

**TEST CHANGE** 

Rapid Whole Genome Sequencing, Familial Control with Report 3005933 RWGS FRPT

3005933, RWGS FRPT	
Specimen Requirements:	
Patient Preparation:	
Collect:	Lavender or pink (EDTA) or yellow (ACD solution A or B).  Peripheral blood required. Contact ARUP's genetic counselor at 800-242-2787 ext. 2141 prior to test submission.
Specimen Preparation:	Transport 2 mL whole blood. (Min: 0.5 mL)
Transport Temperature:	Refrigerated.
Unacceptable Conditions:	
Remarks:	This test is used for parental control samples associated with a proband sample submitted for <a href="Rapid Whole Genome">Rapid Whole Genome</a> <a href="Sequencing">Sequencing</a> (ARUP test code 3005935). <a href="RWGS NGS">RWGS NGS</a> . A report will be provided for samples ordered using this test code. If a report for parental control sample is not desired, order <a href="Rapid Whole Genome Sequencing">Rapid Whole Genome Sequencing</a> , <a href="Familial Control">Familial Control</a> (ARUP test code 3005928). <a american="" and="" college="" genetics="" genomics"="" href="RWGS FAM (3005928)&lt;/a&gt;.&lt;/td&gt;&lt;/tr&gt;&lt;tr&gt;&lt;td&gt;Stability:&lt;/td&gt;&lt;td&gt;Ambient: 72 hours; Refrigerated: 1 week; Frozen: Unacceptable&lt;/td&gt;&lt;/tr&gt;&lt;tr&gt;&lt;td&gt;Methodology:&lt;/td&gt;&lt;td&gt;Massively Parallel Sequencing&lt;/td&gt;&lt;/tr&gt;&lt;tr&gt;&lt;td&gt;Performed:&lt;/td&gt;&lt;td&gt;Varies&lt;/td&gt;&lt;/tr&gt;&lt;tr&gt;&lt;td&gt;Reported:&lt;/td&gt;&lt;td&gt;5-7 days&lt;/td&gt;&lt;/tr&gt;&lt;tr&gt;&lt;td&gt;Note:&lt;/td&gt;&lt;td&gt;Parental samples are used to aid in interpretation of the proband's genome sequencing data. This test is ordered when a report of &lt;a href=" mailto:the="" medical="" of="">the American College of Medical Genetics and Genomics</a> (ACMG)ACMG secondary findings is desired for submitted parental controls. For each parental specimen, please indicate on the intake form that the sample is control and reference the patient's name.
CPT Codes:	NA
New York DOH Approval Status:	Specimens from New York clients will be sent out to a New York DOH approved laboratory, if possible.
Interpretive Data:	

Effective Date: February 20, 2024



Refer to report.

Reference Interval:

By report

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