeldon.wordpress.com/2010/09/06/synthetic-telepathy-also-known-as-techlepathy-or-psychotronics/), describes the process of use of brain-computer interfaces by which human thought (as electromagnetic radiation) is intercepted, processed by computer and a return signal generated that is perceptible by the human brain. Dewan, E. M., “Occipital Alpha Rhythm Eye Position and Lens Accommodation.” *Nature* 214, 975-977 (3 Jun. 1967), demonstrates the mental control of Alpha waves, turning them on and off, to produce Morse code representations of words and phrases by thought alone. U.S. Pat. No. 3,951,134 proposes remotely monitoring and altering brainwaves using radio, and references demodulating the waveform, displaying it to an operator for viewing and passing this to a computer for further analysis. In 1988, Farwell, L. A. & Donchin, E. (1988). Talking off the top of your head: toward a mental prosthesis utilizing event-related brain potentials. *Electroencephalography and Clinical Neurophysiology*, 70(6), 510-523 describes a method of transmitting linguistic information using the P300 response system, which combines matching observed information to what the subject was thinking of. In this case, being able to select a letter of the alphabet that the subject was thinking of. In theory, any input could be used and a lexicon constructed. U.S. Pat. No. 6,011,991 describes a method of monitoring an individual's brainwaves remotely, for the purposes of communication, and outlines a system that monitors an individual's brainwaves via a sensor, then transmits this information, specifically by satellite, to a computer for analysis. This analysis would determine if the individual was attempting to communicate a “word, phrase, or thought corresponding to the matched stored normalized signal.”

(162) Approaches to synthetic telepathy can be categorized into two major groups, passive and active. Like sonar, the receiver can take part or passively listen. Passive reception is the ability to “read” a signal without first broadcasting a signal. This can be roughly equated to tuning into a radio station—the brain generates electromagnetic radiation which can be received at a distance. That distance is determined by the sensitivity of the receiver, the filters used and the bandwidth required. Most universities would have limited budgets, and receivers, such as EEG (and similar devices),

would be used. A related military technology is the surveillance system TEMPEST. Robert G. Malech's approach requires a modulated signal to be broadcast at the target. The method uses an active signal, which is interfered with by the brain's modulation. Thus, the return signal can be used to infer the original brainwave.

(163) Computer mediation falls into two basic categories, interpretative and interactive. Interpretative mediation is the passive analysis of signals coming from the human brain. A computer "reads" the signal then compares that signal against a database of signals and their meanings. Using statistical analysis and repetition, false-positives are reduced overtime. Interactive mediation can be in a passive-active mode or active-active mode. In this case, passive and active denote the method of reading and willing to the brain and whether or not they make use of a broadcast signal. Interactive mediation can also be performed manually or via artificial intelligence. Manual interactive mediation involves a human operator producing return signals such as speech or images. AI mediation leverages the cognitive system of the subject to identify images, pre-speech, objects, sounds and other artifacts, rather than developing AI routines to perform such activities. AI based systems may incorporate natural language processing interfaces that produce sensations, mental impressions, humor and conversation to provide a mental picture of a computerized personality. Statistical analysis and ML techniques, such as neural networks can be used.

(164) ITV News Service (3/1991), reported ultrasound piggybacked on a commercial radio broadcast (100 MHz) aimed at entraining the brains of Iraqi troops and creating feelings of despair. U.S. Pat. No. 5,159,703 that refers to a "silent communications system in which nonaural carriers, in the very low or very high audio frequency range or in the adjacent ultrasonic frequency spectrum, are amplitude or frequency modulated with the desired intelligence and propagated acoustically or vibrationally, for inducement into the brain, typically through the use of loudspeakers, earphones or piezoelectric transducers." See: Dr Nick Begich—Controlling the Human Mind, Earth Pulse Press Anchorage—isbn=1-890693-54-5 cbcg.org/gjcs1.htm %7C God's Judgment Cometh Soon cnslab.ss.uci.edu/muri/research.html, #Dewan, #FarwellDonchin, #ImaginedSpeechProduction, #Overview, MURI: Synthetic Telepathy daprocess.com/01.welcome.html DaProcess of A Federal Investigation deepthought.newsvine.com/_news/2012/01/01/9865851-nsa-disinformation-watch-the-watchers-with-me deepthought.newsvine.com/_news/2012/01/09/10074589-nsa-disinformation-watch-the-watchers-with-me-part-2 deepthought.newsvine.com/_news/2012/01/16/10169491-the-nsa-behind-the-curtain genamason.wordpress.com/2009/10/18/more-on-synthetic-telepathy/ io9.com/5065304/tips-and-tricks-for-mind-control-from-the-us-military newdawnmagazine.com.au/Article/Brain_Zapping_Part_One.html pinktentacle.com/2008/12/scientists-extract-images-directly-from-brain/ Scientists extract images directly from brain timesofindia.indiatimes.com/HealthSci/US_army_developing_synthetic_telepathy/ www.bibliotecapleyades.net/ciencia/ciencia_nonlethalweapons02.htm Eleanor White—New Devices That 'Talk' To The Human Mind Need Debate, Controls www.cbsnews.com/stories/2008/12/31/60_minutes/main4694713.shtml 60 Minutes: Incredible Research Lets Scientists Get A Glimpse At Your Thoughts www.cbsnews.com/video/watch/?id=5119805n&tag=related;photovideo 60 Minutes: Video—Mind Reading www.charlesrehn.com/charlesrehn/books/aconversationwithamerica/essays/myessays/The%20NSA.doc www.govtrack.us/congress/billtext.xpd?bill=h107-2977 Space Preservation Act of 2001 www.informaworld.com/smpp/content?db=all&content=a785359968 Partial Amnesia for a Narrative Following Application of Theta Frequency Electromagnetic Fields www.msnbc.msn.com/id/27162401/ www.psychology.nottingham.ac.uk/staff/lpxdts/TMS%20info.html Transcranial Magnetic Stimulation www.raven1.net/silsoun2.htm Psy-Ops Weaponry Used In The Persian Gulf War www.scribd.com/doc/24531011/Operation-Mind-Control www.scribd.com/doc/6508206/synthetic-telepathy-and-the-early-mind-wars www.slavery.org.uk/Bioeffects_of_Selected_Non-Lethal_Weapons.pdf—Bioeffects of selected non-lethal weapons www.sst.ws/tempstandards.php?pab=1_1 TEMPEST measurement standards www.uwe.ac.uk/hlss/research/cpps/Journal_Psycho-Social_Studies/v2-2/SmithC.shtml Journal of Psycho-Social Studies—Vol 2 (2) 2003—On the Need for New Criteria of Diagnosis of Psychosis in the Light of Mind Invasive Technology by Dr. Carole Smith www.wired.com/dangerroom/2009/05/pentagon-preps-soldier-telepathy-push www.wired.com/wired/archive/7.11/persinger.html This Is Your Brain on God Noah, Shachtman—Pentagon's PCs Bend to Your Brain www.wired.com/dangerroom/2007/03/the_us_military Soldier-Telepathy" Drummond, Katie—Pentagon Preps Soldier Telepathy Push U.S. Pat. No. 3,951,134 U.S. Pat. No. 5,159,703 Silent subliminal presentation system U.S. Pat. No. 6,011,991 U.S. Pat. No. 6,587,729 Apparatus for audibly communicating speech using the radio frequency hearing effect Wall, Judy, "Military Use of Mind Control Weapons", NEXUS, 5/06, October-November 1998

(165) It is known to analyze EEG patterns to extract an indication of certain volitional activity (U.S. Pat. No. 6,011,991). This technique describes that an EEG recording can be matched against a stored normalized signal using a computer. This matched signal is then translated into the corresponding reference. The patent application describes a method "a system capable of identifying particular nodes in an individual's brain, the firings of which affect characteristics such as appetite, hunger, thirst, communication skills" and "devices mounted to the person (e.g. underneath the scalp) may be energized in a predetermined manner or sequence to remotely cause particular identified brain node(s) to be fired in order to cause a predetermined feeling or reaction in the individual" without technical description of implementation. This patent also describes, that "brain activity (is monitored) by way of electroencephalograph (EEG) methods, magnetoencephalograph (MEG) methods, and the like. For example, see U.S. Pat. Nos. 5,816,247 and 5,325,862.

(166) See also, U.S. Pat. Nos. 3,951,134; 4,437,064; 4,591,787; 4,613,817; 4,689,559; 4,693,000; 4,700,135; 4,733,180; 4,736,751; 4,749,946; 4,753,246; 4,761,611; 4,771,239; 4,801,882; 4,862,359; 4,913,152; 4,937,525; 4,940,058; 4,947,480; 4,949,725; 4,951,674; 4,974,602; 4,982,157; 4,983,912; 4,996,479; 5,008,622; 5,012,190; 5,020,538; 5,061,680; 5,092,835; 5,095,270; 5,126,315; 5,158,932; 5,159,703; 5,159,928; 5,166,614; 5,187,327; 5,198,977; 5,213,338; 5,241,967; 5,243,281; 5,243,517; 5,263,488; 5,265,611; 5,269,325; 5,282,474; 5,283,523; 5,291,888; 5,303,705; 5,307,807; 5,309,095; 5,311,129; 5,323,777; 5,325,862; 5,326,745; 5,339,811; 5,417,211; 5,418,512; 5,442,289; 5,447,154; 5,458,142; 5,469,057; 5,476,438; 5,496,798; 5,513,649; 5,515,301; 5,552,375; 5,579,241; 5,594,849; 5,600,243; 5,601,081; 5,617,856; 5,626,145; 5,656,937; 5,671,740; 5,682,889; 5,701,909; 5,706,402; 5,706,811; 5,729,046; 5,743,854; 5,743,860; 5,752,514; 5,752,911; 5,755,227; 5,761,332; 5,762,611; 5,767,043; 5,771,261; 5,771,893; 5,771,894; 5,797,853; 5,813,993; 5,815,413; 5,842,986; 5,857,978; 5,885,976; 5,921,245; 5,938,598; 5,938,688; 5,970,499; 6,002,254; 6,011,991; 6,023,161; 6,066,084; 6,069,369; 6,080,164; 6,099,319; 6,144,872; 6,154,026; 6,155,966; 6,167,298; 6,167,311; 6,195,576; 6,230,037; 6,239,145; 6,263,189; 6,290,638; 6,354,087; 6,356,079; 6,370,414; 6,374,131; 6,385,479; 6,418,344; 6,442,948; 6,470,220; 6,488,617; 6,516,246; 6,526,415; 6,529,759; 6,538,436; 6,539,245; 6,539,263; 6,544,170; 6,547,746; 6,557,558; 6,587,729; 6,591,132; 6,609,030; 6,611,698; 6,648,822; 6,658,287; 6,665,552; 6,665,553; 6,665,562; 6,684,098; 6,687,525; 6,695,761; 6,697,660; 6,708,051; 6,708,064; 6,708,184; 6,725,080; 6,735,460; 6,774,929; 6,785,409; 6,795,724; 6,804,661; 6,815,949; 6,853,186; 6,856,830; 6,873,872; 6,876,196; 6,885,192; 6,907,280; 6,926,921; 6,947,790; 6,978,179; 6,980,863; 6,983,184; 6,983,264; 6,996,261; 7,022,083; 7,023,206; 7,024,247; 7,035,686; 7,038,450; 7,039,266; 7,039,547; 7,053,610; 7,062,391; 7,092,748; 7,105,824; 7,116,102; 7,120,486; 7,130,675; 7,145,333; 7,171,339; 7,176,680; 7,177,675; 7,183,381; 7,186,209; 7,187,169; 7,190,826; 7,193,413; 7,196,514; 7,197,352; 7,199,708; 7,209,787; 7,218,104; 7,222,964; 7,224,282; 7,228,178; 7,231,254; 7,242,984; 7,254,500; 7,258,659; 7,269,516; 7,277,758; 7,280,861; 7,286,871; 7,313,442; 7,324,851; 7,334,892; 7,338,171; 7,340,125; 7,340,289; 7,346,395; 7,353,064; 7,353,065; 7,369,896; 7,371,365; 7,376,459; 7,394,246; 7,400,984; 7,403,809; 7,403,820; 7,409,321; 7,418,290; 7,420,033; 7,437,196; 7,440,789; 7,453,263; 7,454,387; 7,457,653; 7,461,045; 7,462,155; 7,463,024; 7,466,132; 7,468,350; 7,482,298; 7,489,964; 7,502,720; 7,539,528; 7,539,543; 7,553,810; 7,565,200; 7,565,809; 7,567,693; 7,570,054; 7,573,264; 7,573,268; 7,580,798; 7,603,174; 7,608,579; 7,613,502; 7,613,519; 7,613,520; 7,620,456; 7,623,927; 7,623,928; 7,625,340; 7,627,370; 7,647,098; 7,649,351; 7,653,433; 7,672,707; 7,676,263; 7,678,767; 7,697,979; 7,706,871; 7,715,894; 7,720,519; 7,729,740; 7,729,773; 7,733,973; 7,734,340; 7,737,687; 7,742,820; 7,746,979; 7,747,325; 7,747,326; 7,747,551; 7,756,564; 7,763,588; 7,769,424; 7,771,341; 7,792,575; 7,800,493; 7,801,591; 7,801,686; 7,831,305; 7,834,627; 7,835,787; 7,840,039; 7,840,248; 7,840,250; 7,853,329; 7,856,264; 7,860,552; 7,873,411; 7,881,760; 7,881,770; 7,882,135; 7,891,814; 7,892,764; 7,894,903; 7,895,033; 7,904,139; 7,904,507; 7,908,009; 7,912,530; 7,917,221; 7,917,225; 7,929,693; 7,930,035; 7,932,225; 7,933,727; 7,937,152; 7,945,304; 7,962,204; 7,974,787;

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(167) Brain entrainment Brain entrainment, also referred to as brainwave synchronization and neural entrainment, refers to the capacity of the brain to naturally synchronize its brainwave frequencies with the rhythm of periodic external stimuli, most commonly auditory, visual, or tactile.

Brainwave entrainment technologies are used to induce various brain states, such as relaxation or sleep, by creating stimuli that occur at regular, periodic intervals to mimic electrical cycles of the brain during the desired states, thereby “training” the brain to consciously alter states. Recurrent acoustic frequencies, flickering lights, or tactile vibrations are the most common examples of stimuli applied to generate different sensory responses. It is hypothesized that listening to these beats of certain frequencies one can induce a desired state of consciousness that corresponds with specific neural activity. Patterns of neural firing, measured in Hz, correspond with alertness states such as focused attention, deep sleep, etc.

(168) The term “entrainment” has been used to describe a shared tendency of many physical and biological systems to synchronize their periodicity and rhythm through interaction. This tendency has been identified as specifically pertinent to the study of sound and music generally, and acoustic rhythms specifically. The most ubiquitous and familiar examples of neuromotor entrainment to acoustic stimuli is observable in spontaneous foot or finger tapping to the rhythmic beat of a song. Exogenous rhythmic entrainment, which occurs outside the body, has been identified and documented for a variety of human activities, which include the way people adjust the rhythm of their speech patterns to those of the subject with whom they communicate, and the rhythmic unison of an audience clapping. Even among groups of strangers, the rate of breathing, locomotive and subtle expressive motor movements, and rhythmic speech patterns have been observed to synchronize and entrain, in response to an auditory stimulus, such as a piece of music with a consistent rhythm. Furthermore, motor synchronization to repetitive tactile stimuli occurs in animals, including cats and monkeys as well as humans, with accompanying shifts in electroencephalogram (EEG) readings. Examples of endogenous entrainment, which occurs within the body, include the synchronizing of human circadian sleep-wake cycles to the 24-hour cycle of light and dark, and the frequency following response of humans to sounds and music.

(169) Neural oscillations Neural oscillations are rhythmic or repetitive electrochemical activity in the brain and central nervous system. Such oscillations can be characterized by their frequency, amplitude and phase. Neural tissue can generate oscillatory activity driven by mechanisms within individual neurons, as well as by interactions between them. They may also adjust frequency to synchronize with the periodic vibration of external acoustic or visual stimuli. The functional role of neural oscillations is still not fully understood; however, they have been shown to correlate with emotional responses, motor control, and a number of cognitive functions including information transfer, perception, and memory. Specifically, neural oscillations, in particular theta activity, are extensively linked to memory function, and coupling between theta and gamma activity is considered to be vital for memory functions, including episodic memory. Electroencephalography (EEG) has been most widely used in the study of neural activity generated by large groups of neurons, known as neural ensembles, including investigations of the changes that occur in electroencephalographic profiles during cycles of sleep and wakefulness. EEG signals change dramatically during sleep and show a transition from faster frequencies to increasingly slower frequencies, indicating a relationship between the frequency of neural oscillations and cognitive states including awareness and consciousness.

(170) Brainwaves, or neural oscillations, share the fundamental constituents with acoustic and optical waves, including frequency, amplitude and periodicity. The synchronous electrical activity of cortical neural ensembles can synchronize in response to external acoustic or optical stimuli and also entrain or synchronize their frequency and phase to that of a specific stimulus. Brainwave entrainment is a colloquialism for such ‘neural entrainment’, which is a term used to denote the way in which the aggregate frequency of oscillations produced by the synchronous electrical activity in ensembles of cortical neurons can adjust to synchronize with the periodic vibration of an external stimuli, such as a sustained acoustic frequency perceived as pitch, a regularly repeating pattern of intermittent sounds, perceived as rhythm, or of a regularly rhythmically intermittent flashing light.

(171) Changes in neural oscillations, demonstrable through electroencephalogram (EEG) measurements, are precipitated by listening to music, which can modulate autonomic arousal ergotropically and trophotropically, increasing and decreasing arousal respectively. Musical auditory stimulation has also been demonstrated to improve immune function, facilitate relaxation, improve mood, and contribute to the alleviation of stress.

(172) The Frequency following response (FFR), also referred to as Frequency Following Potential (FFP), is a specific response to hearing sound and music, by which neural oscillations adjust their frequency to match the rhythm of auditory stimuli. The use of sound with intent to influence cortical brainwave frequency is called auditory driving, by which frequency of neural oscillation is ‘driven’ to entrain with that of the rhythm of a sound source.

(173) See, en.wikipedia.org/wiki/Brainwave_entrainment;

(174) U.S. Pat. Nos. 5,070,399; 5,306,228; 5,409,445; 6,656,137; 7,749,155; 7,819,794; 7,988,613; 8,088,057; 8,167,784; 8,213,670; 8,267,851; 8,298,078; 8,517,909; 8,517,912; 8,579,793; 8,579,795; 8,597,171; 8,636,640; 8,638,950; 8,668,496; 8,852,073; 8,932,218; 8,968,176; 9,330,523; 9,357,941; 9,459,597; 9,480,812; 9,563,273; 9,609,453; 9,640,167; 9,707,372; 20050153268; 20050182287; 20060106434; 20060206174; 20060281543; 20070066403; 20080039677; 20080304691; 20100010289; 20100010844; 20100028841; 20100056854; 20100076253; 20100130812; 20100222640; 20100286747; 20100298624; 20110298706; 20110319482; 20120003615; 20120053394; 20120150545; 20130030241; 20130072292; 20130131537; 20130172663; 20130184516; 20130203019; 20130234823; 20130338738; 20140088341; 20140107401; 20140114242; 20140154647; 20140174277; 20140275741; 20140309484; 20140371516; 20150142082; 20150283019; 20150296288; 20150313496; 20150313949; 20160008568; 20160019434; 20160055842; 20160205489; 20160235980; 20160239084; 20160345901; 20170034638; 20170061760; 20170087330; 20170094385; 20170095157; 20170099713; 20170135597; and 20170149945. Carter, J., and H. Russell. “A pilot investigation of auditory and visual entrainment of brain wave activity in learning disabled boys.” Texas Researcher 4.1 (1993): 65-75; Casciaro, Francesco, et al. “Alpha-rhythm stimulation using brain entrainment enhances heart rate variability in subjects with reduced HRV.” World J. Neuroscience 3.04 (2013): 213; Helfrich, Randolph F., et al. “Entrainment of brain oscillations by transcranial alternating current stimulation.” Current Biology 24.3 (2014): 333-339; Huang, Tina L., and Christine Charyton. “A comprehensive review of the psychological effects of brainwave entrainment” Alternative therapies in health and medicine 14.5 (2008): 38; Joyce, Michael, and Dave Siever. “Audio-visual entrainment program as a treatment for behavior disorders in a school setting.” J. Neurotherapy 4.2 (2000): 9-25; Keitel, Christian, Clodhna Quigley, and Philipp Ruhnau. “Stimulus-driven brain oscillations in the alpha range: entrainment of intrinsic rhythms or frequency-following response?” J. Neuroscience

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(175) A baseline correction of event-related time-frequency measure may be made to take pre-event baseline activity into consideration. In general, a baseline period is defined by the average of the values within a time window preceding the time-locking event There are at least four common methods for baseline correction in time-frequency analysis. The methods include various baseline value normalizations. See, Spencer K M, Nestor P G, Perlmuter R, et al. Neural synchrony indexes disordered perception and cognition in schizophrenia. *Proc Natl Acad Sci USA*. 2004; 101:17288-17293; Hoogenboom N, Schoffelen J M, Oostenveld R, Parkes L M, Fries P. Localizing human visual gamma-band activity in frequency, time and space. *Neuroimage*. 2006; 29:764-773; Le Van Quyen M, Foucher J, Lachaux J, et al. Comparison of Hilbert transform and wavelet methods for the analysis of neuronal synchrony. *J Neurosci Methods*. 2001; 111:83-98, Lachaux J P, Rodriguez E, Martinerie J, Varela F J. Measuring phase synchrony in brain signals. *Hum Brain Mapp*. 1999; 8:194-208, Rodriguez E, George N, Lachaux J P, Martinerie J, Renault B, Varela F J. Perception's shadow: long-distance synchronization of human brain activity. *Nature*. 1999; 397:430-433., Canolty R T, Edwards E, Dalal S S, et al. High gamma power is phase-locked to theta oscillations in human neocortex. *Science*. 2006; 313:1626-1628.

(176) The question of whether different emotional states are associated with specific patterns of physiological response has long being a subject of neuroscience research See, for example: James W (1884.) What is an emotion? *Mind* 9: 188-205; Lacey J I, Bateman D E, Vanlehn R (1953) Autonomic response specificity; an experimental study. *Psychosom Med* 15: 8-21; Levenson R W, Heider K, Ekman P, Friesen W V (1992) Emotion and Autonomic Nervous-System Activity in the Minangkabau of West Sumatra. *J Pers Soc Psychol* 62: 972-988.

(177) Some studies have indicated that the physiological correlates of emotions are likely to be found in the central nervous system (CNS). See, for example: Buck R (1999) The biological affects: A typology. *Psychological Review* 106: 301-336; Izard C E (2007) Basic Emotions, Natural Kinds, Emotion Schemas, and a New Paradigm. *Perspect Psychol Sci* 2: 260-280; Panksepp J (2007) Neurologizing the Psychology of Affects How Appraisal-Based Constructivism and Basic Emotion Theory Can Coexist *Perspect Psychol Sci* 2: 281-296.

(178) Electroencephalograms (EEG) and functional Magnetic Resonance Imaging, fMRI have been used to study specific brain activity associated with different emotional states. Mauss and Robinson, in their review paper, have indicated that "emotional state is likely to involve circuits rather than any brain region considered in isolation" (Mauss I B, Robinson M D (2009) Measures of emotion: A review. *Cogn Emot* 23: 209-237.)

(179) The amplitude, latency from the stimulus, and covariance (in the case of multiple electrode sites) of each component can be examined in connection with a cognitive task (ERP) or with no task (EP). Steady-state visually evoked potentials (SSVEPs) use a continuous sinusoidally-modulated flickering light, typically superimposed in front of a TV monitor displaying a cognitive task. The brain response in a narrow frequency band containing the stimulus frequency is measured. Magnitude, phase, and coherence (in the case of multiple electrode sites) may be related to different parts of the cognitive task. Brain entrainment may be detected through EEG or MEG activity.

(180) Brain entrainment may be detected through EEG or MEG activity. See: Abeln, Vera, et al. "Brainwave entrainment for better sleep and post-sleep state of young elite soccer players-A pilot study." *European J. Sport science* 14.5 (2014): 393-402; Acton, George. "Methods for independent entrainment of visual field zones." U.S. Pat. No. 9,629,976. 25 Apr. 2017; Albouy, Philippe, et al. "Selective entrainment of theta oscillations in the dorsal stream causally enhances auditory working memory performance." *Neuron* 94.1 (2017): 193-206. Amengual, J., et al. "P018 Local entrainment and distribution across cerebral networks of natural oscillations elicited in implanted epilepsy patients by intracranial stimulation: Paving the way to develop causal connectomics of the healthy human brain." *Clin. Neurophysiology* 128.3 (2017): e18; Argento, Emanuele, et al. "Augmented Cognition via Brainwave Entrainment in Virtual Reality: An Open, Integrated Brain Augmentation in a Neuroscience System Approach." *Augmented Human Research* 2.1(2017): 3; Bello, Nicholas P. "Altering Cognitive and Brain States Through Cortical Entrainment" (2014); Costa-Faidella, Jordi, Elyse S. Sussman, and Caries Escera. "Selective entrainment of brain oscillations drives auditory perceptual organization." *NeuroImage* (2017); Börgers, Christoph. "Entrainment by Excitatory Input Pulses." *An Introduction to Modeling Neuronal Dynamics*. Springer International Publishing, 2017. 183-192; Calderone, Daniel J., et al. "Entrainment of neural oscillations as a modifiable substrate of attention." *Trends in cognitive sciences* 18.6 (2014): 300-309; Casciaro, Francesco, et al. "Alpha-rhythm stimulation using brain entrainment enhances heart rate variability in subjects with reduced HRV." *World J. Neuroscience* 3.04 (2013): 213; Chang, Daniel Wonchul. 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(181) The entrainment hypothesis (Thut and Miniussi, 2009; Thut et al., 2011a, 2012), suggests the possibility of inducing a particular oscillation frequency in the brain using an external oscillatory force (e.g., rTMS, but also tACS). The physiological basis of oscillatory cortical activity lies in the timing of the interacting neurons; when groups of neurons synchronize their firing activities, brain rhythms emerge, network oscillations are generated, and the basis for interactions between brain areas may develop (Buzsáki, 2006). Because of the variety of experimental protocols for brain stimulation, limits on descriptions of the actual protocols employed, and limited controls, consistency of reported studies is lacking, and extrapolability is limited. Thus, while there is various consensus in various aspects of the effects of extra cranial brain stimulation, the results achieved have a degree of uncertainty dependent on details of implementation. On the other hand, within a specific experimental protocol, it is possible to obtain statistically significant and repeatable results. This implies that feedback control might be effective to control implementation of the stimulation for a given purpose; however, studies that employ feedback control are lacking.

(182) Different cognitive states are associated with different oscillatory patterns in the brain (Buzsáki, 2006; Canolty and Knight 2010; Varela et al., 2001). Thut et al. (2011b) directly tested the entrainment hypothesis by means of a concurrent EEG-TMS experiment They first determined the individual source of the parietal-occipital alpha modulation and the individual alpha frequency (magnetoencephalography study). They then applied rTMS at the individual alpha power while recording the EEG activity at rest. The results confirmed the three predictions of the entrainment hypothesis: the induction of a specific frequency after TMS, the enhancement of oscillation during TMS stimulation due to synchronization, and a phase alignment of the induced frequency and the ongoing activity (Thut et al., 2011b).

(183) If associative stimulation is a general principle for human neural plasticity in which the timing and strength of activation are critical factors, it is possible that synchronization within or between areas using an external force to phase/align oscillations can also favor efficient communication and associative plasticity (or alter communication). In this respect associative, cortico-cortical stimulation has been shown to enhance coherence of oscillatory activity between the stimulated areas (Plewnia et al., 2008).

(184) In coherence resonance (Longtin, 1997), the addition of a certain amount of noise in an excitable system results in the most coherent and proficient oscillatory responses. The brain's response to external timing-embedded stimulation can result in a decrease in phase variance and an enhanced alignment (clustering) of the phase components of the ongoing EEG activity (entraining, phase resetting) that can change the signal-to-noise ratio and increase (or decrease) signal efficacy.

(185) If one considers neuron activity within the brain as a set of loosely coupled oscillators, then the various parameters that might be controlled include the size of the region of neurons, frequency of oscillation, resonant frequency or time-constant, oscillator damping, noise, amplitude, coupling to other oscillators, and of course, external influences that may include stimulation and/or power loss. In a human brain, pharmacological intervention may be significant. For example, drugs that alter excitability, such as caffeine, neurotransmitter release and reuptake, nerve conductance, etc. can all influence operation of the neural oscillators. Likewise, sub-threshold external stimulation effects, including DC, AC and magnetic electromagnetic effects, can also influence operation of the neural oscillators.

(186) Phase resetting or shifting can synchronize inputs and favor communication and, eventually, Hebbian plasticity (Hebb, 1949). Thus, rhythmic stimulation may induce a statistically higher degree of coherence in spiking neurons, which facilitates the induction of a specific cognitive process (or hinders that process). Here, the perspective is slightly different (coherence resonance), but the underlining mechanisms are similar to the ones described so far (stochastic resonance), and the additional key factor is the repetition at a specific rhythm of the stimulation.

(187) In the 1970's, the British biophysicist and psychobiologist C. Maxwell Cade, monitored the brainwave patterns of advanced meditators and 300 of his students. Here he found that the most advanced meditators have a specific brainwave pattern that was different from the rest of his students. He noted that these meditators showed high activity of alpha brainwaves accompanied by beta, theta and even delta waves that were about half the amplitude of the alpha waves. See, Cade "The Awakened Mind: Biofeedback and the Development of Higher States of Awareness" (Dell, 1979). Anna Wise extended Cade's studies, and found that extraordinary achievers which included composers, inventors, artists, athletes, dancers, scientists, mathematicians, CEO's and presidents of large corporations have brainwave patterns differ from average performers, with a specific balance between Beta, Alpha, Theta and Delta brainwaves where Alpha had the strongest amplitude. See, Anna Wise, "The High-Performance Mind: Mastering

Brainwaves for Insight, Healing, and Creativity”.

(188) Entrainment is plausible because of the characteristics of the demonstrated EEG responses to a single TMS pulse, which have a spectral composition which resemble the spontaneous oscillations of the stimulated cortex. For example, TMS of the “resting” visual (Rosanova et al., 2009) or motor cortices (Veniero et al., 2011) triggers alpha-waves, the natural frequency at the resting state of both types of cortices. With the entrainment hypothesis, the noise generation framework moves to a more complex and extended level in which noise is synchronized with on-going activity. Nevertheless, the model to explain the outcome will not change, stimulation will interact with the system, and the final result will depend on introducing or modifying the noise level. The entrainment hypothesis makes clear predictions with respect to online repetitive TMS paradigms' frequency engagement as well as the possibility of inducing phase alignment, i.e., a reset of ongoing brain oscillations via external spTMS (Thut et al., 2011a, 2012; Veniero et al., 2011). The entrainment hypothesis is superior to the localization approach in gaining knowledge about how the brain works, rather than where or when a single process occurs. TMS pulses may phase-align the natural, ongoing oscillation of the target cortex. When additional TMS pulses are delivered in synchrony with the phase-aligned oscillation (i.e., at the same frequency), further synchronized phase-alignment will occur, which will bring the oscillation of the target area in resonance with the TMS train. Thus, entrainment may be expected when TMS is frequency-tuned to the underlying brain oscillations (Veniero et al., 2011).

(189) Binaural Beats Binaural beats are auditory brainstem responses which originate in the superior olivary nucleus of each hemisphere. They result from the interaction of two different auditory impulses, originating in opposite ears, below 1000 Hz and which differ in frequency between one and 30 Hz. For example, if a pure tone of 400 Hz is presented to the right ear and a pure tone of 410 Hz is presented simultaneously to the left ear, an amplitude modulated standing wave of 10 Hz, the difference between the two tones, is experienced as the two wave forms mesh in and out of phase within the superior olivary nuclei. This binaural beat is not heard in the ordinary sense of the word (the human range of hearing is from 20-20,000 Hz). It is perceived as an auditory beat and theoretically can be used to entrain specific neural rhythms through the frequency-following response (FFR)—the tendency for cortical potentials to entrain to or resonate at the frequency of an external stimulus. Thus, it is theoretically possible to utilize a specific binaural-beat frequency as a consciousness management technique to entrain a specific cortical rhythm. The binaural-beat appears to be associated with an electroencephalographic (EEG) frequency-following response in the brain.

(190) Uses of audio with embedded binaural beats that are mixed with music or various pink or background sound are diverse. They range from relaxation, meditation, stress reduction, pain management, improved sleep quality, decrease in sleep requirements, super learning, enhanced creativity and intuition, remote viewing, telepathy, and out-of-body experience and lucid dreaming. Audio embedded with binaural beats is often combined with various meditation techniques, as well as positive affirmations and visualization.

(191) When signals of two different frequencies are presented, one to each ear, the brain detects phase differences between these signals. “Under natural circumstances a detected phase difference would provide directional information. The brain processes this anomalous information differently when these phase differences are heard with stereo headphones or speakers. A perceptual integration of the two signals takes place, producing the sensation of a third “beat” frequency. The difference between the signals waxes and wanes as the two different input frequencies mesh in and out of phase. As a result of these constantly increasing and decreasing differences, an amplitude-modulated standing wave—the binaural beat—is heard. The binaural beat is perceived as a fluctuating rhythm at the frequency of the difference between the two auditory inputs. Evidence suggests that the binaural beats are generated in the brainstem's superior olivary nucleus, the first site of contralateral integration in the auditory system. Studies also suggest that the frequency-following response originates from the inferior colliculus. This activity is conducted to the cortex where it can be recorded by scalp electrodes. Binaural beats can easily be heard at the low frequencies (<30 Hz) that are characteristic of the EEG spectrum.

(192) Synchronized brainwaves have long been associated with meditative and hypnagogic states, and audio with embedded binaural beats has the ability to induce and improve such states of consciousness. The reason for this is physiological. Each ear is “hardwired” (so to speak) to both hemispheres of the brain. Each hemisphere has its own olivary nucleus (sound-processing center) which receives signals from each ear. In keeping with this physiological structure, when a binaural beat is perceived there are actually two standing waves of equal amplitude and frequency present, one in each hemisphere. So, there are two separate standing waves entraining portions of each hemisphere to the same frequency. The binaural beats appear to contribute to the hemispheric synchronization evidenced in meditative and hypnagogic states of consciousness. Brain function is also enhanced through the increase of cross-collosal communication between the left and right hemispheres of the brain. [en.wikipedia.org/wiki/Beat \(acoustics\)#Binaural_beats](http://en.wikipedia.org/wiki/Beat_(acoustics)#Binaural_beats). See: Oster, G (October 1973). “Auditory beats in the brain”. *Scientific American*. 229 (4): 94-102. See: Lane, J. D., Kasian, S. J., Owens, J. E., & Marsh, G. R. (1998). Binaural auditory beats affect vigilance performance and mood. *Physiology & behavior*, 63(2), 249-252; Foster, D. S. (1990). EEG and subjective correlates of alpha frequency binaural beats stimulation combined with alpha biofeedback (Doctoral dissertation, Memphis State University); Kasprzak, C. (2011). Influence of binaural beats on EEG signal. *Acta Physica Polonica A*, 119(6A), 986-990; Pratt, H., Starr, A., Michalewski, H. J., Dimitijevic, A., Bleich, N., & Mittelman, N. (2009). Cortical evoked potentials to an auditory illusion: binaural beats. *Clinical Neurophysiology*, 120(8), 1514-1524; Pratt, H., Starr, A., Michalewski, H. J., Dimitijevic, A., Bleich, N., & Mittelman, N. (2010). A comparison of auditory evoked potentials to acoustic beats and to binaural beats. *Hearing research*, 262(1), 34-44; Padmanabhan, R., Hildreth, A. J., & Laws, D. (2005). A prospective, randomised, controlled study examining binaural beat audio and pre-operative anxiety in patients undergoing general anaesthesia for day case surgery. *Anaesthesia*, 60(9), 874-877; Reedijk, S. A., Bolders, A., & Hommel, B. (2013). The impact of binaural beats on creativity. *Frontiers in human neuroscience*, 7; Atwater, F. H. (2001). Binaural beats and the regulation of arousal levels. *Proceedings of the TANS*, 11; Hink, R. F., Koder, K., Yamada, O., Kaga, K., & Suzuki, J. (1980). Binaural interaction of a beating frequency-following response. *Audiology*, 19(1), 36-43; Gao, X., Cao, H., Ming, D., Qi, H., Wang, X., Wang, X., & Zhou, P. (2014). Analysis of EEG activity in response to binaural beats with different frequencies. *International Journal of Psychophysiology*, 94(3), 399-406; Sung, H. C., Lee, W. L., Li, H. M., Lin, C. Y., Wu, Y. Z., Wang, J. J., & Li, T. L. (2017). Familiar Music Listening with Binaural Beats for Older People with Depressive Symptoms in Retirement Homes. *Neuropsychiatry*, 7(4); Colzato, L. S., Barone, H., Sam, R., & Hommel, B. (2017). More attentional focusing through binaural beats: evidence from the global-local task. *Psychological research*, 81(1), 271-277; Mortazavi, S. M. J., Zahraei-Moghadam, S. M., Masoumi, S., Rafati, A., Haghani, M., Mortazavi, S. A. R., & Zehtabian, M. (2017). Short Term Exposure to Binaural Beats Adversely Affects Learning and Memory in Rats. *Journal of Biomedical Physics and Engineering*.

(193) Brain Entrainment Frequency Following Response (or FFR). See, “Stimulating the Brain with Light and Sound,” Transparent Corporation, Neuroprogrammer™ 3, www.transparentcorp.com/products/np/entrainment.php.

(194) Isochronic Tones Isochronic tones are regular beats of a single tone that are used alongside monaural beats and binaural beats in the process called brainwave entrainment. At its simplest level, an isochronic tone is a tone that is being turned on and off rapidly. They create sharp, distinctive pulses of sound. www.livingflow.net/isochronic-tones-work/; Schulze, H. H. (1989). The perception of temporal deviations in isochronic patterns. *Attention, Perception, & Psychophysics*, 45(4), 291-296; Oster, G. (1973). Auditory beats in the brain. *Scientific American*, 229(4), 94-102; Huang, T. L., & Charyton, C. (2008). A comprehensive review of the psychological effects of brainwave entrainment Alternative therapies in health and medicine, 14(5), 38; Trost W., Frühholz, S., Schön, D., Labbé, C., Pichon, S., Grandjean, D., & Vuilleumier, P. (2014). Getting the beat: entrainment of brain activity by musical rhythm and pleasantness. *NeuroImage*, 103, 55-64; Casciaro, F., Laterza, V., Conte, S., Pieralice, M., Federici, A., Todarello, O., . . . & Conte, E. (2013). Alpha-rhythm stimulation using brain entrainment enhances heart rate variability in subjects with reduced HRV. *World Journal of Neuroscience*, 3(04), 213; Conte, Elio, Sergio Conte, Nunzia Santacroce, Antonio Federici, Orlando Todarello, Franco Orsucci, Francesco Casciaro, and Vincenza Laterza. “A Fast Fourier Transform analysis of time series data of heart rate variability during alfa-rhythm stimulation in brain entrainment” *NeuroQuantology* 11, no. 3 (2013); Doherty, C. (2014). A comparison of alpha brainwave entrainment, with and without musical accompaniment Moseley, R. (2015, July). Inducing targeted brain states utilizing merged reality systems. In *Science and Information Conference (SAI)*, 2015 (pp. 657-663). IEEE.

(195) Time-Frequency Analysis Brian J. Roach and Daniel H. Mathalon, "Event-related EEG time-frequency analysis: an overview of measures and analysis of early gamma band phase locking in schizophrenia. Schizophrenia Bull. USA. 2008; 34:5:907-926., describes a mechanism for EEG time-frequency analysis. Fourier and wavelet transforms (and their inverse) may be performed on EEG signals.

(196) See, U.S. Pat. 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(197) There are many approaches to lime-frequency decomposition of EEG data, including the short-term Fourier transform (STFT), (Gabor D. Theory of Communication. J. Inst Elect. Engrs. 1946; 93:429-457) continuous (Daubechies I. Ten Lectures on Wavelets. Philadelphia, Pa: Society for Industrial and Applied Mathematics; 1992:357. 21. Combes J M, Grossmann A, Tchamitchian P. Wavelets: Time-Frequency Methods and Phase Space-Proceedings of the International Conference; Dec. 14-18, 1987; Marseille, France) or discrete (Mallet S G. A theory for multiresolution signal decomposition: the wavelet representation. IEEE Trans Pattern Anal Mach Intell. 1989; 11:674-693) wavelet transforms, Hilbert transform (Lyons R G. Understanding Digital Signal Processing. 2nd ed. Upper Saddle River, NJ: Prentice Hall PTR; 2004:688), and matching pursuits (Mallat S, Zhang Z. Matching pursuits with time-frequency dictionaries. IEEE Trans. Signal Proc. 1993; 41(12):3397-3415). Prototype analysis systems may be implemented using, for example, MatLab with the Wavelet Toolbox, www.mathworks.com/products/wavelet.html.

(198) See, U.S. Pat. Nos. 6,196,972; 6,338,713; 6,442,421; 6,507,754; 6,524,249; 6,547,736; 6,616,611; 6,816,744; 6,865,494; 6,915,241; 6,936,012; 6,996,261; 7,043,293; 7,054,454; 7,079,977; 7,128,713; 7,146,211; 7,149,572; 7,164,941; 7,209,788; 7,254,439; 7,280,867; 7,282,030; 7,321,837; 7,330,032; 7,333,619; 7,381,185; 7,537,568; 7,559,903; 7,565,193; 7,567,693; 7,604,603; 7,624,293; 7,640,055; 7,715,919; 7,725,174; 7,729,755; 7,751,878; 7,778,693; 7,794,406; 7,797,040; 7,801,592; 7,803,118; 7,803,119; 7,879,043; 7,896,807; 7,899,524; 7,917,206; 7,933,646; 7,937,138; 7,976,465; 8,014,847; 8,033,996; 8,073,534; 8,095,210; 8,137,269; 8,137,270; 8,175,696; 8,177,724; 8,177,726; 8,180,601; 8,187,181; 8,197,437; 8,233,965; 8,236,005; 8,244,341; 8,248,069; 8,249,698; 8,280,514; 8,295,914; 8,326,433; 8,335,664; 8,346,342; 8,355,768; 8,386,312; 8,386,313; 8,392,250; 8,392,253; 8,392,254; 8,392,255; 8,396,542; 8,406,841; 8,406,862; 8,412,655; 8,428,703; 8,428,704; 8,463,374; 8,464,288; 8,475,387; 8,483,815; 8,494,610; 8,494,829; 8,494,905; 8,498,699; 8,509,881; 8,533,042; 8,548,786; 8,571,629; 8,579,786; 8,591,419; 8,606,360; 8,628,480; 8,655,428; 8,666,478; 8,682,422; 8,706,183; 8,706,205; 8,718,747; 8,725,238; 8,738,136; 8,747,382; 8,755,877; 8,761,869; 8,762,202; 8,768,449; 8,781,796; 8,790,255; 8,790,272; 8,821,408; 8,825,149; 8,831,731; 8,843,210; 8,849,392; 8,849,632; 8,855,773; 8,858,440; 8,862,210; 8,862,581; 8,903,479; 8,918,178; 8,934,965; 8,951,190; 8,954,139; 8,955,010; 8,958,868; 8,983,628; 8,983,629; 8,989,835; 9,020,789; 9,026,217; 9,031,644; 9,050,470; 9,060,671; 9,070,492; 9,072,832; 9,072,905; 9,078,584; 9,084,896; 9,095,295; 9,101,276; 9,107,595; 9,116,835; 9,125,574; 9,149,719; 9,155,487; 9,192,309; 9,198,621; 9,204,835; 9,211,417; 9,215,978; 9,232,910; 9,232,984; 9,238,142; 9,242,067; 9,247,911; 9,248,286; 9,254,383;

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(199) Single instruction, multiple data processors, such as graphic processing units including the nVidia CUDA environment or AMD Firepro high-performance computing environment are known, and may be employed for general purpose computing, finding particular application in data matrix transformations.

(200) See, U.S. Pat. Nos. 5,273,038; 5,503,149; 6,240,308; 6,272,370; 6,298,259; 6,370,414; 6,385,479; 6,490,472; 6,556,695; 6,697,660; 6,801,648; 6,907,280; 6,996,261; 7,092,748; 7,254,500; 7,338,455; 7,346,382; 7,490,085; 7,497,828; 7,539,528; 7,565,193; 7,567,693; 7,577,472; 7,597,665; 7,627,370; 7,680,526; 7,729,755; 7,809,434; 7,840,257; 7,860,548; 7,872,235; 7,899,524; 7,904,134; 7,904,139; 7,907,998; 7,983,740; 7,983,741; 8,000,773; 8,014,847; 8,069,125; 8,233,682; 8,233,965; 8,235,907; 8,248,069; 8,356,004; 8,379,952; 8,406,838; 8,423,125; 8,445,851; 8,553,956; 8,586,932; 8,606,349; 8,615,479; 8,644,910; 8,679,009; 8,696,722; 8,712,512; 8,718,747; 8,761,866; 8,781,557; 8,814,923; 8,821,376; 8,834,546; 8,852,103; 8,870,737; 8,936,630; 8,951,189; 8,951,192; 8,958,882; 8,983,155; 9,005,126; 9,020,586; 9,022,936; 9,028,412; 9,033,884; 9,042,958; 9,078,584; 9,101,279; 9,135,400; 9,144,392; 9,149,255; 9,155,521; 9,167,970; 9,179,854; 9,179,858; 9,198,637; 9,204,835; 9,208,558; 9,211,077; 9,213,076; 9,235,685; 9,242,067; 9,247,924; 9,268,014; 9,268,015; 9,271,651; 9,271,674; 9,275,191; 9,292,920; 9,307,925; 9,322,895; 9,326,742; 9,330,206; 9,368,265; 9,395,425; 9,402,558; 9,414,776; 9,436,989; 9,451,883; 9,451,899; 9,468,541; 9,471,978; 9,480,402; 9,480,425; 9,486,168; 9,592,389; 9,615,789; 9,626,756; 9,672,302; 9,672,617; 9,682,232; 20020033454; 20020035317; 20020037095; 20020042563; 20020058867; 20020103428; 20020103429; 20030018277; 20030093004; 20030128801; 20040082862; 20040092809; 20040096395; 20040116791; 20040116798; 20040122787; 20040122790; 20040166536; 20040215082; 20050007091; 20050020918; 20050033154; 20050079636; 20050119547; 20050154290; 20050222639; 20050240253; 20050283053; 20060036152; 20060036153; 20060036154; 20060036155; 20060036156; 20060036157; 20060074290; 20060078183; 20060084858; 20060149160; 20060161218; 20060241382; 20060241718; 20070191704; 20070239059; 20080001600; 20080009772; 20080033291; 20080039737; 20080042067; 20080097235; 20080097785; 20080128626; 20080154126; 20080221441; 20080228077; 20080228239; 20080230702; 20080230705; 20080249430; 20080262327; 20080275340; 20090012387; 20090018407; 20090022825; 20090024050; 20090062660; 20090078875; 20090118610; 20090156907; 20090156955; 20090157323; 20090157481; 20090157482; 20090157625; 20090157751; 20090157813; 20090163777; 20090164131; 20090164132; 20090171164; 20090172540; 20090179642; 20090209831; 20090221930; 20090246138; 20090299169; 20090304582; 20090306532; 20090306534; 20090312808; 20090312817; 20090318773; 20090318794; 20090322331; 20090326604; 20100021378; 20100036233; 20100041949; 20100042011; 20100049482; 20100069739; 20100069777; 20100082506; 20100113959; 20100249573; 20110015515; 20110015539; 20110028827; 20110077503; 20110118536; 201101125077; 20110125078; 20110129129; 20110160543; 20110161011; 20110172509; 20110172553; 20110178359; 20110190846; 20110218405; 20110224571; 20110230738; 20110257519; 20110263962; 20110263968; 20110270074; 20110288400; 20110301448; 20110306845; 20110306846; 20110313274; 20120021394; 20120022343; 20120035433; 20120053483; 20120163689; 20120165904; 20120215114; 20120219195; 20120219507; 20120245474; 20120253261; 20120253434; 20120289854; 20120310107; 20120316793; 20130012804; 20130060125; 20130063550; 20130085678; 20130096408; 20130110616; 20130116561; 20130123607; 20130131438; 20130131461; 20130178693; 20130178733; 20130184558; 20130211238; 20130221961; 20130245424; 20130274586; 20130289385; 20130289386;

20130303934; 20140058528; 20140066763; 20140119621; 20140151563; 20140155730; 20140163368; 20140171757; 20140180088; 20140180092; 20140180093; 20140180094; 20140180095; 20140180096; 20140180097; 20140180099; 20140180100; 20140180112; 20140180113; 20140180176; 20140180177; 20140184550; 20140193336; 20140200414; 20140243614; 20140257047; 20140275807; 20140303486; 20140315169; 20140316248; 20140323849; 20140335489; 20140343397; 20140343399; 20140343408; 20140364721; 20140378830; 20150011866; 20150038812; 20150051663; 20150099959; 20150112409; 20150119658; 20150119689; 20150148700; 20150150473; 20150196800; 20150200046; 20150219732; 20150223905; 20150227702; 20150247921; 20150248615; 20150253410; 20150289779; 20150290453; 20150290454; 20150313540; 20150317796; 20150324692; 20150366482; 20150375006; 20160005320; 20160027342; 20160029965; 20160051161; 20160051162; 20160055304; 20160058304; 20160058392; 20160066838; 20160103487; 20160120437; 20160120457; 20160143541; 20160157742; 20160184029; 20160196393; 20160228702; 20160231401; 20160239966; 20160239968; 20160260216; 20160267809; 20160270723; 20160302720; 20160303397; 20160317077; 20160345911; 20170027539; 20170039706; 20170045601; 20170061034; 20170085855; 20170091418; 20170112403; 20170113046; 20170120041; 20170160360; 20170164861; 20170169714; 20170172527; and 20170202475.

(201) Statistical analysis may be presented in a form that permits parallelization, which can be efficiently implemented using various parallel processors, a common form of which is a SIMD (single instruction, multiple data) processor, found in typical graphics processors (GPUs).

(202) See, U.S. Pat. Nos. 8,406,890; 8,509,879; 8,542,916; 8,852,103; 8,934,986; 9,022,936; 9,028,412; 9,031,653; 9,033,884; 9,037,530; 9,055,974; 9,149,255; 9,155,521; 9,198,637; 9,247,924; 9,268,014; 9,268,015; 9,367,131; 9,414,780; 9,420,970; 9,430,615; 9,442,525; 9,444,998; 9,445,763; 9,462,956; 9,474,481; 9,489,854; 9,504,420; 9,510,790; 9,519,981; 9,526,906; 9,538,948; 9,585,581; 9,622,672; 9,641,665; 9,652,626; 9,684,335; 9,687,187; 9,693,684; 9,693,724; 9,706,963; 9,712,736; 20090118622; 20100098289; 20110066041; 20110066042; 20110098583; 20110301441; 20120130204; 20120265271; 20120321759; 20130060158; 20130113816; 20130131438; 20130184786; 20140031889; 20140031903; 20140039975; 20140114889; 20140226131; 20140279341; 20140296733; 20140303424; 20140313303; 20140315169; 20140316235; 20140364721; 20140378810; 20150003698; 20150003699; 20150005640; 20150005644; 20150006186; 20150029087; 20150033245; 20150033258; 20150033259; 20150033262; 20150033266; 20150081226; 20150088093; 20150093729; 20150105701; 20150112899; 20150126845; 20150150122; 20150190062; 20150190070; 20150190077; 20150190094; 20150192776; 20150196213; 20150196800; 20150199010; 20150241916; 20150242608; 20150272496; 20150272510; 20150282705; 20150282749; 20150289217; 20150297109; 20150305689; 20150335295; 20150351655; 20150366482; 20160027342; 20160029896; 20160058366; 20160058376; 20160058673; 20160060926; 20160065724; 20160065840; 20160077547; 20160081625; 20160103487; 20160104006; 20160109959; 20160113517; 20160120048; 20160120428; 20160120457; 20160125228; 20160157773; 20160157828; 20160183812; 20160191517; 20160193499; 20160196185; 20160196635; 20160206241; 20160213317; 20160228064; 20160235341; 20160235359; 20160249857; 20160249864; 20160256086; 20160262680; 20160262685; 20160270656; 20160278672; 20160282113; 20160287142; 20160306942; 20160310071; 20160317056; 20160324/45; 20160324457; 20160342241; 20160360100; 20160361027; 20160366462; 20160367138; 20160367195; 20160374616; 20160378608; 20160378965; 20170000324; 20170000325; 20170000326; 20170000329; 20170000330; 20170000331; 20170000332; 20170000333; 20170000334; 20170000335; 20170000337; 20170000340; 20170000341; 20170000342; 20170000343; 20170000345; 20170000454; 20170000683; 20170001032; 20170007111; 20170007115; 20170007116; 20170007122; 20170007123; 20170007182; 20170007450; 20170007799; 20170007843; 20170010469; 20170010470; 20170013562; 20170017083; 20170020627; 20170027521; 20170028563; 20170031440; 20170032221; 20170035309; 20170035317; 20170041699; 20170042485; 20170046052; 20170065349; 20170086695; 20170086727; 20170090475; 20170103440; 20170112446; 20170113056; 20170128006; 20170143249; 20170143442; 20170156593; 20170156606; 20170164893; 20170171441; 20170172499; 20170173262; 20170185714; 20170188933; 20170196503; 20170205259; 20170206913; and 20170214786.

(203) Artificial neural networks have been employed to analyze EEG signals.

(204) See, U.S. Pat. Nos. 9,443,141; 20110218950; 20150248167; 20150248764; 20150248765; 20150310862; 20150331929; 20150338915; 20160026913; 20160062459; 20160085302; 20160125572; 20160247064; 20160274660; 20170053665; 20170069306; 20170173262; and 20170206691. Amari, S., Natural gradient works efficiently in learning, *Neural Computation* 10:251-276, 1998. Amari S., Cichocki, A. & Yang, H. H., A new learning algorithm for blind signal separation. In: *Advances in Neural Information Processing Systems 8*, MIT Press, 1996. Bandettini P A, Wong E C, Hinks R S, Tikofsky R S, Hyde J S, Time course EPI of human brain function during task activation. *Magn Reson Med* 25:390-7, 1992. Bell A. J. & Sejnowski T. J. An information-maximization approach to blind separation and blind deconvolution. *Neural Comput* 7:1129-59, 1995. Bell, A. J. & Sejnowski, T. J., Learning the higher-order structure of a natural sound, *Network: Computation in Neural Systems* 7, 1996b. 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(205) Principal Component Analysis Principal component analysis (PCA) is a statistical procedure that uses an orthogonal transformation to convert a set of observations of possibly correlated variables into a set of values of linearly uncorrelated variables called principal components. If there are n observations with p variables, then the number of distinct principal components is $\min(n-1, p)$. This transformation is defined in such a way that the first principal component has the largest possible variance (that is, accounts for as much of the variability in the data as possible), and each succeeding component in turn has the highest variance possible under the constraint that it is orthogonal to the preceding components. The resulting vectors are an uncorrelated orthogonal basis set. PCA is sensitive to the relative scaling of the original variables. PCA is the simplest of the true eigenvector-based multivariate analyses. Often, its operation can be thought of as revealing the internal structure of the data in a way that best explains the variance in the data. If a multivariate dataset is visualized as a set of coordinates in a high-dimensional data space (1 axis per variable), PCA can supply the user with a lower-dimensional picture, a projection of this object when viewed from its most informative viewpoint. This is done by using only the first few principal components so that the dimensionality of the transformed data is reduced. PCA is closely related to factor analysis. Factor analysis typically incorporates more domain specific assumptions about the underlying structure and solves eigenvectors of a slightly different matrix. PCA is also related to canonical correlation analysis (CCA). CCA defines coordinate systems that optimally describe the cross-covariance between two datasets while PCA defines a new orthogonal coordinate system that optimally describes variance in a single dataset. See, en.wikipedia.org/wiki/Principal_component_analysis.

(206) A general model for confirmatory factor analysis is expressed as $x = \alpha + \Lambda \xi + \epsilon$. The covariance matrix is expressed as $E[(x - \mu)(x - \mu)'] = \Lambda \Phi \Lambda' + \Theta$. If residual covariance matrix $\Theta = 0$ and correlation matrix among latent factors $\Phi = I$, then factor analysis is equivalent to principal component analysis and the resulting covariance matrix is simplified to $\Sigma = \Lambda \Lambda'$. When there are p number of variables and all p components (or factors) are extracted, this covariance matrix can alternatively be expressed into $\Sigma = D \Lambda D'$, or $\Sigma = \lambda D \Lambda D'$, where $D = n \times p$ orthogonal matrix of eigenvectors, and $\Lambda = \lambda \Lambda$, $p \times p$.

matrix of eigenvalues, where λ is a scalar and A is a diagonal matrix whose elements are proportional to the eigenvalues of Σ . The following three components determine the geometric features of the observed data: λ parameterizes the volume of the observation, D indicates the orientation, and A represents the shape of the observation.

(207) When population heterogeneity is explicitly hypothesized as in model-based cluster analysis, the observed covariance matrix is decomposed into the following general form $\Sigma_{\text{sub.k}} = \lambda_{\text{sub.k}} D_{\text{sub.k}} A_{\text{sub.k}}^T A_{\text{sub.k}}$, where $\lambda_{\text{sub.k}}$ parameterizes the volume of the k -th cluster, $D_{\text{sub.k}}$ indicates the orientation of that cluster, and $A_{\text{sub.k}}$ represents the shape of that cluster. The subscript k indicates that each component (or cluster) can have different volume, shape, and orientation.

(208) Assume a random vector X , taking values in \mathbb{R}^m , has a mean and covariance matrix of $\mu_{\text{sub.X}}$ and $\Sigma_{\text{sub.X}}$, respectively. $\lambda_{\text{sub.1}} > \lambda_{\text{sub.2}} > \dots > \lambda_{\text{sub.m}} > 0$ are ordered eigenvalues of $\Sigma_{\text{sub.X}}$, such that the i -th eigenvalue of $\Sigma_{\text{sub.X}}$ means the i -th largest of them. Similarly, a vector $\alpha_{\text{sub.i}}$ is the i -th eigenvector of $\Sigma_{\text{sub.X}}$ when it corresponds to the i -th eigenvalue of $\Sigma_{\text{sub.X}}$. To derive the form of principal components (PCs), consider the optimization problem of maximizing $\text{var}[\alpha_{\text{sub.1}}^T X] = \alpha_{\text{sub.1}}^T \Sigma_{\text{sub.X}} \alpha_{\text{sub.1}}$, subject to $\alpha_{\text{sub.1}}^T \alpha_{\text{sub.1}} = 1$. The Lagrange multiplier method is used to solve this question.

(209) $L(\alpha_1, \phi_1) = \alpha_1^T \Sigma_{\text{sub.X}} \alpha_1 + \phi_1 (\alpha_1^T \alpha_1 - 1)$

$$\frac{\partial L}{\partial \alpha_1} = 2 \Sigma_{\text{sub.X}} \alpha_1 + 2\phi_1 \alpha_1 = 0 \quad \Sigma_{\text{sub.X}} \alpha_1 = -\phi_1 \alpha_1 \quad \text{var}[\alpha_1^T X] = -\phi_1 \alpha_1^T \alpha_1 = -\phi_1$$

(210) Because $-\phi_{\text{sub.1}}$ is the eigenvalue of $\Sigma_{\text{sub.X}}$, with $\alpha_{\text{sub.1}}$ being the corresponding normalized eigenvector, $\text{var}[\alpha_{\text{sub.1}}^T X]$ is maximized by choosing $\alpha_{\text{sub.1}}$ to be the first eigenvector of $\Sigma_{\text{sub.X}}$. In this case, $z_{\text{sub.1}} = \alpha_{\text{sub.1}}^T X$ is named the first PC of X , $\alpha_{\text{sub.1}}$ is the vector of coefficients for $z_{\text{sub.1}}$, and $\text{var}(z_{\text{sub.1}}) = \lambda_{\text{sub.1}}$.

(211) To find the second PC, $z_{\text{sub.2}} = \alpha_{\text{sub.2}}^T X$, we need to maximize $\text{var}[\alpha_{\text{sub.2}}^T X] = \alpha_{\text{sub.2}}^T \Sigma_{\text{sub.X}} \alpha_{\text{sub.2}}$ subject to $z_{\text{sub.2}}$ being uncorrelated with $z_{\text{sub.1}}$. Because $\text{cov}(\alpha_{\text{sub.1}}^T X, \alpha_{\text{sub.2}}^T X) = 0$, $\alpha_{\text{sub.1}}^T \Sigma_{\text{sub.X}} \alpha_{\text{sub.2}} = 0$, $\alpha_{\text{sub.1}}^T \alpha_{\text{sub.2}} = 0$, this problem is equivalently set as maximizing $\alpha_{\text{sub.2}}^T \Sigma_{\text{sub.X}} \alpha_{\text{sub.2}}$, subject to $\alpha_{\text{sub.1}}^T \alpha_{\text{sub.2}} = 0$, and $\alpha_{\text{sub.2}}^T \alpha_{\text{sub.2}} = 1$. We still make use of the Lagrange multiplier method.

(212) $L(\alpha_2, \phi, \phi_2) = \alpha_2^T \Sigma_{\text{sub.X}} \alpha_2 + \phi_1 \alpha_1^T \alpha_2 + \phi_2 (\alpha_2^T \alpha_2 - 1)$

$$\frac{\partial L}{\partial \alpha_2} = 2 \Sigma_{\text{sub.X}} \alpha_2 + \phi_1 \alpha_1 + 2\phi_2 \alpha_2 = 0 \quad \alpha_1^T (2 \Sigma_{\text{sub.X}} \alpha_2 + \phi_1 \alpha_1 + 2\phi_2 \alpha_2) = 0 \quad \phi_1 = 0 \quad \Sigma_{\text{sub.X}} \alpha_2 = -\phi_2 \alpha_2$$

(213) Because $-\phi_{\text{sub.2}}$ is the eigenvalue of $\Sigma_{\text{sub.X}}$ with $\alpha_{\text{sub.2}}$ being the corresponding normalized eigenvector, $\text{var}[\alpha_{\text{sub.2}}^T X]$ is maximized by choosing $\alpha_{\text{sub.2}}$ to be the second eigenvector of $\Sigma_{\text{sub.X}}$. In this case, $z_{\text{sub.2}} = \alpha_{\text{sub.2}}^T X$ is named the second PC of X , $\alpha_{\text{sub.2}}$ is the vector of coefficients for $z_{\text{sub.2}}$, and $\text{var}(z_{\text{sub.2}}) = \lambda_{\text{sub.2}}$. Continuing in this way, it can be shown that the i -th PC $z_{\text{sub.i}} = \alpha_{\text{sub.i}}^T X$ is constructed by selecting $\alpha_{\text{sub.i}}$ to be the i -th eigenvector of $\Sigma_{\text{sub.X}}$ and has variance of $\lambda_{\text{sub.i}}$. The key result in regards to PCA is that the principal components are the only set of linear functions of original data that are uncorrelated and have orthogonal vectors of coefficients.

(214) For any positive integer $p \leq m$, let $B = [\beta_{\text{sub.1}}, \beta_{\text{sub.2}}, \dots, \beta_{\text{sub.p}}]$ be an $m \times p$ matrix with orthonormal columns, i.e., $\beta_{\text{sub.i}}^T \beta_{\text{sub.j}} = \delta_{\text{sub.ij}}$, and $Y = B \sup T X$. Then the trace of covariance matrix of Y is maximized by taking $B = [\alpha_{\text{sub.1}}, \alpha_{\text{sub.2}}, \dots, \alpha_{\text{sub.p}}]$, where $\alpha_{\text{sub.i}}$ is the i -th eigenvector of $\Sigma_{\text{sub.X}}$. Because $\Sigma_{\text{sub.X}}$ is symmetric with all distinct eigenvalues, so $\{\alpha_{\text{sub.1}}, \alpha_{\text{sub.2}}, \dots, \alpha_{\text{sub.m}}\}$ is an orthonormal basis with $\alpha_{\text{sub.i}}$ being the i -th eigenvector of $\Sigma_{\text{sub.X}}$ and we can represent the columns of B as

$$(215) \beta_i = \sum_{j=1}^m c_{ji} \alpha_j, i = 1, \dots, p,$$

So we have $B = PC$ where $P = [\alpha_{\text{sub.1}}, \dots, \alpha_{\text{sub.m}}]$, $C = \{c_{\text{sub.ij}}\}$ is an $m \times p$ matrix. Then, $P \sup T \Sigma_{\text{sub.X}} P = \Lambda$, with Λ being a diagonal matrix whose k -th diagonal element is $\lambda_{\text{sub.k}}$ and the covariance matrix of Y is,

$\Sigma_{\text{sub.Y}} = B \sup T \Sigma_{\text{sub.X}} B = C \sup T P \sup T \Sigma_{\text{sub.X}} P C = C \sup T \Lambda C = \sum_{i=1}^m \lambda_{\text{sub.i}} c_{\text{sub.i1}} c_{\text{sub.i1}}^T + \dots + \lambda_{\text{sub.m}} c_{\text{sub.m}} c_{\text{sub.m}}^T$ where $c_{\text{sub.i}} \sup T$ is the i -th row of C . So,

$$(216) \text{trace}(\Sigma_{\text{sub.Y}}) = \sum_{i=1}^m \lambda_i \text{trace}(c_i c_i^T) = \sum_{i=1}^m \lambda_i \text{trace}(c_i^T c_i) = \sum_{i=1}^m \lambda_i c_i^T c_i = \sum_{i=1}^m (\sum_{j=1}^p c_{ij}^2) \lambda_i$$

(217) Because $C \sup T C = B \sup T P P \sup T B = B \sup T B = I$, so

$$(218) \text{trace}(C^T C) = \sum_{i=1}^m \sum_{j=1}^p c_{ij}^2 = p,$$

and the columns of C are orthonormal. By the Gram-Schmidt method, C can expand to D , such that D has its columns as an orthonormal basis of \mathbb{R}^m and contains C as its first p columns. D is square shape, thus being an orthogonal matrix and having its rows as another orthonormal basis of \mathbb{R}^m . One row of C is a part of one row of D so

$$(219) \sum_{j=1}^p c_{ij}^2 \leq 1, i = 1, \dots, m.$$

Considering the constraints

$$(220) \sum_{j=1}^p c_{ij}^2 \leq 1, \sum_{i=1}^m \sum_{j=1}^p c_{ij}^2 = p$$

and the objective

$$(221) \sum_{i=1}^m (\sum_{j=1}^p c_{ij}^2) \lambda_i$$

We derive that $\text{trace}(\Sigma_{\text{sub.Y}})$ is maximized if

$$(222) \sum_{j=1}^p c_{ij}^2 = 1$$

for $i=1, \dots, p$, and

$$(223) \sum_{j=1}^p c_{ij}^2 = 0$$

for $i=p+1, \dots, m$. When $B = [\alpha_{\text{sub.1}}, \alpha_{\text{sub.2}}, \dots, \alpha_{\text{sub.p}}]$, straightforward calculation yields that C is an all-zero matrix except $c_{\text{sub.ii}}=1, i=1, \dots, p$. This fulfills the maximization condition. Actually, by taking $B = [\gamma_{\text{sub.1}}, \gamma_{\text{sub.2}}, \dots, \gamma_{\text{sub.p}}]$, where $\{\gamma_{\text{sub.1}}, \gamma_{\text{sub.2}}, \dots, \gamma_{\text{sub.p}}\}$ is any orthonormal basis of the subspace of $\text{span}\{\alpha_{\text{sub.1}}, \alpha_{\text{sub.2}}, \dots, \alpha_{\text{sub.p}}\}$, the maximization condition is also satisfied, yielding the same trace of covariance matrix of Y .

(224) Suppose that we wish to approximate the random vector X by its projection onto a subspace spanned by columns of B , where $B = [\beta_{\text{sub.1}}, \beta_{\text{sub.2}}, \dots, \beta_{\text{sub.p}}]$ is a real $m \times p$ matrix with orthonormal columns, i.e., $\beta_{\text{sub.i}}^T \beta_{\text{sub.j}} = \delta_{\text{sub.ij}}$. If $\sigma_{\text{sub.i}}^2$ is the residual variance for each component of X , then

$$(225) \sum_{i=1}^m \sigma_i^2$$

is minimized if $B = [\alpha_{\text{sub.1}}, \alpha_{\text{sub.2}}, \dots, \alpha_{\text{sub.p}}]$, where $\{\alpha_{\text{sub.1}}, \alpha_{\text{sub.2}}, \dots, \alpha_{\text{sub.p}}\}$ are the first p eigenvectors of $\Sigma_{\text{sub.X}}$. In other words, the trace of covariance matrix of $X - BB \sup T X$ is minimized if $B = [\alpha_{\text{sub.1}}, \alpha_{\text{sub.2}}, \dots, \alpha_{\text{sub.p}}]$. When $E(X) = 0$, which is a commonly applied preprocessing step in data analysis methods, this property is saying that $E\|X - BB \sup T X\|^2$ is minimized if $B = [\alpha_{\text{sub.1}}, \alpha_{\text{sub.2}}, \dots, \alpha_{\text{sub.p}}]$.

(226) The projection of a random vector X onto a subspace spanned by columns of B is $\hat{X} = BB \sup T X$. Then the residual vector is $\epsilon = X - BB \sup T X$, which has a covariance matrix

$$\Sigma_{\text{sub.}\epsilon} = (I - BB \sup T) \Sigma_{\text{sub.X}} (I - BB \sup T)^T.$$

(243) Spatial Principal Component Analysis (Spatial PCA) Let $J(t, i; \alpha, s)$ be the current density in voxel i , as estimated by LORETA, in condition α at

t time-frames after stimulus onset for subject s . Let $fwdarw.fBA$ be a function, which assigns to each voxel $i \in \text{Voxel}$ the corresponding fBA $b \in fBA$. In a first pre-processing step, for each subjects, the value of the current density averaged over each fBA is calculated:

$$(244) \quad x(t, b; \alpha, s) = \frac{1}{N_{\text{sub},b}} \cdot \text{Math.} \int (t, i; \alpha, s) \quad (4) \quad \text{where } N_{\text{sub},b} \text{ is the number of voxels in the } fBA \text{ } b, \text{ in condition } \alpha \text{ for subjects.}$$

(245) In the second analysis stage, the mean current density $x(t, b; \alpha, s)$ from each fBA b , for every subjects and condition α was subjected to spatial PCA analysis of the correlation matrix and varimax rotation

(246) The spatial PCA uses the above-defined $fBAs$ as variables sampled along the time epoch for which EEG has been sampled (e.g., 0-1000 ms; 512 time-frames), and the inverse solution estimated. Spatial matrices (e.g., each matrix was sized $b \times t = 36 \times 512$ elements) for every subject and condition may be collected, and subjected to PCA analyses, including the calculation of the covariance matrix; eigenvalue decomposition and varimax rotation, in order to maximize factor loadings. In other words, the spatial PCA analysis approximates the mean current density for each subject in each condition as

$$(247) \quad 0 \quad x(t; \alpha, s) \approx x_0(\alpha, s) + \sum_k \text{Math.} \cdot c_k(t) x_k(\alpha, s), \quad (5)$$

where here $x(t; \alpha, s) \in \mathbb{R}^{\text{sup.}36}$ is a vector, which denotes the time-dependent activation of the $fBAs$, $x_{\text{sub},0}(\alpha, s)$ is their mean activation, and $x_{\text{sub},k}(\alpha, s)$ and $c_{\text{sub},k}$ are the principal components and their corresponding coefficients (factor loadings) as computed using the principal component analysis. See: Arzouan Y, Goldstein A, Faust M. Brainwaves are stethoscopes: ERP correlates of novel metaphor comprehension. *Brain Res* 2007; 1160: 69-81. Arzouan Y, Goldstein A, Faust M. Dynamics of hemispheric activity during metaphor comprehension: electrophysiological measures. *NeuroImage* 2007; 36: 222-231. Chapman R M, McCrary J W. EP component identification and measurement by principal components analysis. *Brain and cognition* 1995; 27: 288-310. Dien J, Frishkoff G A, Cerbone A, Tucker D M. Parametric analysis of event-related potentials in semantic comprehension: evidence for parallel brain mechanisms. *Brain research* 2003; 15: 137-153. Dien J, Frishkoff G A. Principal components analysis of event-related potential datasets. In: Handy T (ed). *Event-Related Potentials: A Methods Handbook*. Cambridge, Mass MIT Press; 2004. Potts G F, Dien J, Harty-Speiser A L, McDougal L M, Tucker D M. Dense sensor array topography of the event-related potential to task-relevant auditory stimuli. *Electroencephalography and clinical neurophysiology* 1998; 106: 444-456. Roster F, Manzey D. Principal components and varimax-rotated components in event-related potential research: some remarks on their interpretation. *Biological psychology* 1981; 13: 3-26. Ruchkin D S, McCalley M G, Glaser E M. Event related potentials and time estimation. *Psychophysiology* 1977; 14: 451-455. Spencer K M, Dien J, Donchin E. Spatiotemporal analysis of the late ERP responses to deviant stimuli. *Psychophysiology* 2001; 38: 343-358. Squires K C, Squires N K, Hillyard S A. Decision-related cortical potentials during an auditory signal detection task with cued observation intervals. *Journal of experimental psychology* 1975; 1: 268-279. van Boxtel A, Boelhouwer A J, Bos A R. Optimal EMG signal bandwidth and interelectrode distance for the wording of acoustic, electrocutaneous, and photic blink reflexes. *Psychophysiology* 1998; 35: 690-697.

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(248) Nonlinear Dimensionality Reduction High-dimensional data, meaning data that requires more than two or three dimensions to represent, can be difficult to interpret One approach to simplification is to assume that the data of interest lie on an embedded non-linear manifold within the higher-dimensional space. If the manifold is of low enough dimension, the data can be visualized in the low-dimensional space. Non-linear methods can be broadly classified into two groups: those that provide a mapping (either from the high-dimensional space to the low-dimensional embedding or vice versa), and those that just give a visualization. In the context of ML, mapping methods may be viewed as a preliminary feature extraction step, after which pattern recognition algorithms are applied. Typically, those that just give a visualization are based on proximity data—that is, distance measurements. Related Linear Decomposition Methods include Independent component analysis (ICA), Principal component analysis (PCA) (also called Karhunen-Loève transform—KLT), Singular value decomposition (SVD), and Factor analysis.

(249) The self-organizing map (SOM, also called Kohonen map) and its probabilistic variant generative topographic mapping (GTM) use a point representation in the embedded space to form a latent variable model based on a non-linear mapping from the embedded space to the high-dimensional space. These techniques are related to work on density networks, which also are based around the same probabilistic model.

(250) Principal curves and manifolds give the natural geometric framework for nonlinear dimensionality reduction and extend the geometric interpretation of PCA by explicitly constructing an embedded manifold, and by encoding using standard geometric projection onto the manifold. How to define the “simplicity” of the manifold is problem-dependent, however, it is commonly measured by the intrinsic dimensionality and/or the smoothness of the manifold. Usually, the principal manifold is defined as a solution to an optimization problem. The objective function includes a quality of data approximation and some penalty terms for the bending of the manifold. The popular initial approximations are generated by linear PCA, Kohonen's SOM or autoencoders. The elastic map method provides the expectation-maximization algorithm for principal manifold learning with minimization of quadratic energy functional at the “maximization” step.

(251) An autoencoder is a feed-forward neural network which is trained to approximate the identity function. That is, it is trained to map from a vector of values to the same vector. When used for dimensionality reduction purposes, one of the hidden layers in the network is limited to contain only a small number of network units. Thus, the network must learn to encode the vector into a small number of dimensions and then decode it back into the original space. Thus, the first half of the network is a model which maps from high to low-dimensional space, and the second half maps from low to high-dimensional space. Although the idea of autoencoders is quite old, training of deep autoencoders has only recently become possible through the use of restricted Boltzmann machines and stacked denoising autoencoders. Related to autoencoders is the NeuroScale algorithm, which uses stress functions inspired by multidimensional scaling and Sammon mappings (see below) to learn a non-linear mapping from the high-dimensional to the embedded space. The mappings in NeuroScale are based on radial basis function networks.

(252) Gaussian process latent variable models (GPLVM) are probabilistic dimensionality reduction methods that use Gaussian Processes (GPs) to find a lower dimensional non-linear embedding of high dimensional data. They are an extension of the Probabilistic formulation of PCA. The model is defined probabilistically and the latent variables are then marginalized and parameters are obtained by maximizing the likelihood. Like kernel PCA they use a kernel function to form a nonlinear mapping (in the form of a Gaussian process). However, in the GPLVM the mapping is from the embedded(latent) space to the data space (like density networks and GTM) whereas in kernel PCA it is in the opposite direction. It was originally proposed for visualization of high dimensional data but has been extended to construct a shared manifold model between two observation spaces. GPLVM and its many variants have been proposed specially for human motion modeling, e.g., back constrained GPLVM, GP dynamic model (GPDM), balanced GPDM (B-GPDM) and topologically constrained GPDM. To capture the coupling effect of the pose and gait manifolds in the gait analysis, a multi-layer joint gait-pose manifolds was proposed.

(253) Curvilinear component analysis (CCA) looks for the configuration of points in the output space that preserves original distances as much as possible while focusing on small distances in the output space (conversely to Sammon's mapping which focus on small distances in original space). It should be noticed that CCA, as an iterative learning algorithm, actually starts with focus on large distances (like the Sammon algorithm), then gradually change focus to small distances. The small distance information will overwrite the large distance information, if compromises between the two have to be made. The stress function of CCA is related to a sum of right Bregman divergences. Curvilinear distance analysis (CDA) trains a self-organizing neural network to fit the manifold and seeks to preserve geodesic distances in its embedding. It is based on Curvilinear Component Analysis (which extended Sammon's mapping), but uses geodesic distances instead. Diffeomorphic Dimensionality Reduction or Diffeomap learns a smooth diffeomorphic mapping which transports the data onto a lower-dimensional linear subspace. The method solves for a smooth time indexed vector field such that flows along the field which start at the data points will end at a lower-dimensional linear subspace, thereby attempting to preserve pairwise differences under both the forward and inverse mapping.

(254) Perhaps the most widely used algorithm for kernel principal analysis is Kernel PCA. It is a combination of Principal component analysis and the kernel trick. PCA begins by computing the covariance matrix of the $M \times n$ Matrix X . It then projects the data onto the first k eigenvectors of that matrix. By comparison, KPCA begins by computing the covariance matrix of the data after being transformed into a higher-dimensional space. It then projects the transformed data onto the first k eigenvectors of that matrix, just like PCA. It uses the kernel trick to factor away much of the computation, such that the entire process can be performed without actually computing $\phi(x)$. Of course ϕ must be chosen such that it has a known corresponding kernel.

(255) The Fourier transform (FT) decomposes a function of time (a signal) into the frequencies that make it up. The Fourier transform of a function of time is itself a complex-valued function of frequency, whose absolute value represents the amount of that frequency present in the original function, and whose complex argument is the phase offset of the basic sinusoid in that frequency. The Fourier transform is called the frequency domain representation of the original signal. The term Fourier transform refers to both the frequency domain representation and the mathematical operation that associates the frequency domain representation to a function of time. The Fourier transform is not limited to functions of time, but in order to have a unified language, the domain of the original function is commonly referred to as the time domain. For many functions of practical interest, one can define an operation that reverses this: the inverse Fourier transformation, also called Fourier synthesis, of a frequency domain representation combines the contributions of all the different frequencies to recover the original function of time. See, en.wikipedia.org/wiki/Fourier_transform.

(256) The Fourier transform of a finite Borel measure μ on \mathbb{R}^n is given by: $\mu(\xi) = \int_{\mathbb{R}^n} e^{-2\pi i x \cdot \xi} d\mu(x)$. This transform continues to enjoy many of the properties of the Fourier transform of integrable functions. One notable difference is that the Riemann-Lebesgue lemma fails for measures. In the case that $d\mu = f(x) dx$, then the formula above reduces to the usual definition for the Fourier transform of f . In the case that μ is the probability distribution associated to a random variable X , the Fourier-Stieltjes transform is closely related to the characteristic function, but the typical conventions in probability theory take $e^{ix\xi}$ instead of $e^{-2\pi i x \cdot \xi}$. In the case when the distribution has a probability density function this definition reduces to the Fourier transform applied to the probability density function, again with a different choice of constants. The Fourier transform may be used to give a characterization of measures. Bochner's theorem characterizes which functions may arise as the Fourier-Stieltjes transform of a positive measure on the circle. Furthermore, the Dirac delta function, although not a function, is a finite Borel measure. Its Fourier transform is a constant function (whose specific value depends upon the form of the Fourier transform used). See Pinsky, Mark (2002), *Introduction to Fourier Analysis and Wavelets*, Brooks/Cole, ISBN 978-0-534-37660-4; Katznelson, Yitzhak (1976), *An Introduction to Harmonic Analysis*, Dover, ISBN 978-0-486-63331-2.

(257) The Fourier transform is also a special case of Gelfand transform. In this particular context, it is closely related to the Pontryagin duality map. Given an abelian locally compact Hausdorff topological group G , as before we consider space $L^1(G)$, defined using a Haar measure. With convolution as multiplication, $L^1(G)$ is an abelian Banach algebra. Taking the completion with respect to the largest possible C^* -norm gives its enveloping C^* -algebra, called the group C^* -algebra $C^*(G)$ of G . (Any C^* -norm on $L^1(G)$ is bounded by the L^1 norm, therefore their supremum exists.) τ is the involution operator. Given any abelian C^* -algebra A , the Gelfand transform gives an isomorphism between A and $C_0(\hat{A})$, where \hat{A} is the multiplicative linear functionals, i.e. one-dimensional representations, on A with the weak-* topology. The multiplicative linear functionals of $C^*(G)$, after suitable identification, are exactly the characters of G , and the Gelfand transform, when restricted to the dense subset $L^1(G)$ is the Fourier-Pontryagin transform.

(258) The Laplace transform is very similar to the Fourier transform. While the Fourier transform of a function is a complex function of a real variable (frequency), the Laplace transform of a function is a complex function of a complex variable. Laplace transforms are usually restricted to functions $f(t)$ with $t \geq 0$. A consequence of this restriction is that the Laplace transform of a function is a holomorphic function of the variable s . The Laplace transform of a distribution is generally a well-behaved function. As a holomorphic function, the Laplace transform has a power series representation. This power series expresses a function as a linear superposition of moments of the function. The Laplace transform is invertible on a large class of functions. The inverse Laplace transform takes a function of a complex variable s (often frequency) and yields a function of a real variable t (time). Given a simple mathematical or functional description of an input or output to a system, the Laplace transform provides an alternative functional description that often simplifies the process of analyzing the behavior of the system, or in synthesizing a new system based on a set of specifications. So, for example, Laplace transformation from the time domain to the frequency domain transforms differential equations into algebraic equations and convolution into multiplication. See, en.wikipedia.org/wiki/Laplace_transform.

(259) The short-time Fourier transform (STFT), is a Fourier-related transform used to determine the sinusoidal frequency and phase content of local sections of a signal as it changes over time. In practice, the procedure for computing STFTs is to divide a longer time signal into shorter segments of equal length and then compute the Fourier transform separately on each shorter segment. This reveals the Fourier spectrum on each shorter segment. One then usually plots the changing spectra as a function of time. The signal may be windowed using, e.g., a Hann window or a Gaussian window. See, en.wikipedia.org/wiki/Short-time_Fourier_transform.

(260) The fractional Fourier transform (FRFT), is a generalization of the classical Fourier transform. The FRFT of a signal can also be interpreted as a decomposition of the signal in terms of chirps. The FRFT can be used to define fractional convolution, correlation, and other operations, and can also be further generalized into the linear canonical transformation (LCT). See: en.wikipedia.org/wiki/Fractional_Fourier_transform. Almeida, Luis B. "The fractional Fourier transform and time-frequency representations." *IEEE Transactions on signal processing* 42, no. 11 (1994): 3084-3091. Bailey, David H., and Paul N. Swartztrauber. "The fractional Fourier transform and applications." *SIAM review* 33, no. 3 (1991): 389-404. Candan, Cagatay, M. Alper Kutay, and Haldun M. Ozaktas. "The discrete fractional Fourier transform." *IEEE Transactions on signal processing* 48, no. 5 (2000): 1329-1337. Lohmann, Adolf W. "Image rotation, aligner rotation, and the fractional Fourier transform." *JOSA A* 10, no. 10 (1993): 2181-2186. Ozaktas, Haldun M., and David Mendlovic. "Fourier transforms of fractional order and their optical interpretation." *Optics Communications* 101, no. 3-4 (1993): 163-169. Ozaktas, Haldun M., and M. Alper Kutay. "The fractional Fourier transform." In *Control Conference (ECC), 2001 European*, pp. 1477-1483. IEEE, 2001. Ozaktas, Haldun M., Orhan Arikan, M. Alper Kutay, and Gozde Bozdağt "Digital computation of the fractional Fourier transform." *IEEE Transactions on signal processing* 44, no. 9 (1996): 2141-2150. Pei, Soo-Chang, Min-Hung Yeh, and Chien-Cheng Tseng. "Discrete fractional Fourier transform based on orthogonal projections." *IEEE Transactions on Signal Processing* 47, no. 5 (1999): 1335-1348. Qi, Lin, Ran Tao, Siyong Zhou, and Yue Wang. "Detection and parameter estimation of multicomponent LFM signal based on the fractional Fourier transform." *Science in China series F: information sciences* 47, no. 2 (2004): 184. Tao, Ran, Yan-Lei Li, and Yue Wang. "Short-time fractional Fourier transform and its applications." *IEEE Transactions on Signal Processing* 58, no. 5 (2010): 2568-2580. Xia, Xiang-Gen. "On bandlimited signals with fractional Fourier transform." *IEEE Signal Processing Letters* 3, no. 3 (1996): 72-74. Zayed, Ahmed I. "A convolution and product theorem for the fractional Fourier transform." *IEEE Signal processing letters* 5, no. 4 (1998): 101-103. Zayed, Ahmed I. "On the relationship between the Fourier and fractional Fourier transforms." *IEEE signal processing letters* 3, no. 12 (1996): 310-311.

(261) Laplacian Eigenmaps, (also known as Local Linear Eigenmaps, LLE) are special cases of kernel PCA, performed by constructing a data-dependent kernel matrix. KPCA has an internal model, so it can be used to map points onto its embedding that were not available attaining time. Laplacian Eigenmaps uses spectral techniques to perform dimensionality reduction. This technique relies on the basic assumption that the data lies in a low-dimensional manifold in a high-dimensional space. This algorithm cannot embed out of sample points, but techniques based on Reproducing kernel Hilbert space regularization exist for adding this capability. Such techniques can be applied to other nonlinear dimensionality reduction algorithms as well. Traditional techniques like principal component analysis do not consider the intrinsic geometry of the data. Laplacian eigenmaps builds a graph from neighborhood information of the data set. Each data point serves as a node on the graph and connectivity between nodes is governed by the proximity of neighboring points (using e.g. the k-nearest neighbor algorithm). The graph thus generated can be considered as a

discrete approximation of the low-dimensional manifold in the high-dimensional space. Minimizing a cost function based on the graph ensures that points close to each other on the manifold are mapped close to each other in the low-dimensional space, preserving local distances. The eigenfunctions of the Laplace-Beltrami operator on the manifold serve as the embedding dimensions, since under mild conditions this operator has a countable spectrum that is a basis for square integrable functions on the manifold (compare to Fourier series on the unit circle manifold). Attempts to place Laplacian eigenmaps on solid theoretical ground have met with some success, as under certain nonrestrictive assumptions, the graph Laplacian matrix has been shown to converge to the Laplace-Beltrami operator as the number of points goes to infinity. In classification applications, low dimension manifolds can be used to model data classes which can be defined from sets of observed instances. Each observed instance can be described by two independent factors termed 'content' and 'style', where 'content' is the invariant factor related to the essence of the class and 'style' expresses variations in that class between instances. Unfortunately, Laplacian Eigenmaps may fail to produce a coherent representation of a class of interest when training data consist of instances varying significantly in terms of style. In the case of classes which are represented by multivariate sequences, Structural Laplacian Eigenmaps has been proposed to overcome this issue by adding additional constraints within the Laplacian Eigenmaps neighborhood information graph to better reflect the intrinsic structure of the class. More specifically, the graph is used to encode both the sequential structure of the multivariate sequences and, to minimize stylistic variations, proximity between data points of different sequences or even within a sequence, if it contains repetitions. Using dynamic time warping, proximity is detected by finding correspondences between and within sections of the multivariate sequences that exhibit high similarity.

(262) Like LLE, Hessian LLE is also based on sparse matrix techniques. It tends to yield results of a much higher quality than LLE. Unfortunately, it has a very costly computational complexity, so it is not well-suited for heavily sampled manifolds. It has no internal model. Modified LLE (MLLE) is another LLE variant which uses multiple weights in each neighborhood to address the local weight matrix conditioning problem which leads to distortions in LLE maps. MLLE produces robust projections similar to Hessian LLE, but without the significant additional computational cost.

(263) Manifold alignment takes advantage of the assumption that disparate data sets produced by similar generating processes will share a similar underlying manifold representation. By learning projections from each original space to the shared manifold, correspondences are recovered and knowledge from one domain can be transferred to another. Most manifold alignment techniques consider only two data sets, but the concept extends to arbitrarily many initial data sets. Diffusion maps leverages the relationship between heat diffusion and a random walk (Markov Chain); an analogy is drawn between the diffusion operator on a manifold and a Markov transition matrix operating on functions defined on the graph whose nodes were sampled from the manifold. Relational perspective map is a multidimensional scaling algorithm. The algorithm finds a configuration of data points on a manifold by simulating a multi-particle dynamic system on a closed manifold, where data points are mapped to particles and distances (or dissimilarity) between data points represent a repulsive force. As the manifold gradually grows in size the multi-particle system cools down gradually and converges to a configuration that reflects the distance information of the data points. Local tangent space alignment (LTSA) is based on the intuition that when a manifold is correctly unfolded, all of the tangent hyperplanes to the manifold will become aligned. It begins by computing the k-nearest neighbors of every point. It computes the tangent space at every point by computing the d-first principal components in each local neighborhood. It then optimizes to find an embedding that aligns the tangent spaces. Local Multidimensional Scaling performs multidimensional scaling in local regions, and then uses convex optimization to fit all the pieces together.

(264) Maximum Variance Unfolding was formerly known as Semidefinite Embedding. The intuition for this algorithm is that when a manifold is properly unfolded, the variance over the points is maximized. This algorithm also begins by finding the k-nearest neighbors of every point. It then seeks to solve the problem of maximizing the distance between all non-neighboring points, constrained such that the distances between neighboring points are preserved. Nonlinear PCA (NLPCA) uses backpropagation to train a multi-layer perceptron (MLP) to fit to a manifold. Unlike typical MLP training, which only updates the weights, NLPCA updates both the weights and the inputs. That is, both the weights and inputs are treated as latent values. After training, the latent inputs are a low-dimensional representation of the observed vectors, and the MLP maps from that low-dimensional representation to the high-dimensional observation space. Manifold Sculpting uses graduated optimization to find an embedding. Like other algorithms, it computes the k-nearest neighbors and tries to seek an embedding that preserves relationships in local neighborhoods. It slowly scales variance out of higher dimensions, while simultaneously adjusting points in lower dimensions to preserve those relationships.

(265) Ruffini (2015) discusses Multichannel transcranial current stimulation (tCS) systems that offer the possibility of EEG-guided optimized, non-invasive brain stimulation. A tCS electric field realistic brain model is used to create a forward "lead-field" matrix and, from that, an EEG inverter is employed for cortical mapping. Starting from EEG, 2D cortical surface dipole fields are defined that could produce the observed EEG electrode voltages.

(266) Schestatsky et al. (2017) discuss transcranial direct current stimulation (tDCS), which stimulates through the scalp with a constant electric current that induces shifts in neuronal membrane excitability, resulting in secondary changes in cortical activity. Although tDCS has most of its neuromodulatory effects on the underlying cortex, tDCS effects can also be observed in distant neural networks. Concomitant EEG monitoring of the effects of tDCS can provide valuable information on the mechanisms of tDCS. EEG findings can be an important surrogate marker for the effects of tDCS and thus can be used to optimize its parameters. This combined EEG-tDCS system can also be used for preventive treatment of neurological conditions characterized by abnormal peaks of cortical excitability, such as seizures. Such a system would be the basis of a non-invasive closed-loop device. tDCS and EEG can be used concurrently. See: Albert Jacobo, Sara Lopez-Martin, Jose Antonio Hinojosa, and Luis Carretié. "Spatiotemporal characterization of response inhibition." *Neuroimage* 76 (2013): 272-281. Arzouan Y, Goldstein A, Faust M. Brainwaves are stethoscopes: ERP correlates of novel metaphor comprehension. *Brain Res* 2007; 1160: 69-81. Arzouan Y, Goldstein A, Faust M. Dynamics of hemispheric activity during metaphor comprehension: electrophysiological measures. *NeuroImage* 2007; 36: 222-231. Arzy, Shahar, Yossi Arzouan, Esther Adi-Japha, Sorin Solomon, and Olaf Blanke. "The 'intrinsic' system in the human cortex and self-projection: a data driven analysis." *Neuroreport* 21, no. 8 (2010): 569-574. Bao, Xuecai, Jinli Wang, and Jianfeng Hu. "Method of individual identification based on electroencephalogram analysis." In *New Trends in Information and Service Science*, 2009. NISS'09. International Conference on, pp. 390-393. IEEE, 2009. Bhattacharya, Joydeep.

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(267) EEG analysis approaches have emerged, in which event-related changes in EEG dynamics in single event-related data records are analyzed. See Allen D. Malony et al., *Computational Neuroinformatics for Integrated Electromagnetic Neuroimaging and Analysis*, PAR-99-138. Pfurtscheller, reported a method for quantifying the average transient suppression of alpha band (circa 10-Hz) activity following stimulation. Event-related desynchronization (ERD, spectral amplitude decreases), and event-related synchronization (ERS, spectral amplitude increases) are observed in a variety of narrow frequency bands (4-40 Hz) which are systematically dependent on task and cognitive state variables as well as on stimulus parameters. Makeig (1993) was reported event-related changes in the full EEG spectrum, yielding a 2-D time/frequency measure he called the event-related spectral perturbation (ERSP). This method avoided problems associated with analysis of a priori narrow frequency bands, since bands of interest for the analysis could be based on significant features of the complete time/frequency transform. Rappelsburger et al. introduced event-related coherence (ERCOH). A wide variety of other signal processing measures have been tested for use on EEG and/or MEG data, including dimensionality measures based on chaos theory and the bispectrum. Use of neural networks has also been proposed for EEG pattern recognition applied to clinical and practical problems, though usually these methods have not been employed with an aim of explicitly modeling the neurodynamics involved. Neurodynamics is the mobilization of the nervous system as an approach to physical treatment. The method relies on influencing pain and other neural physiology via mechanical treatment of neural tissues and the non-neural structures surrounding the nervous system. The body presents the nervous system with a mechanical interface via the musculoskeletal system. With movement, the musculoskeletal system exerts non-uniform stresses and movement in neural tissues, depending on the local anatomical and mechanical characteristics and the pattern of body movement. This activates an array of mechanical and physiological responses in neural tissues. These responses include neural sliding, pressurization, elongation, tension and changes in intraneural microcirculation, axonal transport and impulse traffic.

(268) The availability of and interest in larger and larger numbers of EEG (and MEG) channels led immediately to the question of how to combine data from different channels. Donchin advocated the use of linear factor analysis methods based on principal component analysis (PCA) for this purpose. Temporal PCA assumes that the time course of activation of each derived component is the same in all data conditions. Because this is unreasonable for many data sets, spatial PCA (usually followed by a component rotation procedure such as Varimax or Promax) is of potentially greater interest. To this end, several variants of PCA have been proposed for ERP decomposition.

(269) Bell and Sejnowski published an iterative algorithm based on information theory for decomposing linearly mixed signals into temporally independent signals by minimizing their mutual information. First approaches to blind source separation minimized third and fourth-order correlations among the observed variables and achieved limited success in simulations. A generalized approach uses a simple neural network algorithm that used joint information maximization or ‘infomax’ as a training criterion. By using a compressive nonlinearity to transform the data and then following the entropy gradient of the resulting mixtures, ten recorded voice and music sound sources were unmixed. A similar approach was used for performing blind deconvolution, and the ‘infomax’ method was used for decomposition of visual scenes.

(270) EEG source analysis may be accomplished using various techniques. Grech, Roberta, Tracey Cassar, Joseph Muscat Kenneth P. Camilleri, Simon G. Fabri, Michalis Zervakis, Petros Xanthopoulos, Vangelis Sakkalis, and Bart Vanrumste. “Review on solving the inverse problem in EEG source analysis.” *Journal of neuroengineering and rehabilitation* 5, no. 1 (2008): 25. De Munck J C, Van Dijk B W, Spekreijse H. Mathematical Dipoles are Adequate to Describe Realistic Generators of Human Brain Activity. *IEEE Transactions on Biomedical Engineering*. 1988; 35:960-966. doi: 10.1109/10.8677. Hallel H, Vanrumste B, Grech R, Muscat J, De Clercq W, Vergult A, D'Asseler Y, Camilleri K P, Fabri S G, Van Huffel S, Lemahieu I. Review on solving the forward problem in EEG source analysis. *J. of NeuroEngineering and Rehabilitation*. 2007; 4 Whittingstall K, Stroink G, Gates L, Connolly J F, Finley A. Effects of dipole position, orientation and noise on the accuracy of EEG source localization. *Biomedical Engineering Online*. 2003; 2 www.biomedical-engineering-online.com/content/2/1/14 Baillet S, Gamero L. A Bayesian Approach to Introducing Anatomic-Functional Priors in the EEG/MEG Inverse Problem. *IEEE Transactions on Biomedical Engineering*. 1997; 44:374-385. doi: 10.1109/10.568913. Pascual-Marqui R D. Review of Methods for Solving the EEG Inverse Problem. *International Journal of Bioelectromagnetism*. 1999; 1:75-86. Baillet S, Mosher J C, Leahy R M. Electromagnetic Brain Mapping. *IEEE Signal Processing Magazine*. 2001; 18:14-30. doi: 10.1109/79.962275. Groetsch W. Inverse Problems in the Mathematical Sciences. Vieweg. 1993. Hansen P C. Rank-Deficient and Discrete III-Posed Problems. SIAM. 1998. Vogel C R. Computational Methods for Inverse Problems. SIAM. 2002. De Munck J C. The estimation of time varying dipoles on the basis of evoked potentials. *Electroencephalography and Clinical Neurophysiology*. 1990; 77:156-160. doi: 10.1016/0168-5597(90)90032-9. Rodriguez-Rivera A, Van Veen B D, Wakai R T. Statistical Performance Analysis of Signal Variance-Based Dipole Models for MEG/EEG Source Localization and Detection. *IEEE Transactions on Biomedical Engineering*. 2003; 50:137-149. doi: 10.1109/TBME.2002.807661. Liu A K, Dale A M, Belliveau J W. Monte Carlo Simulation Studies of EEG and MEG Localization Accuracy. *Human Brain Mapping*. 2002; 16:47-62. doi: 10.1002/hbm.10024. Schmidt D M, George J S, Wood C C. Bayesian Inference Applied to the Electromagnetic Inverse Problem. Progress Report 1997-1998, Physics Division. 2002. Dale A, Sereno M. Improved Localization of Cortical Activity By Combining EEG and MEG with MRI Cortical Surface Reconstruction: A Linear Approach. *Journal of Cognitive Neuroscience*. 1993; 5:162-176. doi: 10.1162/jocn.1993.5.2.162. Gavit L, Baillet S, Mangin J F, Pascatore J, Gamero L. A Multiresolution Framework to MEG/EEG Source Imaging. *IEEE Transactions on Biomedical*

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(271) The first applications of blind decomposition to biomedical time series analysis applied the infomax independent component analysis (ICA) algorithm to decomposition of EEG and event-related potential (ERP) data and reported the use of ICA to monitor alertness. This separated artifacts, and EEG data into constituent components defined by spatial stability and temporal independence. ICA can also be used to remove artifacts from continuous or event-related (single-trial) EEG data prior to averaging. Vigario et al. (1997), using a different ICA algorithm, supported the use of ICA for identifying artifacts in MEG data. Meanwhile, widespread interest in ICA has led to multiple applications to biomedical data as well as to other fields (Jung et al., 2000b). Most relevant to EEG/MEG analysis, ICA is effective in separating functionally independent components of functional magnetic resonance imaging (fMRI) data

(272) Since the publication of the original infomax ICA algorithm, several extensions have been proposed. Incorporation of a 'natural gradient' term avoided matrix inversions, greatly speeding the convergence of the algorithm and making it practical for use with personal computers on large data EEG and fMRI data sets. An initial 'sphering' step further increased the reliability of convergence of the algorithm. The original algorithm assumed that sources have 'sparse' (super-Gaussian) distributions of activation values. This restriction has recently been relaxed in an 'extended-ICA' algorithm that allows both super-Gaussian and sub-Gaussian sources to be identified. A number of variant ICA algorithms have appeared in the signal processing literature. In general, these make more specific assumptions about the temporal or spatial structure of the components to be separated, and typically are more computationally intensive than the infomax algorithm.

(273) Since individual electrodes (or magnetic sensors) each record a mixture of brain and non-brain sources, spectral measures are difficult to interpret and compare across scalp channels. For example, an increase in coherence between two electrode signals may reflect the activation of a strong brain source projecting to both electrodes, or the deactivation of a brain generator projecting mainly to one of the electrodes. If independent components of the EEG (or MEG) data can be considered to measure activity within functionally distinct brain networks, however, event-related coherence between independent components may reveal transient, event-related changes in their coupling and decoupling (at one or more EEG/MEG frequencies). ERCOH analysis has been applied to independent EEG components in a selective attention task.

(274) Relational Database A database management system (DBMS) is the software which controls the storage, retrieval, deletion, security, and integrity of data within a database. A relational database management system (RDBMS) stores data in tables. Tables are organized into columns, and each column stores one type of data (integer, real number, character strings, date, . . .). The data for a single "instance" of a table is stored as a row. For example, an emotional neural correlate table would have columns such as EmotionLabel, NeuralCorrelate1_under_condition1, NeuralCorrelate2_under_condition2, NeuralCorrelate3_under_condition3, NeuralCorrelate4_under_condition4, etc. Tables typically have keys, one or more columns that uniquely identify a row within the table, in the case of the Emotional neural correlate table the key would be EmotionLabel. To improve access time to a data table an index on the table is defined. An index provides a quick way to look up data based on one or more columns in the table. The most common use of RDBMSs is to implement simple Create, Read, Update, and Delete. A relational database may be manipulated using Structured Query Language (SQL) statements. en.wikipedia.org/wiki/Relational_database. The relational database may be a SQL or noSQL database.

SUMMARY OF THE INVENTION

(275) In other embodiments, the processing of the brain activity patterns does not seek to classify or characterize it, but rather to filter and transform

the information to a form suitable for control of the stimulation of the second subject, according to the embodiment, the subtleties that are not yet reliably classified in traditional brain activity pattern analysis are respected. For example, it is understood that all brain activity is reflected in synaptic currents and other neural modulation and, therefore, theoretically, conscious and subconscious information is, in theory, accessible through brain activity pattern analysis. Since the available processing technology generally fails to distinguish a large number of different brain activity patterns, that available processing technology, is necessarily deficient, but improving. However, just because a computational algorithm is unavailable to extract the information, does not mean that the information is absent. Therefore, this embodiment employs relatively raw brain activity pattern data, such as filtered or unfiltered EEGs, to control the stimulation of the second subject, without a full comprehension or understanding of exactly what information of significance is present. In one embodiment, brainwaves are recorded and “played back” to another subject, similar to recoding and playing back music. Such recording-playback may be digital or analog. Typically, the stimulation may include a low dimensionality stimulus, such as stereo-optic, binaural, isotonic tones, tactile, or other sensory stimulation, operating bilaterally, and with control over frequency and phase and/or waveform and/or transcranial stimulation such as TES, tDCS, HD-tDCS, tACS, or TMS. A plurality of different types of stimulation may be applied concurrently, e.g., visual, auditory, other sensory, magnetic, electrical.

(276) Likewise, a present lack of understanding of the essential characteristics of the signal components in the brain activity patterns does not prevent their acquisition, storage, communication, and processing (to some extent). The stimulation may be direct, i.e., a visual, auditory, or tactile stimulus corresponding to the brain activity pattern, or a derivative or feedback control based on the second subject's brain activity pattern.

(277) To address the foregoing problems, in whole or in part, and/or other problems that may have been observed by persons skilled in the art, the present disclosure provides methods, processes, systems, apparatus, instruments, and/or devices, as described by way of example in implementations set forth below.

(278) While mental states are typically considered internal to the individual, and subjective, in fact, such states are common across individuals and have determinable physiological and electrophysiological population characteristics. Further, mental states may be externally changed or induced in a manner that bypasses the normal cognitive processes. In some cases, the triggers for the mental state are subjective, and therefore the particular subject-dependent sensory or excitation scheme required to induce a particular state will differ. For example, olfactory stimulation can have different effects on different people, based on differences in history of exposure, social and cultural norms, and the like. On the other hand, some mental state response triggers are normative, for example “tear jerker” media.

(279) Mental states are represented in brainwave patterns, and in normal humans, the brainwave patterns and metabolic (e.g. blood flow, oxygen consumption, etc.) follow prototypical patterns. Therefore, by monitoring brainwave patterns in an individual, a state or series of mental states in that person may be determined or estimated. However, the brainwave patterns may be interrelated with context, other activity, and past history. Further, while prototypical patterns may be observed, there are also individual variations in the patterns. The brainwave patterns may include characteristic spatial and temporal patterns indicative of mental state. The brainwave signals of a person may be processed to extract these patterns, which, for example, may be represented as hemispheric signals within a frequency range of 3-100 Hz. These signals may then be synthesized or modulated into one or more stimulation signals, which are then employed to induce a corresponding mental state into a recipient, in a manner seeking to achieve a similar brainwave pattern from the source. The brainwave pattern to be introduced need not be newly acquired for each case. Rather, signals may be acquired from one or more individuals, to obtain an exemplar for various respective mental state. Once determined, the processed signal representation may be stored in a non-volatile memory for later use. However, in cases of complex interaction between a mental state and a context or content or activity, it may be appropriate to derive the signals from a single individual whose context or content-environment or activity is appropriate for the circumstances. Further, in some cases, a single mental state, emotion or mood is not described or fully characterized, and therefore acquiring signals from a source is an efficient exercise.

(280) With a library of target brainwave patterns, a system and method is provided in which a target subject may be immersed in a presentation, which includes not only multimedia content, but also a series of defined mental states, emotional states or moods that accompany the multimedia content. In this way, the multimedia presentation becomes fully immersive. The stimulus in this case may be provided through a headset such as a virtual reality or augmented reality headset. This headset is provided with a stereoscopic display, binaural audio, and a set of EEG and transcranial stimulatory electrodes. These electrodes (if provided) typically deliver a subthreshold signal, which is not painful, which is typically an AC signal which corresponds to the desired frequency, phase, and spatial location of the desired target pattern. The electrodes may also be used to counteract undesired signals, by destructively interfering with them while concurrently imposing the desired patterns. The headset may also generate visual and/or auditory signals which correspond to the desired state. For example, the auditory signals may induce binaural beats, which cause brainwave entrainment. The visual signals may include intensity fluctuations or other modulation patterns, especially those which are subliminal, that are also adapted to cause brainwave entrainment or induction of a desired brainwave pattern.

(281) The headset preferably includes EEG electrodes for receiving feedback from the user. That is, the stimulatory system seeks to achieve a mental state, emotion or mood response from the user. The EEG electrodes permit determination of whether that state is achieved, and if not, what the current state is. It may be that achieving a desired brainwave pattern is state dependent and therefore that characteristics of the stimulus to achieve a desired state depend on the starting state of the subject. Other ways of determining mental state, emotion, or mood include analysis of facial expression, electromyography (EMG) analysis of facial muscles, explicit user feedback, etc.

(282) An authoring system is provided which permits a content designer to determine what mental states are desired, and then encode those states into media, which is then interpreted by a media reproduction system in order to generate appropriate stimuli. As noted above, the stimuli may be audio, visual, multimedia, other senses, or electrical or magnetic brain stimulation, and therefore a VR headset with transcranial electrical or magnetic stimulation is not required. Further, in some embodiments, the patterns may be directly encoded into the audiovisual content, subliminally encoded.

(283) In some cases, the target mental state may be derived from an expert, actor or professional exemplar. The states may be read based on facial expressions, EMG, EEG, or other means, from the actor or exemplar. For example, a prototype exemplar engages in an activity that triggers a response, such as viewing the Grand Canyon or artworks within the Louvre. The responses of the exemplar are then recorded or represented, and preferably brainwave patterns recorded that represent the responses. A representation of the same experience is then presented to the target, with a goal of the target also experiencing the same experience as the exemplar. This is typically a voluntary and disclosed process, so the target will seek to willingly comply with the desired experiences. In some cases, the use of the technology is not disclosed to the target, for example in advertising presentations or billboards. In order for an actor to serve as the exemplar, the emotions achieved by that person must be authentic. However, so-called “method actors” do authentically achieve the emotions they convey. However, in some cases, for example where facial expressions are used as the indicator of mental state, an actor can present desired facial expressions with inauthentic mental states. The act of making a face corresponding to an emotion often achieves the targeted mental state.

(284) In order to calibrate the system, the brain pattern of a person may be measured while in the desired state. The brain patterns acquired for calibration or feedback need not be of the same quality, or precision, or data depth, and indeed may represent responses rather than primary indicia. That is, there may be some asymmetry in the system, between the brainwave patterns representative of a mental state, and the stimulus patterns appropriate for inducing the brain state.

(285) The present invention generally relates to achieving a mental state in a subject by conveying to the brain of the subject patterns of brainwaves. These brainwaves may be artificial or synthetic, or derived from the brain of a second subject (e.g., a person experiencing an authentic experience or engaged in an activity). Typically, the wave patterns of the second subject are derived while the second subject is experiencing an authentic experience.

(286) A special case is where the first and second subjects are the same individual. For example, brainwave patterns are recorded while a subject is in a particular mental state. That same pattern may assist in achieving the same mental state at another time. Thus, there may be a time delay between acquisition of the brainwave information from the second subject, and exposing the first subject to corresponding stimulation. The signals may be recorded and transmitted.

(287) The temporal pattern may be conveyed or induced non-invasively via light (visible or infrared), sound (or ultrasound), transcranial direct or alternating current stimulation (tDCS or tACS), transcranial magnetic stimulation (TMS), Deep transcranial magnetic stimulation (Deep TMS, or dTMS), Repetitive Transcranial Magnetic Stimulation (rTMS) olfactory stimulation, tactile stimulation, or any other means capable of conveying frequency patterns. In a preferred embodiment, normal human senses are employed to stimulate the subject, such as light, sound, smell and touch. Combinations of stimuli may be employed. In some cases, the stimulus or combination is innate, and therefore largely pan-subject. In other cases, response to a context is learned, and therefore subject-specific. Therefore, feedback from the subject may be appropriate to determine the triggers and stimuli appropriate to achieve a mental state.

(288) This technology may be advantageously used to enhance mental response to a stimulus or context. Still another aspect provides for a change in the mental state. The technology may be used in humans or animals.

(289) The present technology may employ an event-correlated EEG time and/or frequency analysis performed on neuronal activity patterns. In a time-analysis, the signal is analyzed temporally and spatially, generally looking for changes with respect to time and space. In a frequency analysis, over an epoch of analysis, the data, which is typically a time-sequence of samples, is transformed, using e.g., a Fourier transform (FT, or one implementation, the Fast Fourier Transform, FFT), into a frequency domain representation, and the frequencies present during the epoch are analyzed. The window of analysis may be rolling, and so the frequency analysis may be continuous. In a hybrid time-frequency analysis, for example, a wavelet analysis, the data during the epoch is transformed using a "wavelet transform", e.g., the Discrete Wavelet Transform (DWT) or continuous wavelet transform (CWT), which has the ability to construct a time-frequency representation of a signal that offers very good time and frequency localization. Changes in transformed data over time and space may be analyzed. In general, the spatial aspect of the brainwave analysis is anatomically modelled. In most cases, anatomy is considered universal, but in some cases, there are significant differences. For example, brain injury, psychiatric disease, age, race, native language, training, sex, handedness, and other factors may lead to distinct spatial arrangement of brain function, and therefore when transferring mood from one individual to another, it is preferred to normalize the brain anatomy of both individuals by experiencing roughly the same experiences, and measuring spatial parameters of the EEG or MEG. Note that spatial organization of the brain is highly persistent, absent injury or disease, and therefore this need only be performed infrequently. However, since electrode placement may be inexact, a spatial calibration may be performed after electrode placement.

(290) Different aspects of EEG magnitude and phase relationships may be captured, to reveal details of the neuronal activity. The "time-frequency analysis" reveals the brain's parallel processing of information, with oscillations at various frequencies within various regions of the brain reflecting multiple neural processes co-occurring and interacting. See, Lisman J, Buzsaki G. A neural coding scheme formed by the combined function of gamma and theta oscillations. *Schizophrenia Bull.* Jun. 16, 2008; doi:10.1093/schbul/sbn060. Such a time-frequency analysis may take the form of a wavelet transform analysis. This may be used to assist in integrative and dynamically adaptive information processing. Of course, the transform may be essentially lossless and may be performed in any convenient information domain representation. These EEG-based data analyses reveal the frequency-specific neuronal oscillations and their synchronization in brain functions ranging from sensory processing to higher-order cognition. Therefore, these patterns may be selectively analyzed, for transfer to or induction in, a subject.

(291) A statistical clustering analysis may be performed in high dimension space to isolate or segment regions which act as signal sources, and to characterize the coupling between various regions. This analysis may also be used to establish signal types within each brain region, and decision boundaries characterizing transitions between different signal types. These transitions may be state dependent, and therefore the transitions may be detected based on a temporal analysis, rather than merely a concurrent oscillator state.

(292) The various measures make use of the magnitude and/or phase angle information derived from the complex data extracted from the EEG during spectral decomposition and/or temporal/spatial/spectral analysis. Some measures estimate the magnitude or phase consistency of the EEG within one channel across trials, whereas others estimate the consistency of the magnitude or phase differences between channels across trials. Beyond these two families of calculations, there are also measures that examine the coupling between frequencies, within trials and recording sites. Of course, in the realm of time-frequency analysis, many types of relationships can be examined beyond those already mentioned.

(293) These sensory processing specific neuronal oscillations, e.g., brainwave patterns, e.g., of a subject (a "source") or to a person trained (for example, an actor trained in "the method") to create a desired state, and can be stored on a tangible medium and/or can be simultaneously conveyed to a recipient making use of the brain's frequency following response nature. See, Galbraith, Gary C., Darlene M. Oilman, and Todd M. Huffman. "Selective attention affects human brain stem frequency-following response." *Neuroreport* 14, no. 5 (2003): 735-738, journals.lww.com/neuroreport/Abstract/2003/04150/Selective_attention_affects_human_brain_stem.15.aspx.

(294) Of course, in some cases, one or more components of the stimulation of the target subject (recipient) may be represented as abstract or semantically defined signals, and, more generally, the processing of the signals to define the stimulation will involve high level modulation or transformation between the source signal received from the first subject (donor) or plurality of donors, to define the target signal for stimulation of the second subject (recipient).

(295) Preferably, each component represents a subset of the neural correlates reflecting brain activity that have a high autocorrelation in space and time, or in a hybrid representation such as wavelet. These may be separated by optimal filtering (e.g., spatial PCA), once the characteristics of the signal are known, and bearing in mind that the signal is accompanied by a modulation pattern, and that the two components themselves may have some weak coupling and interaction.

(296) For example, if the first subject (donor) is listening to music, there will be significant components of the neural correlates that are synchronized with the particular music. On the other hand, the music per se may not be part of the desired stimulation of the target subject (recipient). Further, the target subject (recipient) may be in a different acoustic environment, and it may be appropriate to modify the residual signal dependent on the acoustic environment of the recipient, so that the stimulation is appropriate for achieving the desired effect, and does not represent phantoms, distractions, or irrelevant or inappropriate content. In order to perform signal processing, it is convenient to store the signals or a partially processed representation, though a complete real-time signal processing chain may be implemented.

(297) The stimulation may be one or more stimulus applied to the second subject (trainee or recipient), which may be an electrical or magnetic transcranial stimulation (tDCS, HD-tDCS, tACS, osc-tDCS, or TMS), sensory stimulation (e.g., visual, auditory, or tactile), mechanical stimulation, ultrasonic stimulation, etc., and controlled with respect to waveform, frequency, phase, intensity/amplitude, duration, or controlled via feedback, self-reported effect by the second subject, manual classification by third parties, automated analysis of brain activity, behavior, physiological parameters, etc. of the second subject (recipient).

(298) Typically, the first and the second subjects are spatially remote from each other and may be temporally remote as well. In some cases, the first and second subject are the same subject (human or animal), temporally displaced. In other cases, the first and the second subject are spatially proximate to each other. These different embodiments differ principally in the transfer of the signal from at least one first subject (donor) to the second subject (recipient). However, when the first and the second subjects share a common environment, the signal processing of the neural correlates and, especially of real-time feedback of neural correlates from the second subject, may involve interactive algorithms with the neural correlates of the first subject.

(299) According to another embodiment, the first and second subjects are each subject to stimulation. In one particularly interesting embodiment, the

first subject and the second subject communicate with each other in real-time, with the first subject receiving stimulation based on the second subject, and the second subject receiving feedback based on the first subject. This can lead to synchronization of neural correlates (e.g., neuronal oscillations, or brainwaves) and, consequently, of emotional or mental state between the two subjects. The neural correlates may be neuronal oscillations resulting in brainwaves that are detectable as, for example, EEG, qEEG, or MEG signals. Traditionally, these signals are found to have dominant frequencies, which may be determined by various analyses, such as spectral analysis, wavelet analysis, or principal component analysis (PCA), for example. One embodiment provides that the modulation pattern of a brainwave of at least one first subject (donor) is determined independent of the dominant frequency of the brainwave (though, typically, within the same class of brainwaves), and this modulation imposed on a brainwave corresponding to the dominant frequency of the second subject (recipient). That is, once the second subject achieves that same brainwave pattern as the first subject (which may be achieved by means other than electromagnetic, mechanical, or sensory stimulation), the modulation pattern of the first subject is imposed as a way of guiding the emotional or mental state of the second subject.

(300) According to another embodiment, the second subject (recipient) is stimulated with a stimulation signal, which faithfully represents the frequency composition of a defined component of the neural correlates of at least one first subject (donor). The defined component may be determined based on a principal component analysis, independent component analysis (ICI), eigenvector-based multivariable analysis, factor analysis, canonical correlation analysis (CCA), nonlinear dimensionality reduction (NLDR), or related technique.

(301) The stimulation may be performed, for example, by using a TES device, such as a tDCS device, a high-definition tDCS device, an osc-tDCS device, a pulse-tDCS (“electrosleep”) device, an osc-tDCS, a tACS device, a CES device, a TMS device, rTMS device, a deep TMS device, a light source, or a sound source configured to modulate the dominant frequency on respectively the light signal or the sound signal. The stimulus may be a light signal, a sonic signal (sound), an electric signal, a magnetic field, olfactory or a tactile stimulation. The current signal may be a pulse signal or an oscillating signal. The stimulus may be applied via a cranial electric stimulation (CES), a transcranial electric stimulation (TES), a deep electric stimulation, a transcranial magnetic stimulation (TMS), a deep magnetic stimulation, a light stimulation, a sound stimulation, a tactile stimulation, or an olfactory stimulation. An auditory stimulus may be, for example, binaural beats or isochronic tones.

(302) The technology also provides a processor configured to process the neural correlates of emotional or mental state from the first subject (donor), and to produce or define a stimulation pattern for the second subject (recipient) selectively dependent on a waveform pattern of the neural correlates from the first subject. The processor may also perform a PCA, a spatial PCA, an independent component analysis (ICA), eigenvalue decomposition, eigenvector-based multivariate analyses, factor analysis, an autoencoder neural network with a linear hidden layer, linear discriminant analysis, network component analysis, nonlinear dimensionality reduction (NLDR), or another statistical method of data analysis.

(303) A signal is presented to a second apparatus, configured to stimulate the second subject (recipient), which may be an open loop stimulation dependent on a non-feedback-controlled algorithm, or a closed loop feedback dependent algorithm. The second apparatus produces a stimulation intended to induce in the second subject (recipient) the desired emotional or mental state).

(304) A typically process performed on the neural correlates is a filtering to remove noise. In some embodiments, noise filters may be provided, for example, at 50 Hz, 60 Hz, 100 Hz, 120 Hz, and additional overtones (e.g., tertiary and higher harmonics). The stimulator associated with the second subject (recipient) would typically perform decoding, decompression, decryption, inverse transformation, modulation, etc.

(305) Alternately, an authentic wave or hash thereof may be authenticated via a blockchain, and thus authenticatable by an immutable record. In some cases, it is possible to use the stored encrypted signal in its encrypted form, without decryption.

(306) Due to different brain sizes, and other anatomical, morphological, and/or physiological differences, dominant frequencies associated with the same emotional or mental state may be different in different subjects. Consequently, it may not be optimal to forcefully impose on the recipient the frequency of the donor that may or may not precisely correspond to the recipients frequency associated with the same emotional or mental state. Accordingly, in some embodiments, the donor's frequency may be used to start the process of inducing the desired emotional or mental state in a recipient. As some point, when the recipient is close to achieving the desired emotional or mental state, the stimulation is either stopped or replaced with neurofeedback allowing the brain of the recipient to find its own optimal frequency associated with the desired emotional or mental state.

(307) In one embodiment, the feedback signal from the second subject may be correspondingly encoded as per the source signal, and the error between the two minimized. According to one embodiment, the processor may perform a noise reduction distinct from a frequency-band filtering. According to one embodiment, the neural correlates are transformed into a sparse matrix, and in the transform domain, components having a high probability of representing noise are masked, while components having a high probability of representing signal are preserved. That is, in some cases, the components that represent modulation that are important may not be known a priori. However, dependent on their effect in inducing the desired response in the second subject (recipient), the “important” components may be identified, and the remainder filtered or suppressed. The transformed signal may then be inverse-transformed and used as a basis for a stimulation signal.

(308) According to another embodiment, a method of emotional or mental state modification, e.g., brain entrainment, is provided, comprising: ascertaining an emotional or mental state in a plurality of first subjects (donors); acquiring brainwaves of the plurality of first subjects (donors), e.g., using one of EEG and MEG, to create a dataset containing brainwaves corresponding to different emotional or mental states. The database may be encoded with a classification of emotional or mental states, activities, environment, or stimulus patterns, applied to the plurality of first subjects, and the database may include acquired brainwaves across a large number of emotional or mental states, activities, environment, or stimulus patterns, for example. In many cases, the database records will reflect a characteristic or dominate frequency of the respective brainwaves.

(309) The record(s) thus retrieved are used to define a stimulation pattern for the second subject (recipient). As a relatively trivial example, a female recipient could be stimulated principally based on records from female donors. Similarly, a child recipient of a certain age could be stimulated principally based on the records from children donors of a similar age. Likewise, various demographic, personality, and/or physiological parameters may be matched to ensure a high degree of correspondence to between the source and target subjects. In the target subject, a guided or genetic algorithm may be employed to select modification parameters from the various components of the signal, which best achieve the desired target state based on feedback from the target subject.

(310) Of course, a more nuanced approach is to process the entirety of the database and stimulate the second subject based on a global brainwave-stimulus model, though this is not required, and also, the underlying basis for the model may prove unreliable or inaccurate. In fact, it may be preferred to derive a stimulus waveform from only a single first subject (donor), in order to preserve micro-modulation aspects of the signal, which, as discussed above, have not been fully characterized. However, the selection of the donor(s) need not be static and can change frequently. The selection of donor records may be based on population statistics of other users of the records, i.e., whether or not the record had the expected effect, filtering donors whose response pattern correlates highest with a given recipient, etc. The selection of donor records may also be based on feedback patterns from the recipient.

(311) The process of stimulation typically seeks to target a desired emotional or mental state in the recipient, which is automatically or semi-automatically determined or manually entered. In one embodiment, the records are used to define a modulation waveform of a synthesized carrier or set of carriers, and the process may include a frequency domain multiplexed multi-subcarrier signal (which is not necessarily orthogonal). A plurality of stimuli may be applied concurrently, through the different subchannels and/or through different stimulator electrodes, electric current stimulators, magnetic field generators, mechanical stimulators, sensory stimulators, etc. The stimulus may be applied to achieve brain entrainment (i.e., synchronization) of the second subject (recipient) with one or more first subjects (donors). If the plurality of donors is mutually entrained, then each will have a corresponding brainwave pattern dependent on the basis of brainwave entrainment. This link between donors may be helpful in determining compatibility between a respective donor and the recipient. For example, characteristic patterns in the entrained brainwaves may be determined, even for different target emotional or mental states, and the characteristic patterns may be correlated to find relatively close matches and

to exclude relatively poor matches.

(312) This technology may also provide a basis for a social network, dating site, employment, mission (e.g., space or military), or vocational testing, or other interpersonal environments, wherein people may be matched with each other based on entrainment characteristics. For example, people who efficiently entrain with each other may have better compatibility and, therefore, better marriage, work, or social relationships than those who do not. The entrainment effect need not be limited to emotional or mental states, and may arise across any context.

(313) As discussed above, the plurality of first subjects (donors) may have their respective brainwave patterns stored in separate database records. Data from a plurality of first subjects (donors) is used to train the neural network, which is then accessed by inputting the target stage and/or feedback information, and which outputs a stimulation pattern or parameters for controlling a stimulator(s). When multiple first subject (donors) form the basis for the stimulation pattern, it is preferred that the neural network output parameters of the stimulation, derived from and comprising features of the brainwave patterns or other neural correlates of the emotional or mental state from the plurality of first subject (donors), which are then used to control a stimulator which, for example, generates its own carrier wave(s) which are then modulated based on the output of the neural network. A trained neural network need not periodically retrieve records, and therefore may operate in a more time-continuous manner, rather than the more segmented scheme of record-based control.

(314) In any of the feedback dependent methods, the brainwave patterns or other neural correlates of emotional or mental states may be processed by a neural network, to produce an output that guides or controls the stimulation. The stimulation is, for example, at least one of a light signal, a sound signal, an electric signal, a magnetic field, an olfactory signal, a chemical signal, and a vibration or mechanical stimulus. The process may employ a relational database of emotional or mental states and brainwave patterns, e.g., frequencies/neural correlate waveform patterns associated with the respective emotional or mental states. The relational database may comprise a first table, the first table further comprising a plurality of data records of brainwave patterns, and a second table, the second table comprising a plurality of emotional or mental states, each of the emotional or mental states being linked to at least one brainwave pattern. Data related to emotional or mental states and brainwave patterns associated with the emotional or mental states are stored in the relational database and maintained. The relational database is accessed by receiving queries for selected (existing or desired) emotional or mental states, and data records are returned representing the associated brainwave pattern. The brainwave pattern retrieved from the relational database may then be used for modulating a stimulator seeking to produce an effect selectively dependent on the desired emotional or mental state.

(315) A further aspect of the technology provides a computer apparatus for creating and maintaining a relational database of emotional or mental states and frequencies associated with the emotional or mental state. The computer apparatus may comprise a non-volatile memory for storing a relational database of emotional or mental states and neural correlates of brain activity associated with the emotional or mental states, the database comprising a first table comprising a plurality of data records of neural correlates of brain activity associated with the emotional or mental states, and a second table comprising a plurality of emotional or mental states, each of the emotional or mental states being linked to one or more records in the first table; a processor coupled with the non-volatile memory, and being configured to process relational database queries, which are then used for searching the database; RAM coupled with the processor and the non-volatile memory for temporary holding database queries and data records retrieved from the relational database; and an I/O interface configured to receive database queries and deliver data records retrieved from the relational database. A structured query language (SQL) or alternate to SQL (e.g., noSQL) database may also be used to store and retrieve records. A relational database described above maintained and operated by a general-purpose computer, improves the operations of the general-purpose computer by making searches of specific emotional or mental states and brainwaves associated therewith more efficient thereby, inter alia, reducing the demand on computing power.

(316) A further aspect of the technology provides a method of brain entrainment comprising: ascertaining an emotional or mental state in at least one first subject (donor), recording brainwaves of said at least one first subject (donor) using at least one channel of EEG and/or MEG; storing the recorded brainwaves in a physical memory device, retrieving the brainwaves from the memory device, applying a stimulus signal comprising a brainwave pattern derived from at least one-channel of the EEG and/or MEG to a second subject (recipient) via transcranial electrical and/or magnetic stimulation, whereby the emotional or mental state desired by the second subject (recipient) is achieved. The stimulation may be of the same dimension (number of channels) as the EEG or MEG, or a different number of channels, typically reduced. For example, the EEG or MEG may comprise 64, 128 or 256 channels, while the transcranial stimulator may have 32 or fewer channels. The placement of electrodes used for transcranial stimulation may be approximately the same as the placement of electrodes used in recording of EEG or MEG to preserve the topology of the recorded signals and, possibly, use these signals for spatial modulation.

(317) One of the advantages of transforming the data is the ability to select a transform that separates the information of interest represented in the raw data, from noise or other information. Some transforms preserve the spatial and state transition history, and may be used for a more global analysis. Another advantage of a transform is that it can present the information of interest in a form where relatively simple linear or statistical functions of low order may be applied. In some cases, it is desired to perform an inverse transform on the data. For example, if the raw data includes noise, such as 50 or 60 Hz interference, a frequency transform may be performed, followed by a narrow band filtering of the interference and its higher order intermodulation products. An inverse transform may be performed to return the data to its time-domain representation for further processing. (In the case of simple filtering, a finite impulse response (FIR) or infinite impulse response (IIR) filter could be employed). In other cases, the analysis is continued in the transformed domain.

(318) Transforms may be part of an efficient algorithm to compress data for storage or analysis, by making the representation of the information of interest consume fewer bits of information (if in digital form) and/or allow it to be communicated using lower bandwidth. Typically, compression algorithms will not be lossless, and as a result, the compression is irreversible with respect to truncated information.

(319) Typically, the transformation(s) and filtering of the signal are conducted using traditional computer logic, according to defined algorithms. The intermediate stages may be stored and analyzed. However, in some cases, neural networks or deep neural networks may be used, convolutional neural network architectures, or even analog signal processing. According to one set of embodiments, the transforms (if any) and analysis are implemented in a parallel processing environment such as using a SIMD processor such as a GPU (or GPGPU). Algorithms implemented in such systems are characterized by an avoidance of data-dependent branch instructions, with many threads concurrently executing the same instructions.

(320) EEG signals are analyzed to determine the location (e.g., voxel or brain region) from which an electrical activity pattern is emitted, and the wave pattern characterized. The spatial processing of the EEG signals will typically precede the content analysis, since noise and artifacts may be useful for spatial resolution. Further, the signal from one brain region will typically be noise or interference in the signal analysis from another brain region; so the spatial analysis may represent part of the comprehension analysis. The spatial analysis is typically in the form of a geometrically and/or anatomically-constrained statistical model, employing all of the raw inputs in parallel. For example, where the input data is transcutaneous electroencephalogram information, from 32 EEG electrodes, the 32 input channels, sampled at e.g., 500 sps, 1 ksp/s or 2 ksp/s, are processed in a four or higher dimensional matrix, to permit mapping of locations and communication of impulses over time, space and state.

(321) The matrix processing may be performed in a standard computing environment, e.g., an i7-7920HQ, i7-8700K, or i9-7980XE processor, under the Windows 10 operating system, executing MatLab (MathWorks, Woburn MA) software platform. Alternately, the matrix processing may be performed in a computer cluster or grid or cloud computing environment. The processing may also employ parallel processing, in either a distributed and loosely coupled environment, or asynchronous environment. One preferred embodiment employs a single instruction, multiple data processors, such as a graphics processing unit such as the nVidia CUDA environment or AMD Firepro high-performance computing environment. Artificial intelligence (AI) and ML methods, such as artificial neural networks, deep neural networks, etc., may be implemented to extract the signals of interest. Neural networks act as an optimized statistical classifier and may have arbitrary complexity. A so-called deep neural network having multiple

hidden layers may be employed. The processing is typically dependent on labeled training data, such as EEG data, or various processed, transformed, or classified representations of the EEG data. The label represents the emotion, mood, context, or state of the subject during acquisition. In order to handle the continuous stream of data represented by the EEG, a recurrent neural network architecture may be implemented. Depending preprocessing before the neural network, formal implementations of recurrence may be avoided. A four or more dimensional data matrix may be derived from the traditional spatial-temporal processing of the EEG and fed to a neural network. Since the time parameter is represented in the input data, a neural network temporal memory is not required, though this architecture may require a larger number of inputs. Principal component analysis (PCA, en.wikipedia.org/wiki/Principal_component_analysis), spatial PCA (arxiv.org/pdf/1501.03221v3.pdf, adegenetr-forger-project.org/files/tutorial-spca.pdf, www.ncbi.nlm.nih.gov/pubmed/1510870); and clustering analysis may also be employed (en.wikipedia.org/wiki/Cluster_analysis, see U.S. Pat. Nos. 9,336,302, 9,607,023 and cited references).

(322) In general, a neural network of this type of implementation will, in operation, be able to receive unlabeled EEG data, and produce the output signals representative of the predicted or estimated task, performance, context, or state of the subject during acquisition of the unclassified EEG. Of course, statistical classifiers may be used rather than neural networks.

(323) The analyzed EEG, either by conventional processing, neural network processing, or both, serves two purposes. First, it permits one to deduce which areas of the brain are subject to which kinds of electrical activity under which conditions. Second, it permits feedback during training of a trainee (assuming proper spatial and anatomical correlates between the trainer and trainee), to help the system achieve the desired state, or as may be appropriate, desired series of states and/or state transitions. According to one aspect of the technology, the applied stimulation is dependent on a measured starting state or status (which may represent a complex context and history dependent matrix of parameters), and therefore the target represents a desired complex vector change. Therefore, this aspect of the technology seeks to understand a complex time-space-brain activity associated with an activity or task in a trainer, and to seek a corresponding complex time-space-brain activity associated with the same activity or task in a trainee, such that the complex time-space-brain activity state in the trainer is distinct from the corresponding state sought to be achieved in the trainee. This permits transfer of training paradigms from qualitatively different persons, in different contexts, and, to some extent, to achieve a different result.

(324) The conditions of data acquisition from the trainer will include both task data, and sensory-stimulation data. That is, a preferred application of the system is to acquire EEG data from a trainer or skilled individual, which will then be used to transfer learning, or more likely, learning readiness states, to a naïve trainee. The goal for the trainee is to produce a set of stimulation parameters that will achieve, in the trainee, the corresponding neural activity resulting in the EEG state of the trainer at the time of or preceding the learning of a skill or a task, or performance of the task.

(325) It is noted that EEG is not the only neural or brain activity or state data that may be acquired, and of course any and all such data may be included within the scope of the technology, and therefore EEG is a representative example only of the types of data that may be used. Other types include fMRI, magnetoencephalogram, motor neuron activity, PET, etc.

(326) While mapping the stimulus-response patterns distinct from the task is not required in the trainer, it is advantageous to do so, because the trainer may be available for an extended period, the stimulus of the trainee may influence the neural activity patterns, and it is likely that the trainer will have correlated stimulus-response neural activity patterns with the trainee(s). It should be noted that the foregoing has suggested that the trainer is a single individual, while in practice, the trainer may be a population of trainers or skilled individuals. The analysis and processing of brain activity data may, therefore, be adaptive, both for each respective individual and for the population as a whole.

(327) For example, the system may determine that not all human subjects have common stimulus-response brain activity correlates, and therefore that the population needs to be segregated and clustered. If the differences may be normalized, then a normalization matrix or other correction may be employed. On the other hand, if the differences do not permit feasible normalization, the population(s) may be segmented, with different trainers for the different segments. For example, in some tasks, male brains have different activity patterns and capabilities than female brains. This, coupled with anatomical differences between the sexes, implies that the system may provide gender-specific implementations. Similarly, age differences may provide a rational and scientific basis for segmentation of the population. However, depending on the size of the information base and matrices required, and some other factors, each system may be provided with substantially all parameters required for the whole population, with a user-specific implementation based on a user profile or initial setup, calibration, and system training session.

(328) According to one aspect of the present invention, a source subject is instrumented with sensors to determine localized brain activity during experiencing an event. The objective is to identify regions of the brain involved in processing this response.

(329) The sensors will typically seek to determine neuron firing patterns and brain region excitation patterns, which can be detected by implanted electrodes, transcutaneous electroencephalograms, magnetoencephalograms, fMRI, and other technologies. Where appropriate, transcutaneous EEG is preferred, since this is non-invasive and relatively simple.

(330) The source is observed with the sensors in a quiet state, a state in which he or she is experiencing an event, and various control states in which the source is at rest or engaged in different activities resulting in different states. The data may be obtained for a sufficiently long period of time and over repeated trials to determine the effect of duration. The data may also be a population statistical result, and need not be derived from only a single individual at a single time.

(331) The sensor data is then processed using a 4D (or higher) model to determine the characteristic location-dependent pattern of brain activity over time associated with the state of interest. Where the data is derived from a population with various degrees of arousal, the model maintains this arousal state variable dimension.

(332) A recipient is then prepared for receipt of the mental state. The mental state of the recipient may be assessed. This can include responses to a questionnaire, self-assessment, or other psychological assessment method. Further, the transcutaneous EEG (or other brain activity data) of the recipient may be obtained, to determine the starting state for the recipient, as well as activity during experiencing the desired mental state.

(333) In addition, a set of stimuli, such as visual patterns, acoustic patterns, vestibular, smell, taste, touch (light touch, deep touch, proprioception, stretch, hot, cold, pain, pleasure, electric stimulation, acupuncture, etc.), vagus nerve (e.g., parasympathetic), are imposed on the subject, optionally over a range of baseline brain states, to acquire data defining the effect of individual and various combinations of sensory stimulation on the brain state of the recipient. Population data may also be used for this aspect.

(334) The data from the source or population of sources (see above) may then be processed in conjunction with the recipient or population of recipient data, to extract information defining the optimal sensory stimulation over time of the recipient to achieve the desired brain state resulting in the desired emotional or mental state.

(335) In general, for populations of sources and recipients, the data processing task is immense. However, the statistical analysis will generally be of a form that permits parallelization of mathematical transforms for processing the data, which can be efficiently implemented using various parallel processors, a common form of which is a SIMD (single instruction, multiple data) processor, found in typical graphics processors (GPUs). Because of the cost-efficiency of GPUs, it is referred to implement the analysis using efficient parallelizable algorithms, even if the computational complexity is nominally greater than a CISC-type processor implementation.

(336) During emotional arousal of the recipient, the EEG pattern may be monitored to determine if the desired state is achieved through the sensory stimulation. A closed loop feedback control system may be implemented to modify the stimulation seeking to achieve the target. An evolving genetic algorithm may be used to develop a user model, which relates the emotional or mental state, arousal and valence, sensory stimulation, and brain activity patterns, both to optimize the current session of stimulation and learning, as well as to facilitate future sessions, where the emotional or mental states of the recipient have further enhanced, and to permit use of the system for a range of emotional or mental states.

(337) The stimulus may comprise a chemical messenger or stimulus to alter the subject's level of consciousness or otherwise alter brain chemistry or

functioning. The chemical may comprise a hormone or endocrine analog molecule, (such as adrenocorticotrophic hormone (ACTH) (4-11)), a stimulant (such as cocaine, caffeine, nicotine, phenethylamines), a psychoactive drug, psychotropic or hallucinogenic substance (a chemical substance that alters brain function, resulting in temporary changes in perception, mood, consciousness and behavior such as pleasantness (e.g. euphoria) or advantageousness (e.g., increased alertness).

(338) While typically, controlled or “illegal” substances are to be avoided, in some cases, these may be appropriate for use. For example, various drugs may alter the state of the brain to enhance or selectively enhance the effect of the stimulation. Such drugs include stimulants (e.g., cocaine, methylphenidate (Ritalin), ephedrine, phenylpropanolamine, amphetamines), narcotics/opiates (opium, morphine, heroin, methadone, oxycodone, codeine, fentanyl), hallucinogens (lysergic acid diethylamide (LSD), PCP, MDMA (ecstasy), mescaline, psilocybin, magic mushroom (*Psilocybe cubensis*), *Amanita muscaria* mushroom, marijuana/*Cannabis*), *Salvia divinorum*, diphenhydramine (Benadryl), flexed, tobacco, nicotine, bupropion (Zyban), opiate antagonists, depressants, gamma aminobutyric acid (GABA) agonists or antagonists, NMDA receptor agonists or antagonists, depressants (e.g., alcohol, Xanax; Valium; Halcion; Librium; other benzodiazepines, Ativan; Klonopin; Amytal; Nembutal; Seconal; Phenobarbital, other barbiturates), psychedelics, disassociatives, and deliriants (e.g., a special class of acetylcholine-inhibitor hallucinogen). For example, Carhart-Hanis showed using fMRI that LSD and psilocybin caused synchronization of different parts of the brain that normally work separately by making neurons fire simultaneously. This effect can be used to induce synchronization of various regions of the brain to heighten the emotional state.

(339) It is noted that a large number of substances, natural and artificial, can alter mood or arousal and, as a result, may impact emotions or non-target mental states. Typically, such substances will cross the blood-brain barrier, and exert a psychotropic effect. Often, however, this may not be necessary or appropriate. For example, a painful stimulus can alter mood, without acting as a psychotropic drug; on the other hand, a narcotic can also alter mood by dulling emotions. Further, sensory stimulation can induce mood and/or emotional changes, such as smells, sights, sounds, various types of touch and proprioception sensation, balance and vestibular stimulation, etc. Therefore, peripherally acting substances that alter sensory perception or stimulation may be relevant to mood. Likewise, pharmacopsychotropic drugs may alter alertness, perceptiveness, memory, and attention, which may be relevant to task-specific mental state control.

(340) It is an object to provide a method for inducing an emotional state in a subject, comprising: determining a desired emotional state; selecting a profile from a plurality of profiles stored in a memory, the plurality of profiles each corresponding to a brain activity pattern of at least one exemplar subject under a respective emotional state (the “source”); and exposing a target subject (the “recipient”) to a stimulus modulated according to the selected profile, wherein the exposure, stimulus, and modulation are adapted to induce, in the target subject the desired emotional state.

(341) The brain activity pattern may be an electroencephalographic brainwave pattern, a magnetoencephalographic brainwave pattern, an electrical brainwave pattern, or a metabolic rate pattern, for example.

(342) The stimulus comprises may visual stimulus, an auditory stimulus; an olfactory stimulus; a tactile stimulus; a proprioceptive stimulus; an electrical stimulus; or a magnetic stimulus.

(343) The desired emotional state is may be happiness, joy, gladness, cheerfulness, bliss, delight, ecstasy, optimism, exuberance, merriment, joviality; vivaciousness, pleasure, excitement, sexual arousal, relaxation, harmony, or peace, for example.

(344) The exemplar subject and the target subject may be the same human at different times, or different humans, or different species.

(345) The stimulus may comprise an auditory stimulus adapted to induce binaural beats.

(346) The stimulus may comprise a dynamically changing electromagnetic field adapted synchronize brainwave patterns corresponding to the brain activity pattern of at least one exemplar subject under the desired emotional state.

(347) The selected profile may be derived from measurements of brainwave patterns in the exemplar subject selectively acquired during the desired emotional state.

(348) The selected profile may comprise a model derived from at least spatial, frequency and phase analysis of the measured brainwave patterns.

(349) The stimulus may comprise an auditory or visual stimulus frequency corresponding to a frequency pattern in a brainwave pattern of the exemplar subject.

(350) The target subject may be concurrently exposed to the stimulus and a primary audio or visual presentation which does not induce the desired emotional state, wherein the stimulus does not substantially interfere with the target subject appreciation of the audio or visual presentation.

(351) The method may further comprise recording EEG signals of the exemplar subject in the desired emotional state; decoding at least one of a temporal and a spatial pattern from the recorded EEG signals; and storing the decoded at least one of temporal and spatial pattern in a non-volatile memory.

(352) The method may further comprise selectively modifying the pattern based on differences between the exemplar subject and the target subject.

(353) The stimulus may comprise applying a spatial electrical stimulation pattern to the target subject via transcranial electrical stimulation (tES) to induce the desired emotional state. The spatial electrical stimulation pattern comprises a direct current or an alternating current. The transcranial electrical stimulation (tES) may be at least one of a transcranial direct current stimulation (tDCS), a transcranial alternating current stimulation (tACS), a transcranial pulsed current stimulation (tPCS) transcranial pulsed current stimulation (tPCS), and a transcranial random noise stimulation (tRNS).

(354) The brain activity pattern of the at least one exemplar subject may comprise a magnetoencephalogram (MEG), and the stimulus comprises applying a spatial magnetic stimulation pattern to the target subject via transcranial magnetic stimulation (tMS) to induce the desired emotional state.

(355) The stimulus may achieve brain entrainment in the target subject.

(356) The method may further comprise determining a second desired emotional state; selecting a second profile from the plurality of profiles stored in a memory; and exposing a target subject to a stimulus modulated according to the selected second profile, wherein the exposure, stimulus, and modulation are adapted to induce, in the target subject the desired second emotional state, the second emotional state being different from the first subsequent state and being induced in succession after the emotional state.

(357) It is another object to provide a method of brainwave entrainment comprising the steps of recording EEG of the brainwaves of a first subject in an emotional state; decoding at least one of a temporal and a spatial pattern from the EEG; storing a representation of the pattern in a non-volatile memory; retrieving said pattern from the non-volatile memory modulating the temporal and spatial patterns on a stimulus signal; and applying the stimulus signal to a second subject. The stimulus signal may be an alternating current, and said applying comprises applying the alternating current to the second subject via transcranial alternating current stimulation (tACS) to induce the emotional state.

(358) It is a further object to provide a method of brainwave entrainment comprising the steps of recording EEG of the brainwaves of a first subject in a respective emotional state; decoding at least one of temporal and spatial pattern from the recorded EEG; storing said at least one of temporal and spatial pattern in a non-volatile memory; retrieving said at least one of temporal and spatial pattern from the non-volatile memory; modulating the temporal and spatial patterns on a light signal; and projecting the light signal to the second subject to induce the respective emotional state. The light signal may be selected from the group consisting of an ambient light signal, a directional light signal, a laser beam signal, a visible spectrum light signal and an infrared light signal.

(359) It is another object to provide a method of brainwave entrainment comprising the steps of recording EEG of the brainwaves of a first subject in an emotional state; decoding at least one of a temporal and a spatial pattern from the EEG; storing said at least one of the temporal and the spatial pattern in a non-volatile memory; retrieving the at least one of the temporal and the spatial pattern from the non-volatile memory; modulating the temporal and spatial patterns on an isotonic sound signal; and projecting the isotonic sound signal to a second subject to induce the emotional state.

(360) A still further object provides a method of brainwave entrainment comprising the steps of: recording EEG of the brainwaves of a first subject in

an emotional state; decoding temporal frequency pattern from the EEG; storing the decoded temporal frequency pattern in a memory; retrieving the temporal frequency pattern from the memory; computing a first set of frequencies by adding a predetermined delta to the frequencies of the temporal frequency pattern; computing a second set of frequencies by subtracted the delta from the frequencies of the temporal frequency pattern; modulating the first set of frequencies on a first acoustical signal; modulating the second set of frequencies on a second acoustical signal; projecting the first set of frequencies into a first ear of the second subject and projecting the second set of frequencies into a second ear of the second subject, thereby producing binaural stimulation to induce the emotional state.

(361) Another object provides a method for modifying an emotional state or mood in a subject, comprising: selecting an emotional state or mood profile from a memory, corresponding to a brain activity pattern of at least one exemplar subject in a respective emotional state or mood; and exposing a target subject to a stimulus signal modulated according to the selected emotional state or mood profile, to induce, in the target subject the selected emotional state or mood. The brain activity pattern may be acquired through at least one of an electroencephalogram (EEG) and a magnetoencephalogram (MEG). The stimulus signal may be selected from the group consisting of a light, a sound, a touch, a smell, an electric current, and a magnetic field. The emotional state or mood may be selected from the group consisting of a state of happiness, a state of joy, a state of gladness, a state of cheerfulness, a state of bliss, a state of delight, a state of ecstasy, a state of optimism, a state of exuberance, a state of merriment, a jovial state, a state of vivaciousness, a state of pleasure, a state of excitement, a state of relaxation, a state of harmony, and a state of peace. The exemplar subject and the target subject may be the same subject at different times or different subjects.

(362) A further object provides a method of brainwave entrainment comprising the steps of: recording EEG of a first subject in a positive emotional state; storing a spatial-temporal pattern corresponding to the EEG in a memory; modulating a stimulus pattern according to the spatial-temporal pattern; and stimulating a second subject with the modulated stimulus pattern, to induce the positive emotional state. The modulated stimulus pattern may comprise a binaural audio stimulus. The modulated stimulus pattern may comprise a transcranial electrical stimulation, e.g., a direct current stimulus, an alternating current stimulus, a transcranial direct current stimulation (tDCS), a transcranial alternating current stimulation (tACS), a transcranial pulsed current stimulation (tPCS) transcranial pulsed current stimulation (tPCS), or a transcranial random noise stimulation (tRNS).

(363) It is a still further object to provide a method of brainwave entrainment comprising the steps of: modulating a predefined temporal and spatial pattern on a magnetic field; and applying the modulated magnetic field to the brain of a subject via transcranial magnetic stimulation (TMS) to selectively induce an emotional state corresponding to the predefined temporal and spatial pattern.

(364) It is an object to provide a system and method for enhancing emotional response to a stimulus in a subject.

(365) It is another object to provide a system and method for enhancing the experience virtual reality by enhancing the emotional response in a subject.

(366) It is a further object to provide a system and method for enhancing cinematographic experience by enhancing the emotional response in viewers while watching a movie.

(367) It is yet another object to provide a system and method for improving users' interaction with a computer.

(368) It is still another object to provide a system and method for improving users' interaction with a robot.

(369) It is a further object to provide a system and method for accelerating memory-retention and recall by inducing a desired emotional state in a subject.

(370) It is yet another object to provide a system and method for treatment of patients with dementia.

(371) It is an object to provide a system and method for facilitating an emotional state achievement process, comprising: determining a neuronal activity pattern, of a subject while engaged in a respective emotion; processing the determined neuronal activity pattern with at least one automated processor; and subjecting a subject seeking to achieve the respective emotion to a stimulus selected from the group consisting of one or more of a sensory excitation, a peripheral excitation, a transcranial excitation, and a deep brain stimulation, dependent on the processed electromagnetic determined neuronal activity pattern.

(372) It is yet another object to provide a system and method for facilitating a mental process, comprising: determining a neuronal activity pattern of a skilled subject having the mental process; processing the determined neuronal activity pattern with at least one automated processor; and subjecting a subject seeking a corresponding mental process to a stimulus selected from the group consisting of one or more of a sensory excitation, a peripheral excitation, a transcranial excitation, and a deep brain stimulation, dependent on the processed electromagnetic determined neuronal activity pattern.

(373) It is still another object to provide a system and method for improving achieving a mental state, comprising: determining a neuronal activity pattern, of a subject while having the mental state; processing the determined neuronal activity pattern with at least one automated processor; and subjecting a subject seeking to achieve the mental state to a stimulus selected from the group consisting of one or more of a sensory excitation, a peripheral excitation, a transcranial excitation, and a deep brain stimulation, dependent on the processed electromagnetic determined neuronal activity pattern. The mental state is, e.g., an emotional state, a mood, or other subjective state.

(374) It is also an object to provide an apparatus for facilitating control over an emotional state, comprising: an input, configured to receive data representing a neuronal activity pattern of a subject while having an emotional state; at least one automated processor, configured to process the determined neuronal activity pattern, to determine neuronal activity patterns selectively associated with the emotional state, configured to subject a subject emotional arousal in control over the emotional state to a stimulus selected from the group consisting of one or more of a sensory excitation, a peripheral excitation, a transcranial excitation, and a deep brain stimulation, dependent on the processed determined neuronal activity pattern.

(375) It is further an object to provide an apparatus for facilitating an emotional skill or emotional learning process, comprising: an input, configured to receive data representing a neuronal activity pattern of a subject while engaged in an emotional skill or emotional learning process; at least one automated processor, configured to process the determined neuronal activity pattern, to determine neuronal activity patterns selectively associated with successful learning of the emotional skill or emotional learning process; and a stimulator, configured to subject a subject emotional arousal in the respective emotional skill or emotional learning process to a stimulus selected from the group consisting of one or more of a sensory excitation, a peripheral excitation, a transcranial excitation, and a deep brain stimulation, dependent on the processed determined neuronal activity pattern.

(376) It is also an object to provide an apparatus for inducing of a desired emotional state, comprising: an input, configured to receive data representing a neuronal activity pattern of a skilled subject while experiencing the desired emotional state; at least one automated processor, configured to process the determined neuronal activity pattern, to determine neuronal activity patterns selectively associated with the desired emotional state; and a stimulator, configured to subject a recipient desiring to attain the same emotional state to a stimulus selected from the group consisting of one or more of a sensory excitation, a peripheral excitation, a transcranial excitation, and a deep brain stimulation, dependent on the processed determined neuronal activity pattern.

(377) It is a further object to provide a system for influencing a brain electrical activity pattern of a subject during emotional arousal, comprising: an input, configured to determine a target brain activity state for the subject, dependent on the emotional state; at least one processor, configured to generate a stimulation pattern profile adapted to achieve the target brain activity state for the subject, dependent on the emotional state; and a stimulator, configured to output at least one stimulus, proximate to the subject, dependent on the generated stimulation pattern profile.

(378) It is yet a further object to provide a system for influencing a brain electrical activity pattern of a subject during experiencing information, comprising: an input, configured to determine a target brain activity state for the subject, dependent on the nature of the respective information; at least one processor, configured to generate a stimulation pattern profile adapted to achieve the target brain activity state for the subject, dependent on the emotion; and a stimulator, configured to output at least one stimulus, proximate to the subject, dependent on the generated stimulation pattern

profile.

(379) It is still a further object to provide a system for influencing a brain electrical activity pattern of a subject during a state of emotional arousal, comprising: an input, configured to determine a target brain emotional state for the subject, dependent on the desired emotional state; at least one processor, configured to generate a stimulation pattern profile adapted to achieve the target brain emotional state for the subject, dependent on the emotional state; and a stimulator, configured to output at least one stimulus, proximate to the subject, dependent on the generated stimulation pattern profile.

(380) It is a still further object to provide a system for determining a target brain activity state for a subject, dependent on an emotion state, comprising: a first monitor, configured to acquire a brain activity of a first subject during the emotion state; at least one first processor, configured to analyze a spatial brain activity state over time of the first subject; and determine spatial brain activity states of the first subject, which represent readiness for emotion state; a second monitor, configured to acquire a brain activity of a second subject during performance of a variety of activities, under a variety of stimuli; and at least one second processor, configured to: analyze a spatial brain activity state over time of the second subject and translate the determined spatial brain activity states of the first subject which represent readiness for the emotion state, into a stimulus pattern for the second subject to achieve a spatial brain activity state in the second subject corresponding to emotion state.

(381) It is a still further object to provide a system for determining a target brain activity state for a subject, dependent on an emotion or mood, comprising: a first monitor, configured to acquire a brain activity of a first subject during experiencing the emotion or mood; at least one first processor, configured to analyze a spatial brain activity state over time of the first subject and determine spatial brain activity states of the first subject, which represent the emotion or mood; a second monitor, configured to acquire a brain activity of a second subject during the emotion or mood, under a variety of stimuli; and at least one second processor, configured to: analyze a spatial brain activity state over time of the second subject and translate the determined spatial brain activity states of the first subject which represent the emotion or mood, into a stimulus pattern for the second subject to achieve a spatial brain activity state in the second subject corresponding to the emotion or mood.

(382) It is a further object to provide a method of enhancing an emotional state of a first subject the method comprising: recording a second subject's brainwaves EEG while at rest having the second subject experience or enact an emotionally charged experience to induce an emotional state or mood; recording the second subject's brainwaves EEG while experiencing or enacting said emotionally charged experience; extracting a predominant temporal pattern associated with said emotional state from the recorded brainwaves by comparing them with the brainwaves at rest encoding said temporal pattern as a digital code stored in a tangible media; and using said digital code to modulate the temporal pattern on a signal perceptible to the first subject while said first subject is trying to attain the said emotional state, whereby said perceptible signal stimulates in the second subject brainwaves having said temporal pattern to induce the emotional state or mood.

(383) It is still a further object to provide a method of enhancing an emotional state of a first person, the method comprising: recording a second person's brainwaves or EEG while at rest or prior to achieving a desired emotional state; subjecting having the second person to the performance; recording the second person's brainwaves or EEG while subject to the performance; extracting a predominant temporal pattern associated with said performance from the recorded brainwaves or EEG by comparing them with the brainwaves or EEG at rest or prior to achieving the desired emotional state; encoding said temporal pattern as a digital code stored in a tangible media; and using said digital code to modulate the temporal pattern on a signal perceptible to the first person while said first person is seeking to achieve said desired emotional state, whereby said light signal stimulates in the first subject brainwaves or EEG having said temporal pattern to enhance the achievement of the desired emotional state.

(384) A still further object provides a method of assisted appreciation of art by a first subject, the method comprising: recording a second subject's brainwaves EEG while at rest, wherein the second subject is knowledgeable in the art; having the second subject experience the art; recording the second subject's brainwaves (e.g., EEG, or MEG) while experiencing the art; extracting a predominant temporal pattern associated with appreciating the art from the recorded brainwaves by comparing them with the brainwaves at rest encoding said temporal pattern as a digital code stored in a tangible media; and using said digital code to modulate the temporal pattern on a signal perceptible to the first subject while the first subject is seeking to appreciate the art, whereby said signal stimulates in the first subject brainwaves having said temporal pattern.

(385) It is another object to provide a computer readable medium, storing therein non-transitory instructions for a programmable processor to perform a process, comprising the computer-implemented steps: synchronizing brain activity data of a subject with at least one event involving the subject analyzing the brain activity data to determine a selective change in the brain activity data corresponding to an emotional correlate of the event and determine a stimulation pattern adapted to induce a brain activity having a correspondence to the brain activity data associated with the emotion, based on at least a brain activity model.

(386) The at least one of a sensory excitation, peripheral excitation, and transcranial excitation may be generated based on a digital code. The subjecting of the subject having the emotion or mood to the sensory excitation increases a rate of achieving the emotion in the target subject. Similarly, the subjecting of the subject seeking to achieve the emotion or mood to the sensory excitation increases a rate of achieving the emotion or mood in the target. Likewise, the subjecting of the subject seeking to achieve the respective emotional state to the sensory excitation improves the quality or intensity of the emotional state in the subject.

(387) The method may further comprise determining a neuronal baseline activity of the skilled subject while not engaged in the emotion, a neuronal baseline activity of the subject, a neuronal activity of the skilled subject while engaged in the emotion, and/or a neuronal activity of the subject while engaged in the emotion.

(388) The representation of the processed the determined neuronal activity pattern may be stored in memory. The storage could be on a tangible medium as an analog or digital representation. It is possible to store the representation in a data storage and access system either for a permanent backup or further processing the respective representation. The storage can also be in a cloud storage and/or processing system.

(389) The neuronal activity pattern may be obtained by electroencephalography, magnetoencephalography, MRI, fMRI, PET, low-resolution brain electromagnetic tomography, or other electrical or non-electrical means.

(390) The neuronal activity pattern may be obtained by at least one implanted central nervous system (cerebral, spinal) or peripheral nervous system electrode. An implanted neuronal electrode can be either within the peripheral nervous system or the central nervous system. The recording device could be portable or stationary. Either with or without onboard electronics such as signal transmitters and/or amplifiers, etc. The at least one implanted electrode can consist of a microelectrode array featuring more than one recording site. Its main purpose can be for stimulation and/or recording.

(391) The neuronal activity pattern may be obtained by at least a galvanic skin response. Galvanic skin response or resistance is often also referred as electrodermal activity (EDA), psychogalvanic reflex (PGR), skin conductance response (SCR), sympathetic skin response (SSR) and skin conductance level (SCL) and is the property of the human body that causes continuous variation in the electrical characteristics of the skin.

(392) The stimulus may comprise a sensory excitation. The sensory excitation may by either sensible or insensible. It may be either peripheral or transcranial. It may consist of at least one of a visual, an auditory, a tactile, a proprioceptive, a somatosensory, a cranial nerve, a gustatory, an olfactory, a pain, a compression and a thermal stimulus or a combination of the aforesaid. It can, for example, consist of light flashes either within ambient light or aimed at the subject's eyes, 2D or 3D picture noise, modulation of intensity, within the focus of the subjects eye the visual field or within peripheral sight Within a video presentation, intensity variations may be provided around a periphery of the presentation, globally throughout a presentation (i.e., modulating a backlight or display intensity), or programmed to modulate a brightness of individual objects.

(393) The stimulus may comprise a peripheral excitation, a transcranial excitation, a sensible stimulation of a sensory input, an insensible stimulation of a sensory input, a visual stimulus, an auditory stimulus, a tactile stimulus, a proprioceptive stimulus, a somatosensory stimulus, a cranial nerve stimulus, a gustatory stimulus, an olfactory stimulus, a pain stimulus, an electric stimulus, a magnetic stimulus, or a thermal stimulus.

(394) The stimulus may comprise transcranial magnetic stimulation (TMS), cranial electrotherapy stimulation (CES), transcranial direct current stimulation (tDCS), comprise transcranial alternating current stimulation (tACS), transcranial random noise stimulation (tRNS), comprise transcranial pulsed current stimulation (tPCS), pulsed electromagnetic field, or noninvasive or invasive deep brain stimulation (DBS), for example. The stimulus may comprise transcranial pulsed ultrasound (TPU). The stimulus may comprise a cochlea implant stimulus, spinal cord stimulation (SCS) or a vagus nerve stimulation (VNS), or other direct or indirect cranial or peripheral nerve stimulus. The stimulus may comprise or achieve brainwave entrainment. The stimulus may comprise electrical stimulation of the retina, a pacemaker, a stimulation microelectrode array, electrical brain stimulation (EBS), focal brain stimulation (FBS), light, sound, vibrations, an electromagnetic wave. The light stimulus may be emitted by at least one of a light bulb, a light emitting diode (LED), and a laser. The signal may be one of a ray of light, a sound wave, and an electromagnetic wave. The signal may be a light signal projected onto the first subject by one of a smart bulb generating ambient light, at least one LED position near the eyes of the first subject and laser generating low-intensity pulses.

(395) The mental state may be associated with learning or performing a skill. The skill may comprise a mental skill, e.g., cognitive, alertness, concentration, attention, focusing, memorization, visualization, relaxation, meditation, speedreading, creative skill, “whole-brain-thinking”, analytical, reasoning, problem-solving, critical thinking, intuitive, leadership, learning, speedreading, patience, balancing, perception, linguistic or language, language comprehension, quantitative, “fluid intelligence”, pain management, skill of maintaining positive attitude, a foreign language, musical, musical composition, writing, poetry composition, mathematical, science, art, visual art, rhetorical, emotional control, empathy, compassion, motivational skill, people, computational, science skill, or an inventorship skill. See, U.S. Pat. Nos. 6,435,878, 5,911,581, and 20090069707. The skill may comprise a motor skill, e.g., fine motor, muscular coordination, walking, running, jumping, swimming, dancing, gymnastics, yoga; an athletic or sports, massage skill, martial arts or fighting, shooting, self-defense; speech, singing, playing a musical instrument, penmanship, calligraphy, drawing, painting, visual, auditory, olfactory, game-playing, gambling, sculptor's, craftsman, massage, or assembly skill. Where a skill is to be enhanced, and an emotion to be achieved (or suppressed), concurrently, the stimulus to the recipient may be combined in such a way as to achieve the result. In some cases, the component is universal, while in others, it is subjective. Therefore, the combination may require adaptation based on the recipient characteristics.

(396) The technology may be embodied in apparatuses for acquiring the brain activity information from the source, processing the brain activity information to reveal a target brain activity state and a set of stimuli, which seek to achieve that state in a recipient, and generating stimuli for the recipient to achieve and maintain the target brain activity state over a period of time and potential state transitions. The generated stimuli may be feedback controlled. A general-purpose computer may be used for the processing of the information, a microprocessor, a FPGA, an ASIC, a system-on-a-chip, or a specialized system, which employs a customized configuration to efficiently achieve the information transformations required. Typically, the source and recipient act asynchronously, with the brain activity of the source recorded and later processed. However, real-time processing and brain activity transfer are also possible. In the case of a general purpose programmable processor implementation or portions of the technology, computer instructions may be stored on a nontransient computer readable medium. Typically, the system will have special-purpose components, such as a transcranial stimulator, or a modified audio and/or display system, and therefore the system will not be a general purpose system. Further, even in a general purpose system the operation per se is enhanced according to the present technology.

(397) It is another object to provide a method of teaching one of an emotion-dependent mental skill and a motor skill to a first subject, the method comprising: recording a second subject's brainwaves EEG while at rest having the second subject perform said one of a mental skill and a motor skill; recording the second subject's brainwaves while performing said one of a mental skill and a motor skill; extracting a predominant temporal pattern associated with said one of a mental skill and a motor skill from the recorded brainwaves by comparing them with the brainwaves at rest encoding said temporal pattern together with an emotional state targeting stimulus pattern, as a digital code stored in a tangible media; and using said digital code to modulate the temporal pattern on a signal perceptible to the first subject while the first subject is learning said one of a mental and a motor skill, whereby said light signal stimulates in the first subject brainwaves having said temporal pattern to accelerate learning of said one if a mental skill and a motor skill. The emotional state targeting stimulus pattern may be derived from the first subject, the second subject, or a one or more different subjects. The stimulation pattern may thus be modified from the second subject pattern to bias the first subject toward a desired emotional state.

(398) It is a further object to provide a high-definition transcranial alternating current stimulation (HD-tACS) stimulation of a target, having a stimulation frequency, amplitude pattern, spatial pattern, dependent on an existing set of states in the target, and a set of brainwave patterns from a target engaged in a mood, adapted to improve an emotional state or mood of the recipient

(399) It is yet another object to provide a system and method for facilitating a mental process, comprising: determining a neuronal activity pattern, of a subject while engaged in an emotional process; processing the determined neuronal activity pattern with at least one automated processor, and subjecting a subject targeting the emotional process to a stimulus selected from the group consisting of one or more of a sensory excitation, a peripheral excitation, a transcranial excitation, and a deep brain stimulation, dependent on the processed electromagnetic determined neuronal activity pattern while the subject is subjected to tES, a psychedelic and/or other pharmaceutical agents.

(400) It is a still further object to provide a method of facilitating a skill learning process, comprising: determining a neuronal activity pattern of a skilled subject while engaged in a respective skill; processing the determined neuronal activity pattern with at least one automated processor; modifying the determined neuronal activity pattern according to an emotional state neuronal activity pattern; and subjecting a subject training in the respective skill to a stimulus selected from the group consisting of one or more of a sensory excitation, a peripheral excitation, a transcranial excitation, and a deep brain stimulation, dependent on the modified processed determined neuronal activity pattern. The transcranial electric stimulation (tES) may be one of transcranial direct current stimulation (tDCS), transcranial alternative current stimulation (tACS), and high-definition transcranial alternative current stimulation (tES). The emotional state neuronal activity pattern may be a pattern that increases alertness and focus, for example.

(401) Another object provides a method of facilitating a skill learning process, comprising: determining a respective neuronal activity pattern of a skilled subject while engaged in a respective skill and having an emotional state appropriate for learning the skill and while engaged in the respective skill and not having the emotional state appropriate for learning the skill; processing the determined neuronal activity pattern with at least one automated processor; subjecting a subject training in the respective skill to one of a pharmaceutical agent and a psychedelic agent and subjecting a subject training in the respective skill to a stimulus selected from the group consisting of one or more of a sensory excitation, a peripheral excitation, a transcranial excitation, and a deep brain stimulation, dependent on the processed determined neuronal activity pattern while engaged in a respective skill and having an emotional state appropriate for learning the skill, and adapting the stimulus based on feedback based on a measurement of a neuronal activity pattern of the subject training in the respective skill to determine an emotional state of the subject training in the respective skill.

(402) It is another object to provide a method of inducing an emotional state in a target subject, comprising determining a desired emotional state; selecting a profile from a plurality of profiles stored in a memory, the plurality of profiles each corresponding to a brain activity pattern of a donor subject having a respective emotional state; and exposing the target subject to at least one stimulus modulated according to the selected profile representing and being adapted to induce, in the target subject, the desired emotional state. The brain activity pattern may be at least one of an electroencephalographic brainwave pattern and a magnetoencephalographic brainwave pattern. The at least one stimulus may stimulate a cranial nerve of the target subject. The at least one stimulus may comprise at least one of a visual stimulus, and an auditory stimulus, a two-channel auditory stimulus adapted to induce binaural beats, at least one of a tactile stimulus and a proprioceptive stimulus, an at least one of a direct electrical current and an alternating electrical current, and/or a magnetic field. The stimulus may comprise at least one of an auditory stimulus and a visual stimulus with a frequency corresponding to at least a frequency pattern in a brainwave pattern of the donor subject.

(403) The desired emotional state may be one of happiness, joy, gladness, cheerfulness, bliss, delight, ecstasy, optimism, exuberance, merriment, joviality; vivaciousness, pleasure, excitement, sexual arousal, relaxation, harmony, and peace.

(404) The target subject may be the same as or different from the donor subject. The target subject may be identical with the donor subject, wherein the brain activity pattern of the donor subject was recorded prior to the exposing the target subject to at least one stimulus.

(405) The at least one stimulus may comprise a dynamically changing electromagnetic field adapted to synchronize the target subject's brainwave pattern with a brainwave pattern of the donor subject having the desired emotional state.

(406) The selected profile may be derived from recording of brainwave patterns of the donor subject selectively acquired during the desired emotional state. The selected profile may comprise a model derived from at least one of a spatial, a frequency and a phase analysis of the recorded brainwave patterns.

(407) The method may further comprise recording EEG signals of the donor subject in the desired emotional state; decoding at least one of a temporal and a spatial pattern from the recorded EEG signals; and storing the decoded at least one of temporal and spatial pattern in a non-volatile memory as at least one profile.

(408) The method may comprise selectively modifying stimulus based on differences between the donor subject, from which the profile may be derived, and the target subject.

(409) The stimulus may comprise applying at least one of a temporal and a spatial electrical stimulation pattern to the target subject via transcranial electrical stimulation (TES) to induce the desired emotional state. The transcranial electrical stimulation (TES) may be at least one of a transcranial direct current stimulation (tDCS), an oscillating transcranial direct current stimulation (osc-tDCS), a transcranial alternating current stimulation (tACS), a transcranial pulsed current stimulation (tPCS), and a transcranial random noise stimulation (tRNS).

(410) The profile may be derived from brain activity pattern of the donor subject comprising a magnetoencephalogram (MEG), and the stimulus may comprise applying a spatial magnetic stimulation pattern to the target subject via transcranial magnetic stimulation (TMS) to induce the desired emotional state.

(411) The stimulus may achieve brain entrainment in the target subject.

(412) The method may further comprise determining a second desired emotional state; selecting a second profile from the plurality of profiles stored in a memory; and exposing the target subject to a stimulus modulated according to the selected second profile, representing and being adapted to induce, in the target subject, the second desired emotional state, the second emotional state being different from the emotional state and being induced in succession after the emotional state.

(413) At least one profile may correspond to consensus brain activity pattern of a plurality of donor subjects, each of the plurality of donor subjects having the respective emotional state

(414) It is a further object to provide a method of brainwave entrainment comprising: recording brainwaves of a first subject in a desired emotional state; decoding at least one of a temporal and a spatial pattern from the brainwaves; storing a representation of the pattern in a memory; retrieving said pattern from the memory; modulating the decoded at least one of the temporal and the spatial pattern on at least one stimulus signal; and applying said at least one stimulus signal to a second subject, to induce the second subject to assume the emotional state. The step of recording brainwaves comprise recording of at least one of electroencephalogram and magnetoencephalogram of the brainwaves. The stimulus signal may be at least one of a direct current and an alternating current, and said applying may comprise applying said at least one of a direct current and an alternating current to the second subject via respectively a transcranial direct current stimulation (tDCS) or transcranial alternating current stimulation (tACS) to induce the desired emotional state.

(415) It is a still further object to provide a method of brainwave entrainment comprising: recording the brainwaves of a first subject in a desired emotional state; decoding at least one of temporal and spatial pattern from the recorded brainwaves; storing said at least one of the temporal and spatial pattern in a memory; retrieving said at least one of the temporal and spatial pattern from the memory; modulating the at least one of the temporal and spatial pattern on at least one of a current, a magnetic field, a light signal, and an acoustic signal; and exposing the second subject to the at least one of the current, the magnetic field, the light signal, and the acoustic signal, to induce the desired emotional state.

(416) The step of recording the brainwaves may comprise recording of at least one of an electroencephalogram and a magnetoencephalogram of the brainwaves.

(417) Another object provides a method of recording a desired emotional state from a donor, comprising: determining an emotional state of the donor, if the donor may be in the desired emotional state, recording neural correlates of the emotional state of the donor, analyzing neural correlates of the desired emotional state of the donor to decode at least one of a temporal and a spatial pattern corresponding to the desired emotional state; converting said at least one of a temporal and a spatial pattern corresponding to the desired emotional state into a neurostimulation pattern; and storing the neurostimulation pattern in the nonvolatile memory. The neural correlates may be brainwaves of the donor.

(418) The step of analyzing neural correlates may comprise identifying principal components of the brainwaves. The identifying of principal components may comprise performing one of a principal component analysis (PCA), a curvilinear principal component analysis, an independent component analysis (ICA), a Karhunen-Loève transform (KLT), a singular value decomposition (SVD), and a Factor analysis. The step of analyzing neural correlates may comprise performing a frequency domain analysis.

(419) The step of performing the frequency analysis may comprise performing one of a Fourier Transform, a Laplace Transform, a Fourier-Stieltjes transform, a Gelfand transform, time-frequency analysis, a short-time Fourier transform, and a fractional Fourier transform.

(420) The desired emotional state may be one of of happiness, joy, gladness, cheerfulness, bliss, delight, ecstasy, optimism, exuberance, merriment, joviality; vivaciousness, pleasure, excitement, sexual arousal, relaxation, harmony, and peace.

(421) The method may further comprise retrieving the neurostimulation pattern from the nonvolatile memory; and stimulating the recipients brain with at least one stimulus modulated with the neurostimulation pattern to induce the desired emotional state in the recipient.

(422) The at least one stimulus may be one of a direct current, an alternating current, a magnetic field, a light, a sound, a tactile signal and an olfactory signal.

(423) The recipient may be the donor at a point in time subsequent to the time of recording the neural correlates of the emotional state of the donor.

(424) The method may further comprise determining an emotional state of the recipient to confirm that the recipient may be in the desired emotional state. The method may further comprise developing a brain model of the recipient and adjusting said at least one stimulus in accordance with the model to adjust for the differences between the recipients brain and the donor's brain. The method may further comprise the step of administering a pharmacological agent to the recipient to facilitate response of the recipient to the at least one stimulus to induce the desired emotional state. The method may further comprise performing, by the recipient, a physical exercise in conjunction with the at least one stimulus.

(425) It is another object of provide a relational database of neural correlates of emotional states, comprising a first table storing a plurality of respective emotional states, linked with a second table storing information associated with respective emotional states obtained by recording neural correlates of the respective emotional state of each of a plurality of donors while in the respective emotional state; decoding from the recorded neural correlates at least one of a temporal and a spatial pattern corresponding to the plurality of respective emotional states; and storing information selectively derived from the at least one of the temporal and the spatial pattern corresponding to the plurality of respective emotional states in the second table. The neural correlates of each respective emotional state may be brainwaves. The recording of neural correlates may be done by using one of an electroencephalogram and a magnetoencephalogram. The relational database may be accessible by receipt of a respective emotional state and responsive by providing information linked to the respective emotional state.

(426) Another object provides a method of increasing emotional emersion in a presentation, comprising: defining a target emotional state associated

with at least a portion of the presentation; retrieving a record from a database associated with the target emotional state, derived from recorded neural correlates of donors engaged in the target emotional state; defining a neurostimulation pattern based on the record retrieved from the database; and subjecting a recipient to the defined neurostimulation pattern concurrent with being presented with the at least a portion of the presentation.

(427) The defining a target emotional state associated with at least a portion of the presentation may comprise defining a series of emotional states synchronized with activity or objects depicted in the presentation. The retrieving of the record from the database associated with the target emotional state may comprise retrieving a plurality of records corresponding to the series of emotional states. The defining of the neurostimulation pattern may comprise defining a series of neurostimulation patterns based on the retrieved plurality of records. The subjecting the recipient to the defined neurostimulation pattern concurrent with being presented with the at least a portion of the presentation may comprise subjecting the recipient to the defined series of neurostimulation patterns, temporally synchronized with the portions of presentation, in an order defined by the presentation.

(428) The target emotional state may be defined by an author of the presentation, or automatically derived from the presentation.

(429) The database may be a relational database, having a first table of respective emotional states, and a second table of information relating to neural correlates of the respective emotional states, the first table and the second table being linked together and searchable based on respective emotional state.

(430) At least one record of the database may be derived from recorded neural correlates of a plurality of different donors engaged in a common respective target emotional state. The at least one record may comprise a consensus of the plurality of different donors. The at least one record may comprise a plurality of sub-records, each sub-record being derived from a distinct subpopulation of the plurality of different donors, further comprising determining a characteristic of the recipient, and selecting a respective sub-record from the record based on the determined characteristic.

(431) The neurostimulation pattern may be at least one of an electrical current, a magnetic field, a light signal, and an acoustic signal. The neurostimulation pattern may be encoded in the record and/or may be defined by at least one automated processor after retrieving the record, and in selective dependence on at least one characteristic of the recipient. The presentation may comprise an audiovisual presentation, e.g., a virtual reality presentation. The defined neurostimulation pattern may be encoded as at least one of an audio and a visual stimulus within the audiovisual presentation. The defined neurostimulation pattern may be encoded as the at least one of the audio and the visual stimulus within the audiovisual presentation dependent on at least one characteristic of the recipient. The defined neurostimulation pattern may be dependent on automatically generated or manual feedback from the recipient.

(432) Another object provides a system for increasing emotional response to a presentation, comprising: a database comprising a record associated with a target emotional state, the record being derived from recorded neural correlates of at least one donor engaged in the respective target emotional state; at least one input configured to receive an association of the target emotional state with a portion of a presentation; at least one automated processor configured to define a neurostimulation pattern based on the record retrieved from the database; and a neurostimulator, configured to emit the defined neurostimulation pattern concurrent with presentation of the portion of the presentation.

(433) The input may be configured to receive a series of associations of respective target emotional states with respective portions of the presentation, and the neurostimulator may be configured to emit a series of the defined neurostimulation patterns synchronized with the received series of associations of the respective target emotional states with the respective portions of the presentation. The database may be a relational database, having a first table of respective emotional states, and a second table of information relating to neural correlates of the respective emotional states, the first table and the second table being linked together and searchable based on respective emotional state. At least one record may be derived from recorded neural correlates of a plurality of different donors engaged in a common respective target emotional state. The at least one record may comprise a consensus of the plurality of different donors. The at least one record may comprise a plurality of sub-records, each sub-record being derived from a distinct subpopulation of the plurality of different donors, a respective sub-record being selectable from the record based on the determined characteristic. The neurostimulator may be at least one of an electrical current stimulator, a magnetic field stimulator, a light signal stimulator, and an acoustic signal stimulator. The neurostimulation pattern may be encoded in the record, and/or may be defined by the at least one automated processor dependent on the record, and in selective dependence on at least one characteristic of the recipient. The presentation may comprise an audiovisual presentation, e.g., a virtual reality presentation, and optionally the defined neurostimulation pattern may be encoded as at least one of an audio and a visual stimulus within the audiovisual presentation. The defined neurostimulation pattern may be encoded as the at least one of the audio and the visual stimulus within the audiovisual presentation dependent on at least one characteristic of the recipient. The defined neurostimulation pattern may be dependent on automatically or manually generated feedback from the recipient.

(434) Other objects will become apparent from a review of disclosure hereof.

Description

BRIEF DESCRIPTION OF THE DRAWINGS

- (1) The detailed description is described with reference to the accompanying figures. In the figures, the left-most digit(s) of a reference number identifies the figure in which the reference number first appears. The use of the same reference number in different figures indicates similar or identical items.
- (2) FIG. 1 shows the electric activity of a neuron contributing to a brainwave.
- (3) FIG. 2 shows transmission of an electrical signal generated by a neuron through the skull, skin and other tissue to be detectable by an electrode transmitting this signal to EEG amplifier.
- (4) FIG. 3 shows an illustration of a typical EEG setup with a subject wearing a cap with electrodes connected to the EEG machine, which is, in turn, connected to a computer screen displaying the EEG.
- (5) FIG. 4 shows a typical EEG reading.
- (6) FIG. 5 shows one second of a typical EEG signal.
- (7) FIG. 6 shows main brainwave patterns.
- (8) FIGS. 7-13 shows a flowchart according to embodiments of the invention.
- (9) FIG. 14 shows a schematic representation of an apparatus according to one embodiment of the invention.
- (10) FIG. 15 shows brainwave real-time BOLD (Blood Oxygen Level Dependent) fMRI studies acquired with synchronized stimuli.
- (11) FIG. 16 shows Brain Entrainment Frequency Following Response (or FFR).
- (12) FIG. 17 shows brainwave entrainment before and after synchronization.
- (13) FIG. 18 shows brainwaves during inefficient problem solving and stress.
- (14) FIGS. 19 and 20 show how binaural beats work.
- (15) FIG. 21 shows Functional Magnetic Resonance Imaging (Mental states may be induced in a subject)
- (16) FIG. 22 shows a photo of a brain forming a new idea.
- (17) FIG. 23 shows 3D T2 CUBE (SPACENISTA) FLAIR & DSI tractography
- (18) FIG. 24 shows an EEG tracing.
- (19) FIGS. 25-29 show flowcharts according to embodiments of the invention.
- (20) FIG. 30 shows human brain anatomy.
- (21) FIG. 31 shows a brain map.
- (22) FIG. 32 shows an image depicting neuron anatomy.

- (23) FIG. 33 shows graphs representing a dimensional view of emotions.
(24) FIG. 34 shows a representation of neural activity with respect to emotional state.
(25) FIGS. 35-41 show flowcharts according to embodiments of the invention.
(26) FIG. 42 shows graphs of tDCS, tRNS, and tACS stimulation patterns.
(27) FIGS. 43 and 44 show representations of tDCS neural stimulation.
(28) FIG. 45 shows a representation of tACS or tRNS neural stimulation.
(29) FIG. 46 shows a representation of intracranial electrode implantation.
(30) FIG. 47 shows a representation of tDCS electrode location.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

- (31) Hereinafter, embodiments of the present disclosure will be described in detail with reference to the accompanying drawings so that the present disclosure may be readily implemented by those skilled in the art. However, it is to be noted that the present disclosure is not limited to the embodiments but can be embodied in various other ways. In drawings, parts irrelevant to the description are omitted for the simplicity of explanation, and like reference numerals denote like parts through the whole document.
- (32) Through the whole document, the term “connected to” or “coupled to” that is used to designate a connection or coupling of one element to another element includes both a case that an element is “directly connected or coupled to” another element and a case that an element is “electronically connected or coupled to” another element via still another element. Further, it is to be understood that the term “comprises or includes” and/or “comprising or including” used in the document means that one or more other components, steps, operation and/or existence or addition of elements are not excluded in addition to the described components, steps, operation and/or elements unless context dictates otherwise.
- (33) Through the whole document, the term “unit” or “module” includes a unit implemented by hardware or software and a unit implemented by both of them. One unit may be implemented by two or more pieces of hardware, and two or more units may be implemented by one piece of hardware.
- (34) Other devices, apparatus, systems, methods, features and advantages of the invention will be or will become apparent to one with skill in the art upon examination of the following figures and detailed description. It is intended that all such additional systems, methods, features and advantages be included within this description, be within the scope of the invention, and be protected by the accompanying claims.
- (35) The present invention generally relates to enhancing emotional response by a subject in connection with the received information by conveying to the brain of the subject temporal patterns of brainwaves of a second subject who had experienced such emotional response, said temporal pattern being provided non-invasively via light, sound, transcranial direct current stimulation (tDCS), transcranial alternating current stimulation (tDAS) or HD-tACS, transcranial magnetic stimulation (TMS) or other means capable of conveying frequency patterns.
- (36) The transmission of the brainwaves can be accomplished through direct electrical contact with the electrodes implanted in the brain or remotely employing light, sound, electromagnetic waves and other non-invasive techniques. Light, sound, or electromagnetic fields may be used to remotely convey the temporal pattern of prerecorded brainwaves to a subject by modulating the encoded temporal frequency on the light, sound or electromagnetic field signal to which the subject is exposed.
- (37) Every activity, mental or motor, and emotion is associated with unique brainwaves having specific spatial and temporal patterns, i.e., a characteristic frequency or a characteristic distribution of frequencies over time and space. Such waves can be read and recorded by several known techniques, including electroencephalography (EEG), magnetoencephalography (MEG), exact low-resolution brain electromagnetic tomography (eLORETA), sensory evoked potentials (SEP), fMRI, functional near-infrared spectroscopy (fNIRS), etc. The cerebral cortex is composed of neurons that are interconnected in networks. Cortical neurons constantly send and receive nerve impulses-electrical activity-even during sleep. The electrical or magnetic activity measured by an EEG or MEG (or another device) device reflects the intrinsic activity of neurons in the cerebral cortex and the information sent to it by subcortical structures and the sense receptors.
- (38) An EEG electrode mainly detects the neuronal activity in the brain region just beneath it. However, the electrodes receive the activity from thousands of neurons. One square millimeter of cortex surface, for example, has more than 100,000 neurons. It is only when the input to a region is synchronized with electrical activity occurring at the same time that simple periodic waveforms in the EEG become distinguishable.
- (39) The spatial and temporal pattern associated with specific brainwaves can be digitized and encoded in software code. It has been observed that “playing back the brainwaves” to another animal or person by providing decoded temporal pattern through transcranial direct current stimulation (tDCS), transcranial alternating current stimulation (tACS), high definition transcranial alternating current stimulation (HD-tDCS), transcranial magnetic stimulation (TMS), or through electrodes implanted in the brain allows the recipient to achieve the emotional or mental state at hand or to increase a speed of achievement. For example, if the brainwaves of a mouse navigated a familiar maze are decoded (by EEG or via implanted electrodes), playing this temporal pattern to another mouse unfamiliar with this maze will allow it to learn to navigate this maze faster.
- (40) Similarly, recording brainwaves associated with a specific emotional or mental response of one subject and later “playing back” this response to another subject will induce a similar emotional or mental response in the second subject. More generally, when one animal assumes an emotional or mental state, parts of the brain will have characteristic activity patterns. Further, by “artificially” inducing the same pattern in another animal, the other animal will have the same emotional or mental state, or more easily be induced into that state. The pattern of interest may reside deep in the brain, and thus be overwhelmed in an EEG signal by cortical potentials and patterns. However, techniques other than surface electrode EEG may be used to determine and spatially discriminate deep brain activity, e.g., from the limbic system. For example, various types of magnetic sensors may sense deep brain activity. See, e.g., 9,618,591; 9,261,573; 8,618,799; and 8,593,141.
- (41) In some cases, EEGs dominated by cortical excitation patterns may be employed to sense the emotional or mental state, since the cortical patterns may correlate with lower-level brain activity. Note that the determination of a state representation of an emotional or mental need not be performed each time the system is used; rather, once the brain spatial and temporal activity patterns and synchronization states associated with a particular emotional or mental states are determined, those patterns may be used for multiple targets and over time.
- (42) Similarly, while the goal is, for example, to trigger the target to assume the same brain activity patterns as the exemplar, this can be achieved in various ways, and these methods of inducing the desired patterns need not be invasive. Further, user feedback, especially in the case of a human emotional or mental state transferee, may be used to tune the process. Finally, using the various senses, especially sight, sound, vestibular, touch, proprioception, taste, smell, vagus afferent other cranial nerve afferent, etc. can be used to trigger high level mental activity, that in a particular subject achieves the desired mental state, emotion or mood.
- (43) Thus, in an experimental subject, which may include laboratory scale and/or invasive monitoring, a set of brain electrical activity patterns that correspond to particular emotions or emotional or mental states is determined. Preferably, these are also correlated with surface EEG findings. For the transferee, a stimulation system is provided that is non-hazardous and non-invasive. For example, audiovisual stimulation may be exclusively used. A set of EEG electrodes is provided to measure brain activity, and an adaptive or genetic algorithm scheme is provided to optimize the audiovisual presentation, seeking to induce in the transferee the target pattern found in the experimental subject. After the stimulation patterns, which may be path dependent, are determined, it is likely that these patterns will be persistent, though over longer time periods, there may be some desensitization to the stimulation pattern(s). In some cases, audiovisual stimulation is insufficient, and TMS or other electromagnetic stimulation (superthreshold, or preferably subthreshold) is employed to assist in achieving the desired state and maintaining it for the desired period.
- (44) Such technology can be used to significantly enhance the emotional response to viewing photos, reproduction of art, virtual reality, TV, listening to music, reading a book, etc. The user's emotional state may be primed for the secondary stimulation, to enhance the results.
- (45) For example, when a movie is filmed, actors get into their roles and experience real emotions. If we record these emotions by recording their

brainwaves during acting and later playing them back to viewers or otherwise induce in the viewers the same emotional states, while they are watching the film, this would significantly enhance the experience. As discussed above, the emotional state of an actor may be determined based on a script, facial recognition, explicit statement of the actor, etc., and need not be deciphered from the EEG.

(46) Similarly, while producing virtual reality, we can couple digital files containing video with files of brainwaves of people present during the recording, who see the nature in real time and experience emotions first hand, which would dramatically enhance VR experience.

(47) In another example, a book or an eBook can be coupled with a file of recorded brainwaves of the writer or an experienced actor who is trained to evoke an emotional response while reading a script may provide the stimulus.

(48) One of the challenges of adapting robotic technology and artificial intelligence (AI) is a typical lack of an emotional response by a human subject to a robot or an AI software agent Using brainwaves can help evoke a positive emotional response in humans while interacting with robots and/or AI agents.

(49) One purpose of this invention is to enhance an emotional response by a subject while engaged in mood. Yet another purpose of this invention is to enhance an emotional response by a subject while engaged in entertainment Still another purpose of this invention is to enhance an emotional response by a subject while engaged with a robot or an artificial intelligence, another purpose of this invention is to assist a person with recalling a past experience, still another purpose of this invention is to assist a person suffering from a form of dementia to recognize the person's family members and friends.

(50) It may be difficult for many to experience the emotional response to a representation of an experience as to the genuine experience. Looking at a photograph of a Grand Canyon does not elicit the same emotional response as seeing the Grand Canyon itself. Looking at a reproduction of Mona Lisa does not elicit the same emotional response as seeing the original painting in Louvre. An immersive experience achieved through virtual reality (VR) applications goes a long way in simulating the reality, but still falls short of eliciting the emotional response comparable with the one associated with real experience.

(51) Elderly people suffering from Alzheimer's disease or other forms of dementia have difficult recalling their past experiences and recognized family members and friends. While in the early stages of the disease they may have difficulty recalling the person's name or identity, but they still recognize a family member as a loved one responding to seeing a family member with a positive emotion. In later stages, however, the patients no longer feel the emotional response upon seeing a family member and are frightened as if seeing a total stranger.

(52) Recording brainwaves while a person is experiencing a strong emotional response to a genuine experience and later transmitting these recorded brainwaves to another or same individual may help experience stronger emotional response. For example, recording brainwaves of a person seeing for the first time the Grand Canyon and transmitting these brainwaves to another (or the same) person who is viewing a photograph of the Grand Canyon or viewing it through VR glasses would enhance the emotional response of that person and help create more genuine immersive experience. Similarly, recording brainwaves of a person seeing for the first time the original painting of Mona Lisa in the Louvre and transmitting these brainwaves to another (or the same) person who is viewing a reproduction of this painting or on a virtual museum tour of the Louvre viewing it through VR glasses would enhance the emotional response of that person and help create more genuine immersive experience.

(53) In another example, recording brainwaves of a musician playing the music in a concert and transmitting these brainwaves to another person who is listening to a recording of this music would enhance the emotional response of that person and help create more genuine immersive experience.

(54) In a further example, recording brainwaves of actors while acting in movie and transmitting these brainwaves to viewers who are watching the movie in a theater, on a television, on a computer, or through VR glasses would enhance the emotional response of that person and help create more genuine immersive experience.

(55) A further example provides that brainwaves associated with specific emotions may be recorded from actors asked to experience these emotions. A library of brainwaves corresponding to specific emotions can be assembled and used to enhance emotional response, for example, of a gamer playing a computer game, with sequences of emotions triggered in the gamer according to the context or paradigm of the game. There are many applications where such library of brainwaves can be use. Examples include use by law enforcement in helping deescalate a conflict or diffuse a situation by calming down people invoked in the conflict or situation. It can be used by health care providers in the hospitals to help patients maintain positive attitude so important to their recovery. It can be used by personnel in psychiatric wards in calming down psychiatric patient without the use of psychotropic medications. It can be used in spas and meditation retreats or by individuals wishing to achieve the relaxation response to induce feeling of peace and calm or, perhaps, even the altered state of consciousness. It can be used by athletes, creative people, scientists and other wishing to get into the "zone" to achieve pick performance or creative inspiration.

(56) In another example, recording brainwaves of a passionate teacher enthusiastically explaining a difficult subject and transmitting these brainwaves to a student who is studying the same subject would enhance the emotional response of that person and help maintain focus, concentration, interest and may even help understand the subject of study.

(57) In a further example, recording brainwaves associated with the emotional response of a person to his family members or friends while in the initial stages of the Alzheimer's disease or another form of dementia and later transmitting these brainwaves to the same person while in a later stages of the disease may help the patient recognize the familiar faces or, at least, create a positive emotional response upon seeing family members reducing the fear and anxiety associate with inability to recognize familiar faces typical for the later stages of Alzheimer's disease and dementia.

(58) The transmission of the brainwaves can be accomplished through direct electrical contact with the electrodes implanted in the brain or remotely employing light, sound, electromagnetic waves and other non-invasive techniques.

(59) Light, sound or invisible electromagnetic fields may be used to remotely convey the temporal pattern of prerecorded brainwaves to a subject, by modulating the encoded temporal frequency on the light, sound or electromagnetic filed signal to which the subject is exposed.

(60) Another embodiment is combining a text with the code encoding the temporal pattern of brainwaves of a person reading the text who has normal or accentuated affect Say a user is reading a lengthy text (a legal brief or an eBook) on a computer screen. While displaying the text computer monitor (or another light source) generates light frequency corresponding to the temporal pattern of brainwaves of another person reading the same text, prerecorded and embedded with the text. The result is speed reading and improved comprehension and retention of the information while achieving the same emotional states as the other person. This may have use in persons with abnormal psyche, who fail to achieve normal emotional response to media.

(61) Employing light, sound or electromagnetic field to remotely convey the temporal pattern of brainwaves (which may be prerecorded) to a subject by modulating the encoded temporal frequency on the light, sound or electromagnetic filed signal to which the subject is exposed.

(62) When a group of neurons fires simultaneously, the activity appears as a brainwave. Different brainwave-frequencies are linked to different emotional or mental states in the brain.

(63) The EEG pattern may be derived from another individual or individuals, the same individual at a different time, or an in vivo animal model of the desired metal state. The method may therefore replicate a mental state of a first subject in a second subject. The mental state typically is not a state of consciousness or an idea, but rather a subconscious (in a technical sense) state, representing an emotion, readiness, receptivity, or other state, often independent of particular thoughts or ideas. In essence, a mental state of the first subject (a "trainer" or "donor" who is in a desired mental state) is captured by recording neural correlates of the mental state, e.g., as expressed by brain activity patterns, such as EEG or MEG signals. The neural correlates of the first subject, either as direct or recorded representations, may then be used to control a stimulation of the second subject (a "trainee" or "recipient"), seeking to induce the same brain activity patterns in the second subject (recipient/trainee) as were present in the first subject (donor/trainer) to assist the second subject (recipient/trainee) to attain the desired mental state that had been attained by the donor/trainer. In an alternative embodiment, the signals from the first subject (donor/trainer) being in the first mental state are employed to prevent the second subject

(recipient/traine) from achieving a second mental state, wherein the second mental state is an undesired/one.

(64) The source brain wave pattern may be acquired through multichannel EEG or MEG, from a human in the desired brain state. A computational model of the brain state is difficult to create. However, such a model is not required according to the present technology. Rather, the signals may be processed by a statistical process (e.g., PCA or a related technology), or a statistically trained process (e.g., a neural network). The processed signals preferably retain information regarding signal source special location, frequency, and phase. In stimulating the recipient's brain, the source may be modified to account for brain size differences, electrode locations, etc. Therefore, the preserved characteristics are normalized spatial characteristics, frequency, phase, and modulation patterns.

(65) The normalization may be based on feedback from the target subject, for example based on a comparison of a present state of the target subject and a corresponding state of the source subject, or other comparison of known states between the target and source. Typically, the excitation electrodes in the target subject do not correspond to the feedback electrodes or the electrodes on the source subject. Therefore, an additional type of normalization is required, which may also be based on a statistical or statistically trained algorithm.

(66) According to one embodiment, the stimulation of the second subject is associated with a feedback process, to verify that the second subject has appropriately responded to the stimulation, e.g., has a predefined similarity to the mental state as the first subject, has a mental state with a predefined difference from the first subject, or has a desire change from a baseline mental state. Advantageously, the stimulation may be adaptive to the feedback. In some cases, the feedback may be functional, i.e., not based on brain activity per se, or neural correlates of mental state, but rather physical, psychological, or behavioral effects that may be reported or observed.

(67) The feedback typically is provided to a computational model-based controller for the stimulator, which alters stimulation parameters to optimize the stimulation in dependence on a brain and brain state model applicable to the target.

(68) For example, it is believed that brainwaves represent a form of resonance, where ensembles of neurons interact in a coordinated fashion as a set of coupled or interacting oscillators. The frequency of the wave is related to neural responsivity to neurotransmitters, distances along neural pathways, diffusion limitations, etc., and perhaps pacemaker neurons or neural pathways. That is, the same mental state may be represented by different frequencies in two different individuals, based on differences in the size of their brains, neuromodulators present, physiological differences, etc. These differences may be measured in microseconds or less, resulting in fractional changes in frequency. However, if the stimulus is different from the natural or resonant frequency of the target process, the result may be different from that expected. Therefore, the model-based controller can determine the parameters of neural transmission and ensemble characteristics, vis-à-vis stimulation, and resynthesize the stimulus wave to match the correct waveform, with the optimization of the waveform adaptively determined. This may not be as simple as speeding up or slowing down playback of the signal, as different elements of the various waveforms representing neural correlates of mental state may have different relative differences between subjects. Therefore, according to one set of embodiments, the stimulator autocalibrates for the target, based on a correspondence (error) of a measured response to the stimulation and the desired mental state sought by the stimulation. In cases where the results are chaotic or unpredictable based on existing data, a genetic algorithm may be employed to explore the range of stimulation parameters, and determine the response of the target. In some cases, the target has an abnormal or unexpected response to stimulation based on a model maintained within the system. In this case, when the deviance from the expected response is identified, the system may seek to new model, such as from a model repository that may be on-line, such as through the Internet. If the models are predictable, a translation may be provided between an applicable model of a source or trainer, and the applicable model of the target, to account for differences. In some cases, the desired mental state is relatively universal, such as sleep and awake. In this case, the brain response model may be a statistical model, rather than a neural network or deep neural network type implementation.

(69) Thus, in one embodiment, a hybrid approach is provided, with use of donor-derived brainwaves, on one hand, which may be extracted from the brain activity readings (e.g., EEG or MEG) of the first at least one subject (donor), preferably processed by principal component analysis, or spatial principal component analysis, autocorrelation, or other statistical processing technique (clustering, PCA, etc.) or statistically trained technique (backpropagation of errors, etc.) that separates components of brain activity, which can then be modified or modulated based on high-level parameters, e.g., abstractions. See, [ml4a.github.io/ml4a/how_neural_networks_are_trained/](https://github.com/ml4a/ml4a/how_neural_networks_are_trained/). Thus, the stimulator may be programmed to induce a series of brain states defined by name (e.g., emotional or mental state 1, emotional or mental state 2, etc.) or as a sequence of "abstract" semantic labels, icons, or other representations, each corresponding to a technical brain state or sequence of sub-states. The sequence may be automatically defined, based on biology and the system training, and thus relieve the programmer of low-level tasks. However, in a general case, the present technology maintains use of components or subcomponents of the donor's brain activity readings, e.g., EEG or MEG, and does not seek to characterize or abstract them to a semantic level.

(70) According to the present technology, a neural network system or statistical classifier may be employed to characterize the brain wave activity and/or other data from a subject. In addition to the classification or abstraction, a reliability parameter is presented, which predicts the accuracy of the output. Where the accuracy is high, a model-based stimulator may be provided to select and/or parameterize the model, and generate a stimulus for a target subject. Where the accuracy is low, a filtered representation of the signal may be used to control the stimulator, bypassing the model(s). The advantage of this hybrid scheme is that when the model-based stimulator is employed, many different parameters may be explicitly controlled independent of the source subject. On the other hand, where the data processing fails to yield a highly useful prediction of the correct model-based stimulator parameters, the model itself may be avoided, in favor of a direct stimulation type system.

(71) Of course, in some cases, one or more components of the stimulation of the target subject may be represented as abstract or semantically defined signals, and more generally the processing of the signals to define the stimulation will involve high level modulation or transformation between the source signal received from the first subject, to define the target signal for stimulation of the second subject.

(72) Preferably, each component represents a subset of the neural correlates reflecting brain activity that have a high spatial autocorrelation in space and time, or in a hybrid representation such as wavelet. For example, one signal may represent a modulated 10.2 Hz signal, while another signal represents a superposed modulated 15.7 Hz signal, with respectively different spatial origins. These may be separated by optimal filtering, once the spatial and temporal characteristics of the signal are known, and bearing in mind that the signal is accompanied by a modulation pattern, and that the two components themselves may have some weak coupling and interaction.

(73) In some cases, the base frequency, modulation, coupling, noise, phase jitter, or other characteristic of the signal may be substituted. For example, if the first subject is listening to music, there will be significant components of the neural correlates that are synchronized with the particular music. On the other hand, the music per se may not be part of the desired stimulation of the target subject. Therefore, though signal analysis and decomposition, the components of the signal from the first subject, which have a high temporal correlation with the music, may be extracted or suppressed from the resulting signal. Further, the target subject may be in a different acoustic environment, and it may be appropriate to modify the residual signal dependent on the acoustic environment of the target subject, so that the stimulation is appropriate for achieving the desired effect, and does not represent phantoms, distractions, or irrelevant or inappropriate content. In order to perform processing, it is convenient to store the signals or a partially processed representation, though a complete real-time signal processing chain may be implemented. Such a real-time signal processing chain is generally characterized in that the average size of a buffer remains constant, i.e., the lag between output and input is relatively constant, bearing in mind that there may be periodicity to the processing.

(74) The mental state of the first subject may be identified, and the neural correlates of brain activity captured. The second subject is subject to stimulation based on the captured neural correlates and the identified mental state. The mental state may be represented as a semantic variable, within a limited classification space. The mental state identification need not be through analysis of the neural correlates signal, and may be a volitional self-identification by the first subject, a manual classification by third parties, or an automated determination. The identified mental state is useful, for

example, because it represents a target toward (or against) which the second subject can be steered.

(75) The stimulation may be one or more inputs to the second subject, which may be an electrical or magnetic transcranial stimulation, sensors stimulation, mechanical stimulation, ultrasonic stimulation, etc., and controlled with respect to waveform, intensity/amplitude, duration, feedback, self-reported effect by the second subject, manual classification by third parties, automated analysis of brain activity, behavior, physiological parameters, etc. of the second subject.

(76) The process may be used to induce in the target subject neural correlates of the desired mental state, which are derived from a different time for the same person, or a different person at the same or a different time. For example, one seeks to induce the neural correlates of the first subject in a desired mental state in a second subject, through the use of stimulation parameters comprising a waveform over a period of time derived from the neural correlates of mental state of the first subject.

(77) The first and second subjects may be spatially remote from each other, and may be temporally remote as well. In some cases, the first and second subject are the same animal (e.g., human), temporally displaced. In other cases, the first and second subject are spatially proximate to each other. In some cases, neural correlates of a desired mental state are derived from a mammal having a simpler brain, which are then extrapolated to a human brain. (Animal brain stimulation is also possible, for example to enhance training and performance). When the first and second subjects share a common environment, the signal processing of the neural correlates, and especially of real-time feedback of neural correlates from the second subject may involve interactive algorithms with the neural correlates of the first subject.

(78) The first and second subjects may each be subject to stimulators. The first subject and the second subject may communicate with each other in real-time, with the first subject receiving stimulation based on the second subject, and the second subject receiving feedback based on the first subject. This can lead to synchronization of mental state between the two subjects. However, the first subject need not receive stimulation based on real-time signals from the second subject, as the stimulation may derive from a third subject, or the first or second subjects at different points in time.

(79) The neural correlates may be, for example, EEG, qEEG, or MEG signals. Traditionally, these signals are found to have dominant frequencies, which may be determined by various analyses. One embodiment provides that the modulation pattern of a brainwave of the first subject is determined independent of the dominant frequency of the brainwave (though typically within the same class of brainwaves), and this modulation imposed on a wave corresponding to the dominant frequency of the second subject. That is, once the second subject achieves that same brainwave pattern as the first subject (which may be achieved by means other than electromagnetic, mechanical, or sensors stimulation), the modulation pattern of the first subject is imposed as a way of guiding the mental state of the second subject.

(80) The second subject may be stimulated with a stimulation signal which faithfully represents the frequency composition of a defined component of the neural correlates of the first subject.

(81) The stimulation may be performed, for example, by using a tDCS device, a high-definition tDCS device, a tACS device, a TMS device, a deep TMS device, and a source of one of a light signal and a sound signal configured to modulate the dominant frequency on the one of a light signal and a sound signal. The stimulus may be at least one of a light signal, a sound signal, an electric signal, and a magnetic field. The electric signal may be a direct current signal or an alternating current signal. The stimulus may be a transcranial electric stimulation, a transcranial magnetic stimulation, a deep magnetic stimulation, a light stimulation, or a sound stimulation. A visual stimulus may be ambient light or a direct light. An auditory stimulus may be binaural beats or isochronic tones.

(82) The technology may also provide a processor configured to process the neural correlates of mental state from the first subject, and to produce or define a stimulation pattern for the second subject selectively dependent on a waveform pattern of the neural correlates from the first subject. Typically, the processor performs signal analysis and calculates at least a dominant frequency of the brainwaves of the first subject, and preferably also spatial and phase patterns within the brain of the first subject.

(83) A signal is presented to a second apparatus, configured to stimulate the second subject, which may be an open loop stimulation dependent on a non feedback controlled algorithm, or a closed loop feedback dependent algorithm. In other cases, Do analog processing is employed in part or in whole, wherein the algorithm comprises an analog signal processing chain. The second apparatus receives information from the processor (first apparatus), typically comprising a representation of a portion of a waveform represented in the neural correlates. The second apparatus produces a stimulation intended to induce in the second subject the desired mental state, e.g., representing the same mental state as was present in the first subject.

(84) A typical process performed on the neural correlates is a filtering to remove noise. For example, notch filters may be provided at 50 Hz, 60 Hz, 100 Hz, 120 Hz, and additional overtones. Other environmental signals may also be filtered in a frequency-selective or waveform-selective (temporal) manner. Higher level filtering may also be employed, as is known in the art. The neural correlates, after noise filtering, may be encoded, compressed lossy or losslessly, encrypted, or otherwise processed or transformed. The stimulator associated with the second subject would typically perform decoding, decompression, decryption, inverse transformation, etc.

(85) Information security and copy protection technology, similar to that employed for audio signals, may be employed to protect the neural correlate signals from copying or content analysis before use. In some cases, it is possible to use the stored encrypted signal in its encrypted form, without decryption. For example, with an asymmetric encryption scheme, which supports distance determination. See U.S. Pat. No. 7,269,277; Sahai and Waters (2005) Annual International Conference on the Theory and Applications of Cryptographic Techniques, pp. 457-473. Springer, Berlin, Heidelberg; Bringer et al. (2009) IEEE International Conference on Communications, pp. 1-6; Juels and Sudan (2006) Designs, Codes and Cryptography 2:237-257; Thaker et al. (2006) IEEE International Conference on Workload Characterization, pp. 142-149; Galil et al. (1987) Conference on the Theory and Application of Cryptographic Techniques, pp. 135-155.

(86) Because the system may act intrusively, it may be desirable to authenticate the stimulator or parameters employed by the stimulator before use. For example, the stimulator and parameters it employs may be authenticated by a distributed ledger, e.g., a blockchain. On the other hand, in a closed system, digital signatures and other hierarchical authentication schemes may be employed. Permissions to perform certain processes may be defined according to smart contracts, which automated permissions (i.e., cryptographic authorization) provided from a blockchain or distributed ledger system. Of course, centralized management may also be employed.

(87) In practice, the feedback signal from the second subject may be correspondingly encoded as per the source signal, and the error between the two minimized. In such an algorithm, the signal sought to be authenticated is typically brought within an error tolerance of the encrypted signal before usable feedback is available. One way to accomplish this is to provide a predetermined range of acceptable authenticatable signals which are then encoded, such that an authentication occurs when the putative signal matches any of the predetermined range. In the case of the neural correlates, a large set of digital hash patterns may be provided representing different signals as hash patterns. The net result is relatively weakened encryption, but the cryptographic strength may still be sufficiently high to abate the risks.

(88) The processor may perform a noise reduction distinct from a frequency-band filtering. The neural correlates may be transformed into a sparse matrix, and in the transform domain, components representing high probability noise are masked, while components representing high probability signal are preserved. The distinction may be optimized or adaptive. That is, in some cases, the components which represent modulation that are important may not be known a priori. However, dependent on their effect in inducing the desired response in the second subject, the "important" components may be identified, and the remainder filtered or suppressed. The transformed signal may then be inverse-transformed, and used as a basis for a stimulation signal.

(89) A mental state modification, e.g., brain entrainment, may be provided, which ascertains a mental state in a plurality of first subjects; acquires brainwaves of the plurality of first subjects, e.g., using one of EEG and MEG, to create a dataset containing representing brainwaves of the plurality of first subjects. The database may be encoded with a classification of mental state, activities, environment, or stimulus patterns, applied to the

plurality of first subjects and the database may include acquired brainwaves across a large number of mental states, activities, environments, or stimulus patterns, for example. In many cases, the database records will reflect a characteristic or dominate frequency of the respective brainwaves. As discussed above, the trainer or first subject is a convenient source of the stimulation parameters, but is not the sole available source. The database may be accessed according to its indexing, e.g., mental states, activities, environment, or stimulus patterns, for example, and a stimulation pattern for a second subject defined based on the database records of one or more subjects.

(90) The record(s) thus retrieved are used to define a stimulation pattern for the second subject. The selection of records, and their use, may be dependent on the second subject and/or feedback from the second subject. As a relatively trivial example, a female second subject could be stimulated principally dependent on records from female first subjects. Of course, a more nuanced approach is to process the entirety of the database and stimulate the second subject based on a global brain wave-stimulus model, though this is not required, and also, the underlying basis for the model may prove unreliable or inaccurate. In fact, it may be preferred to derive a stimulus waveform from only a single first subject, in order to preserve micro-modulation aspects of the signal, which as discussed above have not been fully characterized. However, the selection of the first subject(s) need not be static, and can change frequently. The selection of first subject records may be based on population statistics of other users of the records (i.e., collaborative filtering, i.e., whose response pattern do I correlate highest with? etc.). The selection of first subject records may also be based on feedback patterns from the second user.

(91) The process of stimulation may seek to target a desired mental state in the second subject, which is automatically or semi-automatically determined or manually entered. That target then represents a part of the query against the database to select the desired record(s). The selection of records may be a dynamic process, and reselection of records may be feedback dependent.

(92) The records may be used to define a modulation waveform of a synthesized carrier or set of carriers, and the process may include a frequency domain multiplexed multi-subcarrier signal (which is not necessarily orthogonal). A plurality of stimuli may be applied concurrently, through the suffered subchannels and/or through different stimulator electrodes, magnetic field generators, mechanical stimulators, sensory stimulators, etc. The stimuli for the different subchannels or modalities need not be derived from the same records.

(93) The stimulus may be applied to achieve the desired mental state, e.g., brain entrainment of the second subject with one or more first subjects. Brain entrainment is not the only possible outcome of this process. If the plurality of first subjects are mutually entrained, then each will have a corresponding brain wave pattern dependent on the basis of brainwave entrainment. This link between first subject may be helpful in determining compatibility between a respective first subject and the second subject. For example, characteristic patterns in the entrained brainwaves may be determined, even for different target mental states, and the characteristic patterns correlated to find relatively close matches and to exclude relatively poor matches.

(94) This technology may also provide a basis for a social network, dating site, employment or vocational testing, or other interpersonal environments, wherein people may be matched with each other based on entrainment characteristics. For example, people who efficiently entrain with each other may have better social relationships than those who do not. Thus, rather than seeking to match people based on personality profiles, the match could be made based on an ability of each party to efficiently entrain the brainwave pattern of the other party. This enhances non-verbal communication, and assists in achieving corresponding states during activities. This can be assessed by monitoring neural responses of each individual to video, and also by providing a test stimulation based on the other party's brainwave correlates of mental state, to see whether coupling is efficiently achieved. On the other hand, the technology could be used to assist in entrainment when natural coupling is inefficient or to block coupling where the coupling is undesirable. An example of the latter is hostility; when two people are entrained in a hostile environment, emotional escalation ensues. However, if the entrainment is attenuated, undesired escalation may be impeded.

(95) As discussed above, the plurality of first subjects may have their respective brain wave patterns stored in association with separate database records. However, they may also be combined into a more global model. One such model is a neural network or deep neural network. Typically, such a network would have recurrent features. Data from a plurality of first subjects is used to train the neural network, which is then accessed by inputting the target state and/or feedback information, and which outputs a stimulation pattern or parameters for controlling a stimulator. When multiple first subjects form the basis for the stimulation pattern, it is preferred that the neural network output parameters of the stimulation, derived from and comprising features of the brain wave patterns or other neural correlates of mental state from the plurality of first subjects, which are then used to control a stimulator which, for example, generates its own carrier wave(s) which are then modulated based on the output of the neural network. The neural network need not periodically retrieve records, and therefore may operate in a more time-continuous manner, rather than the more segmented scheme of record-based control.

(96) In any of the feedback dependent methods, the brainwave patterns or other neural correlates of mental state may be processed by a neural network, to produce an output that guides or controls the stimulation. The stimulation, is, for example, at least one of a light (visual) signal, a sound signal, an electric signal, a magnetic field, and a vibration or mechanical stimulus, or other sensory input. The fields may be static or dynamically varying.

(97) The process may employ a relational database of mental states and brainwave patterns, e.g., frequencies/neural correlate waveform patterns associated with the respective mental states. The relational database may comprise a first table, the first table further comprising a plurality of data records of brainwave patterns, and a second table, the second table comprising a plurality of mental states, each of the mental states being linked to at least one brainwave pattern. Data related to mental states and brainwave patterns associated with the mental states are stored in the relational database and maintained. The relational database is accessed by receiving queries for selected mental states, and data records are returned representing the associated brainwave pattern. The brainwave pattern retrieved from the relational database may then be used for modulating a stimulator seeking to produce an effect selectively dependent on the mental state at issue.

(98) A computer apparatus may be provided for creating and maintaining a relational database of mental states and frequencies associated with the mental states, the computer apparatus comprising: a non-volatile memory for storing a relational database of mental states and neural correlates of brain activity associated with the mental states, the database comprising a first table, the first table further comprising a plurality of data records of neural correlates of brain activity associated with the mental states, and a second table, the second table comprising a plurality of mental states, each of the mental states being linked to one or more records in the first table; a processor coupled with the non-volatile memory, configured to process relational database queries, which are then used for searching the database; RAM coupled with the processor and the non-volatile memory for temporary holding database queries and data records retrieved from the relational database; and an I/O interface configured to receive database queries and deliver data records retrieved from the relational database. A SQL or noSQL database may also be used to store and retrieve records.

(99) A further aspect of the technology provides a method of brain entrainment comprising: ascertaining a mental state in a first subject recording brainwaves of the plurality of subjects using at least one channel one of EEG and MEG; storing the recorded brainwaves in a physical memory device; retrieving the brainwaves from the memory device; applying a stimulus signal comprising a brainwave pattern derived from at least one-channel one of the EEG and MEG to a second subject via transcranial stimulation, whereby the mental state desired by the second subject is achieved. The stimulation may be of the same order (number of channels) as the EEG or MEG, or a different number of channels, typically reduced. For example, the EEG or MEG may comprise 128 or 256 channels, while the transcranial stimulator may have 8 or fewer channels. Sensory stimulation of various modalities and patterns may accompany the transcranial stimulation.

(100) The at least one channel may be less than six channels and the placement of electrodes used for transcranial stimulation may be approximately the same as the placement of electrodes used in recording of said one of EEG and MEG.

(101) The present technology may be responsive to chronobiology, and in particular to the subjective sense of time. For a subject, this may be determined volitionally subjectively, but also automatically, for example by judging attention span, using e.g., eye movements, and analyzing

the persistence of brainwave patterns or other physiological parameters over a discrete stimulus. Further, time-constants of the brain, reflected by delays and phase may also be analyzed. Further, the contingent negative variation (CNV) preceding a volitional act may be used, both to determine (or measure) conscious action timing, and also the time relationships between thought and action more generally.

(102) Typically, brainwave activity is measured with a large number of EEG electrodes, which each receive signals from a small area on the scalp, or in the case of a MEG, by a number of sensitive magnetic field detectors, which are responsive to local field differences. Typically, the brainwave capture is performed in a relatively high number of spatial dimensions, e.g., corresponding to the number of sensors. It is often unfeasible to process the brainwave signals to create a source model, given that the brainwaves are created by billions of neurons, connected through axons, which have long distances. Further, the neurons are generally non-linear, and interconnected. However, a source model is not required.

(103) Various types of artificial intelligence techniques may be exploited to analyze the neural correlates of an emotional or mental state represented in the brain activity data of both the first subject (donor) (or plurality of donors) and the second subject (recipient). The algorithm or implementation need not be the same, though in some cases, it is useful to confirm the approach of the source processing and feedback processing so that the feedback does not achieve or seek a suboptimal target emotional or mental state. However, given the possible differences in conditions, resources, equipment, and purpose, there is no necessary coordination of these processes. The artificial intelligence may take the form of neural networks or deep neural networks, though rule/expert-based systems, hybrids, and more classical statistical analysis may be used. In a typical case, an artificial intelligence process will have at least one aspect, which is non-linear in its output response to an input signal, and thus at least the principle of linear superposition is violated. Such systems tend to permit discrimination, since a decision and the process of decision-making are, ultimately, non-linear. An artificially intelligent system requires a base of experience or information upon which to train. This can be a supervised (external labels applied to data), unsupervised (self-discrimination of classes), or semi-supervised (a portion of the data is externally labelled).

(104) A self-learning or genetic algorithm may be used to tune the system, including both or either the signal processing at the donor system and the recipient system. In a genetic algorithm feedback-dependent self-learning system, the responsivity of a subject, e.g., the target, to various kinds of stimuli may be determined over a stimulus space. This stimulation may be in the context of use, with a specific target emotional or mental state provided, or unconstrained. The stimulator may operate using a library of stimulus patterns, or seek to generate synthetic patterns or modifications of patterns. Over a period of time, the system will learn to map a desired emotional or mental state to optimal context-dependent parameters of the stimulus pattern.

(105) In some cases it may be appropriate to administer a drug or pharmacological agent, such as melatonin, hypnotic or soporific drug, a sedative (e.g., barbiturates, benzodiazepines, nonbenzodiazepine hypnotics, orexin antagonists, antihistamines, general anesthetics, *Cannabis* and other herbal sedatives, methaqualone and analogues, muscle relaxants, opioids) that assists in achieving the target emotional or mental state, and for emotional states and/or dreams, this may include certain psychotropic drugs, such as epinephrine, norepinephrine reuptake inhibitors, serotonin reuptake inhibitors, peptide endocrine hormones, such as oxytocin, ACTH fragments, insulin, etc. Combining a drug with stimulation may reduce the required dose of the drug and the associated side effects of the drug.

(106) The technology may be used to modify or alter a mental state (e.g., from sleep to waking and vice versa) in a subject. Typically, the starting mental state, brain state, or brainwave pattern is assessed, such as by EEG, MEG, observation, stimulus-response amplitude and/or delay, or the like. Of particular interest in uncontrolled environments are automated mental state assessments, which do not rely on human observation or EEG signals, and rather may be acquired through MEG (e.g., SQID, optically-pumped magnetometer), EMG, MMG (magnetomyogram), mechanical (e.g., accelerometer, gyroscope, etc.), data from physiological sensors (e.g., AKG, heartrate, respiration rate, temperature, galvanic skin potential, etc.), or automated camera sensors.

(107) For example, cortical stimulus-response pathways and reflexes may be exercised automatically, to determine their characteristics on a generally continuous basis. These characteristics may include, for example, a delay between stimulus and the observed central (e.g., EEG) or peripheral response (e.g., EMG, limb accelerometer, video). Typically, the same modality will be used to assess the pre-stimulation state, stimulus response, and post-stimulation state, though this is not a limitation.

(108) In order to change the mental state, a stimulus is applied in a way designed to alter the mental state in a desired manner. A state transition table, or algorithm, may be employed to optimize the transition from a starting mental state to a desired mental state. The stimulus may be provided in an open loop (predetermined stimulus protocol) or closed loop (feedback adapted stimulus protocol), based on observed changes in a monitored variable.

(109) Advantageously, a characteristic delay between application of stimulus and determination of response varies with the brain or mental state. For example, some mental states may lead to increased delay or greater variability in delay, while others may lead to decreased or lower variability. Further, some states may lead to attenuation of response, while others may lead to exaggerated response. In addition, different mental states can be associated with qualitatively different responses. Typically, the mere assessment of the brain or mental state should not itself alter the state, though in some cases the assessment and transition influence may be combined. For example, in seeking to assist in achieving a deep sleep state, excitation that disturbs sleep is contraindicated.

(110) In cases where a brainwave pattern is itself determined by EEG (which may be limited to relatively controlled environments), brainwaves representing that pattern represent coherent firing of an ensemble of neurons, defining a phase. One way to change the state is to advance or retard the triggering of the neuronal excitation, which can be a direct or indirect excitation or inhibition, caused, for example, by electrical, magnetic, mechanical, or sensory stimulation. This stimulation may be time-synchronized with the detected (e.g., by EEG) brainwaves, for example with a phase lead or lag with respect to the detected pattern. Further, the excitation can steer the brainwave signal by continually advancing to a desired state, which through the continual phase rotation represents a different frequency. After the desired new state is achieved, the stimulus may cease, or be maintained in a phase-locked manner to hold the desired state.

(111) A predictive model may be used to determine the current mental state, optimal transition to a desired mental state, when the subject has achieved the desired mental state, and how to maintain the desired mental state. The desired mental state itself may represent a dynamic sequence (e.g., stage 1.fwdarw.stage 2.fwdarw.stage 3, etc.), such that the subject's mental state is held for a desired period in a defined condition. Accordingly, the stimulus may be time-synchronized with respect to the measured brainwave pattern.

(112) Direct measurement or determination of brainwaves or their phase relationships is not necessarily required. Rather, the system may determine tremor or reflex patterns. Typically, the reflex patterns of interest involve central pathways, and more preferably brain reflex pathways, and not spinal cord mediated reflexes, which are less dependent on instantaneous brain state. The central reflex patterns can reflect a time delay between stimulation and motor response, an amplitude of motor response, a distribution of response through various afferent pathways, variability of response, tremor or other modulation of motor activity, etc. Combinations of these characteristics may be employed, and different subsets may be employed at different times or to reflect different states. Similar to evoked potentials, the stimulus may be any sense, especially sight, sound, touch/proprioception/pain/etc., though the other senses, such as taste, smell, balance, etc., may also be exercised. A direct electrical or magnetic excitation is also possible. As discussed, the response may be determined through EEG, MEG, or peripheral afferent pathways.

(113) Normalization of brain activity information may be spatial and/or temporal. For example, the EEG electrodes between sessions or for different subject may be in different locations, leading to a distortion of the multichannel spatial arrangement. Further, head size and shape of different individuals is different, and this needs to be normalized and/or encoded as well. The size and shape of the head/skull and/or brain, may also lead to temporal differences in the signals, such as characteristic time delays, resonant or characteristic frequencies, etc.

(114) One way to account for these effects is through use of a time-space transform, such as a wavelet-type transform. It is noted that, in a corresponding way that statistical processes are subject to frequency decomposition analysis through Fourier transforms, they are also subject to

time-frequency decomposition using wavelet transform (DWT), though more complex and less regular transforms may be employed. As discussed above, principal component analysis (PCA) and spatial PCA may be used to analyze signals, presuming linearity (linear superposition) and statistical independence of components. However, these presumptions technically do not apply to brainwave data, and practically, one would normally expect interaction between brain wave components (non-independence) and lack of linearity (since “neural networks” by their nature are non-linear), defeating use of PCA or spatial PCA unmodified. However, a field of nonlinear dimensionality reduction provides various techniques to permit corresponding analyses under presumptions of non-linearity and non-independence. 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(115) Therefore, statistical approaches are available for separating EEG signals from other signals, and for analyzing components of EEG signals themselves. According to the present invention, various components that might be considered noise in other contexts, e.g., according to prior technologies, such as a modulation pattern of a brainwave, are preserved. Likewise, interactions and characteristic delays between significant brainwave events are preserved. This information may be stored either integrated with the brainwave pattern in which it occurs, or as a separated modulation pattern that can then be recombined with an unmodulated brainwave pattern to approximate the original subject.

(116) According to the present technology, lossy “perceptual” encoding (i.e., functionally optimized with respect to subjective response) of the brainwaves may be employed to process, store and communicate the brainwave information. In a testing scenario, the “perceptual” features may be tested, so that important information is preserved over information that does not strongly correspond to the effective signal. Thus, while one might not know a priori which components represent useful information, a genetic algorithm may empirically determine which features or data reduction algorithms or parameter sets optimize retention of useful information vs. information efficiency. It is noted that subjects may differ in their response to signal components, and therefore the “perceptual” encoding may be subjective with respect to the recipient. On the other hand, different donors may have different information patterns, and therefore each donor may also require individual processing. As a result, pairs of donor and recipient may require optimization, to ensure accurate and efficient communication of the relevant information. According to the present invention, sleep/wake mental states and their corresponding patterns are sought to be transferred. In the recipient, these patterns have characteristic brainwave patterns. Thus, the donor may be used, under a variety of alternate processing schemes, to stimulate the recipient, and the sleep/wake response of the recipient determined based on objective criteria, such as resulting brainwave patterns or expert observer reports, or subjective criteria, such as recipient self-reporting, survey or feedback. Thus, after a training period, an optimized processing of the donor, which may include filtering, dominant frequency resynthesis, feature extraction, etc., may be employed, which is optimized for both donor and recipient. In other cases, the donor characteristics may be sufficiently normalized, that only recipient characteristics need be compensated. In a trivial case, there is only one exemplar donor, and the signal is oversampled and losslessly recorded, leaving only recipient variation as a significant factor.

(117) Because dominant frequencies tend to have low information content (as compared to the modulation of these frequencies and interrelation of various sources within the brain), one efficient way to encode the main frequencies is by location, frequency, phase, and amplitude. The modulation of a wave may also be represented as a set of parameters. By decomposing the brainwaves according to functional attributes, it becomes possible, during stimulation, to modify the sequence of “events” from the donor, so that the recipient need not experience the same events, in the same order, and in the same duration, as the donor. Rather, a high-level control may select states, dwell times, and transitions between states, based on classified patterns of the donor brainwaves. The extraction and analysis of the brainwaves of the donors, and response of the recipient, may be performed using statistical processes, such as principle components analysis (PCA), independent component analysis (ICA), and related techniques; clustering, classification, dimensionality reduction and related techniques; neural networks and other known technologies. These algorithms may be implemented on general purpose CPUs, array processors such as GPUs, and other technologies.

(118) In practice, a brainwave pattern of the first subject may be analyzed by a PCA technique that respects the non-linearity and non-independence

of the brainwave signals, to extract the major cyclic components, their respective modulation patterns, and their respective interrelations. The major cyclic components may be resynthesized by a waveform synthesizer, and thus may be efficiently coded. Further, a waveform synthesizer may modify frequencies or relationships of components from the donor based on normalization and recipient characteristic parameters. For example, the brain of the second subject (recipient) may have characteristic classified brainwave frequencies 3% lower than the donor (or each type of wave may be separately parameterized), and therefore the resynthesis may take this difference into account. The modulation patterns and interrelations may then be reimposed onto the resynthesized patterns. The normalization of the modulation patterns and interrelations may be distinct from the underlying major cyclic components, and this correction may also be made, and the normalized modulation patterns and interrelations included in the resynthesis. If the temporal modifications are not equal, the modulation patterns and interrelations may be decimated or interpolated to provide a correct continuous time sequence of the stimulator. The stimulator may include one or more stimulation channels, which may be implemented as electrical, magnetic, auditory, visual, tactile, or other stimulus, and/or combinations.

(119) The stimulator is preferably feedback controlled. The feedback may relate to the brainwave pattern of the recipient, and/or context or ancillary biometric basis. For example, if the second subject (recipient) begins to awaken from sleep, which differs from the first subject (donor) sleep pattern, then the stimulator may resynchronize based on this finding. That is, the stimulator control will enter a mode corresponding to the actual state of the recipient, and seek to guide the recipient to a desired state from a current state, using the available range and set of stimulation parameters. The feedback may also be used to tune the stimulator, to minimize error from a predicted or desired state of the recipient subject based on the prior and current stimulation.

(120) The control for the stimulator is preferably adaptive, and may employ a genetic algorithm to improve performance over time. For example, if there are multiple first subjects (donors), the second subject (recipient) may be matched with those donors from whose brainwave signals (or algorithmically modified versions thereof) the predicted response in the recipient is best, and distinguished from those donors from whose brainwave signals the predicted response in the recipient subject poorly corresponds. Similarly, if the donors have brainwave patterns determined over a range of time and context and stored in a database, the selection of alternates from the database may be optimized to ensure best correspondence of the recipient subject to the desired response.

(121) It is noted that a resynthesizer-based stimulator is not required, if a signal pattern from a donor is available that properly corresponds to the recipient and permits a sufficiently low error between the desired response and the actual response. For example, if a donor and a recipient are the same subject at different times, a large database may be unnecessary, and the stimulation signal may be a minimally processed recording of the same subject at an earlier time. Likewise, in some cases, a deviation is tolerable, and an exemplar signal may be emitted, with relatively slow periodic correction. For example, a sleep signal may be derived from a single subject, and replayed with a periodicity of 90 minutes or 180 minutes, such as a light or sound signal, which may be useful in a dormitory setting, where individual feedback is unavailable or unhelpful.

(122) In some cases, it is useful to provide a stimulator and feedback-based controller on the donor. This will better match the conditions of the donor and recipient, and further allow determination of not only the brainwave pattern of the donor, but also responsiveness of the donor to the feedback. One difference between the donors and the recipients is that in the donor, the natural sleep pattern is sought to be maintained and not interrupted. Thus, the adaptive multi-subject database may include data records from all subject, whether selected ab initio as a useful exemplar or not. Therefore, the issue is whether a predictable and useful response can be induced in the recipient from the database record, and if so, that record may be employed. If the record would produce an unpredictable result, or a non-useful result, the use of that record should be avoided. The predictability and usefulness of the responses may be determined by a genetic algorithm, or other parameter-space searching technology.

(123) FIG. 1 shows the electric activity of a neuron contributing to a brainwave.

(124) FIG. 2 shows transmission of an electrical signal generated by a neuron through the skull, skin and other tissue to be detectable by an electrode transmitting this signal to EEG amplifier.

(125) FIG. 3 shows an illustration of a typical EEG setup with a subject wearing a cap with electrodes connected to the EEG machine, which is, in turn, connected to a computer screen displaying the EEG. FIG. 4 shows a typical EEG reading. FIG. 5 shows one second of a typical EEG signal. FIG. 6 shows main brainwave patterns.

(126) FIG. 7 shows a flowchart according to one embodiment of the invention. Brainwaves from a subject who is in an emotional state are recorded. Brainwaves associated with the emotion are identified. A temporal pattern in the brainwave associated with the emotion is decoded. The decoded temporal pattern is used to modulate the frequency of at least one stimulus. The temporal pattern is transmitted to the second subject by exposing the second subject to said at least one stimulus.

(127) FIG. 8 shows a flowchart according to one embodiment of the invention. Brainwaves in a subject at rest and in an emotional state are recorded, and a brainwave characteristic associated with the emotion is separated by comparing with the brainwaves at rest. A temporal pattern in the brainwave associated with the emotion is decoded and stored. The stored code is used to modulate the temporal pattern on a stimulus, which is transmitted to the second subject by exposing the second subject to the stimulus.

(128) FIG. 9 shows a flowchart according to one embodiment of the invention. Brainwaves in a subject in an emotional state are recorded, and a Fourier Transform analysis performed. A temporal pattern in the brainwave associated with the emotion is then decoded and stored. The stored code is then used to modulate the temporal pattern on a stimulus, which is transmitted to the second subject by exposing the second subject to the stimulus.

(129) FIG. 10 shows a flowchart according to one embodiment of the invention. Brainwaves in a plurality of subjects in a respective emotional state are recorded. A neural network is trained on the recorded brainwaves associated with the emotion. After the neural network is defined, brainwaves in a first subject engaged in the emotion are recorded. The neural network is used to recognize brainwaves associated with the emotion. A temporal pattern in the brainwaves associated with the emotion is decoded and stored. The code is used to modulate the temporal pattern on a stimulus. Brainwaves associated with the emotion in a second subject are induced by exposing the second subject to the stimulus.

(130) FIG. 11 shows a flowchart according to one embodiment of the invention. Brainwaves in a subject both at rest and in an emotional state are recorded. A brainwave pattern associated with the emotion is separated by comparing with the brainwaves at rest. For example, a filter or optimal filter may be designed to distinguish between the patterns. A temporal pattern in the brainwave associated with the emotion is decoded, and stored in software code, which is then used to modulate the temporal pattern of light, which is transmitted to the second subject, by exposing the second subject to the source of the light.

(131) FIG. 12 shows a flowchart according to one embodiment of the invention. Brainwaves in a subject at rest and in an emotion are recorded. A brainwave pattern associated with the emotion is separated by comparing with the brainwaves at rest. A temporal pattern in the brainwave associated with the emotion is decoded and stored as a temporal pattern in software code. The software code is used to modulate the temporal pattern on a sound signal. The temporal pattern is transmitted to the second subject by exposing the second subject to the sound signal.

(132) FIG. 13 shows a flowchart according to one embodiment of the invention. Brainwaves in a subject in an emotional state are recorded, and brainwaves selectively associated with the emotion are identified. A pattern, e.g., a temporal pattern, in the brainwave associated with the emotion, is decoded and used to entrain the brainwaves of the second subject.

(133) FIG. 14 shows a schematic representation of an apparatus according to one embodiment of the invention.

(134) FIG. 15 shows brainwave real time BOLD (Blood Oxygen Level Dependent) fMRI studies acquired with synchronized stimuli.

(135) FIG. 16 shows that a desired mental state may be induced in a target individual (e.g., human, animal), by providing selective stimulation according to a temporal pattern, wherein the temporal pattern is correlated with an EEG pattern of the target when in the desired mental state, or represents a transition which represents an intermediate toward achieving the desired mental state. The temporal pattern may be targeted to a discrete

spatial region within the brain, either by a physical arrangement of a stimulator, or natural neural pathways through which the stimulation (or its result) passes.

(136) FIG. 17 shows brainwave entrainment before and after synchronization. See, Understanding Brainwaves to Expand our Consciousness, fractalenlightenment.com/14794/spirituality/understanding-brainwaves-to-expand-our-consciousness FIG. 18 shows brainwaves during inefficient problem solving and stress.

(137) FIGS. 19 and 20 show how binaural beats work. Binaural beats are perceived when two different pure-tone sine waves, both with frequencies lower than 1500 Hz, with less than a 40 Hz difference between them, are presented to a listener dichotically (one through each ear). See, for example, if a 530 Hz pure tone is presented to a subject's right ear, while a 520 Hz pure tone is presented to the subjects left ear, the listener will perceive the auditory illusion of a third tone, in addition to the two pure-tones presented to each ear. The third sound is called a binaural beat, and in this example would have a perceived pitch correlating to a frequency of 10 Hz, that being the difference between the 530 Hz and 520 Hz pure tones presented to each ear. Binaural-beat perception originates in the inferior colliculus of the midbrain and the superior olivary complex of the brainstem, where auditory signals from each ear are integrated and precipitate electrical impulses along neural pathways through the reticular formation up the midbrain to the thalamus, auditory cortex, and other cortical regions.

(138) FIG. 21 shows Functional Magnetic Resonance Imaging (fMRI)

(139) FIG. 22 shows a photo of a brain forming a new idea.

(140) FIG. 23 shows 3D T2 CUBE (SPACENISTA) FLAIR & DSI tractography.

(141) FIG. 24 shows The EEG activities for a healthy subject during a working memory task.

(142) FIG. 25 shows a flowchart according to one embodiment of the invention. Brainwaves in a subject in an emotional state are recorded.

Brainwaves associated with the emotion are identified. A temporal pattern in the brainwave associated with the emotion is extracted. First and second dynamic audio stimuli are generated, whose frequency differential corresponds to the temporal pattern. Binaural beats are provided using the first and the second audio stimuli to stereo headphones worn by the second subject to entrain the brainwaves of the second subject.

(143) FIG. 25 shows a flowchart according to one embodiment of the invention. Brainwaves of a subject engaged in an emotional state are recorded, and brainwaves associated with the emotion identified. A pattern in the brainwave associated with the emotion is identified, having a temporal variation. Two dynamic audio stimuli whose frequency differential corresponds to the temporal variation are generated, and applied as a set of binaural bits to the second subject, to entrain the brainwaves of the second subject.

(144) FIG. 26 shows a flowchart according to one embodiment of the invention. Brainwaves of a subject in an emotional state are recorded, and brainwaves associated with the emotion identified. A pattern in the brainwave associated with the emotion is identified, having a temporal variation. A series of isochronic tones whose frequency differential corresponds to the temporal variation is generated and applied as a set of stimuli to the second subject, to entrain the brainwaves of the second subject See:

(145) FIG. 27 shows a flowchart according to one embodiment of the invention. Brainwaves of a subject in an emotional state are recorded, and brainwaves associated with the emotion identified. A pattern in the brainwave associated with the emotion is identified, having a temporal variation. Two dynamic light stimuli whose frequency differential corresponds to the temporal variation are generated, and applied as a set of stimuli to the second subject, wherein each eye sees only one light stimulus, to entrain the brainwaves of the second subject.

(146) FIG. 28 shows a flowchart according to one embodiment of the invention. Brainwaves of a subject in an emotional state are recorded, and brainwaves associated with the emotion identified. A pattern in the brainwave associated with the emotion is identified, having a temporal variation. Two dynamic electric stimuli whose frequency differential corresponds to the temporal variation are generated, and applied as transcranial stimulation to the second subject, wherein each electric signal is applied to the opposite side of the subject's head, to entrain the brainwaves of the second subject.

(147) FIG. 29 shows a flowchart according to one embodiment of the invention. Brainwaves of a subject are recorded at rest, and in an emotional state. A brainwave associated with the emotion is separated from the remainder of the signal by comparison with the brainwaves at rest A temporal pattern if the brainwave associated with the emotion is decoded, and stored in software code, in a memory. The software code is then used to modulate a temporal pattern in light, which is transmitted to a second subject, who is exposed to the light.

(148) FIG. 30 shows picture of brain anatomy. FIG. 31 shows a brain map. FIG. 32 shows an image depicting neuron anatomy. FIG. 33 shows graphs representing a dimensional view of emotions. FIG. 34 shows a representation of neural activity with respect to emotional state.

(149) In one embodiment, as shown in FIG. 35, brainwaves of the first subject (donor) being in a positive emotional state are recorded 10. A temporal and spatial patterns are decoded from the recorded brainwaves 20 and stored in a non-volatile memory 30. At a later time, the temporal and spatial patters are retrieved from the non-volatile memory 40 and modulated on at least one stimulus 50, which is applied to the first subject via non-invasive brain stimulation technique 60 to induce the positive emotional state. The positive emotional state may be one of or a combination of the state of happiness, joy, gladness, cheerfulness, delight, optimism, merriment, jovialness, vivaciousness, pleasure, excitement, sexual arousal, exuberance, bliss, ecstasy, relaxation, harmony peacefulness.

(150) In another embodiment, as shown in FIG. 36, brainwaves of the first subject being in a positive emotional state are recorded using EEG 80. A temporal and spatial patterns are decoded from the EEG 70 and stored in a non-volatile memory 90.

(151) At a later time, the temporal and spatial patters are retrieved from the non-volatile memory 100 and modulated on a direct current 110, which is applied to the first subject via transcranial direct current stimulation (tDCS) 120 to induce the positive emotional state. See FIG. 42.

(152) In further embodiment, as shown in FIG. 37, brainwaves of the first subject being in a positive emotional state are recorded using EEG 130. A temporal and spatial patterns are decoded from the EEG 140 and stored in a non-volatile memory 150. At a later time, the temporal and spatial patters are retrieved from the non-volatile memory 160 and modulated on an alternating current 170, which is applied to the first subject via transcranial alternating current stimulation (tACS) 180 to induce the positive emotional state. It will be understood by a person skilled in the art that transcranial pulsed current stimulation (tPCS), transcranial random noise stimulation (tRNS), or any other type of transcranial electrical stimulation (tES) may be used. See FIGS. 43-47.

(153) In certain embodiments, as shown in FIG. 38, brainwaves of the first subject being in a positive emotional state are recorded using magnetoencephalogram (MEG) 190. A temporal and spatial patterns are decoded from the MEG 200 and stored in a non-volatile memory 210. At a later time, the temporal and spatial patters are retrieved from the non-volatile memory 220 and modulated on a magnetic field 230, which is applied to the second subject via transcranial magnetic stimulation (tMS) 240 to induce the positive emotional state.

(154) In certain embodiments, as shown in FIG. 39, brainwaves of the first subject being in a positive emotional state are recorded using electroencephalogram (EEG) 250. A temporal and spatial patterns are decoded from the EEG 260 and stored in a non-volatile memory 270. At a later time, the temporal and spatial patters are retrieved from the non-volatile memory 280 and modulated on a light signal 290, which is projected to the second subject 300 to induce the positive emotional state. The light signal may be an ambient light, a directed light or a laser beam. The light may be in a visible spectrum or an infrared light In all embodiments the second subject may the same as the first subject.

(155) In certain embodiments, as shown in FIG. 40, brainwaves of the first subject being in a positive emotional state are recorded using electroencephalogram (EEG) 310. A temporal pattern is decoded from the EEG 320 and stored in a non-volatile memory 330. At a later time, the temporal patter is retrieved from the non-volatile memory 340 and modulated on an isotonic sound signal 350, which is projected to the second subject 360 to induce the positive emotional state. The isotonic sound signal may be imbedded in a music or an ambient noise. The sound may be in an audible spectrum, infrasound or ultrasound.

(156) In certain embodiments, as shown in FIG. 41, brainwaves of the first subject being in a positive emotional state are recorded using

electroencephalogram (EEG) **370**. A temporal spatial pattern is decoded from the EEG **380** and stored in a non-volatile memory **390**. The first set of frequencies is computed by adding a predetermined delta to the frequencies of the temporal frequency pattern **400**. The second set of frequencies is computed by subtracting the delta from the frequencies of the temporal frequency pattern **410**. The first set of frequencies is modulated on the first acoustical signal **420**. The second set of frequencies is modulated on the second acoustical signal **430**. The first acoustic signal is played into an ear of the second subject **440**. The second acoustic signal is played into another ear of the second subject **450** thereby producing binaural stimulation to induce the positive emotional state. The isotonic sound signal may be imbedded in a music or an ambient noise. The sound may be in an audible spectrum, infrasound or ultrasound.

(157) FIG. **42** shows graphs of tDCS, tRNS, and tACS stimulation patterns.

(158) FIGS. **43** and **44** show representations of tDCS neural stimulation.

(159) FIG. **45** shows a representation of tACS or tRNS neural stimulation.

(160) FIG. **46** shows a representation of intracranial electrode implantation.

(161) FIG. **47** shows a representation of tDCS electrode location.

Example 1

(162) We record EEG of a first person (source) experiencing an emotional arousal while seeing an authentic scenic view of nature (e.g., standing in front of the Grand Canyon, or Niagara Falls, or Giza Pyramids); then decode the dynamic spatial and/or temporal patterns of the EEG and encode them in software. If a second person (recipient) wants to experience the same emotional arousal while viewing a representation (e.g., a painting, a photograph or a video) of the same scenic view, the software with an encoded dynamic temporal pattern is used to drive “smart bulbs” or another source of light and/or sound while the second person is viewing the representation of the scenic view. The result is an enhanced emotional response and a deeper immersive experience. See FIG. **1**.

Example 2

(163) We record EEG of an actor (or actress) while the actor (or actress) is playing a particular role in a film or theatrical production; we then decode the temporal patterns of the EEG and encode them in software. If another person wants to experience enhanced emotional state while watch the same film or a recording of the theatrical production, the software with encoded temporal pattern is used to drive smart bulbs or another source of light and/or sound while the second person is watching the same film or a recording of the theatrical production. The result is an enhanced emotional response and a deeper immersive experience.

Example 3

(164) We record EEG of a first person (source) experiencing an emotional arousal while engaged in an activity (playing a game, sports, etc.); then decode the dynamic spatial and/or temporal patterns of the EEG and encode them in software coupled with the virtual reality representation of the activity. If a second person (recipient) wants to experience the same emotional arousal while viewing the virtual reality representation of the activity, the software with an encoded dynamic temporal pattern is used to drive a current used in transcranial electric or magnetic brain stimulation. The result is an enhanced emotional response and a deeper immersive experience.

Example 4

(165) A person is reading a book, and during the course of the reading, brain activity, including electrical or magnetic activity, and optionally other measurements, is acquired. The data is processed to determine the frequency and phase, and dynamic changes of brainwave activity, as well as the spatial location of emission. Based on a brain model, a set of non-invasive stimuli, which may include any and all senses, magnetic nerve or brain stimulation, ultrasound, etc., is devised for a subject who is to read the same book. The set of non-invasive stimuli includes not only content-based components, but also emotional response components. The subject is provided with the book to read, and the stimuli are presented to the subject synchronized with the progress through the book. Typically, the book is presented to the subject through an electronic reader device, such as a computer or computing pad, to assist in synchronization. The same electronic reader device may produce the temporal pattern of stimulation across the various stimulus modalities. The result is that the subject will be guided to the same emotional states as the source of the target brain patterns.

(166) In this description, several preferred embodiments were discussed. Persons skilled in the art will, undoubtedly, have other ideas as to how the systems and methods described herein may be used. It is understood that this broad invention is not limited to the embodiments discussed herein.

Rather, the invention is limited only by the following claims.

(167) The aspects of the invention are intended to be separable and may be implemented in combination, sub-combination, and with various permutations of embodiments. Therefore, the various disclosure herein, including that which is represented by acknowledged prior art, may be combined, sub-combined and permuted in accordance with the teachings hereof, without departing from the spirit and scope of the invention.

(168) All references and information sources cited herein are expressly incorporated herein by reference in their entirety.

Claims

1. A method of neurostimulation, comprising: receiving a plurality of signals, each respective signal representing an encephalographically-derived brainwave pattern of a donor associated with a respective time-varying brain state of the donor; storing the plurality of the signals in conjunction with the respective time-varying brain state of the donor for each respective signal, over a range of the respective time-varying brain state of the donor; analyzing the plurality of signals according to an eigenvector-based multivariate analysis, to define a reduced dimensionality time-varying brain state-dependent pattern associated with the respective time-varying brain state of the donor; defining a neurostimulation pattern for a selected time-varying brain state of the donor dependent on a deconvolution of the reduced dimensionality pattern for the selected time-varying brain state of the donor; and stimulating a recipient with the defined neurostimulation pattern to entrain the recipient's brain with the neurostimulation pattern.

2. The method of claim 1, wherein the respective time-varying brain state of the donor comprises an emotional state, further comprising determining the emotional state of the donor with a biometric sensor.

3. The method of claim 2, further comprising determining an electroencephalographic pattern of the recipient's brain, and controlling a phase of the neurostimulation pattern dependent on the determined electroencephalographic pattern of the recipient's brain.

4. A method defining a neurostimulation pattern for a recipient dependent on an encephalographically derived brainwave pattern of a donor, comprising: analyzing, with at least one automated processor, a time varying brain state-dependent brainwave pattern of the donor using an eigenvector-based multivariate analysis, to determine a reduced dimensionality brainwave pattern; defining, with the at least one automated processor, the neurostimulation pattern for the recipient dependent on a deconvolution of the reduced dimensionality brainwave pattern; and stimulating the recipient with the neurostimulation pattern using a central nervous system stimulator or peripheral nervous system stimulator, wherein the neurostimulation pattern is selectively dependent on the time varying brain state-dependent brainwave pattern of the donor.

5. The method of claim 4, wherein the analyzing comprises performing at least one of a principal component analysis (PCA), an independent component analysis (ICA), a factor analysis (FA), a Fourier Transform, and a Laplace Transform.

6. The method of claim 4, wherein defining comprises adapting the neurostimulation pattern dependent on a frequency and a phase of a brainwave pattern of the recipient.

7. The method of claim 4, wherein the defining comprises adapting the neurostimulation pattern dependent on a computational model of the recipient's brain.

8. A method neurostimulation, comprising: recording a brainwave pattern derived from an encephalographic recorder, associated with an emotional state of a donor, and storing the recorded brainwave pattern having a first dimensionality and associated emotional state in a memory; automatically

analyzing the stored brainwave pattern associated with the emotional state of the donor using a multivariate analysis in a multivariate analysis process, to determine a reduced dimensionality pattern having a second dimensionality, the second dimensionality being less than the first dimensionality; automatically defining a neurostimulation pattern dependent on the reduced dimensionality pattern; and stimulating a target subject with a neurostimulator according to the neurostimulation pattern.

9. The method according to claim 8, wherein the multivariate analysis comprises an eigenvector-based multivariate analysis.

10. The method according to claim 8, wherein the eigenvector-based multivariate analysis is selected from the group consisting of a principal component analysis (PCA), an independent component analysis (ICA), a factor analysis (FA), a Fourier Transform, and a Laplace Transform.

11. The method according to claim 8, wherein the neurostimulation pattern is defined in a blind deconvolution process of the reduced dimensionality pattern.

12. The method according to claim 8, further comprising producing the reduced dimensionality pattern using a Local Linear Eigenmaps (LLE) technique to approximate a low-dimensional manifold in a high-dimensional space having a dimensionality in excess of the second dimensionality, wherein the target subject is a human subject.

13. The method according to claim 8, further comprising producing the reduced dimensionality pattern using a nonlinear dimensionality reduction technique.

14. The method according to claim 8, further comprising reading an electroencephalographic pattern of the target subject, and synchronizing the neurostimulation pattern with the read electroencephalographic pattern.

15. The method according to claim 8, wherein the memory comprises a database, the method further comprising storing a plurality of different brainwave patterns, each associated with a respective different emotional state of a donor in the database.

16. The method according to claim 8, further determining the emotional state of the donor with a biometric sensor.

17. The method according to claim 8, further comprising entraining a brain of the target subject with the neurostimulation pattern using a neurostimulator.

18. The method according to claim 17, further comprising producing a visual output with the neurostimulator.

19. The method according to claim 17, further comprising producing an auditory output with the neurostimulator.

20. The method according to claim 17, further comprising producing a haptic or vibrational output with the neurostimulator.

21. The method according to claim 8, further comprising modelling a brain of the target subject, and adapting the neurostimulation pattern in real time in accordance with the model of the brain of the target subject.

22. The method according to claim 8, further comprising controlling the stimulation of the target subject with electroencephalographic waveform feedback from a brain of the target subject to adaptively optimize a waveform of the stimulation.

23. The method according to claim 8, further comprising producing by the neurostimulator, an auditory stimulus comprising binaural beats, which entrain brainwaves of a brain of the target subject.
