Diagnosing Frontotemporal Lobar Degeneration

NCT02964637

Status	RECRUITING
Sponsor	University Health Network, Toronto
Enrollment	100 participants

Key Eligibility Criteria

Inclusion (3)

- Participant must have a reliable study partner who can provide an independent evaluation of functioning.
- Able to read, understand and speak English for neuropsychological testing.
- Control subjects must have a normal neurological exam, a CDR sum of boxes = 0, and MMSE score equal to or greater than 28

Exclusion (4)

- Patients with clinical, imaging or CSF A beta/ tau profile consistent with AD
- History of traumatic brain injury, brain tumors, stroke or other neurological or psychiatric disorders that can explain symptoms will be excluded.
- Premenopausal women will be asked to consent to a pregnancy test prior to each scan as pregnant women will be excluded from study because of potential harm to fetus from PET study.
- Presence of pacemakers, aneurysm clips, artificial heart valves, ear implants, metal fragments or foreign objects in the eyes, skin or body.

Locations (1 total)

Toronto Western Hospital, University Health Network, Toronto, Ontario, Canada