Area of study	Cognition and Activities of Daily Living	
Name of trial	Tele-rehabilitation for cognitive disability post-stroke: Enhancing function in the face of geographical disparities	
Study Type	Intervention	
Description of trial	This study is a single-blind (assessor), randomized wait-list control trial with a one-month follow-up. All testing and intervention sessions will be delivered online. Stroke results in long-term disability for many people, and particularly for those with cognitive impairment. Yet, access to rehabilitation services for survivors is declining: the 2014 Ontario Stroke Evaluation Report identified that the proportion of stroke patients receiving publicly funded rehabilitation declined to 51.3% in 2011-13. Data from the Cardiovascular Health Study (n=5,888) demonstrates an annual increase in the trajectory of disability post-stroke. The Ontario Stroke Network reported that to achieve best practice standards, 2-3 outpatient or community-based health professional visits for 8-12 weeks should be provided: 16-36 visits and not the current average of 5.8. To rectify the situation, an estimated 1,700 additional stroke patients require these services. It behooves us to consider cost-effective ways to deliver these services particularly in geographically remote areas.	
Objective of the Study	The objectives of the study are to investigate the feasibility and preliminary efficacy of delivering a rehabilitation intervention shown to reduce disability post-stroke via tele-rehabilitation.	
Eligibility requirements	**Who can apply** (inclusion criteria) • Community-dwelling adults at least three months post-stroke • Fluent in written and spoken English • Impairment of cognitive functions • Ability to self-identify specific areas of difficulty in their everyday life that they would like to improve • Access to a computer or tablet with a high-speed internet connection and webcam	**Who can not apply** (exclusion criteria) • Presence of dementia • Severe concurrent depression • Severe aphasia • Concurrent substance abuse

Study Commitment	Participants will receive 16 one-hour therapy sessions over a period of 10 weeks. Prior to and following the therapy sessions participants will participate in test sessions involving answering questionnaires and some thinking tests. These will each last about 2 hours and will be done over the phone and online. The total commitment for the study is 3-5 months. Participants will be reimbursed for time spent in testing.
Location of trial	Therapy and testing sessions are conducted online via the computer program SKYPE™
Funder	Heart and Stroke Foundation Canadian Partnership for Stroke Recovery (CPSR)
Contact information	Study Contact Information: **PI: Deirdre R. Dawson, PhD, OT Reg (ON)** Senior Scientist, Rotman Research Institute, Baycrest 3560 Bathurst St., Toronto, ON, M6A 2E1 Phone: 416-785-2500 x 2136 Email: ddawson@research.baycrest.org **Study coordinator: Yael Bar MSW, RSW** Lab Manager, Rotman Research Institute 3560 Bathurst St. Toronto, ON, M6A 2E1 Phone: 416 785 2500, ext 3377 Email: ybar@research.baycrest.org General Information: If you are interested in participating in or learning more about this study, please email research@canadianstroke.ca or call 416-480-6100 ext. 80941