

**PSYCHIATRY ACADEMY** 

# Targeting brain circuits with non-invasive brain stimulation

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# Outline

- rTMS for Depression
- rTMS for OCD
- Combining Therapies to Improve Outcomes
- Theta Burst Stimulation
- Accelerated Protocols
- Transcranial Direct Current Stimulation



#### Disclosures

Neither I nor my spouse/partner has a relevant financial relationship with a commercial interest to disclose.





N ACCESS



#### <mark>eural circuits</mark>

Dysconnectivity of Multiple Brain Networks in Schizophrenia: A Meta-Analysis of Resting-State Functional Connectivity



Large-scale network dysfunction in Major Depressive Disorder: Meta-analysis of resting-state functional connectivity

Roselinde H. Kaiser, Ph.D.<sup>1,\*</sup>, Jessica R. Andrews-Hanna, Ph.D.<sup>2</sup>, Tor D. Wager, Ph.D.<sup>2</sup>, and Diego A. Pizzagalli, Ph.D.<sup>1</sup>

Frontoparietal areas link impairments of large-scale intrinsic brain networks with aberrant fronto-striatal interactions in OCD: a meta-analysis of resting-state functional connectivity

Deniz A. Gürsel<sup>a,b,\*,1</sup>, Mihai Avram<sup>a,b,1</sup>, Christian Sorg<sup>a,b,c</sup>, Felix Brandl<sup>a,b,2</sup>, Kathrin Koch<sup>a,b,2</sup>





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# Circuit-based Interventions: need to know...





# **Transcranial Magnetic Stimulation**

1831 Faraday's Electromagnetic Induction







# **Transcranial Magnetic Stimulation**

1831 Faraday's Electromagnetic Induction







#### What is Transcranial Magnetic Stimulation (TMS)?

- Safe
- Noninvasive
- Nonconvulsive
- Neuromodulation therapy
  - Changes neural excitability and activity



### **TMS** Theory

- Target treatment to a specific, affected region
- Changes spread to other regions
- Effects are network wide
- Repeated treatments lead to lasting effects



Liston 2014

### TMS Parameters

- 1) Location (low tech vs. neuronavigation)
- 2) Focality & Depth (coil selection)
- 3) Frequency (up- or downregulate)
- 4) Intensity (relative to stimulator or subject)
- 5) Duration (number of pulses / sessions)



#### **Current Therapeutic Uses**

#### **FDA Approved**

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- Unipolar Depression
  - Obsessive Compulsive Disorder

#### **Investigative**

- Auditory Hallucinations
- Post Traumatic Stress Disorder
- Generalized Anxiety Disorder
- Tourette Syndrome
- Bipolar Depression
- Autism

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• Neurorehabilitation

Migraine with Aura

- Parkinson Disease
- Alzheimer Disease
- Epilepsy
- Focal Dystonia
- Chronic Pain

#### TMS – Basic Equipment





MagVenture © System

Brainsway © System

### **Treatment Logistics**

- Remain **awake** during treatment
- No restrictions on activity
- Initial treatment course: five daily treatment per week (M-F) for 4-6 weeks
- •
- Taper period: 1-3 treatments per week
- Daily treatment duration: 3 30 minutes
- A tapping sensation is experienced
- A clicking noise accompanies each electromagnetic pulse

#### Magventure ©



#### Neural Networks Associated with Depression



### **Therapeutic applications: MDD**

• Early PET data argued for an overall hypofrontality and metabolic asymmetry in the two frontal areas

Depression Rx Strategy:

Left DLPFC: High Frequency (5-20 Hz)



Right DLPFC: Low Frequency (1 Hz)



### TMS Clinical Trials in MDD

- Multiple small single center trials since 1996
- Large multicenter trials in US leading to FDA approval in 2008 (O'Reardon et al., 2007)
- Follow up large NIMH trial confirms (George et al. 2010).
- Deep TMS (dTMS) system was granted FDA approval in 2013, after showing response rate of 38.4 % and remission rate of 32.6 % after 20 sessions.
- 7 companies have FDA-cleared devices for the treatment of MDD (6 Conventional rTMS systems and 1 dTMS system)



### TMS in the Treatment of Depression

• Conventional rTMS was FDA approved for the treatment of unipolar depression in 2008

#### Open label study of 10 Hz rTMS using conventional TMS device



 The H1 coil (deep TMS) was FDA approved for treatment of depression in 2013



#### Why Consider TMS treatment for Depression?

STAR\*D Study: Depression Treatment Outcomes



Likelihood of achieving remission drops with each subsequent medication trial

Rush AJ et al. Am J Psych 163:1905-1917, 2006

#### Potential Side Effects of TMS



# **TMS Safety**



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#### **Current Therapeutic Uses**

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### **OCD** Targets

#### OCD has a well-defined neurologic basis:

- The Cortical Striatal Thalamic

   Cortical pathway is a brain
   circuit that controls movement
   execution, habit formation, and
   reward.
- OCD is associated with hyperactivity of this pathway
- Poor thalamic gating may increase anterior cingulate cortex activity
- Medial prefrontal stimulation decreases anterior cingulate cortex activity



#### Medial prefrontal cortex/Anterior Cingulate Cortex

#### dTMS outcomes for OCD after 6 weeks of treatment



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Unpublished Information From Brainsway Website:

https://www.brainsway.com/treatments/obsessive-compulsive-disorder

www.mghcme.org

### OCD Symptoms Must Be Provoked!

- Provocation consists of internal or external stimuli which will provoke or induce typical OCD symptoms and distress the subject – lasts up to 5 minutes
- The goal is to induce a moderate-to-major distress immediately before initiating TMS

How much does the script/photo cause you distress right now?



### Enhancing the effect of TMS?

![](_page_24_Figure_1.jpeg)

Idea: TMS + Second Therapy = **Synergistic Effects** 

Activating a network with a task -> Increases susceptibility of network to the changes introduced by TMS

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www.mghcme.org

#### **State Effects: Simultaneous TMS + Therapy**

![](_page_25_Picture_1.jpeg)

Donse et al. 2018

-None

![](_page_25_Figure_3.jpeg)

#### **State Effects: Mood Alteration + TMS**

Prior to dTMS for depression subjects randomized to:

–Positive cognitive emotional reactivation

–Negative cognitive emotional reactivation

![](_page_25_Figure_8.jpeg)

![](_page_25_Picture_9.jpeg)

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Isserles et al, 2011

### Medications

- Alter physiology:
  - − Excitability → Affects Motor Threshold!
  - − Plasticity → Affect Treatment Efficacy

Might continuing medication help the efficacy of treatment?

![](_page_26_Figure_5.jpeg)

Augmentation of medication management with rTMS in treatment-

MASSACHUSETTS GENERAL HOSPITAL resistant depression leads to significant symptom improvement

### **Medications Effects on Treatment Outcomes**

![](_page_27_Figure_1.jpeg)

**Medications May Impact Response** 

![](_page_27_Picture_3.jpeg)

Hunter et al (2019) Brain and Behavior

### **Medication Effects on Treatment Outcome**

Patients taking anticonvulsants had a *faster* rate of response than those not taking anticonvulsants.

There was not significant difference between response and remission rates between those taking anticonvulsants and those not taking anticonvulsants.

![](_page_28_Figure_3.jpeg)

![](_page_28_Figure_4.jpeg)

Unpublished data from our clinic

![](_page_28_Picture_6.jpeg)

### **Theta Burst Stimulation**

![](_page_29_Figure_1.jpeg)

- Shorter duration
- May allow more sessions per day
- Longer-lasting physiological and cognitive effects are established in mechanistic studies

![](_page_29_Picture_5.jpeg)

#### Standard 10 Hz vs iTBS

#### TBS is an FDA approved treatment protocol that takes ~3 minutes to administer!

Parameters	10 Hz	iTBS
Train Duration	4 seconds	2 seconds
Inter-Train Interval	26 seconds	8 seconds
Total Pulses	3000	600
Total Treatment Duration	27 min 30 sec	3 min 9 sec
Frequency	120% resting MT	120% resting MT

N = 385

![](_page_30_Figure_4.jpeg)

#### Adapted from Blumberger et al. 2018; The Lancet www.mghcme.org

#### **Accelerated Protocols**

Accelerated iTBS treatment of depression in an inpatient setting

![](_page_31_Figure_2.jpeg)

- Each patient received 10 iTBS treatments per day
- Number of pulses delivered to in 1 day of treatment = standard treatment course.

HAMD-6

- Response Rate = 87.1%
- Remission Rate = 83.9%

#### MADRS

- Response Rate = 90.3%
- Remission Rate = 90.3%

Are safe and can shorten the duration of treatment!

![](_page_32_Figure_0.jpeg)

Patients with more treatment resistant depression may need more time to achieve response

Cole et al, 2019 (unpublished)

### **Transcranial Direct Current Stimulation**

![](_page_33_Picture_1.jpeg)

- Continuous low amplitude electrical current is delivered to a specified cortical regions using scalp electrodes
- <u>Anodal Stimulation</u>: Increases cortical excitability via depolarization of neuronal membrane potential
- <u>Cathodal Stimulation</u>: Decreases cortical excitability via hyperpolarization of neuronal membrane potential
- Repeated use may lead to neural plasticity
- Voltage: 2 mA over 30 minutes
- NOT FDA APROVED

![](_page_33_Picture_8.jpeg)

### **Transcranial Direct Current Stimulation**

![](_page_34_Picture_1.jpeg)

#### Advantages:

- Easy to use
- Inexpensive
- Safe
- Potential for Home Use

#### Recent meta-analysis of 7 studies in Bipolar Depression

- Standardized Mean Difference after acute phase: 0.71
- Standardized Mean Difference after furthest endpoint from treatment: 1.97

#### May be good option for bipolar depression

Donde et al. 2017

![](_page_34_Picture_12.jpeg)

### Thank you for your attention!

![](_page_35_Picture_1.jpeg)