Abstracts

Decreased occipital lobe metabolism by FDG-PET/CT: An anti-NMDA receptor encephalitis biomarker

Objective To compare brain metabolism patterns on FDG-PET/CT in anti-NMDAR and other definite autoimmune encephalitides (AE) and assess how these patterns differ between anti-NMDAR neurologic disability groups.

Methods This was a retrospective review of clinical data and initial dedicated brain FDG-PET/CT studies for neurology inpatients with definite AE, per published consensus criteria, treated at a single academic medical center over a 10-year period. Z score maps of FDG-PET/CT scans were made using 3D stereotactic surface projections with comparison to age group–matched controls. Brain region mean Z scores with magnitudes >2.00 were interpreted as significant. Comparisons were made between anti-NMDAR and other definite AE patients as well as among anti-NMDAR patients based on modified Rankin Scale scores (mRS) at time of FDG-PET/CT.

Results The medial occipital lobes were markedly hypometabolic in 6 of 8 anti-NMDAR patients and as a group (Z = −4.02, interquartile range [IQR] 2.14) relative to those with definite AE (Z = −2.32, 1.46; p = 0.004). Among anti-NMDAR patients, the lateral and medial occipital lobes were markedly hypometabolic for patients with mRS 4–5 (lateral occipital lobe Z = −3.69, IQR 1; medial occipital lobe Z = −4.08, 1) compared to those with mRS 0–3 (lateral occipital lobe Z = −0.83, 2; p < 0.0005; medial occipital lobe Z = −1.07, 2; p = 0.001).

Conclusions Marked medial occipital lobe hypometabolism by dedicated brain FDG-PET/CT may serve as an early biomarker for discriminating anti-NMDAR encephalitis from other AE. Resolution of lateral and medial occipital hypometabolism may correlate with improved neurologic status in anti-NMDAR encephalitis.

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Safety and preliminary efficacy of deep transcranial magnetic stimulation in MS-related fatigue

Objective This was a randomized, sham-controlled phase I/IIa study investigating the safety and preliminary efficacy of deep brain H-coil rTMS over the prefrontal cortex (PFC) and the primary motor cortex (MC) in patients with multiple sclerosis (MS) with fatigue or depression (NCT01106365).

Methods Thirty-three patients with MS were recruited to undergo 18 consecutive rTMS sessions over 6 weeks, followed by follow-up assessments over 6 weeks. Patients were randomized to receive high-frequency stimulation of the left PFC, MC, or sham stimulation. Primary endpoint was the safety of stimulation. Preliminary efficacy was assessed based on changes in Fatigue Severity Scale (FSS) and Beck Depression Inventory scores. Randomization allowed analysis of preliminary efficacy for fatigue only.

Results No serious adverse events were observed. Five patients terminated participation during treatment due to mild side effects. Treatment resulted in a significant median FSS decrease of 1.0 point (95% confidence interval 0.45, 1.65), which was sustained during follow-up.

Conclusion H-coil rTMS is safe and well tolerated in patients with MS. The observed sustained reduction in fatigue after subthreshold motor cortex stimulation warrants further investigation.

Classification of evidence This study provides Class III evidence that rTMS of the prefrontal or primary motor cortex is not associated with serious adverse effects, although the study is underpowered to state this with any precision.

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