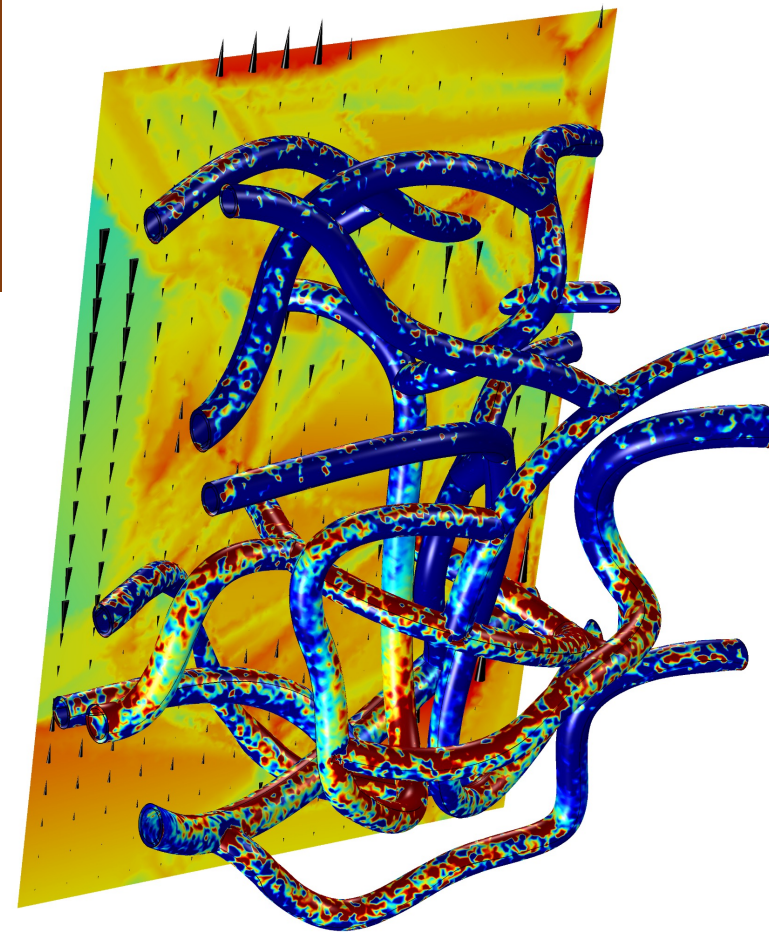


Neurovascular-Modulation: How Brain Stimulation Techniques Like tDCS, TMS and ECT May Activate the Blood-Brain-Barrier?

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Disclosure

The City University of New York: Patents on brain stimulation.

Soterix Medical: Produces tDCS and High-Definition tDCS.

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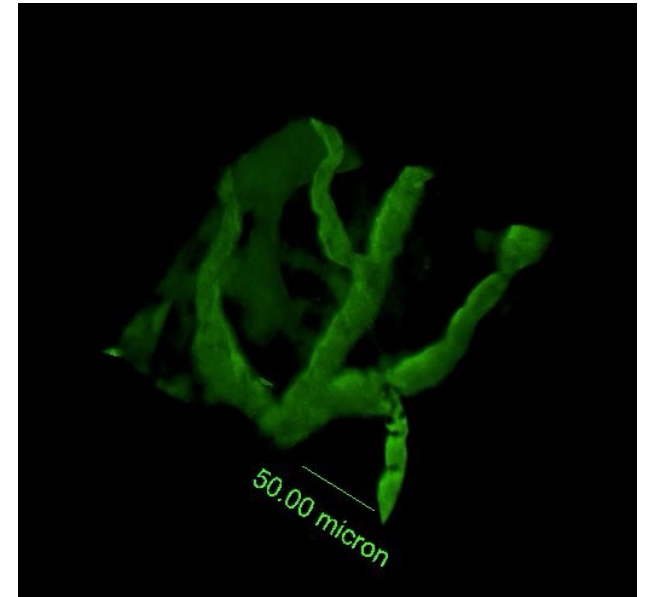


Neurovascular-modulation

First aspect: Brain stimulation of neuro-vascular coupling.

Second aspect: Direct stimulation of brain vascular/ blood-brain-barrier function.

- **Neurovascular coupling (unit):** Coupling between neuronal activity with vascular flow and blood-brain barrier (BBB) permeability.
- **Two-way interaction.** Neuronal activity activates vascular (eg. fMRI), Transport across BBB tightly controlled to regulate brain function.



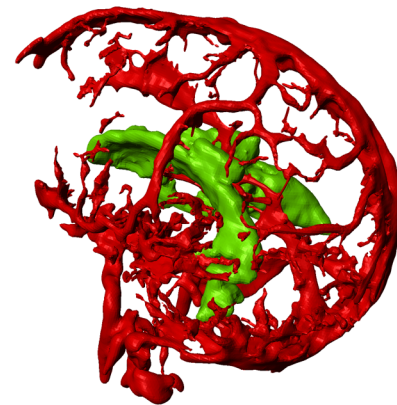
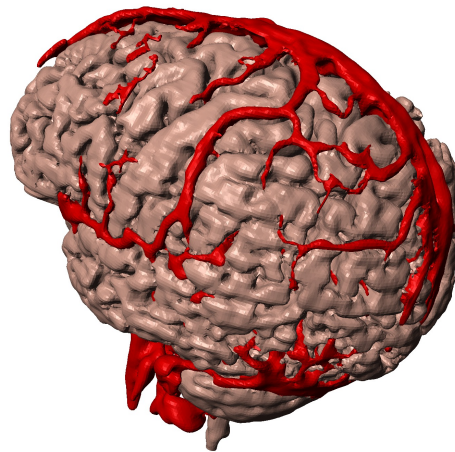
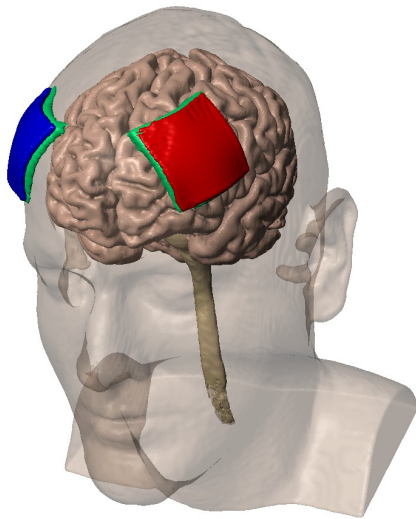
Stimulation of neurovascular unit:

FIRST ASPECT: Brain vasculature changes inevitable **secondary** to neuronal stimulation (eg. fMRI changes after brain stimulation).

SECOND ASPECT: Can neuromodulation **directly** activate endothelial cells of the BBB, leading to secondary neuronal changes.

Transcranial Direct Current Stimulation (tDCS) of the BBB [second aspect]

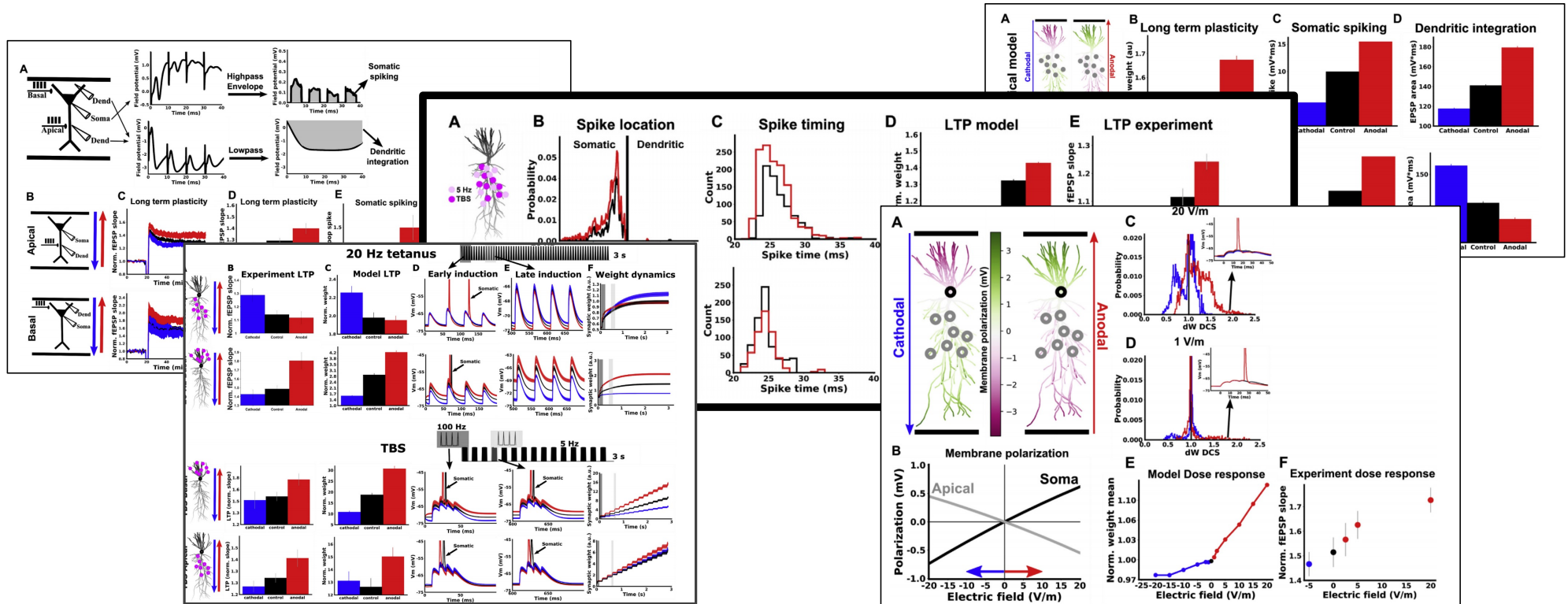
Neuronal response to DC fields extensively characterized.
Including in brain slices (where vasculature is absent)



Can tDCS (DC fields)
directly activate the
BBB, which in turn
modulates neurons?

Vascular response to tDCS established (eg. fMRI, fNIRS)
but considered epiphenomena !

Well established: Direct Current stimulation is a “direct” modulator of ongoing neuronal plasticity



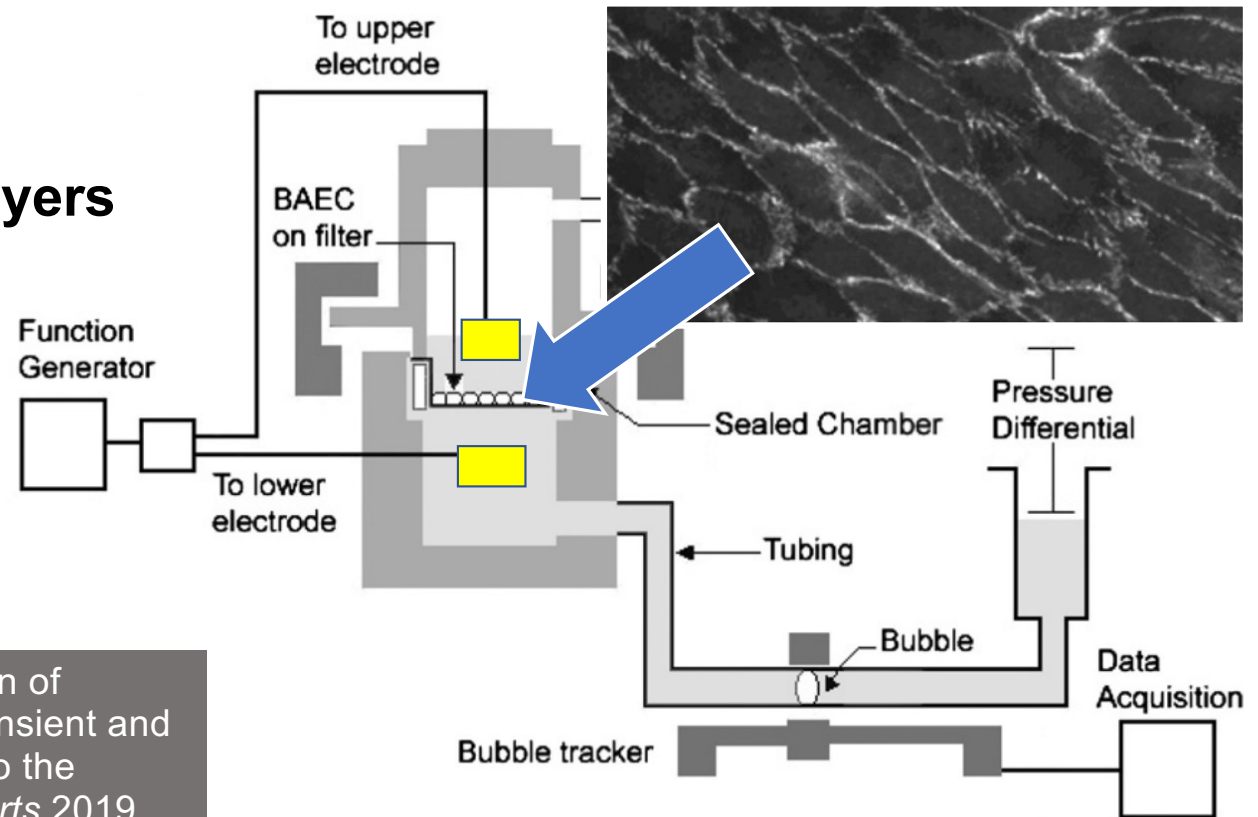
Jackson et al. Animal models of transcranial direct current stimulation. *Clin Neurophys* 2016

Kronberg et al. Direct current stimulation boosts Hebbian plasticity in vitro. *Brain Stim* 2020

“Primacy” of neurons as targets of neuromodulation means any changes in vascular function assumed secondary to neuron stimulation.

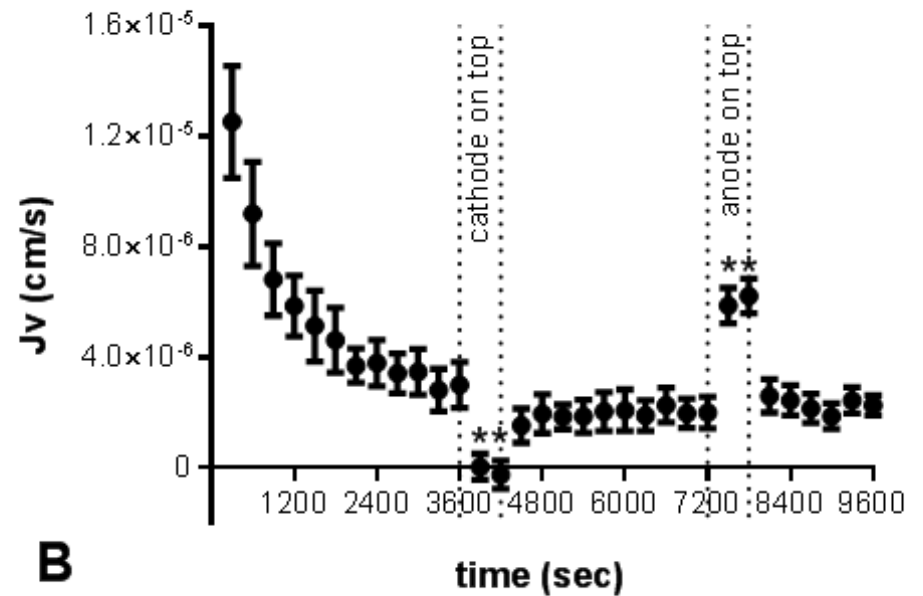
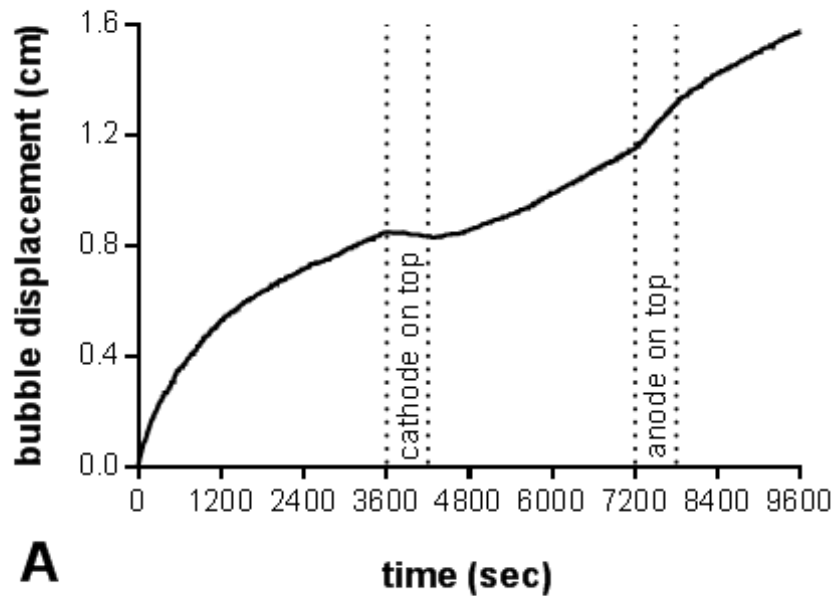
Isolated BBB stimulation established direct neuromodulation.

BBB model: **cultured endothelium monolayers**



Cancel et al. Direct current stimulation of endothelial monolayers induces a transient and reversible increase in transport due to the electroosmotic effect. *Scientific Reports* 2019

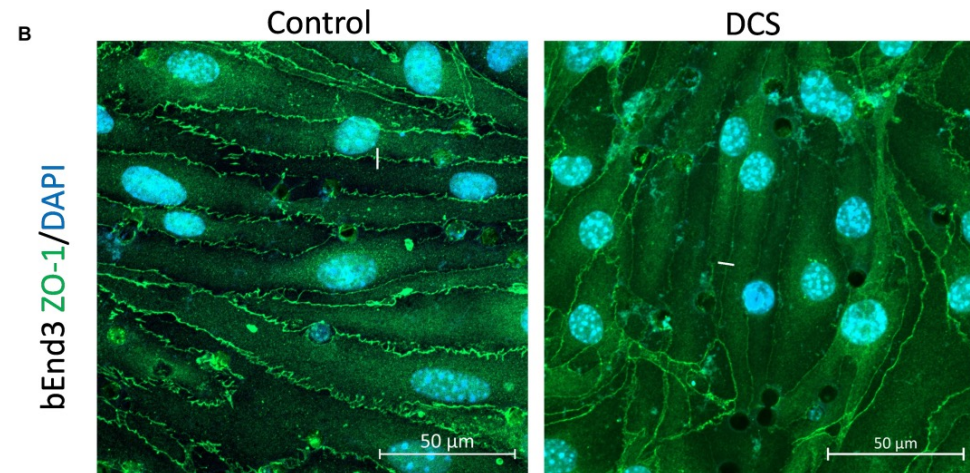
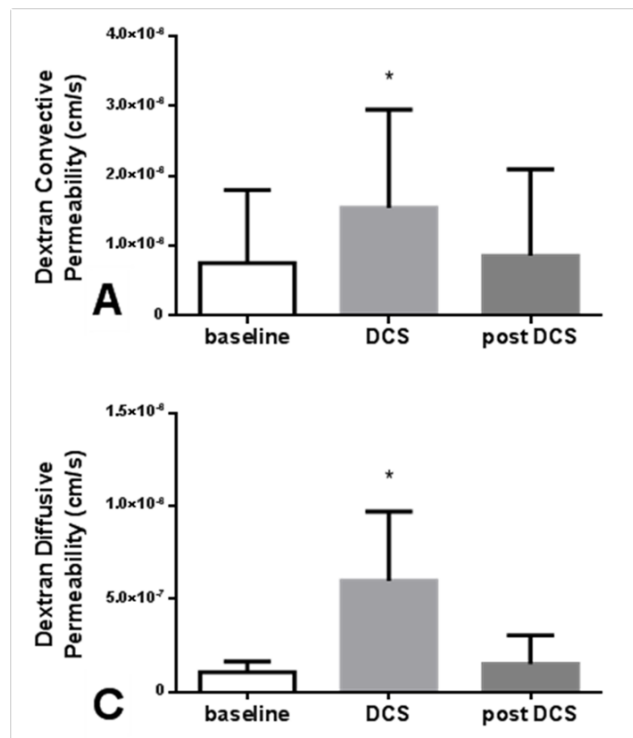
Direct Current stimulation produced an acute, polarity specific change in water transport across BBB model



Electroosmosis: Current will drag water through a (charged) barrier, proportional to tightness of barrier.

Cancel et al. Direct current stimulation of endothelial monolayers induces a transient and reversible increase in transport due to the electroosmotic effect. *Scientific Reports* 2019

Direct Current stimulation enhances specific molecule transport across the BBB and activates structural (tight junction) / molecular (eNOS) /early gene expression (VEGF).

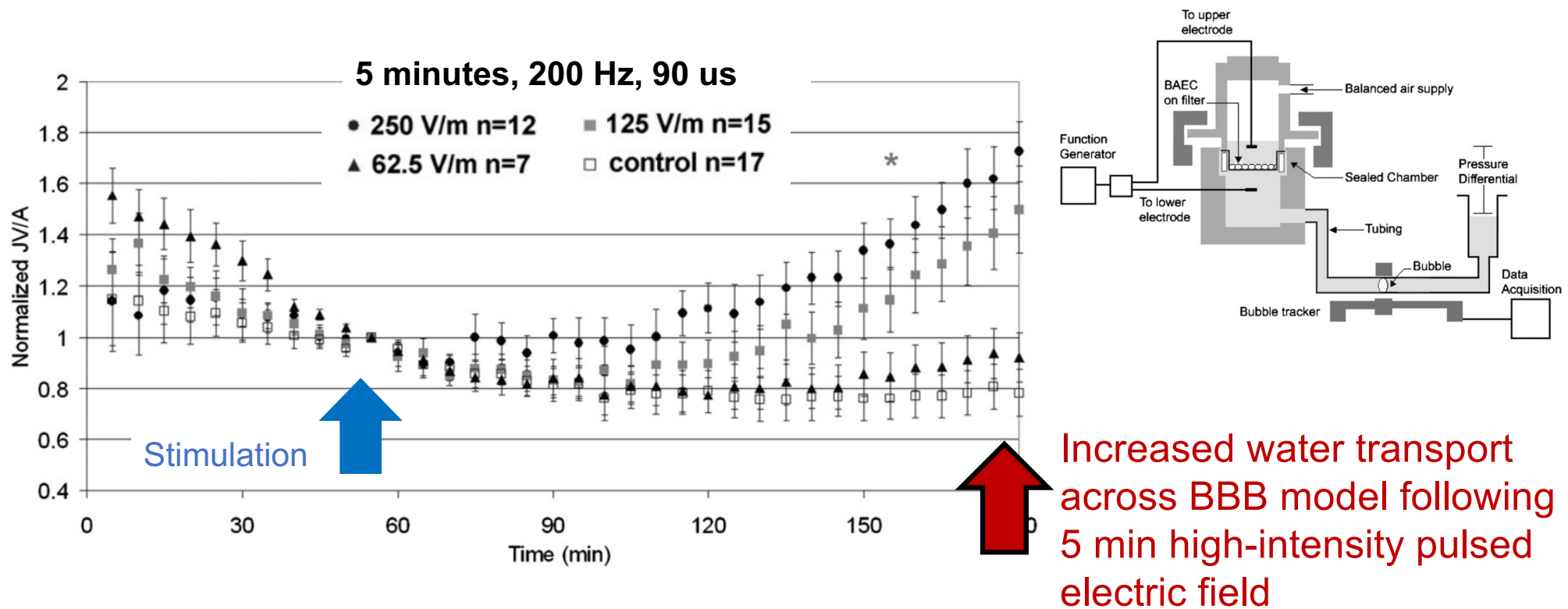


Xia et al. Direct Current Stimulation Disrupts Endothelial Glycocalyx and Tight Junctions of the Blood-Brain Barrier in vitro. *Frontiers cell and developmental biology* 2021

Plasticity from brain vasculature stimulation.

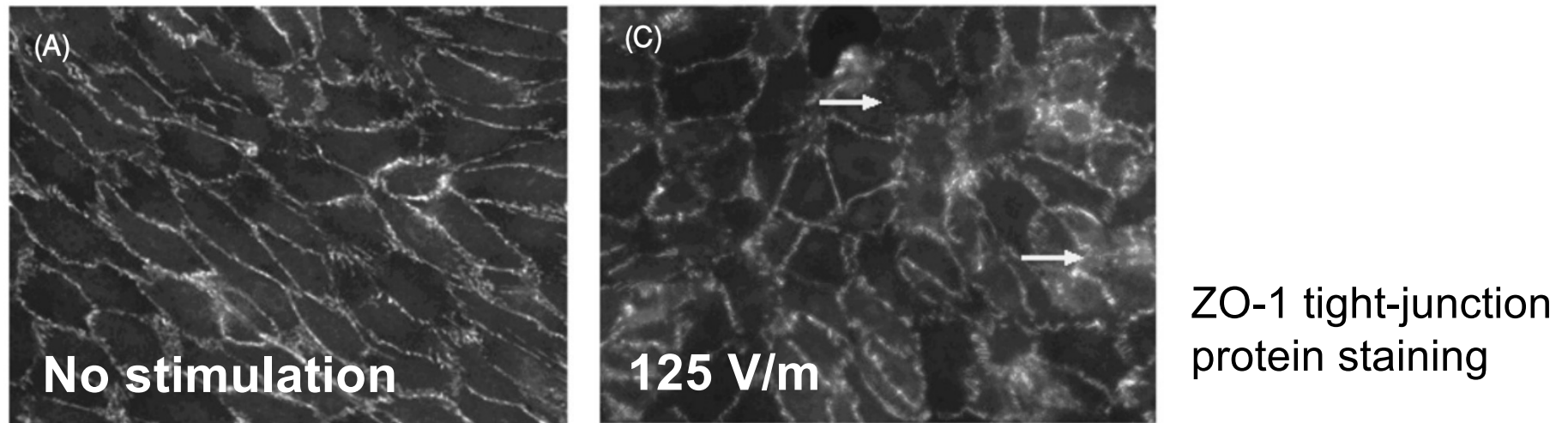
Cancel et al. Direct current stimulation of endothelial monolayers induces a transient and reversible increase in transport due to the electroosmotic effect. *Scientific Reports* 2019

High-intensity pulsed electric fields (DBS, ECT, TMS like) modulate isolated endothelial cells (BBB) including water and transport flux.



Cancel et al. DBS-relevant electric fields increases hydraulic conductivity of in vitro endothelial monolayers. *J Neural Engr* 2010

High-intensity pulsed electric fields (DBS, ECT, TMS like) induce lasting (plastic) changes in endothelial cells (BBB) function.

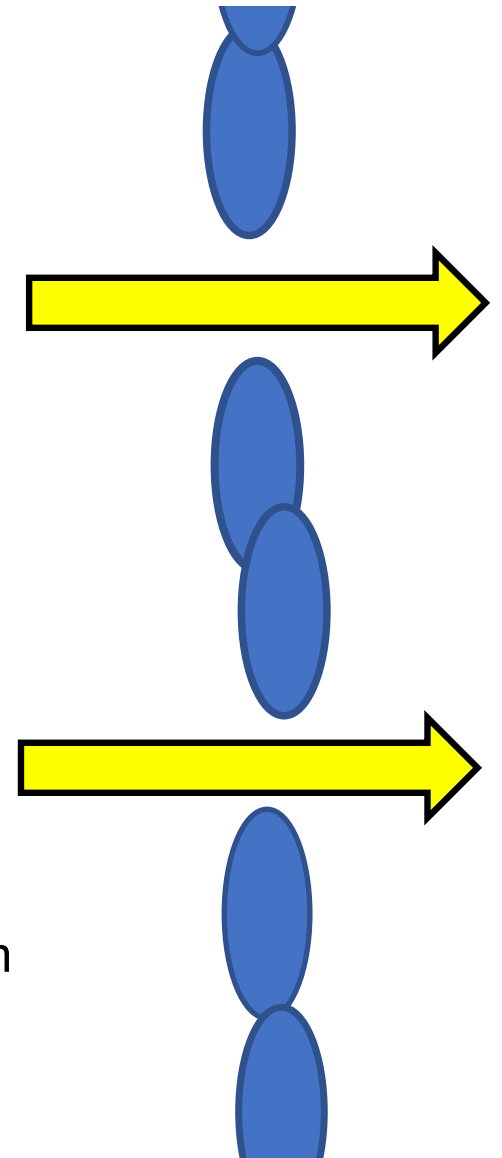


ZO-1 tight junction protein surrounds endothelial cells in control. Pulses electric fields modify continuity (arrows).

Cancel et al. DBS-relevant electric fields increases hydraulic conductivity of in vitro endothelial monolayers. *J Neural Engr* 2010

Pulsed electric fields increase hydraulic conductivity of in vitro endothelial monolayers

- Increased water transport likely enhances solute transport, impacting neurons
- Dose dependent increase in BBB transport
- **Via opening peri-cellular Tight Junctions.**
- No evidence for cell electroporation / transmembrane transport.
- Plausible in any brain / spinal structures / peripheral
- In vitro BBB system may not be good model for long-term (reversible) changes



Neurovascular Modulation: Direct effects on brain vasculature suggest unique therapeutic strategies (pathways)

”Boosting” of brain function (transport) / neurorehabilitation efficacy

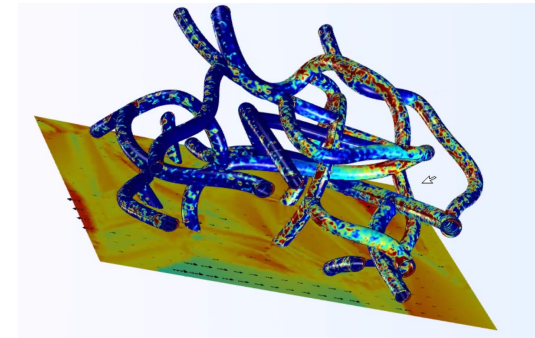
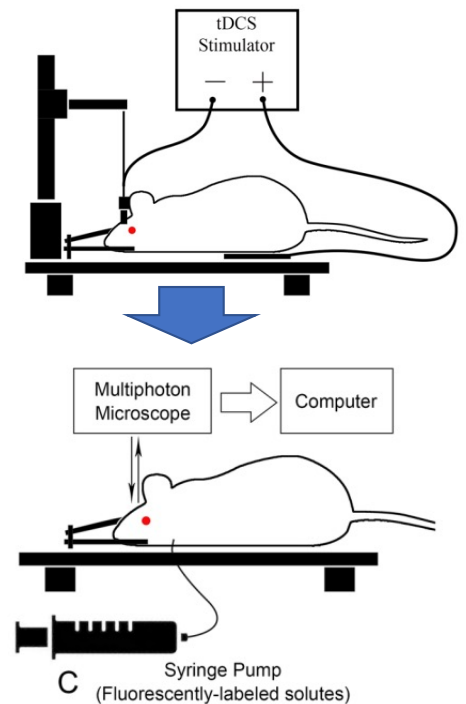
- Cancel et al. DCS of endothelial monolayers induces a transient and reversible increase in transport due to electroosmotic. *Sci Reports* 2019
- Shin et al. In Vivo Modulation of the Blood-Brain Barrier Permeability by tDCS. *Ann Biomed Eng.* 2020

Drive brain clearance (eg. dementia)

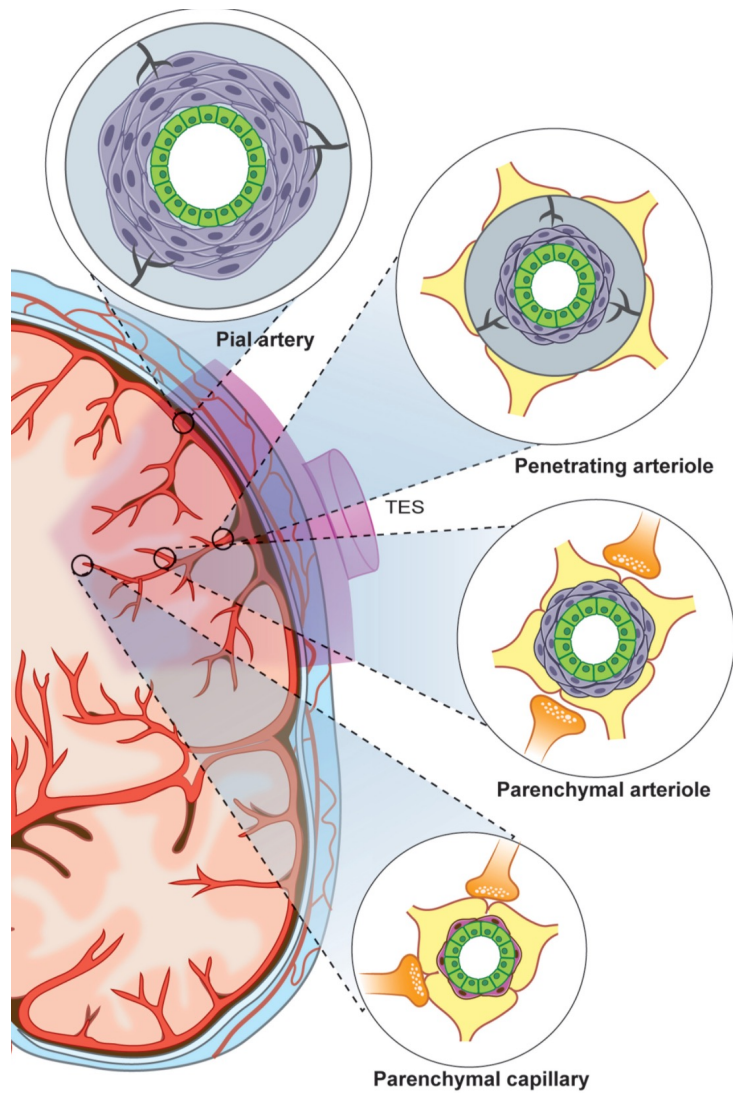
- Khadka et al. Neurocapillary-modulation. *Neuromodulation.* 2020
- Xia et. al Modulation of solute diffusivity in brain tissue as a novel mechanism of transcranial direct current stimulation (tDCS). *Sci Rep* 2020

Neuro-protective role (acute stroke)

- Bahr Hosseini et al. Central Nervous System Electrical Stimulation for Neuroprotection in Acute Cerebral Ischemia: Meta-Analysis of Preclinical Studies. *Stroke* 2019



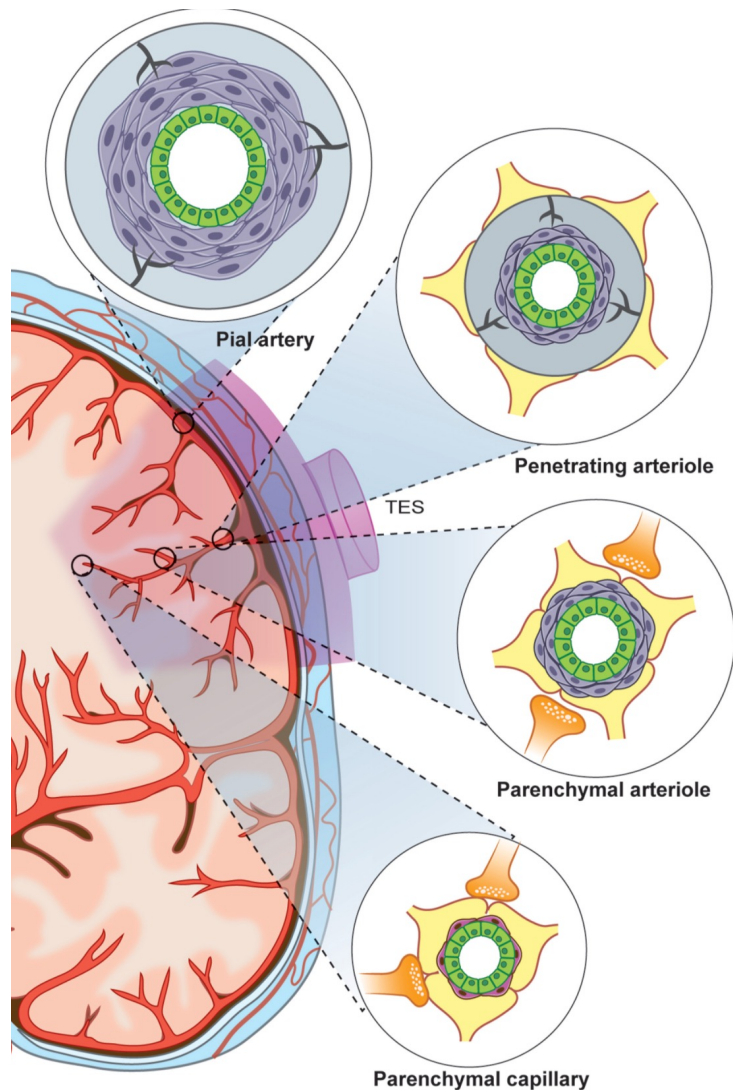
Why neurovascular modulation?



Bahr-Hosseini et al. Neurovascular-modulation. *Brain Stim* 2021

Why neurovascular modulation?

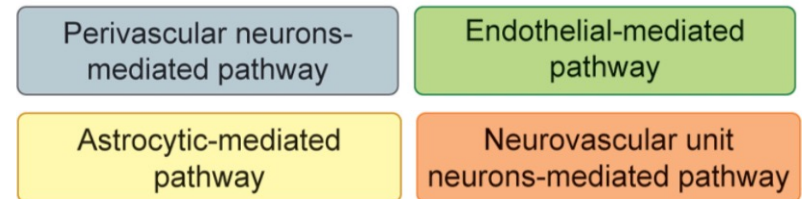
- Neurons are not alone in the brain. And are not functional without cells supporting transport.



● Dural Surface/Arteries
 ● Perivascular nerves
 ● Perivascular Space
 ● Smooth muscle cells
● Endothelium
 ● Astrocytic end feet
 ● Neurons of neurovascular unit
 ● Pericytes

FIRST ASPECT: Neuronal stimulation must consider neurovascular coupling.

SECOND ASPECT: Direct stimulation of brain vasculature.



Neurovascular-modulation

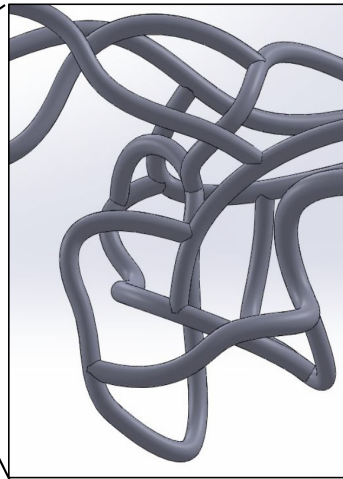
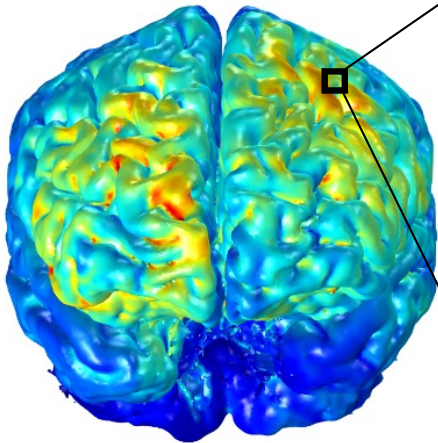
Third aspect: Fundamentally changing how neurons are directly stimulated

Endothelial-mediated pathway

Neurovascular modulation

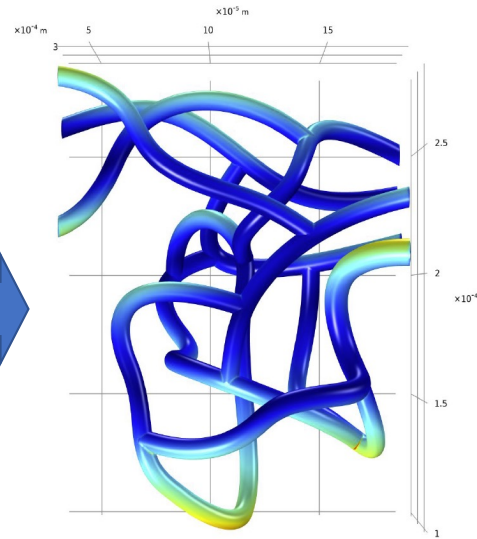
Macroscale (anatomy based) current flow models.

Brain parenchyma Electric Field :
(0.4 V/m at 1 mA tDCS)



Multi-scale models with brain vasculature structure.

Microscale current flow models. BBB Electric Field :
(160 V/m at 1 mA tDCS)



The structure of capillaries (extremely resistive wall, conductive interior) change microscopic current flow.

Electric fields are magnified across the Blood-Brain-Barrier (>400x of brain parenchyma).

For DBS /SCS /TMS/ ECT/ VNS: BBB Electric Fields >10,000 V/m

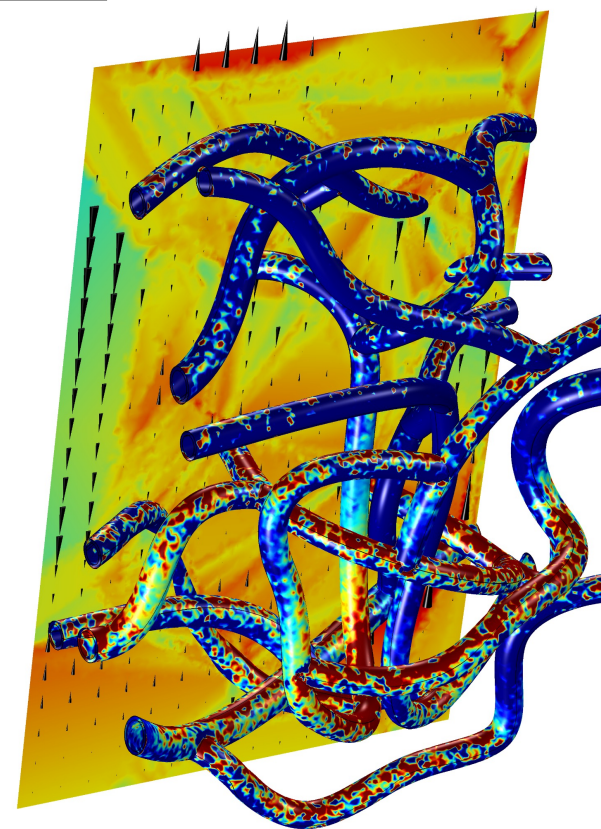
Multi-physics models couple to treatment mechanisms (eg. fluid clearance).

Things **Neuro-vascular Modulation** can explain

FIRST ASPECT: Brain stimulation (tDCS, TMS, ECT, DBS....) cannot significantly modulate neuronal function without engaging neuro-vascular coupling. Imaging based on hemodynamic coupling (fMRI...) **measure changes in neuro-vascular coupling**.

SECOND ASPECT: A direct vascular stimulation also likely. Isolated vascular (BBB) systems respond to stimulation in a dose / mechanisms / time-course specific manner. These dependencies can predict system / behavioral scale outcomes. And suggests **unique therapy strategies** (brain “flushing...”)

THIRD ASPECT: Reconsider how neuronal compartments or polarized. Impacts **neuronal sensitivity** (can provide “super-sensitivity” above traditional theory) and spatial distribution.



Multi-scale multi-physics model predict fluid “push” around brain during stimulation.