6 KEY POINTS FROM THE HUWE1 DATA COLLECTION PROGRAM 2025

HUWE1 + RARE-X

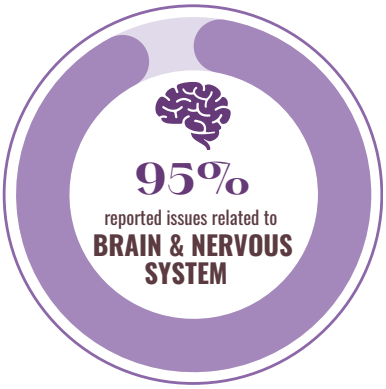
WHAT IS THE HUWE1 RARE-X DATA COLLECTION PROGRAM?

Louie's Huwe, the 501c3 non-profit dedicated to furthering research into HUWE1-related genetic conditions, has partnered with RARE-X to build a Data Collection Program for the HUWE1 community. Patients and caregivers have taken part in **self-report surveys** to help understand symptoms, provide opportunities to participate in research/clinical trials and pave the way to identifying treatments. As of **January 2025**, we currently have **78 enrolled** and **27 who have provided full genetic details** in reports/clinic notes*. Thank you to everyone who has participated! To learn more visit huwe1.rare-x.org.

LOUIE'S HUWE – FOR HUWE1-RELATED GENETIC CONDITIONS

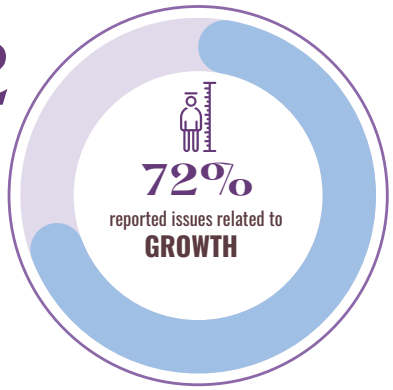
* **Note**: this data is self-reported by participants and the genetic data/variants for each participant have not been verified or classified by researchers or clinicians.

1



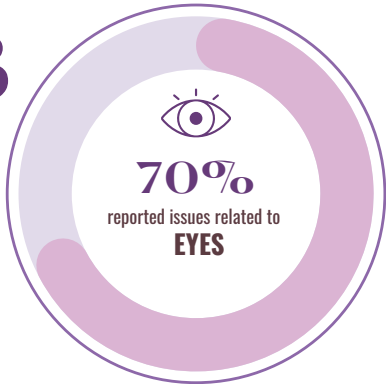
MOST COMMON ISSUES: COGNITIVE IMPAIRMENT, HYPOTONIA, COORDINATION, ABNORMAL EEG, SEIZURES

2



MOST COMMON ISSUES: SHORT STATURE, UNDERGROWTH

3

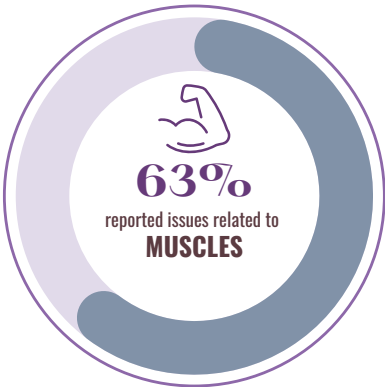


MOST COMMON ISSUES: ABNORMAL EYE MOVEMENT, FARSIGHTEDNESS, VISUAL IMPAIRMENT

4

MOST COMMON ISSUES: REPETITIVE BEHAVIOUR, UNUSUAL EYE GAZE, DIFFICULTY WITH NONVERBAL BEHAVIOR, AUTISM SPECTRUM, SHORT ATTENTION SPAN, MELTDOWNS, SELF INJURIOUS BEHAVIOR

5



MOST COMMON ISSUES: ABNORMAL MUSCLE FUNCTION, ARM/LEG MUSCLE ISSUES, NECK MUSCLE ISSUES

6

MOST COMMON ISSUES: CONSTIPATION, FEEDING DIFFICULTIES