Characterization and Natural History of Williams Syndrome and Other Chromosome 7q11.23 Variants

NCT06930417

Status	RECRUITING
Sponsor	University of Pennsylvania
Enrollment	2,000 participants

Key Eligibility Criteria

Inclusion (4)

- clinical and/or molecular diagnosis of Williams syndrome (WS)
- biological parents or siblings of individuals diagnosed with WS
- molecular diagnosis of 7q11.23 duplication syndrome (Dup7)
- molecular diagnosis of another abnormality in the 7q11.23 region

Exclusion (1)

- \- No diagnosis of abnormalities in the 7q11.23 region, while not being a biological relative of affected individuals

Locations (1 total)

University of Pennsylvania, Philadelphia, Pennsylvania, United States

DISCLAIMER: This document is for informational purposes only and does not constitute medical advice. Always consult your healthcare provider before enrolling in any clinical trial. Information may not be up to date — verify details at ClinicalTrials.gov. Generated by ClinicalTrialsFinder.org.