INTERNATIONAL FOXP1 FOUNDATION™

RareX Data Analysis ~ Dec 2023

In 2023, thanks to the FOXP1 community's involvement in the RARE-X data collection program, we were able to participate in an initial data analysis effort to identify symptoms that are commonly reported by families but not represented in the literature as of mid-2023. See the graphic below. Although an excellent review published in late 2023 provides an updated, comprehensive review of FOXP1 (https://www.ncbi.nlm.nih.gov/books/NBK594825/), this data analysis nevertheless shows how our own lived experiences can improve the "official" understanding of FOXP1 syndrome.

Again, thank you for your participation in the data collection program; as always, the more we all participate (including with longitudinal surveys over time) the richer the dataset and more impactful the insights will be. If you have not yet registered for the RARE-X data collection program, or you have not checked your dashboard lately for new surveys to complete, please follow this link: https://foxp1.rare-x.org/