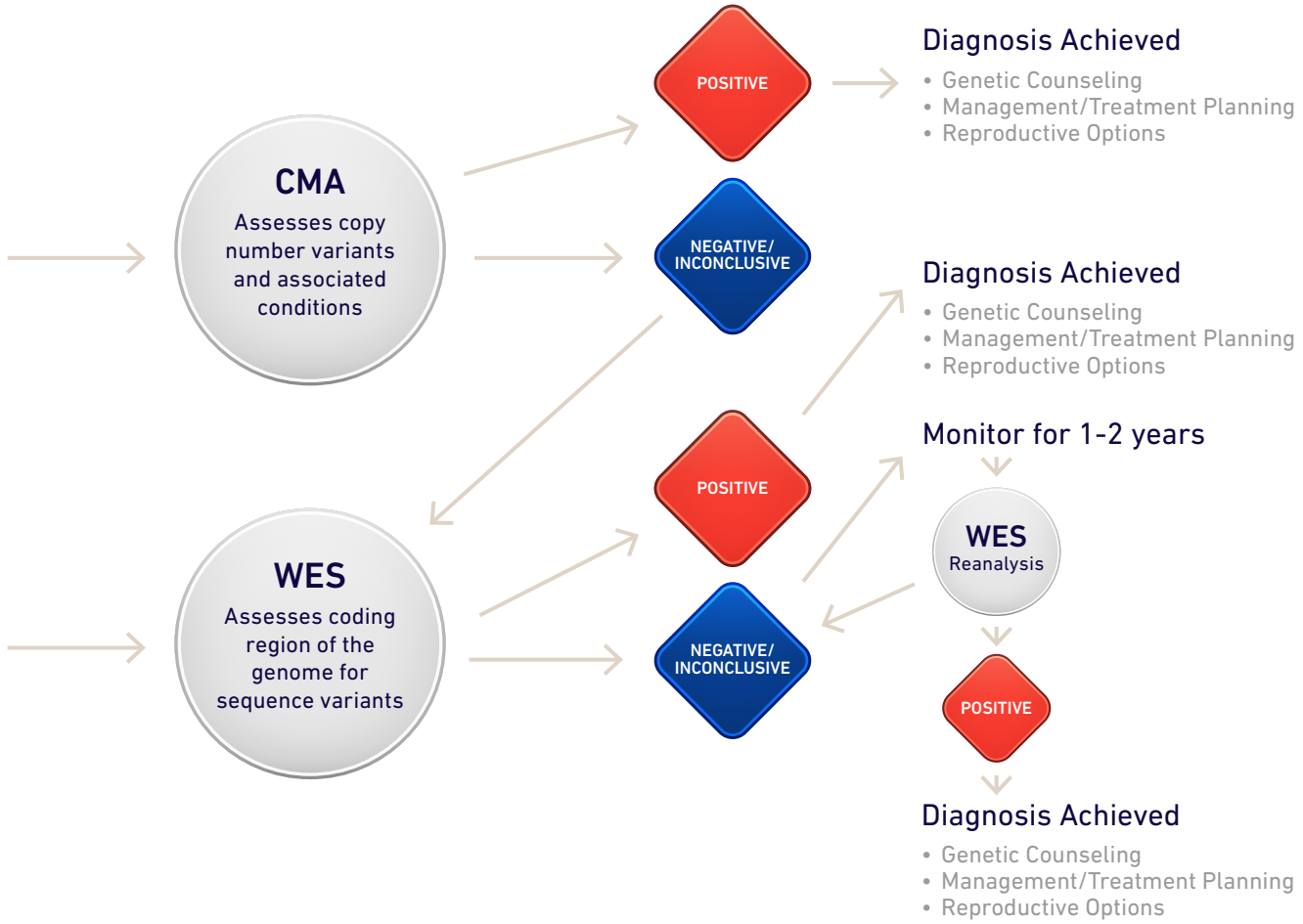
Rare Disease Diagnosis Decision Support Tool:

Chromosomal Microarray Analysis (CMA) & Whole Exome Sequencing (WES)

TO FIND A UNIFYING DIAGNOSIS AND GUIDE GENETIC TESTING STRATEGY FOR PATIENTS WITH MULTIPLE MEDICAL ISSUES

Multiple Congenital Anomalies
Failure To Thrive
Dysmorphic Features
Autism Spectrum Disorders
Neurodevelopmental Disorders



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.

BAYLOR GENETICS