

Familial Arrhythmia Panel

Order Name: Fam Arrhythmia

Test Number: 5194954
Revision Date: 03/21/2023

TEST NAME		METHO	DOLOGY	LOINC CODE	
Familial Arrhythmia Panel		Polymerase Chain Reaction			
SPECIMEN REQUIREMENTS					
SPECIMEN REQUIREM	MEN 13				
Specimen	Specimen Volume (min)	Specimen Type	Specimen Container	Transport Environment	
Preferred	8.5 mL (3 mL)	Whole Blood	ACD Solution A or B (Yellow Top)	Room Temperature	
Alternate 1	8.5 mL (3 mL)	Whole Blood	EDTA (Lavender Top)	Room Temperature	
Alternate 2	1	Saliva	Oragene Dx saliva kit	Room Temperature	
Alternate 3	1	Buccal swab	PurFlock buccal swab kit	Room Temperature	
Instructions	Specimen Type: Whole blood or PurFlock buccal swab kit or Oragene Dx saliva kit				
	Specimen Volume: 8.5 mL whole blood or PurFlock buccal swab kit or Oragene Dx saliva kit				
	Mininum Volume: 3 mL whole blood or PurFlock buccal swab kit or Oragene Dx saliva kit				
	Collection: Standard phlebotomy. Follow PurFlock buccal swab kit or Oragene Dx 500 saliva kit collection instructions. Do not eat, drink, smoke, or				
	chew gum 30 min prior to collection.				
	Specimen Storage: Maintain specimen at room temperature or refrigerate at 4C Do not freeze.				
	Special Instructions: In cases in which there is a known variant documented in the family, the physician may prefer to order Targeted Variant				

GENERAL INFORMATION			
Expected TAT	14 - 21 days In some cases, additional time may be required for confirmatory or reflex tests.		
Clinical Use	This test includes the following genes: AKAP9, ANK2, CACNA1C, CACNB2, CALM1, CALM2, CALM3, CASQ2, CAV3, CTNNA3, DES, DSC2, DSG2, DSP, FLNC, GJA5, GPD1L, HCN4, JUP, KCNA5, KCND2, KCND3, KCNE1, KCNE2, KCNE3, KCNE5, KCNH2, KCNJ2, KCNJ5, KCNJ8, KCNQ1, LIG3, NPPA, PKP2, PLN, PRKAG2, RANGRF, RYR2, SCN10A, SCN1B, SCN2B, SCN3B, SCN4B, SCN5A, SLMAP, SNTA1, TECRL, TGFB3, TMEM43, TRDN and TRPM4.		
Notes	Labcorp Test Code: 482225		
Service Provided By	Oklahoma, Inc.		

Analysis, test code 5194970. Test orders must include an attestation that the provider has the patient's informed consent for genetic testing.