ARTICLE IN PRESS

YNIMG-12503; No. of pages: 14; 4C: 3, 5, 8, 9, 10

NeuroImage xxx (2015) xxx-xxx



Contents lists available at ScienceDirect

NeuroImage

journal homepage: www.elsevier.com/locate/ynimg



The functional significance of EEG microstates—Associations with modalities of thinking

- P. Milz ^{a,*}, P.L. Faber ^a, D. Lehmann ^{a,1}, T. Koenig ^b, K. Kochi ^a, R.D. Pascual-Marqui ^a
- a The KEY Institute for Brain-Mind Research, Department of Psychiatry, Psychotherapy and Psychosomatics, University Hospital of Psychiatry, P.O. Box 1931, CH-8032 Zurich, Switzerland
- ^b Department of Psychiatric Neurophysiology, University Hospital of Psychiatry, University of Bern, Bern, Switzerland

6 ARTICLE INFO

Article history:

5

- Received 4 March 2015
- Accepted 10 August 2015
- 10 Available online xxxx
- 11 Keywords:
- 12 Cognitive style
- 13 eLORETA
- 14 Object
- 15 Spatial
- 16 Verbal
- 17 Visual

ABSTRACT

The momentary, global functional state of the brain is reflected by its electric field configuration. Cluster analytical approaches consistently extracted four head-surface brain electric field configurations that optimally explain 19 the variance of their changes across time in spontaneous EEG recordings. These four configurations are referred 20 to as EEG microstate classes A, B, C, and D and have been associated with verbal/phonological, visual, attention 21 reorientation, and subjective interoceptive-autonomic processing, respectively. The present study tested these 22 associations via an intra-individual and inter-individual analysis approach. The intra-individual approach tested 23 the effect of task-induced increased modality-specific processing on EEG microstate parameters. The inter- 24 individual approach tested the effect of personal modality-specific parameters on EEG microstate parameters. 25 We obtained multichannel EEG from 61 healthy, right-handed, male students during four eyes-closed conditions: 26 object-visualization, spatial-visualization, verbalization (6 runs each), and resting (7 runs). After each run, we 27 assessed participants' degrees of object-visual, spatial-visual, and verbal thinking using subjective reports. Before 28 and after the recording, we assessed modality-specific cognitive abilities and styles using nine cognitive tests and 29 two questionnaires. The EEG of all participants, conditions, and runs was clustered into four classes of EEG micro- 30states (A, B, C, and D). RMANOVAs, ANOVAs and post-hoc paired t-tests compared microstate parameters be- 31 tween conditions. TANOVAs compared microstate class topographies between conditions. Differences were 32 localized using eLORETA. Pearson correlations assessed interrelationships between personal modality-specific 33 parameters and EEG microstate parameters during no-task resting.

As hypothesized, verbal as opposed to visual conditions consistently affected the duration, occurrence, and coverage of microstate classes A and B. Contrary to associations suggested by previous reports, parameters were increased for class A during visualization, and class B during verbalization. In line with previous reports, microstate 37 D parameters were increased during no-task resting compared to the three internal, goal-directed tasks. Topographic differences between conditions concerned particular sub-regions of components of the metabolic default 39 mode network. Modality-specific personal parameters did not consistently correlate with microstate parameters 40 except verbal cognitive style which correlated negatively with microstate class A duration and positively with 41 class C occurrence.

This is the first study that aimed to induce EEG microstate class parameter changes based on their hypothesized 43 functional significance. Beyond, the associations of microstate classes A and B with visual and verbal processing, 44 respectively and microstate class D with interoceptive-autonomic processing, our results suggest that a finely-45 tuned interplay between all four EEG microstate classes is necessary for the continuous formation of visual and 46 verbal thoughts, as well as interoceptive-autonomic processing. Our results point to the possibility that the 47 EEG microstate classes may represent the head-surface measured activity of intra-cortical sources primarily 48 exhibiting inhibitory functions. However, additional studies are needed to verify and elaborate on this 49 hypothesis.

© 2015 Elsevier Inc. All rights reserved. 51

59

* Corresponding author at: The KEY Institute for Brain-Mind Research, University Hospital of Psychiatry, P.O. Box 1931, CH-8032 Zurich, Switzerland.

E-mail addresses: patricia.milz@key.uzh.ch (P. Milz), pfaber@key.uzh.ch (P.L. Faber), dlehmann@key.uzh.ch (D. Lehmann), thomas.koenig@puk.unibe.ch (T. Koenig), osakachochiku@gmail.com (K. Kochi), pascualm@key.uzh.ch (R.D. Pascual-Marqui).

¹ Deceased.

Introduction 56

The momentary, global functional state of the brain is reflected by its 57 electric field configuration. The time-course of brain electric field config-58 urations can be analyzed with high resolution, "millisecond by millisec-59 ond" using head-surface EEG recordings (Koenig et al., 2002). Inspecting 60 these time-courses reveals that these electric field configurations 61

http://dx.doi.org/10.1016/j.neuroimage.2015.08.023 1053-8119/© 2015 Elsevier Inc. All rights reserved.

63

64 65

66

67

68 69

70

71

72

73

74

75

76 77

78

79

80

81

82

83

84

85

86

87

90 91

92

93

94

95

96

97 98

gg

100

101

102 103

104 105

106

107 108

109

110

111

113

114

115

116

117 118

119

120

121

122

123

124

125

126

change discontinuously, i.e. particular configurations remain quasistable for a duration of approximately 100 ms and then transition abruptly to another configuration (Lehmann et al., 1987, 2009; Michel et al., 2009). Cluster analytical approaches consistently extracted four head-surface brain electric field configurations that explain approximately 80% of the variance of these changes across time in spontaneous EEG recordings (Koenig et al., 2002; Wackermann et al., 1993). These four states are referred to as EEG microstate classes and are conventionally labeled from A through D (Koenig et al., 2002). The particular instances of these four states as quasi-stable configurations observed over the time-course of brain electric field configuration changes are referred to as "brain electric microstates" (Lehmann, 1990). The previous literature suggests that these brain electric microstates qualify as basic building blocks of mentation and may be considered candidates for conscious or non-conscious "atoms of thought and emotion" (Lehmann, 1990; Lehmann et al., 2004, 2005; Lehmann and Michel, 2011; Strik et al 1998)

Changes in topography, and other parameters such as mean duration, occurrence, coverage, and syntax (microstate sequence) of the four EEG microstate classes have been reported across stages of development (Koenig et al., 2002), in psychopathology compared to health (disorders of consciousness: Fingelkurts et al., 2012; schizophrenia: Andreou et al., 2014; Kikuchi et al., 2007; Kindler et al., 2011; Koenig et al., 1999; Lehmann et al., 2005; Nishida et al., 2013; Strelets et al., 2003; Tomescu et al., 2014, 2015; depression: Strik et al., 1995; dementia: Strik et al., 1997), in states of sleep (Brodbeck et al., 2012; Wehrle et al., 2007), in altered states such as hypnosis (Katayama et al., 2007) and meditation (Faber et al., 2005, 2014), and for particular personality profiles compared to others (Schlegel et al., 2012).

Despite the obvious need to understand the functional significance of these four EEG microstate classes to interpret the above changes, the functional significance of these four EEG microstate classes was only directly targeted in very few studies.

Britz et al. (2010) carried out a simultaneous EEG-fMRI study where the four EEG microstate classes were spatially correlated with four of the fMRI resting-state networks that had previously been attributed to phonological processing (microstate class A), visual imagery (microstate class B), attention reorientation (microstate class C), and subjective interoceptive-autonomic processing (microstate class D). Associations of EEG microstates with visual as opposed to non-visual processing are also supported by reports that showed different microstate topographies for visual as opposed to abstract thoughts (Koenig et al., 1998; Lehmann et al., 1998, 2010).

However, a more recent study suggested that the four EEG microstate classes may reflect the temporally distinct electrophysiological activation of four main components of the default mode network (Pascual-Marqui et al., 2014). The correspondence of the intra-cortical sources of the four EEG microstate classes with the four main components of the default mode network suggests that the metabolic DMN as measured with fMRI may reflect a time averaged version of distinct electrical networks which only become identifiable with the increased time resolution available in EEG recordings. An increase of metabolic DMN activity is associated with no-task resting, as well as a wide range of internal, goal-directed tasks (Andrews-Hanna, 2012). Possibly, the increased time resolution of the EEG will allow us to obtain a more detailed account of the roles of the four temporally distinct electrophysiological DMN components by assessing the effects of internal, goaldirected tasks as compared to no-task resting on the four EEG microstate classes.

The present study used four conditions, three internal, goaldirected tasks (object-visualization, spatial-visualization, verbalization) and no-task resting, to induce increased visual, verbal, and subjective interoceptive-autonomic processing, respectively. Subjective reports after conditions assessed participants' degree of object-visual, spatial-visual, and verbal thinking; cognitive tests assessed participants' object-visual, spatial-visual, and verbal abilities; and cognitive style

questionnaires assessed participants' object-visual, spatial-visual, and 128 verbal cognitive style. An intra-individual analysis approach tested the 129 success of the induction of the particular thinking modality, and the effects of the four conditions on the occurrence, duration, coverage, and 131 topography of the four EEG microstate classes. An inter-individual anal- 132 ysis approach correlated EEG microstate parameters during resting 133 with the subjective degree of modality-specific thinking, as well as 134 modality-specific cognitive abilities and style.

Firstly, we expected intra-individual differences between conditions 136 in the degree of subjectively reported thinking modality and EEG 137 microstate properties (occurrence, duration, coverage, and topogra- 138 phy). Secondly, we expected inter-individual differences reflected by 139 correlations of the degree of subjectively reported thinking modality, 140 modality-specific cognitive abilities, and style with EEG microstate pa- 141 rameters (occurrence, duration, coverage) during resting. Thirdly, we 142 expected correspondences between these intra- and inter-individual 143 differences.

Methods 145

Participants 146

Seventy healthy right-handed male volunteers from the University 147 of Zurich and the Swiss Federal Institute of Technology Zurich were 148 invited via flyers and through the Participant Server of the Institute of 149 Psychology of the University of Zurich. Applicants were screened with 150 an online questionnaire and excluded when they reported a history of 151 head trauma, brain disease, or current drug usage. The first 70 qualifying 152 men were invited to participate in the EEG experiment. Only male 153 participants were considered because of the effects of the menstrual 154 cycle and hormonal contraceptives on EEG in women (Becker et al., 155 1982; Krug et al., 1999). Each participant received 40 Swiss Francs for 156 their participation. The Ethics Committee of the Canton of Zurich ap- 157 proved the experimental protocol (reference number: KEK-ZH-Nr. 158 2011-0278). All participants gave their written informed consent. 159 After EEG pre-processing, EEG data from 61 participants was available 160 (mean age 24.2 years, SD = 3.3, range = 18-34). Behavioral data 161 were available for 60 of these 61 participants. All of these participants 162 were right-handed (German version of the Edinburgh handedness in- 163 ventory by Oldfield, 1971: mean = 4.48, SD = 0.38 where 1 indicates 164 left-handedness and 5 right-handedness). The reasons for participant 165 rejection are listed in the data pre-processing section.

Questionnaires 167

166

Modality of Thinking Questionnaire (MOTQ)

The MOTQ (Milz et al., submitted for publication) is a self-rating 169 questionnaire based on the object-spatial-verbal cognitive style model 170 (Blazhenkova and Kozhevnikov, 2009). It assesses modality-specific 171 cognitive style in three dimensions: object-visualization, spatial- 172 visualization, and verbalization. It comprises 36 items: 12 items to assess each style dimension. Each item asks respondents to rate their 174 degree of agreement with a modality-specific statement on a 5-point 175 Likert-scale ranging from 1 (complete disagreement) to 5 (absolute 176 agreement). Participants' scores were obtained separately for each 177 dimension by computing the mean of the items of the respective scale. 178 The reliabilities of the three scales are high (internal consistency: 179 0.81-0.86; re-test reliability: 0.85-0.87; Milz et al., submitted for 180 publication).

Cognitive tests 182

Object-visual ability

To assess object-visual ability, we used the Snowy Pictures Test Part 184 1 and 2 (Ekstrom et al., 1976), and the Incomplete Pictures Test (Horn, 185 1962). The former two ask participants to identify objects hidden in 186

an image of short unconnected lines (3 min time for twelve images each). The latter asks participants to identify the first letter of the name of small images of incomplete drawings of objects from a selection of five letters (one minute time for 40 images). Each correctly identified solution was added up to retrieve a sum score for each of the three

Spatial-visual ability

187

188

189 190

191

192

193 194

195 196

197

198

199

200

201

202

203

204

205

206 207

208

209

210

211 212

213

214

215 216

217

218

219

220

221

222

223

224

225

226

227

228 229

230

231 232

To assess spatial-visual ability, we used the Paper Folding Test Part 1 and 2 (PFT: Ekstrom et al., 1976), and the "Cube Test" (Wuerfelabwicklungen: Meili, 1955). The former ask participants to identify an unfolded paper which had previously been folded in a series of steps and then punched, by its holes from a selection of five solutions (3 min time for ten images each). The latter asks participants to label top and connecting edges of unfolded cubes by imagining what they would look like if folded (10 min time for eleven images). Each correctly identified solution was added up to retrieve a sum score for each of the three tests.

Verbal ability

To assess verbal ability, we used the Masselon, Part-Whole, and the Insight Test from the verbal dimension of the Berlin Intelligence Structure Test (Jäger et al., 1997). The Masselon Test asks participants to write down as many sentences possible containing three defined nouns (two min time), the Part-Whole Test asks participants to identify wholes if they are followed by their part (e.g. "year" if followed by "month") in four columns of nouns (40 s time). The Insight Test asks participants to write down as many keywords possible to explain a given statement (e.g. why to a person fashion may be important) (2 min time). Correct sentences for the Masselon Test, correctly identified wholes for the Part-Whole Test, and reasonable keywords for the Insight Test were added up to retrieve a sum score for each of the three tests.

Recording

EEG recordings were done at the KEY Brain Mapping Laboratory at the University Hospital of Psychiatry, Zurich, using a 64-channel BioSemi Recording System. Sixty-four electrodes were attached using a Biosemi Headcap at the following positions: Fp1/2, Fpz, AF7/8, AF3/ 4, AFz, F7/8, F5/6, F3/4, F1/2, Fz, FT7/8, FC5/6, FC3/4, FC1/2, FCz, T7/8, C5/6, C3/4, C1/2, Cz, TP7/8, CP5/6, CP3/4, CP1/2, CPz, P7/8, P5/6, P3/4, P1/2, Pz, PO7/8, PO3/4, POz, O1/2, Oz, P9/10, and Iz (according to the International "10-10 System", Chatrian et al., 1985; Nuwer, 1987). Eye movements were tracked using 2 additional electrodes, one above the left and one below the right eye. Breathing was recorded with a strain gauge chest band. Recordings were done with a sampling frequency of 2048 Hz.

Protocol

Prior to the experiment, participants completed an online version of the Modality of Thinking Questionnaire (MOTQ: Milz et al., submitted for publication) and the Vividness of Visual Imagery Questionnaire 233 (VVIQ: Marks, 1973). On arrival at the EEG laboratory, participants 234 were informed and gave their consent to their participation in the ex- 235 periment. Then, they completed the Edinburgh handedness test 236 (Oldfield, 1971) and a computerized version of the Object-Spatial Imag- 237 ery and Verbal Questionnaire (OSIVQ: Blazhenkova and Kozhevnikov, 238 2009, purchased via MM Virtual Design). During electrode attachment, 239 the experimental design and the upcoming tasks were thoroughly 240 explained and participants were informed that during each of the 241 upcoming recording runs they should sit relaxed with arms and legs 242 in a comfortable position. Participants were then seated in an electrical- 243 ly, acoustically and light-shielded chamber on an armchair, viewing a 244 computer monitor at 1 m. Each participant was recorded during four 245 eyes-closed conditions: no-task resting (7 recording runs), object- 246 visualization, spatial-visualization, and verbalization (6 recording runs 247 each). Each recording run lasted 50 s and conditions were presented 248 in pseudo-randomized order (no-task resting always initiated a permu- 249 tation of the other three tasks, and ended the recording).

Instructions were again presented on the monitor prior to each run 251 using Presentation® (www.neurobs.com). Before no-task resting runs, 252 participants were asked to just relax and close their eyes until a tone 253 would indicate the end of that run. Before visualization runs, partici- 254 pants were asked to memorize an upcoming image and to then visualize 255 it internally with their eyes closed for the whole period of the 50 s re- 256 cording. The image was visible for 7 s. For the six object-visual runs, 257 images were six photos containing many visual details such as colors, 258 textures, and shapes (objects were foods: apple bowl, bread loaf; furni- 259 ture: bed, lamp; and animals: deer, tigers). For the six spatial-visual 260 runs, images were six spatial configurations of black circles. Before ver-261 balization runs, participants were asked to read an upcoming noun to 262 then think of a verbal definition for that noun to internally explain its 263 meaning to an imaginary other person. The noun was visible for 2 s. 264 For the six verbalization runs, six nouns of lowest visually imaginable 265 ranking according to Baschek et al. (1977) were displayed. Stimulus ex-266 amples are illustrated in Fig. 1.

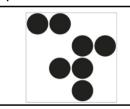
After each run, participants reported their degree of object-visual, 268 spatial-visual, and verbal thinking during task execution by specifying 269 their agreement to three corresponding statements (for object-visual: 270 "My thinking was comprised of concrete, detailed, realistic visual 271 images."; for spatial-visual: "My thinking was comprised of images 272 depicting spatial relations between objects."; for verbal: "My thinking 273 was very verbal in nature.") on a continuous rating scale ranging from 274 0 (not at all) to 100 (very much so). Moreover, for object-visual tasks, 275 they answered a three-option multiple choice question regarding the 276 photography they had perceived (e.g. "How were the dots on the deer 277 aligned?"). For the spatial-visual tasks, they chose the spatial configura- 278 tion they had just visualized from three possible configurations (only 279 one of which was correct). For the verbalization task, participants en- 280 tered their definition within 30 s using a keyboard on a table in front 281 of them.

It was emphasized to participants that their success in answering 283 these questions and the quality of their definition was of minor 284

Object-Visualization



Spatial-Visualization



Verbalization

Define: Familiarization"

Fig. 1. Example stimuli for the three tasks: object-visualization, spatial-visualization, and verbalization. Image source for the object-visualization example: http://www.public-domainimage.com/fauna-animals-public-domain-images-pictures/deers-public-domain-images-pictures/white-tailed-deer-pictures/white-tailed-deer-fawn-in-grass-odocoileus-virginianus.ipg.html

Please cite this article as: Milz, P., et al., The functional significance of EEG microstates—Associations with modalities of thinking, NeuroImage (2015), http://dx.doi.org/10.1016/j.neuroimage.2015.08.023

287

289

290

291

292 293

294

295

296

297

298

299

300

301 302

303

304 305

306

307

308

311 312

313

314

315

316

317

318 319

320

321

322

323 324

325

326

327

328 329

330

331

332

333

334

335

336

337

338

339

340

341

342

343

344

345

importance. Instead, their main focus should be to visualize as well as possible during the visualization tasks and to verbalize as well as possible during the verbalization task. Regarding their answer to the degree of object-visual, spatial-visual, and verbal thinking during tasks, they were told that it would be best for the experiment if they answered as truthfully as possible even if their thinking may not have been in line with what they had been asked to do. After the EEG recording, participants' abilities in the three thinking modalities were assessed using three cognitive tasks each for object-, spatial- visual, and verbal ability. Two weeks after the experiment, participants completed the MOTQ and OSIVQ a second time.

The answers to multiple choice questions after the visualization runs and the definitions written after the verbalization runs clearly indicated that participants had successfully remembered the images they had perceived (88.4 \pm 9.35% correct answers) and were able to form a definition within the given time period. Of the 61 participants, 59 were able to form a coherent definition (sometimes simply not yet complete) across the six runs as assessed by two independent raters.

Data pre-processing

EEG data were pre-processed using Brain Vision Analyzer 2 (Brain Products, Munich, Germany). Eve movement artifacts were removed using the software's ocular correction ICA. Sweat, muscle, movement, and electrode artifacts were discarded through visual inspection. Channels with severe artifacts across the whole recording run (mean number of channels across participants = 6.7, SD = 3.8) were interpolated. Interpolations were done separately for each affected run. All channels directly surrounding the affected channel were used for interpolation unless they were contaminated with severe artifacts themselves. Three channels (P9, P10, and Iz) were discarded for the analysis because they were spatial outliers relative to the other 61 electrodes that cover the scalp in an approximate uniformly distributed manner (see also: Pascual-Marqui et al., 2014). The data was then segmented into 2-s epochs, down-sampled to 256 Hz, re-referenced to average reference, and FFT filtered from 2-20 Hz (no windowing). After data conditioning, the mean number of available data epochs per run (average across the 25 runs: 6 runs each for object-visualization, spatial-visualization, and verbalization, 7 runs for resting) ranged between 3.36 and 21.76 (S.D. between 2.48 and 6.56) across participants. On average, 12.88 epochs (SD = 4.14) were available for each participant in each run. The EEG of five participants had to be rejected because of strong eye rolling artifacts across the whole recording which could not be satisfactorily removed with ocular correction ICA approaches. Another four participants had to be rejected because of severe local frontal and temporal muscular artifacts across the whole recording which could not be satisfactorily compensated for with interpolation procedures since they affected up to eight surrounding channels. Due to technical problems with the Presentation software, behavioral data for one participant were not available.

Microstate analysis

Microstate class computation

The microstate class computations were done following the procedure described in (Koenig et al., 1999) using the implementation in the microstate package of the KEY EEG Python library (Milz, in preparation). This open source library provides EEG pre-processing and analysis routines implemented in the Python programming language. For each participant, each condition, and each run, the EEG map topographies at time points of global field power peaks (GFPL1 Norm: Lehmann and Skrandies, 1980) were collected and fed into a modified k-means clustering algorithm (see also Pascual-Marqui et al., 1995) to deduce the four classes of map topographies that maximally explain the variance of the map topographies (see also Koenig et al., 1999; Koenig et al., 2002). These four classes of map topographies were then used to compute mean classes across runs, which were then 347 used to compute mean classes across participants, which were finally 348 used to compute mean classes across conditions. The following descrip- 349 tion depicts how the mean classes across runs were computed. Howev- 350 er, we applied the same procedure to compute mean classes across 351 participants, and across conditions, thereafter. Mean classes across 352 runs were computed by using a full permutation procedure that deter- 353 mines the solution of maximal mean correlation across the four classes 354 across all runs. The means across runs were computed as the first principle component of all maps that were assigned to each other in this 356 solution.

357

358

381

382

Microstate parameter computation

Using the mean microstate classes across conditions as templates, for 359 all participants, all conditions, all runs, and each two-second epoch, the 360 EEG map topographies at global field power peaks were assigned to one 361 of these 4 microstate classes based on maximal Pearson correlation. Successive GFP peak maps (maps) assigned to the same class were recog- 363 nized as belonging to one microstate. The duration of a microstate was 364 computed as the time period during which maps were successively 365 assigned to the same class, starting at the midpoint between the last 366 map of the preceding microstate and the first map of the following microstate. Based on a given 2-s epoch, we cannot reconstruct when the 368 first microstate began (previous epoch) and when the last microstate 369 ended (next epoch). Therefore, their duration cannot conclusively be 370 determined. For this reason, the first and last microstate of each 2-s 371 epoch were ignored for the following parameter computations.

Three microstate parameters were computed: mean duration (aver- 373 age time in ms covered by a given microstate class), occurrence (mean 374 number of distinct microstates of a given class occurring within a 1 s 375 window), and coverage (percentage of time covered by a given micro- 376 state class). Computations were done for each participant, each condition, each run, and each microstate class by averaging the respective 378 parameter values across all 2-s epochs of a given run (see also Strelets 379 et al., 2003).

Analysis methods

Intra-individual analysis

Where not otherwise specified, statistics were computed using SPSS Version 22.0 (IBM Corporation, 2014).

Success of modality induction. Subjective reports on the degree of object- 385 visual, spatial-visual, and verbal thinking during the four conditions 386 were averaged across repeated runs. A repeated measures MANOVA 387 (independent variable: condition, dependent variable: degree of ob- 388 ject-visual thinking, spatial-visual thinking, and verbal thinking), post- 389 hoc ANOVAs, and paired t-tests tested whether the three tasks induced 390 thinking in the attempted modality.

Mauchly's Test of Sphericity indicated that the assumption of sphe- 392 ricity had been violated for object-visual thinking ($\chi^2(5) = 24.65$, 393 p = 0.0001), spatial-visual thinking ($\chi^2(5) = 19.47$, p = 0.0016), and 394 verbal thinking ($\chi^2(5) = 20.24$, p = 0.0011). Therefore, results are reported with Greenhouse-Geisser correction.

Task effects on microstate parameters. Microstate duration, occurrence, 397 and coverage during the four conditions were averaged across repeated 398 runs. For each microstate parameter (duration, occurrence, and cover- 399 age), a repeated measures MANOVA (independent variable: condition, 400 dependent variable: parameter values of classes A, B, C, D), post-hoc 401 ANOVAs and paired t-tests tested whether the three tasks affected the 402 three microstate parameters.

Mauchly's Test of Sphericity indicated that the assumption of 404 sphericity had been violated for duration, occurrence, and coverage for 405 all four EEG microstate classes ($\chi^2(5) > 11.91, p < 0.05$). Therefore, re- 406 sults are reported with Greenhouse-Geisser correction. 407

t1.1 t1.2

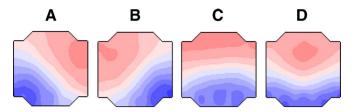


Fig. 2. The topographies of the four microstate classes retrieved from the clustering algorithm, sorted and labeled according to the normative classes described in Koenig et al. (2002). Note that only the map's topography is important, whereas polarity is disregarded in the spontaneous EEG clustering algorithm.

Task effects on microstate topographies. The EEG map topographies of the four classes for the four conditions were tested for global differences using TANOVA (Strik et al., 1998) implemented in RAGU (Koenig et al., 2011). TANOVA computes the Global Map Dissimilarity (Lehmann and Skrandies, 1980) between two map topographies and the probability of the observed differences using a randomization procedure. In a first step, a two-factor TANOVA (independent variables: class, condition; dependent variable: map topography) was computed. In a second step, class-wise one-factor TANOVAs (independent variable: condition; dependent variable: map topography) were computed. Channel-wise paired *t*-tests specified the spatial distribution of observed differences. The intra-cortical sources of significant differences were localized using exact low resolution electromagnetic tomography (eLORETA: Pascual-Marqui, 2007).

Inter-individual analysis

408

409

410

411

412

413

414

415

416 417

418

419

420

421

422

423

424

425

426

427

428

Correlations between microstate parameters across conditions. We computed Pearson correlations between the microstate parameters of the four conditions (object-visualization, spatial-visualization, verbalization, and no-task resting) for each condition pair (means across runs).

Correlations between microstate parameters and subjectively reported thinking modality. We computed Pearson correlations between the

Table 1
Multivariate effects of condition on self-reported degree of thinking in the three modali-
ties (object-visual, spatial-visual, and verbal).

Dependent variable	df	df error	F	p	Wilks' A	Partial η^2	t1.4
Degree of thinking	9	51	100.362	< 0.001	0.053	0.95	t1.5

duration, occurrence, and coverage of the four EEG microstate classes 429 and the reported degree of object-visual, spatial-visual, and verbal 430 thinking during resting (means across runs, and separately for each 431 resting run).

Correlations between microstate parameters and modality-specific ability. 433
We computed Pearson correlations between modality-specific abilities 434
and the duration, occurrence, and coverage of the four EEG microstate 435
classes during resting (means across runs). Correlations were computed 436
separately for an object-visual, spatial-visual, and verbal ability factor 437
derived from a principal component analysis of the nine cognitive 438
tests (see Inline Supplementary Table S1).

Inline Supplementary Table S1 can be found online at http://dx.doi. 440 org/10.1016/j.neuroimage.2015.08.023. 441

Correlations between microstate parameters and modality-specific style. Q3
We computed Pearson correlations between modality-specific cogni- 443
tive style and the duration, occurrence, and coverage of the four EEG mi- 444
crostate classes during resting (means across runs). Correlations were 445
computed separately for object-visual, spatial-visual, and verbal style 446
based on the MOTQ (means across the two assessment time points). 447

Results 448

Microstate classes 449

Across participants, the topographies of the individual four micro- 450 state classes explained $84.0\pm4.8\%$ of the total variance of all GFP 451 peaks, and $77.0\pm5.1\%$ of the variance of all EEG time frames. The four 452 mean microstate classes across conditions are illustrated in Fig. 2. The 453

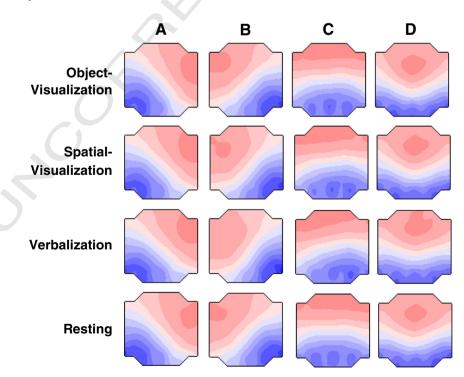


Fig. 3. The four microstate classes retrieved from the clustering algorithm for the four conditions (object-visualization, spatial-visualization, verbalization, and no-task resting) separately. They were sorted and labeled according to the normative classes described in Koenig et al. (2002).

Please cite this article as: Milz, P., et al., The functional significance of EEG microstates—Associations with modalities of thinking, NeuroImage (2015), http://dx.doi.org/10.1016/j.neuroimage.2015.08.023

t2.2

t2.3

t2.6 t2.7

454

455

456

457

458

459

460

461

462

463 464

465

466

467

468 469

470

471

472

473

474

Table 2 Univariate effects of condition on self-reported degree of object-visual, spatial-visual, and verbal thinking (values based on Greenhouse-Geisser correction).

Dependent variable	df	df error	F	p	Partial η ²
Object-visual thinking	2	136	87.922	< 0.001	0.60
Spatial-visual thinking	3	149	319.619	< 0.001	0.84
Verbal thinking	3	150	172.594	< 0.001	0.75

four mean microstate classes across participants, separately for the four conditions are illustrated in Fig. 3. The topographies were labeled as classes A, B, C, and D based on their best fit with the four normative classes described in Koenig et al. (2002).

Intra-individual analysis

Success of modality induction

Analyses of variance revealed a significant effect of condition on object-visual, spatial-visual, and verbal thinking (Tables 1, 2). The self-reported degrees of object-visual, spatial-visual, and verbal thinking during the four conditions are illustrated in Fig. 4. Post-hoc ttests revealed that during each task (object-visualization, spatialvisualization, and verbalization) participants were thinking more strongly in the modality that was attempted to be induced than in other modalities (Inline Supplementary Table S2). Moreover, they were thinking most strongly in the modality that was attempted to be induced in the task which attempted to induce it compared to the other tasks and also compared to resting (Inline Supplementary Table S3). This was true for all three tasks but most prominent for the verbalization task (see Fig. 4).

Inline Supplementary Tables S2 and S3 can be found online at http:// dx.doi.org/10.1016/j.neuroimage.2015.08.023.

Multivariate effects of condition on duration, occurrence, and coverage of the four microstate classes A. B. C. and D. t3.3

Dependent variable	df	df error	F	р	Wilks' Λ	Partial η^2	t3.4
Duration	12	49	2.734	0.007	0.599	0.40	t3.5
Occurrence	12	49	3.126	0.002	0.566	0.43	t3.6
Coverage	12	49	3.914	<0.001	0.511	0.49	t3.7

Task effects on EEG microstate parameters

Analyses of variance revealed a significant effect of condition on microstate duration, occurrence, and coverage for microstate classes A and 477 B (Tables 3 and 4).

475

483

Post-hoc paired t-tests revealed a similar pattern of differences be- 479 tween conditions for the three parameters (Inline Supplementary 480 Tables S4-S7). 481

Inline Supplementary Tables S4–S7 can be found online at http://dx. 482 doi.org/10.1016/j.neuroimage.2015.08.023.

Table 5 illustrates all significant differences (p < 0.05) and trends 484 (p < 0.10). For microstate class A, parameters were increased during 485 the visualization compared to the other conditions. For microstate 486 class B, parameters were increased during the verbalization compared 487 to the other conditions. Microstate class C was shorter during visualiza- 488 tion compared to resting. Microstate class D lasted longer during resting 489 compared to verbalization, occurred more often during resting com- 490 pared to visualization, and showed increased coverage compared to 491 both object-visualization and verbalization. There were no significant 492 differences between the two visualization tasks.

Fig. 5 illustrates the absolute magnitude of coverage (in percent) ob-494 served for the four microstate classes. We note lower mean coverage 495 (20–24%) for microstate classes A and B as compared to microstate 496 classes C and D (27–30%) during all conditions. The absolute magnitude 497 of significant (p < 0.05) coverage differences between conditions for 498

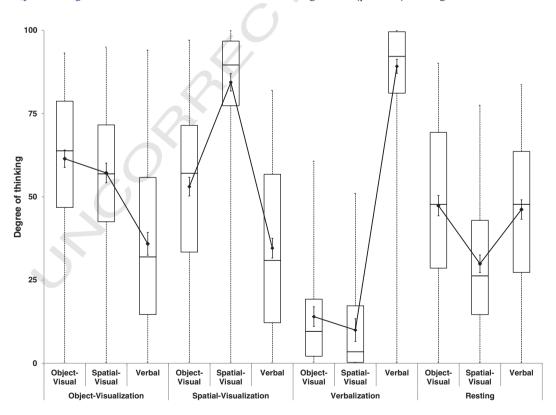


Fig. 4. Self-reported degree of object-visual, spatial-visual, and verbal thinking during the three tasks (object-visualization, spatial-visualization, and verbalization) and during no-task resting. The end of the lower whisker of the boxes represents the minima, the boxes lower bound the first quartiles, the midline the medians, the boxes upper bound the third quartiles, and the end of the upper whisker the maxima of the distributions of participants' degree of thinking modality, respectively. Distribution means are indicated by the small circles within the boxes, standard errors are indicated by the error bars starting from these circles. The means are connected with lines within each condition (N = 60).

t6.1

t6.2

±6.11

P. Milz et al. / NeuroImage xxx (2015) xxx-xxx

Table 4Univariate effects of condition on duration, occurrence, and coverage of the four microstate classes A, B, C, and D (Greenhouse–Geisser corrected values).

t4.2

t43

t5.1

t5.2

t5.4

t5.5

t5.6

t5.7

t5.8

t5.9

t5.10t5.11

t5.12

t5.13

Dependent variable		df	df error	F	p	Partial η ²
Duration	Α	2.5	151	4.164	0.011	0.06
	В	2.1	128	7.020	0.001	0.10
	C	1.7	102	1.923	0.157	0.03
	D	2.4	144	1.200	0.309	0.02
Occurrence	Α	2.7	160	7.510	< 0.001	0.11
	В	2.4	143	5.172	0.004	0.08
	C	2.2	134	0.103	0.920	0.00
	D	2.6	156	1.940	0.134	0.03
Coverage	Α	2.5	147	11.432	< 0.001	0.16
	В	2.1	125	10.518	< 0.001	0.15
	C	2.2	132	0.442	0.663	0.01
	D	2.3	139	2.092	0.120	0.03

Table 5Microstate parameter changes between the four conditions: resting (Re), object-visualization (Ov), spatial-visualization (Sv), and verbalization (Ve).

	Dur	Duration				Occurrence				Coverage				
Conditions	A	В	С	D	A	В	С	D	A	В	С	D		
$Ov \rightarrow Ve$	7	1			7	1			7	1				
$Sv \rightarrow Ve$ $Ov \rightarrow Sv$	1	1			7	1			7	1				
$Re \to Ov$	7		1		1			\	1			\		
$Re \to Sv$			1		1			/	1					
$Re \rightarrow Ve$		1		7		1			7	1		/		

Differences at p < 0.10 are depicted by arrows; arrows for p < 0.05 are in bold (no correction for multiple testing).

Table 6Paired TANOVA *p*-values for each condition pair.

Conditions			Microstat	Microstate classes							
			A	В	С	D					
Ov	-	Ve	0.048	< 0.001	< 0.001	< 0.001					
Sv	-	Ve	0.407	0.177	0.013	0.004					
Ov	-	Sv	0.202	0.015	0.125	0.482					
Re	-	Ov	0.145	0.593	0.165	0.096					
Re	-	Sv	0.485	0.098	0.004	0.100					
Re	-	Ve	0.415	0.003	< <u>0.001</u>	0.002					

Re: resting, Ov: object-visualization, Sv: spatial-visualization, Ve: verbalization. Significant differences (*p* < 0.05, no cmt) are bold, italic, and underlined.

microstate classes A and B was of moderate effect size (absolute effect 499 sizes ranging from 0.38 to 0.60, see also Inline Supplementary Table S7). 500

Task effects on EEG microstate topographies

The two-factor TANOVA revealed a significant effect of microstate class (p < 0.001) and the interaction between condition and microstate 503 class (p < 0.001) on microstate map topographies. Separate one-factor 504 TANOVAs for the four EEG microstate classes revealed a significant 505 effect of condition on microstate map topographies for Map B 506 (p < 0.001), Map C (p < 0.001), and Map D (p < 0.001). Post-hoc paired 507 TANOVAs revealed significant topographic differences between several 508 condition pairs (see Table 6). Topographic differences of these significant differences are illustrated in Fig. 6.

Multidimensional scaling (Koenig et al., 2011) of the maps for all 511 four conditions revealed that object- and spatial-visualization deviated 512 similarly from verbalization on the left-right and anterior-posterior 513 map topography for microstate classes C and D, respectively. Moreover, 514

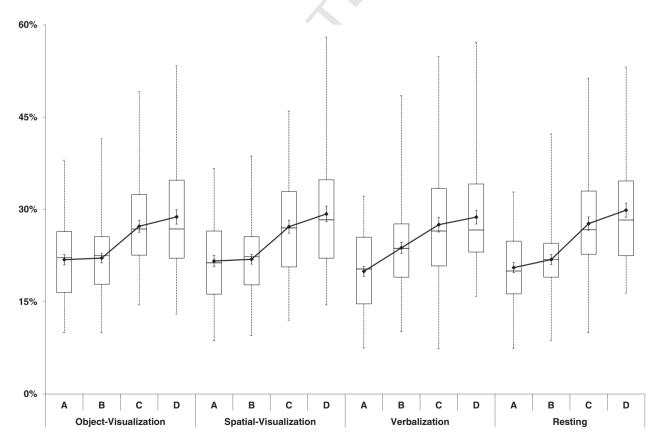


Fig. 5. Box-plots illustrating the coverage of the four microstate classes A, B, C, and D during the four conditions (object-visualization, spatial-visualization, verbalization, and no-task resting). The end of the lower whisker of the boxes represents the minima, the boxes lower bound the first quartiles, the midline the medians, the boxes upper bound the third quartiles, and the end of the upper whisker the maxima of the distributions of participants' degree of thinking modality, respectively. Distribution means are indicated by the small circles within the boxes, standard errors are indicated by the error bars starting from these circles. The means are connected with lines within each condition (N = 60).

Please cite this article as: Milz, P., et al., The functional significance of EEG microstates—Associations with modalities of thinking, NeuroImage (2015), http://dx.doi.org/10.1016/j.neuroimage.2015.08.023

516 517

518

520

521 522

523

524

525

526

527

528

529

530

O10

P. Milz et al. / NeuroImage xxx (2015) xxx-xxx

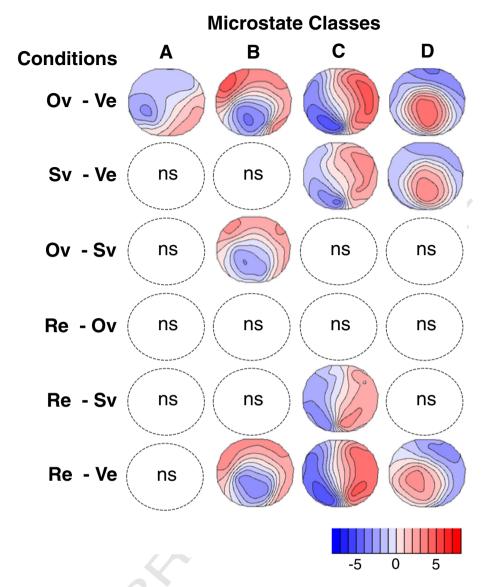


Fig. 6. Maps visualizing channel-wise paired *t*-test differences for condition pairs significantly different at *p* < 0.05 (two-tailed) for the four EEG microstate classes A, B, C, and D. Positive *t*-values are depicted in red, negative *t*-values in blue. For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.

the three tasks deviated similarly from no-task testing. Therefore, we computed the source of the mean difference between visualization (object- and spatial-visualization) and verbalization, and between tasks (object-, spatial-visualization and verbalization) and no-task resting using eLORETA (Pascual-Marqui, 2007) (Fig. 7).

During visualization compared to verbalization, source localizations revealed increased activity in left parietal areas for Map B (insula, BA 13) and Map C (temporal lobe, BA 39) and in midline posterior areas (posterior cingulate, BA23) for Map D. Increased activity during verbalization was observed in right occipito-parietal areas (posterior cingulate, BA31) for Map B, right occipito-parietal areas (insula, BA13) for Map C, and midline anterior areas (anterior cingulate, BA25) for Map D (see Fig. 6).

During tasks compared to no-task resting, source localizations revealed increased activity in left Broca's area (BA44) for Map B, the precuneus (BA7) for Map C, and the medial frontal gyrus (BA9) for

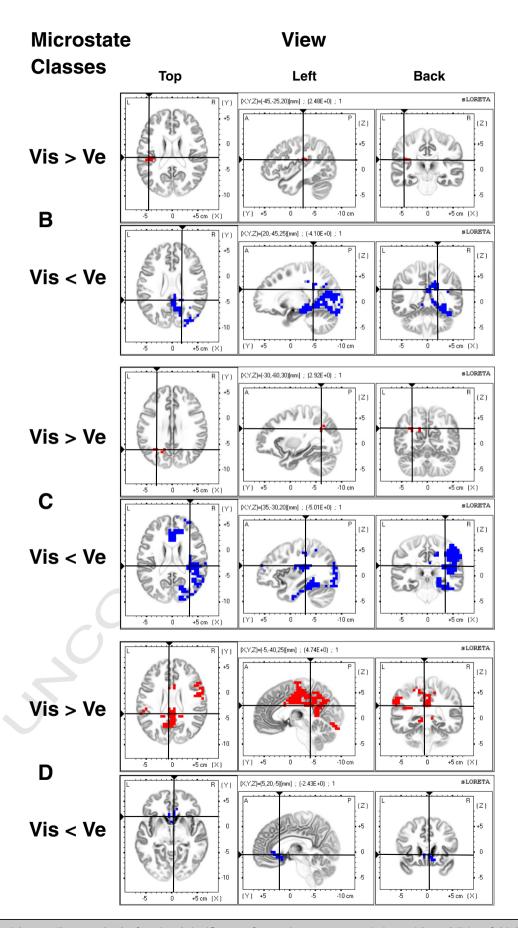
Map D. Increased activity during resting was observed in the right pri- 531 mary somatosensory cortex (BA3) for Map B, left visual association 532 areas (BA18) for Map C, and the left auditory association area (BA21) 533 for Map D (see Fig. 8).

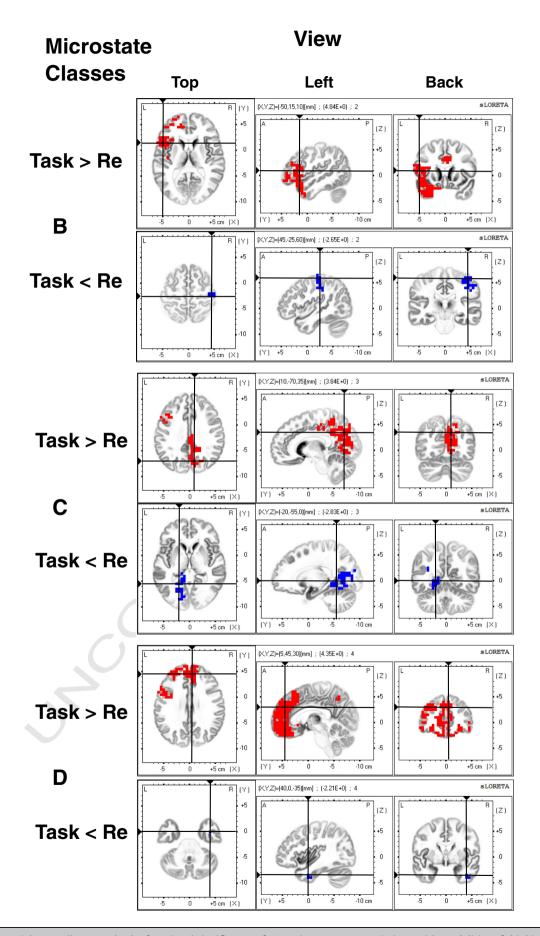
535

Inter-individual analysis

Pearson correlations between EEG microstate parameters across 536 conditions revealed strong associations (all r > 0.75) between duration, 537 occurrence, and coverage for all microstate classes and all condition 538 pairs (Inline Supplementary Table S8). These results suggest that the 539 relative duration, occurrence, and coverage of the four EEG microstate classes of the same participant compared to others are very similar during all conditions. A participant with a comparably long duration of EEG 542 microstate class A during resting, will also exhibit a comparably long 543 duration of microstate class A during tasks, and vice versa.

Fig. 7. eLORETA brain images of the statistical difference in cortical distribution of electric sources of the three microstate classes B, C, and D between visualization (Vis) and verbalization (Ve). Left to right: slices in axial (from top, nose up), sagittal (from left) and coronal (from back) view in MNI space. Images depict all t-values with p < 0.05 (two-tailed), t = 2.00 (df = 60). Red = stronger activity during visualization, blue = stronger activity during verbalization. Vis > Ve: slices through the voxel (indicated by arrowheads) of maximally stronger activity during verbalization. The MNI coordinates (X, Y, Z) of these voxels are reported in millimeters. For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.





t7.1 **Table 7**t7.2 Correlations (Pearson's r) between the microstate parameters (duration, occurrence, coverage) during no-task resting of the four EEG microstate classes (A, B, C, D) with three modality-t7.3 related person parameters: modality-specific ability (Ability), modality-specific style (Style), and self-reported thinking modality (Thinking).

t7.4			Modality-related personal parameters											
t7.5	Microstate parameters	Modalities	Ability	Ability			Style				Thinking			
t7.6			A	В	С	D	A	В	С	D	A	В	С	D
t7.7	Duration	Object-visual	0.04	-0.03	0.07	0.06	0.07	0.20	0.10	-0.12	0.01	-0.01	0.28	-0.21
t7.8		Spatial-visual	0.08	0.05	0.01	0.00	0.17	0.06	0.08	-0.13	0.11	0.14	0.25	0.04
t7.9		Verbal	-0.13	0.14	0.00	-0.25	-0.34	-0.22	-0.08	-0.09	-0.05	-0.18	-0.09	-0.02
t7.10	Occurrence	Object-visual	-0.14	-0.16	0.01	0.08	-0.11	0.07	-0.04	-0.12	-0.03	-0.11	0.21	-0.15
t7.11		Spatial-visual	-0.04	-0.01	-0.16	0.11	-0.02	-0.05	0.03	0.09	-0.11	-0.10	-0.12	-0.27
t7.12		Verbal	0.02	0.19	0.15	-0.10	0.02	0.08	0.28	0.24	0.04	0.04	0.15	0.04
t7.13	Coverage	Object-visual	-0.08	-0.13	0.07	0.09	-0.07	0.19	0.05	-0.13	-0.02	-0.07	0.32	-0.25
t7.14		Spatial-visual	0.02	0.01	-0.08	0.05	0.05	-0.01	0.05	-0.07	-0.04	0.01	0.11	-0.08
t7.15		Verbal	-0.02	0.24	0.10	-0.25	-0.14	-0.05	0.11	0.04	0.04	-0.07	0.04	-0.01

R-values of significant (p < 0.05 no correction for multiple testing) correlations are highlighted in bold, italic, and underlined.

Inline Supplementary Table S8 can be found online at http://dx.doi.org/10.1016/j.neuroimage.2015.08.023.

Pearson correlations between EEG microstate classes and behavioral variables (Table 7) inspected whether these relatively stable interindividual differences in the EEG microstate class parameter distributions might be associated with modality-related personal parameters.

However, only four of the 108 possible correlations between the three EEG microstate parameters and the three modality-related personal parameters were significant at p < 0.05 (not corrected for multiple testing). The first two significant correlations concerned object-visual thinking during resting. Object-visual thinking was positively associated with the duration and coverage of microstate class C. However, post-hoc run-wise analyses revealed that these significant correlations did not occur consistently across runs (Inline Supplementary Tables S9 and S10). The other two significant correlations concerned verbal cognitive style. Verbal style was negatively associated with the duration of microstate class A, and positively associated with the occurrence of microstate class C.

Inline Supplementary Tables S9 and S10 can be found online at http://dx.doi.org/10.1016/j.neuroimage.2015.08.023.

Discussion

t7.16

579

580

581

583 584

585

586

587

589

590

591

592 593

594

595

596

597

598

599

600 601

602

603

604

605

606

607

608 609

610

611 612

613 614

615

616

012

As hypothesized, EEG microstate parameters and topographies were affected by an individual's degree of visual and verbal thinking, and differed between internal, goal-directed tasks and no-task resting.

Intra-individual differences

EEG microstate parameters

Participants' subjective ratings suggest that our tasks successfully induced object-visual, spatial-visual, and verbal thinking using three corresponding tasks. EEG microstate parameter differences between the three tasks were compared to each other and to resting. These comparisons revealed an association of microstate class A with visualization and microstate class B with verbalization. Microstate class D was associated with no-task resting, possibly reflecting the increase of the degree of subjective interoceptive-autonomic processing particular to no-task resting compared to the three goal-directed tasks.

Our results are in line with the findings of Britz et al. (2010) with regard to microstate class D. In their combined fMRI EEG study, microstate class D was also associated with subjective interoceptive-autonomic processing, namely an fMRI resting-state network that had previously

been attributed to this processing mode. However, our associations of 618 microstate classes A and B at first sight appear to conflict with their find-619 ings. Whereas we found increased duration, occurrence, and coverage of 620 microstate class A during visualization, and of microstate class B during 621 verbalization, Britz et al. (2010) associated microstate class A with fMRI 622 resting-state networks that had previously been attributed to phono-623 logical processing, and microstate class B with fMRI resting-state networks previously attributed to visual imagery.

According to Pascual-Marqui et al. (2014), the four EEG microstate 626 classes are produced by a strong common generator in the posterior 627 cortex, with an additional generator in left occipital areas for micro- 628 state class A, right occipital areas for microstate class B, and anterior 629 cingulate areas for microstate class C. Again our results at first sight ap- 630 pear to conflict with this finding. Why would we find increased dura- 631 tion, occurrence, and coverage of left-lateralized vs. right-lateralized 632 microstates during visualization compared to verbalization and vice 633 versa when in the literature language processing has primarily been 634 associated (though clearly not uniquely, see e.g. Jonides et al., 1998) 635 with a large-scale left-hemispheric network (Buchsbaum et al., 2001; 636 Vigneau et al., 2006), whereas secondary visual processing has been 637 primarily associated (though again clearly not uniquely, see e.g. Mehta 638 and Newcombe, 1991; Sergent, 1990) with right-lateralized regions, 639 particularly the right-posterior parietal cortex (Malhotra et al., 2009; 640 Newcombe et al., 1987; Weiss et al., 2006)?

A possible answer to this question may be associated with the EEG 015 frequency band range and the computation of the four EEG microstate 643 classes is based on our knowledge of the location of their intra-cortical 644 sources (Pascual-Marqui et al., 2014). The computation of EEG micro- 645 state topographies is conventionally performed on EEG data band-pass 646 filtered from 2 to 20 Hz (e.g. Koenig et al., 2002; Lehmann et al., 2005; 647 Pascual-Marqui et al., 2014). However, the frequency band with the 648 strongest power in this frequency range is the alpha band. The function- 649 al significance of EEG alpha activity has been a matter of debate 650 (Bazanova and Vernon, 2014). It may depend on the specific frequency 651 range (upper vs. lower alpha) and its cortical sources. However, the pri- 652 mary cortical sources of the EEG microstate classes are posterior where 653 alpha exhibits inhibitory rather than excitatory functions on modality- 654 specific processing (Harmony, 2013; Niedermeyer and da Silva, 2005; 655 O'gorman et al., 2013; Pfurtscheller, 2003; Pfurtscheller and Da Silva, 656 1999; Pfurtscheller et al., 1996). Consequently, we suggest that microstate class A with its left-posterior source is observed more frequently 658 during visualization because it reflects activation in a left-posterior 659 hub which triggers inhibition of the connected left-hemispheric areas 660

Fig. 8. eLORETA brain images of the statistical difference in cortical distribution of electric sources of the three microstate classes B, C, and D between tasks (Task) and no-task resting (Re). Left to right: slices in axial (from top, nose up), sagittal (from left) and coronal (from back) view in MNI space. Images depict all t-values with p < 0.05 (two-tailed), t = 2.00 (df = 60). Red = stronger activity during tasks, blue = stronger activity during no-task resting. Task > Re: slices through the voxel (indicated by arrowheads) of maximally stronger activity during tasks. Task < Re: slices through the voxel (indicated by arrowheads) of maximally stronger activity during resting. The MNI coordinates (X, Y, Z) of these voxels are reported in millimeters. For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.

661

662

663 664

665

666

667

668

669

670

671

672

673

674

675

676

677

678

679

680

681

682

691

692

693

694

695

696

697

698

699

700

701

702

703

704

705

706

707

708

709

710

711

712

713

714

715

716

717

718

719

720

721

related to language processing, whereas microstate class B with its right-posterior source is observed more frequently during verbalization because it reflects activation in a right-posterior hub which triggers inhibition of the connected right-hemispheric areas related to visuospatial processing. Studies on the effective connectivity of the brain sources of the four EEG microstates are needed to shed more light on this issue.

EEG microstate topographies

Comparing the EEG microstate topographies between the four conditions revealed significant topographic differences for microstate classes B, C, and D. These differences varied across classes. This suggests that the biological correlates of mental states representing modalities such as visual vs. verbal/abstract are themselves (micro-) state dependent, which is something that, to our knowledge, has not been taken into account previously (e.g. Koenig et al., 1998; Lehmann et al., 1998,

Intra-cortical source localizations revealed stronger left parietal (BA13, BA39) and weaker right occipito-parietal (BA31, BA13) activation for visualization compared to verbalization for microstate classes B and C respectively. These left-right topography differences for microstate classes B and C might again reflect increased inhibitory activity in left-lateralized areas of verbal processing (Buchsbaum et al., 2001; LLoyd, 2007; Vigneau et al., 2006) and decreased inhibitory activity in right-lateralized areas of spatial-visual processing (LLoyd, 2007; Malhotra et al., 2009; Newcombe et al., 1987; Weiss et al., 2006).

For microstate class D, posterior cingulate activity (BA23) was stronger and anterior cingulate (BA25) activity weaker during visualization compared to verbalization. Unlike posterior areas associated with modality-specific processing, the anterior cingulate cortex (ACC) is not primarily regarded as a source of alpha but as a source of EEG theta activity. EEG theta induced by activity in the ACC is referred to as frontal midline theta. This distinct rhythm occurs during working-memory tasks (Asada et al., 1999; Ishii et al., 1999) and correlates positively with demands on attention and executive processing (Gevins et al., 1997). These findings suggest that increased ACC activity during verbalization may be interpreted as increased demands on attention and executive processing compared to visualization. The interpretation of increased posterior cingulate activity during visualization is less straightforward. Its maxima was localized in an area previously associated with spatial orientation and memory (Vogt et al., 1992). However, since it extends beyond the posterior cingulate cortex to the motor- and somatosensory cortices, it may also reflect increased alpha inhibition of information processing in competing modalities. In line with this hypothesis, opposing alpha effects in visual as opposed to motor- and somatosensory cortical areas have frequently been reported (Pfurtscheller, 2003).

The areas of increased activity during no-task resting compared to the three tasks largely overlap with areas previously reported to be associated with the dorsal medial prefrontal cortex subsystem, a default mode network subsystem associated with introspection about mental states. The areas of increased activity during tasks largely overlap with areas associated with the medial temporal lobe subsystem and primary default mode network hubs which have been associated with memory-based construction/ simulation and valuation of motivationally-salient/personally-significant information respectively (Andrews-Hanna, 2012). Future studies must reveal the interplay of the cortical neuronal rhythms that these networks rely on.

Inter-individual differences

EEG microstate parameters

Pearson correlations revealed strong associations between the distributions of the three parameters of the four EEG microstate classes across conditions (all r > 0.75). These results suggest that beyond task-specificity, the EEG microstate parameters also exhibit strong 723 person-specificity.

Pearson correlations between behavioral variables and EEG micro- 725 state parameters during resting revealed no consistent associations 726 with participants' degree of object-visual, spatial-visual, and verbal 727 thinking during resting or participants' modality-specific cognitive 728 abilities. However, verbal cognitive style was negatively associated 729 with the duration of microstate class A, and positively associated with 730 the occurrence of microstate class C. The negative association with 731 class A is in line with our observed task differences which suggest that 732 class A duration is shortened during verbalization compared to visualization. However, due to the exploratory nature of this investigation, 734 many correlations had to be computed and significant results (not 735 corrected for multiple testing) only concerned singular parameter- 736 modality pairs. Hypothesis-driven studies or studies of larger sample 737 sizes are needed to validate our results.

The lack of associations between EEG microstate parameters 739 with reported thinking modality and modality-specific cognitive abili-740 ties may be due to a number of reasons. Firstly, other factors such as 741 alertness, wakefulness, and/or other personality traits (Knyazev et al., 742 2004; Schmidtke and Heller, 2004; Stenberg, 1992) which are known 743 to affect EEG parameters and differ between participants may play a 744 confounding role. Moreover, with regard to reported thinking modality, 745 participants may struggle with reporting their degree of object-, spatial-746 visual, and verbal thinking with the necessary precision and similar 747 point of reference to allow comparing these reports between individ- 748 uals. Furthermore, we note that participants' reported thinking mo- 749 dality referred to their perception of the whole duration of the 50 s 750 recording run. However, on average only 25.76 s of clean EEG for each 751 run was available for analysis. This limitation of overlap between the 752 time period rated and the time period analyzed may also contribute to 753 the lack of consistent inter-individual associations between EEG microstate parameters and subjectively reported thinking modality.

755

756

The functional significance of the EEG microstate classes

Our inter-individual results showed clear associations of the left- 757 lateralized microstate class A with visual, and the right-lateralized mi- 758 crostate class B with verbal processing. The increased duration, occur- 759 rence, and coverage of these microstate classes during the respective 760 tasks might reflect increased inhibitory activity in default mode 761 network hubs associated with areas associated with language and 762 visuo-spatial processing, respectively. Complementary to this effect, 763 topographic analyses suggested a shift of two microstate classes 764 (B and C) towards increased left-hemispheric and decreased right- 765 hemispheric source activity for visualization compared to verbalization. 766 Apparently, when the same four microstate classes are enforced for vi- 767 sualization and verbalization conditions, the lateralization difference 768 shows in the predominance of the two lateralized microstate classes A 769 and B, whereas when the microstate classes for the tasks are inspected 770 separately, microstate classes B and C appear more right- respectively 771 left-lateralized.

Conclusions 773

Our results revealed associations of microstate classes A and B with 774 visual and verbal processing respectively, and microstate class D with 775 interoceptive-autonomic processing. However, all microstate classes 776 occurred in all four conditions and the absolute differences of coverage 777 for microstate classes A and B for visualization compared to verbalization conditions, and the absolute differences of coverage for microstate 779 class D for no-task resting compared to goal-directed tasks are consider- 780 ably small. Therefore, we suggest that while increased occurrence, dura-781 tion, and coverage of EEG microstate classes A, B, and D can clearly be 782 associated with visual, verbal, and interoceptive-autonomic processing 783 respectively, they can certainly not be reduced to these functions. 784

875

883

884

889

892

896

899

901

905

909

910

913

017

915

922

931

933

939

Instead, a finely-tuned interplay between the four EEG microstate classes is necessary for the continuous formation of visual and verbal thoughts, as well as interoceptive-autonomic processing. Our results point to the possibility that the EEG microstate classes may represent the head-surface measured activity of intra-cortical sources primarily exhibiting inhibitory functions. However, additional studies are needed to verify and elaborate on this hypothesis.

Acknowledgments

785

786

787

788

790

791

792

793

794

795

796

797

798

799

800

801

802

803

804

805

806

807

808

809

810

811 812

813

814

815

816

817

818

819

820

821 822

823

825

826 827

829

830 831

832

833

834

835

836

837 838

839

840

841

842

843

844

845

846

847

848 849

850

851

852

853

854

855

856

857

858 859

860

861

We thank Daniel Kiper, Stephan Gerhard, and Christian Oberholzer for suggesting valuable improvements to this manuscript. P.M. is grateful to Dietrich Lehmann, Klaus Oberauer, and Daniel Kiper for their supervision and support during this project.

References

- Andreou, C., Faber, P.L., Leicht, G., Schoettle, D., Polomac, N., Hanganu-Opatz, I.L. Lehmann, D., Mulert, C., 2014. Resting-state connectivity in the prodromal phase of schizophrenia: insights from EEG microstates, Schizophr, Res. 152, 513-520
 - Andrews-Hanna, J.R., 2012. The brain's default network and its adaptive role in internal mentation. Neuroscientist 18, 251-270.
 - Asada, H., Fukuda, Y., Tsunoda, S., Yamaguchi, M., Tonoike, M., 1999. Frontal midline theta rhythms reflect alternative activation of prefrontal cortex and anterior cingulate cortex in humans. Neurosci. Lett. 274, 29-32.
 - Baschek, I.L., Bredenkamp, J., Oehrle, B., Wippich, W., 1977. Assessment of imagery (I), concreteness (C) and meaningfulness (m1) of 800 nouns. Z. Exp. Angew. Psychol. 24, 353-396.
 - Bazanova, O.M., Vernon, D., 2014. Interpreting EEG alpha activity. Neurosci. Biobehav. Rev. 44, 94-110
 - Becker, D., Creutzfeldt, O., Schwibbe, M., Wuttke, W., 1982. Changes in physiological, EEG and psychological parameters in women during the spontaneous menstrual cycle and following oral contraceptives. Psychoneuroendocrinology 7, 75-90.
 - Blazhenkova, O., Kozhevnikov, M., 2009. The new object-spatial-verbal cognitive style model: theory and measurement. Appl. Cogn. Psychol. 23, 638-663.
 - Britz, J., Van De Ville, D., Michel, C.M., 2010. BOLD correlates of EEG topography reveal rapid resting-state network dynamics. NeuroImage 52, 1162-1170.
 - Brodbeck, V., Kuhn, A., von Wegner, F., Morzelewski, A., Tagliazucchi, E., Borisov, S., Michel, C.M., Laufs, H., 2012. EEG microstates of wakefulness and NREM sleep. NeuroImage 62, 2129-2139.
 - Buchsbaum, B.R., Hickok, G., Humphries, C., 2001. Role of left posterior superior temporal gyrus in phonological processing for speech perception and production. Cogn. Sci. 25, 663-678.
 - Chatrian, G., Lettich, E., Nelson, P., 1985. Ten percent electrode system for topographic studies of spontaneous and evoked EEG activities. Am. J. EEG Technol. 25, 83-92.
 - Ekstrom, R.B., French, J.W., Harman, H.H., Derman, D., 1976. Manual for kit of factorreferenced cognitive tests. Educational Testing Service, Princeton, NJ.
- Faber, P.L., Lehmann, D., Barendregt, H., Kaelin, M., Gianotti, L.R., 2005. Increased duration of EEG microstates during meditation. Brain Topogr. 18, 131.
- Faber, P., Lehmann, D., Milz, P., Travis, F., Parim, N., 2014. EEG microstates differ between transcending and mind wandering. ZNZ Symposium 2014, Zurich.
- Fingelkurts, A.A., Fingelkurts, A.A., Bagnato, S., Boccagni, C., Galardi, G., 2012. EEG oscillatory states as neuro-phenomenology of consciousness as revealed from patients in vegetative and minimally conscious states. Conscious. Cogn. 21, 149-169.
- Gevins, A., Smith, M.E., McEvoy, L., Yu, D., 1997. High-resolution EEG mapping of cortical activation related to working memory: effects of task difficulty, type of processing, and practice. Cereb. Cortex 7, 374-385.
- Harmony, T., 2013. The functional significance of delta oscillations in cognitive processing. Front. Integr. Neurosci. 7.
- Horn, W., 1962. Leistungsprüfsystem, LPS: Handanweisung für die Durchführung, Auswertung und Interpretation.
- IBM Corporation, 2014. IBM SPSS Statistics for Windows, Version 22.0. IBM Corp., Armonk, NY
- Ishii, R., Shinosaki, K., Ukai, S., Inouye, T., Ishihara, T., Yoshimine, T., Hirabuki, N., Asada, H., Kihara, T., Robinson, S.E., Takeda, M., 1999. Medial prefrontal cortex generates frontal midline theta rhythm. Neuroreport 10, 675-679.
- Jäger, A.O., Süß, H.-M., Beauducel, A., 1997. Berliner Intelligenzstruktur-Test, BIS-Test. Form 4. Handanweisung (The Berlin Intelligence Structure Test, BIS test. 4. Form. Test Manual), Hogrefe, Göttingen,
- Jonides, J., Schumacher, E.H., Smith, E.E., Koeppe, R.A., Awh, E., Reuter-Lorenz, P.A., Marshuetz, C., Willis, C.R., 1998. The role of parietal cortex in verbal working memory. I. Neurosci. 18, 5026-5034.
- Katayama, H., Gianotti, L.R., Isotani, T., Faber, P.L., Sasada, K., Kinoshita, T., Lehmann, D., 2007. Classes of multichannel EEG microstates in light and deep hypnotic conditions. Brain Topogr. 20, 7–14.
- Kikuchi, M., Koenig, T., Wada, Y., Higashima, M., Koshino, Y., Strik, W., Dierks, T., 2007. Native EEG and treatment effects in neuroleptic-naive schizophrenic patients: time and frequency domain approaches. Schizophr. Res. 97, 163-172.
- Kindler, J., Hubl, D., Strik, W., Dierks, T., Koenig, T., 2011. Resting-state EEG in schizophrenia: auditory verbal hallucinations are related to shortening of specific microstates. Clin. Neurophysiol. 122, 1179-1182.

- Knyazev, G.G., Savostvanov, A.N., Levin, E.A., 2004, Alpha oscillations as a correlate of trait 862 anxiety. Int. J. Psychophysiol. 53, 147-160.
- Koenig, T., Kochi, K., Lehmann, D., 1998. Event-related electric microstates of the brain dif-864 fer between words with visual and abstract meaning. Electroencephalogr. Clin. 865 Neurophysiol, 106, 535-546. 866
- Koenig, T., Lehmann, D., Merlo, M.C., Kochi, K., Hell, D., Koukkou, M., 1999. A deviant EEG 867 brain microstate in acute, neuroleptic-naive schizophrenics at rest. Eur. Arch. Psychi-868 atry Clin. Neurosci, 249, 205-211. 869
- Koenig T Prichen I. Jehmann D Sosa P.V. Braeker F. Kleinlogel H. Jsenhart R. John. 870 E.R., 2002. Millisecond by millisecond, year by year: normative EEG microstates and 871 developmental stages, NeuroImage 16, 41-48, 872
- Koenig, T., Kottlow, M., Stein, M., Melie-García, L., 2011. Ragu: a free tool for the analysis 873 of EEG and MEG event-related scalp field data using global randomization statistics. 874 Comput. Intell. Neurosci. 2011. 4.
- Krug, R., Moelle, M., Fehm, H., Born, J., 1999. Variations across the menstrual cycle in EEG 876 activity during thinking and mental relaxation. J. Psychophysiol, 13, 163-172. 877
- Lehmann, D., 1990. Brain electric microstates and cognition: the atoms of thought. Ma-878 chinery of the Mind. Springer, pp. 209-224. 879
- Lehmann, D., Michel, C.M., 2011. EEG-defined functional microstates as basic building 880 blocks of mental processes. Clin. Neurophysiol. 122, 1073-1074. 881
- Lehmann, D., Skrandies, W., 1980. Reference-free identification of components of 882 checkerboard-evoked multichannel potential fields. Electroencephalogr. Clin. Neurophysiol. 48, 609-621.
- Lehmann, D., Ozaki, H., Pal, I., 1987. EEG alpha map series: brain micro-states by space-885 oriented adaptive segmentation. Electroencephalogr. Clin. Neurophysiol. 67, 271–288. 886
- Lehmann, D., Strik, W., Henggeler, B., Koenig, T., Koukkou, M., 1998. Brain electric micro-887 states and momentary conscious mind states as building blocks of spontaneous 888 thinking: I. Visual imagery and abstract thoughts. Int. J. Psychophysiol. 29, 1-11.
- Lehmann, D., Koenig, T., Henggeler, B., Strik, W., Kochi, K., Koukkou, M., Pascual-Marqui, 890 R., 2004. Brain areas activated during electric microstates of mental imagery versus 891 abstract thinking. Klin. Neurophysiol. 35, 160.
- Lehmann, D., Faber, P.L., Galderisi, S., Herrmann, W.M., Kinoshita, T., Koukkou, M., Mucci, 893 A., Pascual-Marqui, R.D., Saito, N., Wackermann, J., 2005. EEG microstate duration and 894 syntax in acute, medication-naive, first-episode schizophrenia: a multi-center study. 895 Psychiatry Res. Neuroimaging 138, 141-156.
- Lehmann, D., Pascual-Marqui, R.D., Michel, C., 2009. EEG microstates. Scholarpedia 4, 897 898
- Lehmann, D., Pascual-Marqui, R.D., Strik, W.K., Koenig, T., 2010. Core networks for visualconcrete and abstract thought content; a brain electric microstate analysis. NeuroImage 900 49, 1073-1079
- LLoyd, D., 2007. "What Do Brodmann Areas Do?," Or: Scanning the Neurocracy. CT: Pro-902 gram in Neuroscience, Hartford, 903
- Malhotra, P., Coulthard, E.J., Husain, M., 2009. Role of right posterior parietal cortex in 904 maintaining attention to spatial locations over time. Brain awn350
- Marks, D.F., 1973. Visual imagery differences in the recall of pictures. Br. J. Psychol. 64, 906
- Mehta, Z., Newcombe, F., 1991. A role for the left hemisphere in spatial processing. Cortex 908 27, 153-167.
- Meili, R., 1955. Würfelabwicklungen. Huber, Bern.
- Michel, C., Brandeis, D., Koenig, T., 2009. Electrical neuroimaging in the time domain. In: 911 Michel, C., Koenig, T., Brandeis, D., Gianotti, L., Wackermann, J. (Eds.), Electrical Neu-912 roimaging. Cambridge University Press, Cambridge, pp. 111-143.
- Milz, P., 2015n. The KEY EEG Python Library: keypy (in preparation for upload to public github repository).
- Milz, P., Faber, P., Lehmann, D., 2015. The Modality of Thinking Questionnaire—A New Self-Report Measure of Object-Spatial-Verbal Cognitive Style (submitted for publication). 917
- Newcombe, F., Ratcliff, G., Damasio, H., 1987. Dissociable visual and spatial impairments 918 following right posterior cerebral lesions: clinical, neuropsychological and anatomical 919 evidence. Neuropsychologia 25, 149-161. 921
- Niedermeyer, E., da Silva, F.L., 2005. Electroencephalography: basic principles, clinical applications, and related fields. Lippincott Williams & Wilkins.
- Nishida, K., Morishima, Y., Yoshimura, M., Isotani, T., Irisawa, S., Jann, K., Dierks, T., Strik, 923 W., Kinoshita, T., Koenig, T., 2013. EEG microstates associated with salience and 924 frontoparietal networks in frontotemporal dementia, schizophrenia and Alzheimer's 925 disease. Clin. Neurophysiol. 124, 1106-1114. 926
- Nuwer, M.R., 1987. Recording electrode site nomenclature. J. Clin. Neurophysiol. 4, 927 121-133. 928
- O'gorman, R., Poil, S.-S., Brandeis, D., Klaver, P., Bollmann, S., Ghisleni, C., Lüchinger, R., 929 Martin, E., Shankaranarayanan, A., Alsop, D., 2013. Coupling between resting cerebral 930 perfusion and EEG. Brain Topogr. 26, 442-457.
- Oldfield, R.C., 1971. The assessment and analysis of handedness: the Edinburgh inventory. 932 Neuropsychologia 9, 97-113.
- Pascual-Marqui, R.D., 2007. Discrete, 3D distributed, linear imaging methods of electric 934 neuronal activity. Part 1: exact, zero error localization (arXiv preprint arXiv: 935 0710.3341). 936
- Pascual-Marqui, R.D., Michel, C.M., Lehmann, D., 1995, Segmentation of brain electrical activity into microstates: model estimation and validation, IEEE Trans. Biomed. Eng. 42. 938 658-665
- Pascual-Marqui, R.D., Lehmann, D., Faber, P., Milz, P., Kochi, K., Yoshimura, M., Nishida, K., 940 Isotani, T., Kinoshita, T., 2014. The resting microstate networks (RMN): cortical distributions, dynamics, and frequency specific information flow (arXiv preprint arXiv: 942 1411.1949) 943
- Pfurtscheller, G., 2003. Induced oscillations in the alpha band: functional meaning. 944 Epilepsia 44, 2–8. 945
- Pfurtscheller, G., Da Silva, F.L., 1999. Event-related EEG/MEG synchronization and 946 desynchronization: basic principles. Clin. Neurophysiol. 110, 1842-1857. 947

P. Milz et al. / NeuroImage xxx (2015) xxx-xxx

948

949

950

951

952

953

954

955

956

957

958

959

960 961

962

963

964

965

966 967

968

969

970

971

972

973

998

Pfurtscheller, G., Stancak, A., Neuper, C., 1996, Event-related synchronization (ERS) in the alpha band-an electrophysiological correlate of cortical idling: a review. Int. I. Psychophysiol, 24, 39-46.

Schlegel, F., Lehmann, D., Faber, P.L., Milz, P., Gianotti, L.R., 2012, EEG microstates during resting represent personality differences. Brain Topogr. 25, 20–26.

- Schmidtke, I.I., Heller, W., 2004, Personality, affect and EEG; predicting patterns of regional brain activity related to extraversion and neuroticism, Personal, Individ, Differ, 36. 717-732.
- Sergent, J., 1990. The neuropsychology of visual image generation: data, method, and theory. Brain Cogn. 13, 98-129.
- Stenberg, G., 1992. Personality and the EEG: arousal and emotional arousability. Personal. Individ. Differ. 13, 1097-1113.
- Strelets, V., Faber, P., Golikova, J., Novototsky-Vlasov, V., Koenig, T., Gianotti, L., Gruzelier, J., Lehmann, D., 2003. Chronic schizophrenics with positive symptomatology have shortened EEG microstate durations. Clin. Neurophysiol. 114, 2043–2051.
- Strik, W., Dierks, T., Becker, T., Lehmann, D., 1995. Larger topographical variance and decreased duration of brain electric microstates in depression. J. Neural Transm. Gen. Sect_INT 99_213-222
- Strik, W.K., Chiaramonti, R., Muscas, G.C., Paganini, M., Mueller, T.J., Fallgatter, A.J., Versari, A., Zappoli, R., 1997. Decreased EEG microstate duration and anteriorisation of the brain electrical fields in mild and moderate dementia of the Alzheimer type. Psychiatry Res. Neuroimaging 75, 183-191.
- Strik, W., Fallgatter, A., Brandeis, D., Pascual-Marqui, R., 1998. Three-dimensional tomography of event-related potentials during response inhibition: evidence for phasic frontal lobe activation. Electroencephalogr. Clin. Neurophysiol. Evoked Potentials Sect. 108, 406-413.

- Tomescu, M.I., Rihs, T.A., Becker, R., Britz, I., Custo, A., Grouiller, F., Schneider, M., Debbané, 974 M., Eliez, S., Michel, C.M., 2014. Deviant dynamics of EEG resting state pattern in 975 22q11.2 deletion syndrome adolescents: a vulnerability marker of schizophrenia? 976 Schizophr. Res. 157, 175-181.
- Tomescu, M.I., Rihs, T.A., Roinishvili, M., Karahanoglu, F.I., Schneider, M., Menghetti, S., 978 Van De Ville, D., Brand, A., Chkonia, E., Eliez, S., Herzog, M.H., Michel, C.M., Cappe, 979 C., 2015. Schizophrenia patients and 22q11.2 deletion syndrome adolescents at risk 980 express the same deviant patterns of resting state EEG microstates; a candidate 981 endophenotype of schizophrenia. Schizophr. Res. Cogn.
- Vigneau, M., Beaucousin, V., Herve, P.-Y., Duffau, H., Crivello, F., Houde, O., Mazoyer, B., 983 Tzourio-Mazoyer, N., 2006. Meta-analyzing left hemisphere language areas: phonol- 984ogy, semantics, and sentence processing. NeuroImage 30, 1414-1432.
- Vogt, B.A., Finch, D.M., Olson, C.R., 1992. Functional heterogeneity in cingulate cortex: the 986 anterior executive and posterior evaluative regions. Cereb. Cortex 2, 435-443.

985

- Wackermann, J., Lehmann, D., Michel, C., Strik, W., 1993. Adaptive segmentation of spontaneous EEG map series into spatially defined microstates. Int. J. Psychophysiol. 14, 989
- Wehrle, R., Kaufmann, C., Wetter, T.C., Holsboer, F., Auer, D.P., Pollmächer, T., Czisch, M., 991 2007. Functional microstates within human REM sleep: first evidence from fMRI of $\,992$ a thalamocortical network specific for phasic REM periods. Eur. J. Neurosci. 25, 993 863-871
- Weiss, P.H., Rahbari, N.N., Lux, S., Pietrzyk, U., Noth, J., Fink, G.R., 2006. Processing the spa-995 tial configuration of complex actions involves right posterior parietal cortex: an fMRI 996 study with clinical implications. Hum. Brain Mapp. 27, 1004–1014.

Please cite this article as: Milz, P., et al., The functional significance of EEG microstates—Associations with modalities of thinking, NeuroImage (2015), http://dx.doi.org/10.1016/j.neuroimage.2015.08.023