





# Speech Tracker: Research into speech, language, and non-verbal communication development over time in individuals with *FOXP1*- related disorder

This study builds on our previous research showing that speech and language is a core developmental challenge for children with *FOXP1*-related disorder (Braden et al., 2021, Developmental Medicine & Child Neurology)

In this new study, we are examining communication development <u>over time</u>. This data will help us better understand prognosis and will help us develop more targeted speech therapies.

# We are looking for individuals:

- Confirmed to have a FOXP1-related disorder by a genetic test
- Aged 6 months to adulthood
- Who are verbal or non-verbal
- Who speak English, German, Dutch, Italian, French, Portuguese, Spanish

## What is involved?

- Enrol using the QR code or email us at: speechtracker@mcri.edu.au
- Completing brief online surveys of language skills once a year (families who have taken part in our previous research may have already completed some of the required surveys, and these can be reused for this study)
- Completing a quick 15 minute speech task from home twice a year

### **About us**

The Translational Centre of Research Excellence for Speech Disorders examines speech and language in individuals with rare genetic conditions.

We are trying to better understand the strengths and difficulties of communication in children with rare genetic syndromes. Our longer-term aim is to help develop better targeted therapies.

### **GET IN TOUCH!**

If you or someone you know may be interested in helping with this research, we would love to hear from you. Contact us at:

speechtracker@mcri.edu.au

Or scan the QR code to register!