

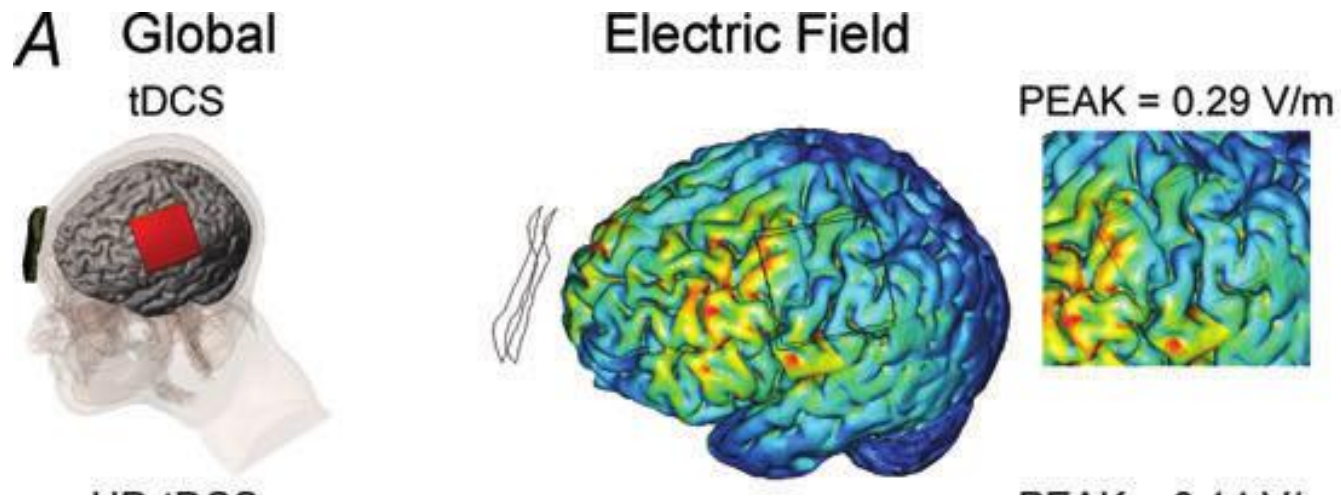
Modeling the effects of DC stimulation on presynaptic vesicle release

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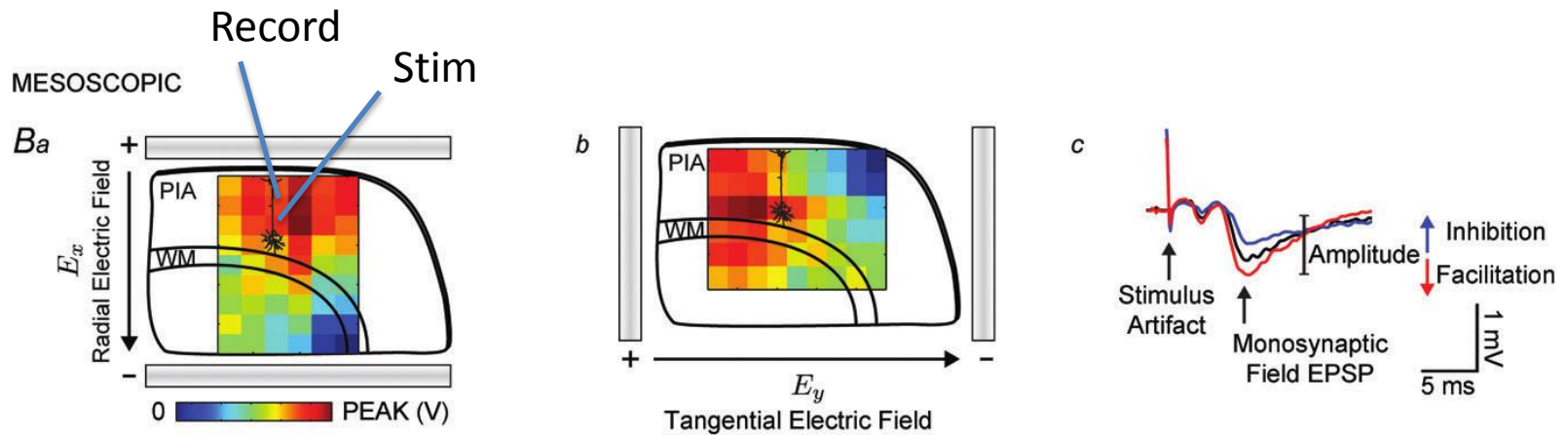
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Transcranial Direct Current Stimulation

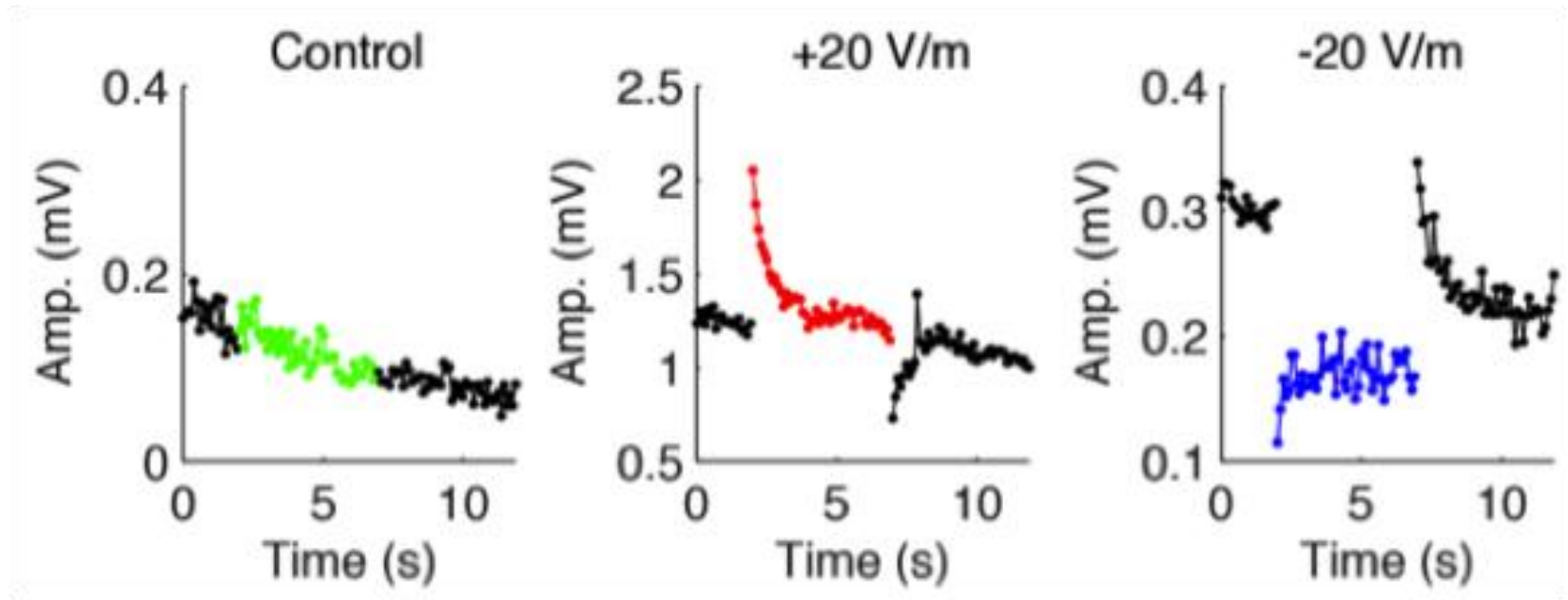


Used to treat: depression, stroke rehab, alzheimer's, epilepsy, addiction, many more

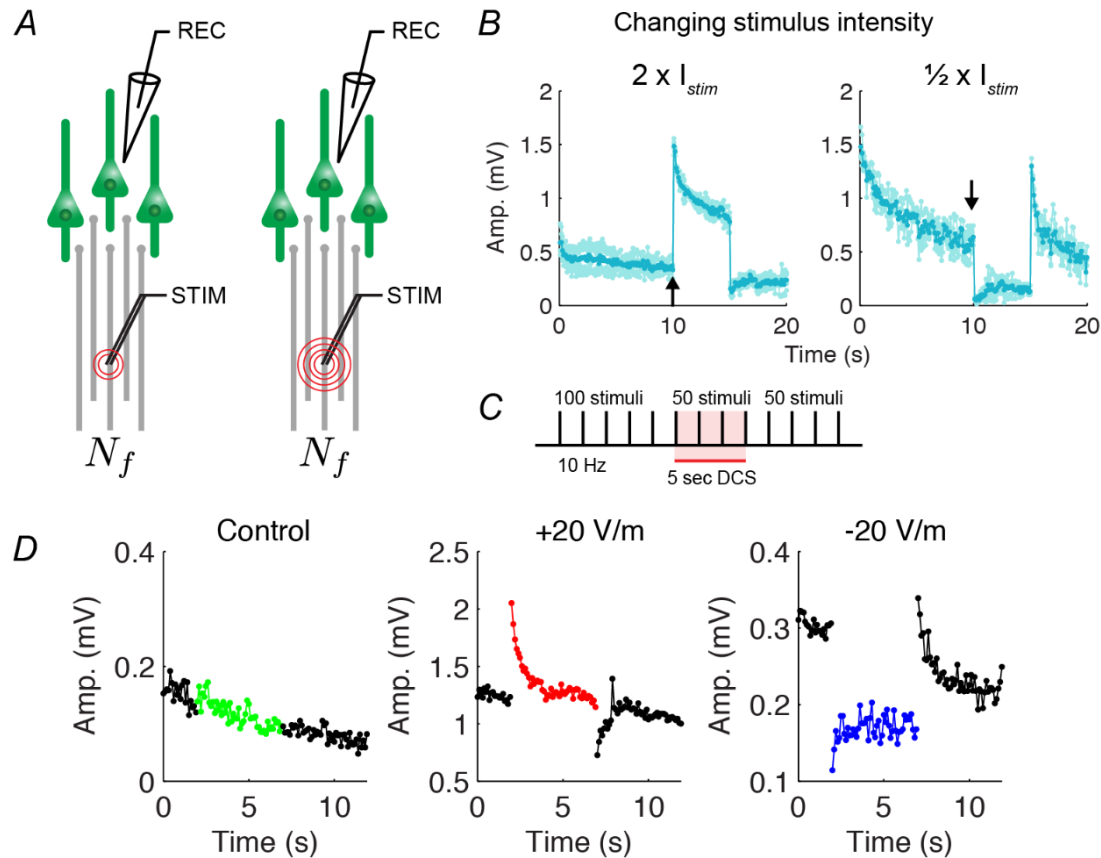
In vitro model of tDCS



Adaptation during DCS



Adaptation during DCS



Results are consistent with presynaptic effects

Modeling presynaptic release

- A group of $nsyn$ synapses
- Given stimuli at time t , terminal i has probability p_{ap_i} of firing
- All terminals have poisson docking/undocking (mean rates α/β) at finite number of release sites ns with initial release probability p_0

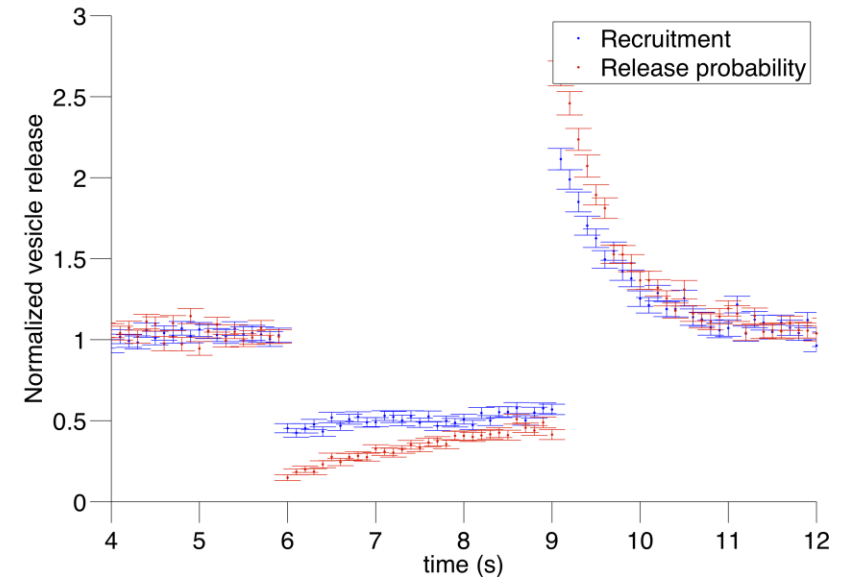
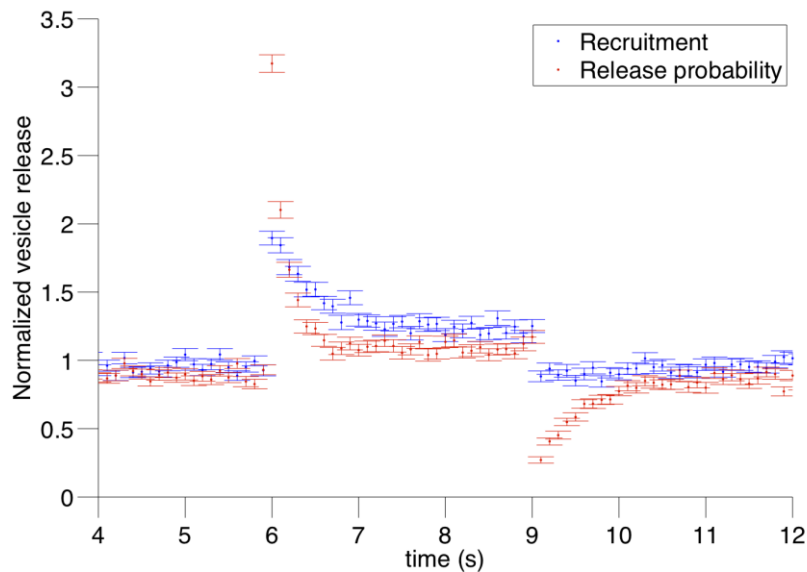
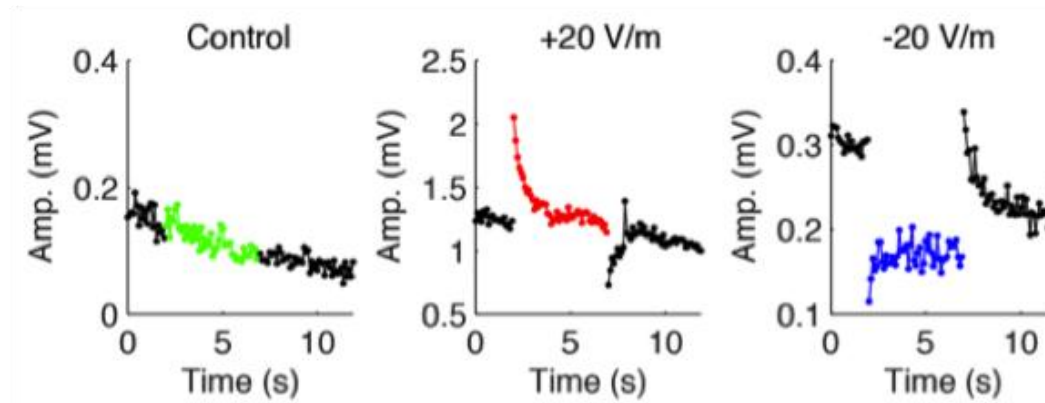
$$\sum_{i=1}^{nsyn} \overline{N^i} = \sum_{i=1}^{nsyn} \frac{p_{ap}^i p_0 ns_* (1 - e^{-gDt})}{1 - (1 - p_{ap} p_0) e^{-gDt}} \quad \begin{aligned} g &= a + b \\ ns_* &= \frac{ans}{a + b} \end{aligned}$$

Modeling presynaptic release with DCS

- DCS is known to modulate membrane potential at axon terminals
- Here it is modeled as having either of two effects
 - Recruitment
 - Modulate p_{ap}
 - Release
 - Modulate p_0

$$\sum_{i=1}^{nsyn} \overline{N^i} = \sum_{i=1}^{nsyn} \frac{\overset{\downarrow \text{blue}}{p_{ap}^i} \overset{\downarrow \text{red}}{p_0} ns_* (1 - e^{-g_D t})}{1 - \underset{\uparrow \text{blue}}{(1 - p_{ap})} \underset{\uparrow \text{red}}{p_0} e^{-g_D t}}$$

Modeling presynaptic release with DCS



Modeling presynaptic release with DCS

- Modulation of p_0 better reproduces edge detect when DCS is turned off
- Results depend on initial distribution of p_0 and p_{ap}

