

Lipid Profiles and Associated Symptoms and Outcomes in Acute Ischemic Stroke Patients
Sarah Martha, PhD, RN
University of Illinois at Chicago, Chicago, Illinois, USA
Sigma/Western Institute of Nursing Research Grant

Aim/Purpose/Objective: The purpose of this study was to determine the feasibility of a 1-month protocol and to describe self-reported symptoms and their association to cognitive and functional outcomes in persons with acute ischemic stroke (AIS) who did and did not receive reperfusion intervention (e.g., tPA, mechanical thrombectomy). We explored these factors and their association to underlying lipid biomarker signatures.

Sample: We enrolled 30 participants post-AIS categorized into reperfusion intervention and non-reperfusion intervention groups (n=15/each) during hospital admission (inpatient). Peripheral plasma samples were collected for lipidomic analysis (study day 5 and 1-month). Self-reported symptoms, cognitive and functional outcome data were collected (study day 5 and 1-month) from patients.

Setting: For this 1-month prospective study, a convenience sample of 30 AIS survivors were recruited from UI Health Hospital in the acute and critical-care units.

Methodology: Quantitative, Observational/Participant Observation, Physiological Data, a Prospective Cohort Study

Results: This protocol was feasible as evidence by 70% of participants completed the questionnaires and blood draws. Higher fatigue and anger symptoms were associated with poorer cognitive outcomes. Oxidative stress pathways were associated with higher symptoms and poorer cognitive outcomes in AIS participants. Information on the participants' characteristics are summarized in Table 1.

Conclusions: Analyzing plasma lipid markers may provide lipidomic signatures useful in predicting the development of symptoms as well as cognitive and functional outcomes following acute ischemic stroke.

Implications: This pilot study evaluated the feasibility and acceptability of our research protocol and will lead to the submission of an R-series grant. The long-term goal is to understand the underlying pathophysiology of AIS, in order to develop additional interventions (e.g., dietary, physical activity) to improve symptoms and outcomes for AIS survivors.

References:

1. Benjamin EJ, Muntner P, Alonso A, et al. Heart Disease and Stroke Statistics-2019 Update: A Report From the American Heart Association. *Circulation*. 2019;139(10):e56-e528.
2. Doyle KP, Simon RP, Stenzel-Poore MP. Mechanisms of ischemic brain damage. *Neuropharmacology*. 2008;55(3):310-318.

3. Adibhatla RM, Hatcher JF. Role of Lipids in Brain Injury and Diseases. *Future Lipidol.* 2007;2(4):403-422.
4. Muralikrishna Adibhatla R, Hatcher JF. Phospholipase A2, reactive oxygen species, and lipid peroxidation in cerebral ischemia. *Free Radic Biol Med.* 2006;40(3):376-387.
5. Phillis JW, O'Regan MH. The role of phospholipases, cyclooxygenases, and lipoxygenases in cerebral ischemic/traumatic injuries. *Crit Rev Neurobiol.* 2003;15(1):61-90.
6. Tobin MK, Bonds JA, Minshall RD, Pelligrino DA, Testai FD, Lazarov O. Neurogenesis and inflammation after ischemic stroke: what is known and where we go from here. *J Cereb Blood Flow Metab.* 2014;34(10):1573-1584.
7. Fiorella DJ, Fargen KM, Mocco J, et al. Thrombectomy for acute ischemic stroke: an evidence-based treatment. *J Neurointerv Surg.* 2015;7(5):314-315.
8. Campbell BC, Donnan GA, Lees KR, et al. Endovascular stent thrombectomy: the new standard of care for large vessel ischaemic stroke. *Lancet Neurol.* 2015;14(8):846-85.
9. Tissue plasminogen activator for acute ischemic stroke. The National Institute of Neurological Disorders and Stroke rt-PA Stroke Study Group. *N Engl J Med.* 1995;333(24):1581-1587.
10. Singh A. Tools for metabolomics. *Nature Methods.* 2020;17(1):24-24.