HOTLINE: Effective May 16, 2022

2013661 Cystic Fibrosis (*CFTR*) Expanded Variant Panel CF VAR

Specimen Required: Collect: Lavender (EDTA), pink (K_2EDTA).
Specimen Preparation: Transport 3 mL whole blood. (Min: 1 mL)
Storage/Transport Temperature: Refrigerated.
Unacceptable Conditions: Plasma or serum. Specimens collected in sodium heparin, yellow (ACD solution), or lithium heparin tubes. Frozen specimens in glass collection tubes.
Stability (collection to initiation of testing): Ambient: 72 hours; Refrigerated: 1 week; Frozen: 1 month

Interpretive Data:
Refer to report.

This test was developed and its performance characteristics determined by ARUP Laboratories. It has not been cleared or approved by the US Food and Drug Administration. This test was performed in a CLIA certified laboratory and is intended for clinical purposes.

Counseling and informed consent are recommended for genetic testing. Consent forms are available online.

Note: The Cystic Fibrosis (*CFTR*) Expanded Variant Panel includes the 23 pathogenic CF variants recommended by the American College of Medical Genetics for carrier screening as well as many more.

HOTLINE NOTE: There is a clinically significant charting name change associated with this test.
Change the charting name for component 2013677, CF Common Variants Interp from CF Common Variants Interp to CF Expanded Variant Panel Interp.