



**THIS STUDY IS BEING LED BY  
DR. RUJUTA B. WILSON, MD, MS AT UCLA**

**FOXP1 SYNDROME:  
UNDERSTANDING THROUGH NEURODEVELOPMENTAL DEEP  
PHENOTYPING TO SUPPORT CLINICAL TRIALS**

# THE FOUND STUDY



We are seeking to fill important gaps in the knowledge about the neurodevelopmental manifestations of FOXP1 syndrome and how they change in early childhood.

**YOUR FAMILY MAY BE ELIGIBLE FOR THIS STUDY IF YOU OR YOUR CHILD HAS A DIAGNOSIS OF FOXP1 SYNDROME AND:**

- Is between 1 to 20 years old
- Live anywhere in North America

**PARTICIPATION INCLUDES**

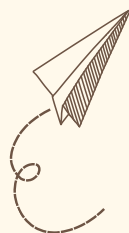
- Behavioral/Cognitive Assessments
- Neurological/Physical Exam
- Gait/Upper Extremity Assessments
- Parent Interview/Questionnaires



Opportunity  
for a 6-month  
and 12-month  
follow up

**YOU WILL RECIEVE**

- \$50 gift card upon baseline completion
- Clinical feedback



**CONTACT US**

[found-study@mednet.ucla.edu](mailto:found-study@mednet.ucla.edu)