Systemic Biomarkers of Brain Injury From Hyperammonemia

NCT04602325

Status	RECRUITING
Sponsor	Children's National Research Institute
Enrollment	24 participants

Key Eligibility Criteria

Inclusion (25)

- Inherited Hyperammonemias:
- A clinical diagnosis of 1 of 7 diagnosed urea cycle disorders:
- N-acetylglutamate Synthetase Deficiency (NAGS)
- Carbamyl Phosphate Synthetase Deficiency (CPSD)
- Ornithine Transcarbamylase Deficiency (OTCD)

... and 20 more (see full listing online)

Exclusion (2)

- Prior Solid-Organ Transplant
- Use of any other investigational drug, biologic, or therapy or any clinical or laboratory abnormality or medical condition that, as determined by the investigator, may interfere with or obscure the biomarker measurements

Locations (1 total)

Children's National Research Institute, Washington D.C., District of Columbia, United States