MAYO CLINIC LABORATORIES

Targeted Genes and Methodology Details for Congenital Fibrinogen Disorders, *FGA*, *FGB*, and *FGG* Genes

The following applies to GNFIB / Congenital Fibrinogen Disorders, *FGA*, *FGB*, and *FGG* Genes, Next-Generation Sequencing. Testing is performed to evaluate for the presence of variants in coding regions and extending to +/- 10 base pairs of adjacent intronic sequence on either side of the coding exons of the genes analyzed. In addition, the analysis will cover select non-coding variants. Next-generation sequencing and/or a polymerase chain reaction-based quantitative method is performed to test for the presence of copy number variants in the genes analyzed. Confirmation of select reportable variants may be performed by alternate methodologies based on internal laboratory criteria.

This list is current from June 2023 to the present. This document is intended to highlight additional evaluations for variants of high clinical interest as well as technical limitations. However, this document does not comprehensively reflect all genomic regions covered by this test. For questions regarding transcripts, genes, or regions covered, contact the laboratory at 800-533-1710.

Genomic Build: GRCh37 (hg19) unless otherwise specified

Gene	Reference Transcript	Additional Evaluations	Technical Limitations
FGA	NM_000508.5	c.-202 to c.-1, c.1891+1 to c.1891+54	-
FGB	NM_005141.4	-	-
FGG	NM_000509.5	-	-
FGG	NM_021870.3	-	-