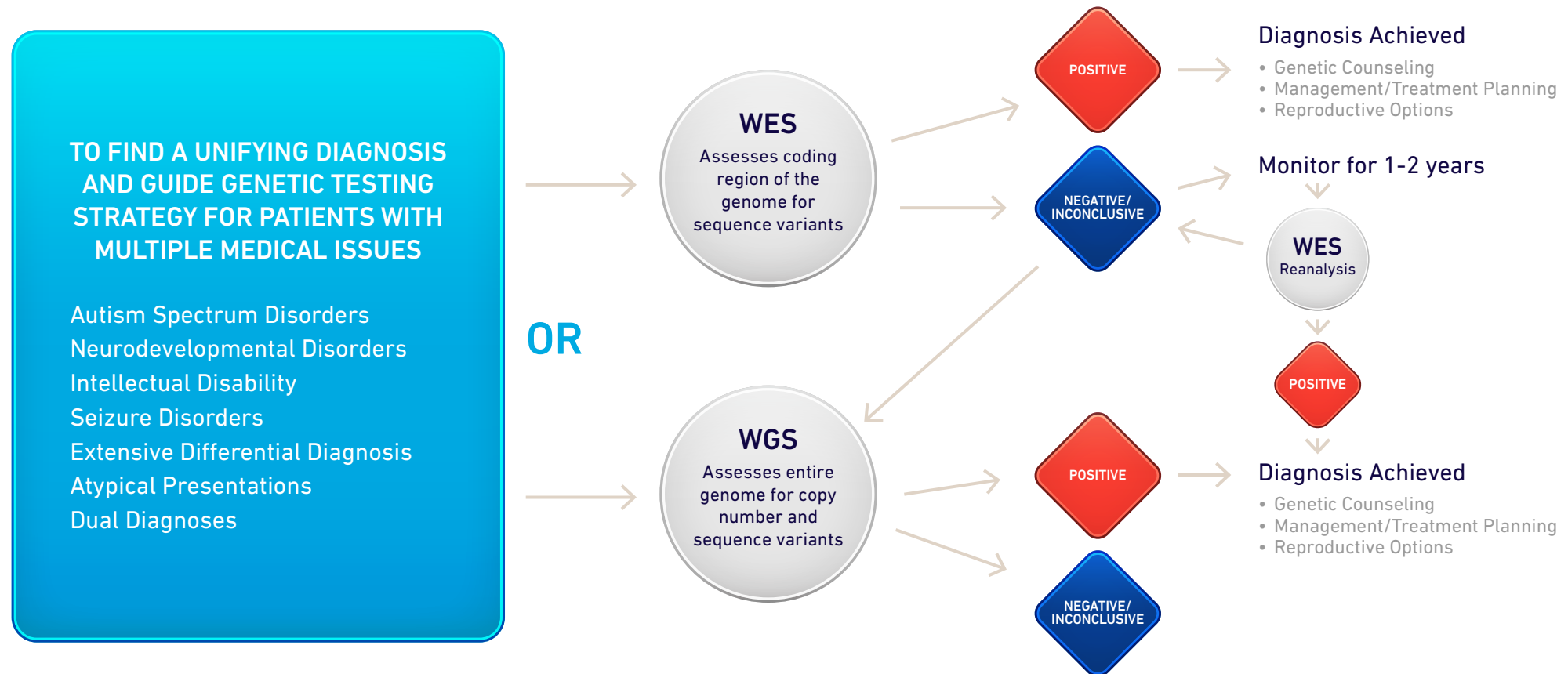


# Rare Disease Diagnosis Decision Support Tool:

## Whole Exome Sequencing (WES) and Whole Genome Sequencing (WGS)



Disclaimer: The above provides general recommendations for appropriate testing to consider based on presenting phenotype. Optimal test selection and strategy is best determined by the clinical health care provider. Metabolic, mitochondrial, and/or repeat expansion disorders may need an additional separate assessment.