Active Research Studies at Corewell Health West

Neurology	
Epilepsy	
RD2022-0122 NAUTILUS NCT05147571	RNS® System Responsive Thalamic Stimulation for Primary Generalized Seizures Study (NAUTILUS Study)
Neurodegenerative	
Healey ALS	Platform Trial for the Treatment of Amyotrophic Lateral Sclerosis (ALS): A perpetual multi-center, multi-regimen clinical trial evaluating the safety and efficacy of investigational products for the treatment of ALS
Neurosurgery	
Impact of BDNF Variants on Deep Brain Stimulation outcomes	TRAC - This study is designed to assess the safety and effectiveness of the ThrombX Retriever in revascularizing the site of primary occlusion in patients presenting with acute ischemic stroke secondary to intracranial, large vessel, occlusive disease
Neurovascular	
Distal Ischemic Stroke Treatment with Adjustable Low-profile Stentriever (DISTALS Study)	The objective of the randomized DISTALS Study is to evaluate the safety and effectiveness of the Tigertriever 13 Revascularization Device in restoring blood flow in the neurovasculature by removing thrombus in subjects presenting within 24 hours of onset with an ischemic stroke with disabling neurological deficits due to a primary distal vessel occlusion (DVO), as compared to medical management.
Stroke	
RP2021-0488 Genentech HAStE	Increase stroke awareness among the at-risk population as defined by validated risk score. Stroke awareness will significantly increase following the implementation of stroke risk intervention program(s) of this project, in regional communites.
Imperative	A prospective, multi-center, open label and single arm clinical investigation to evaluate the safety and efficacy of using the Zoom Reperfusion System in thrombectomy procedures to treat acute ischemic stroke patients
EMBOLISE NCT04402632	Embolization of the Middle Meningeal Artery with ONYXTM Liquid Embolic System In the Treatment of Subacute and Chronic Subdural HEmatoma (EMBOLISE)

For information about research studies listed please use these contacts below: Email: researchassist@corewellhealth.org

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RAGE	Second-generation hydrogel coils (HydroFrame®, HydroFill®, HydroSoft®, HydroSoft® 3D) used in any combination comprising a minimum of 90% total coil length used. Future hydrogel coils approved by the FDA (or applicable regulatory body) may also be included upon availability. Objective To determine occlusion rates and safety when hydrogel coils are used in the treatment of ruptured intracranial aneurysms.
Timeless	This study will evaluate the efficacy and safety of tenecteplase compared with placebo inpatients with acute ischemic stroke (Al.S)