

Congenital Heart Disease Gene Panel, Varies

Reporting Title: Congenital Heart Disease Gene Panel

Performing Location: Rochester

#### **Ordering Guidance:**

Chromosomal microarray is often used as a first-tier test in the setting of congenital heart disease. If chromosomal microarray testing is desired, order either CMACB / Chromosomal Microarray, Congenital, Blood or CMAP / Chromosomal Microarray, Prenatal, Amniotic Fluid/Chorionic Villus Sampling.

Customization of this panel and single gene analysis for any gene present on this panel are available. For more information see CGPH / Custom Gene Panel, Hereditary, Next-Generation Sequencing, Varies.

Targeted testing for familial variants (also called site-specific or known mutations testing) is available for the genes on this panel. See FMTT / Familial Variant, Targeted Testing, Varies. To obtain more information about this testing option, call 800-533-1710.

#### **Additional Testing Requirements:**

All prenatal specimens must be accompanied by a maternal blood specimen; order MATCC / Maternal Cell Contamination, Molecular Analysis, Varies on the maternal specimen as this must be a different order number than the prenatal specimen.

#### **Shipping Instructions:**

Specimen preferred to arrive within 96 hours of collection.

#### **Necessary Information:**

<u>Prior Authorization</u> is available, **but not required**, for this test. If proceeding with the prior authorization process, submit the required form with the specimen.

#### **Specimen Requirements:**

**Patient Preparation:** A previous bone marrow transplant from an allogenic donor will interfere with testing. For instructions for testing patients who have received a bone marrow transplant, call 800-533-1710.

#### Submit only 1 of the following specimens:

Specimen Type: Whole blood

Container/Tube:

Preferred: Lavender top (EDTA) or yellow top (ACD)

Acceptable: Any anticoagulant

Specimen Volume: 3 mL Collection Instructions:

1. Invert several times to mix blood.

2. Send whole blood specimen in original tube. **Do not aliquot.** 

Specimen Stability Information: Ambient (preferred)/Refrigerated

Due to its complexity, consultation with the laboratory is required for all prenatal testing; call 800-533-1710 to speak to a genetic counselor.



#### Congenital Heart Disease Gene Panel, Varies

Specimen Type: Amniotic fluid

Container/Tube: Amniotic fluid container

Specimen Volume: 20 mL

Specimen Stability Information: Refrigerated (preferred)/Ambient

**Additional information:** 

1. If amniotic fluid or nonconfluent cultures are received, CULAF / Culture for Genetic Testing, Amniotic Fluid will be added at an additional charge.

2. **All prenatal specimens must be accompanied by a maternal blood specimen;** order MATCC / Maternal Cell Contamination, Molecular Analysis, Varies on the maternal specimen.

Specimen Type: Chorionic villi

Container/Tube: 15-mL tube containing 15 mL of transport media

Specimen Volume: 20 mg

Specimen Stability Information: Refrigerated

**Additional Information:** 

- 1. If nonconfluent cultures are received, CULFB / Fibroblast Culture for Biochemical or Molecular Testing will be added at an additional charge.
- 2. **All prenatal specimens must be accompanied by a maternal blood specimen;** order MATCC / Maternal Cell Contamination, Molecular Analysis, Varies on the maternal specimen.

#### Acceptable:

Specimen Type: Confluent cultured cells

Container/Tube: T-25 flask Specimen Volume: 2 Flasks

**Collection Instructions:** Submit confluent cultured cells from another laboratory.

Specimen Stability Information: Ambient (preferred)/Refrigerated

Additional Information: All prenatal specimens must be accompanied by a maternal blood specimen; order MATCC /

Maternal Cell Contamination, Molecular Analysis, Varies on the maternal specimen.

#### Forms:

- 1. **New York Clients-Informed consent is required.** Document on the request form or electronic order that a copy is on file. The following documents are available:
- -Informed Consent for Genetic Testing (T576)
- -Informed Consent for Genetic Testing (Spanish) (T826)
- 2. Congenital Heart Disease Genetic Testing Patient Information
- 3. Congenital Heart Disease Gene Panel (CHDGG) Prior Authorization Ordering Instructions
- 4. If not ordering electronically, complete, print, and send a <u>Cardiovascular Test Request</u> (T724) with the specimen.

Specimen Type	Temperature	Time	Special Container
Varies	Varies		

#### **Result Codes:**

Result ID Reporting Name	Туре	Unit	LOINC®
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## Congenital Heart Disease Gene Panel, Varies

617198	Test Description	Alphanumeric	62364-5
617199	Specimen	Alphanumeric	31208-2
617200	Source	Alphanumeric	31208-2
617201	Result Summary	Alphanumeric	50397-9
617202	Result	Alphanumeric	82939-0
617203	Interpretation	Alphanumeric	69047-9
617204	Additional Results	Alphanumeric	82939-0
617205	Resources	Alphanumeric	99622-3
617206	Additional Information	Alphanumeric	48767-8
617207	Method	Alphanumeric	85069-3
617208	Genes Analyzed	Alphanumeric	48018-6
617209	Disclaimer	Alphanumeric	62364-5
617210	Released By	Alphanumeric	18771-6

LOINC® and CPT codes are provided by the performing laboratory.

#### **Supplemental Report:**

Supplemental

#### **CPT Code Information:**

81404

81405 x 3

81406 x 6

81407 x 3

81408

81479

81479 (if appropriate for government payers)

81265-Maternal cell contamination (if appropriate)

88233-Tissue culture, skin, solid tissue biopsy (if appropriate)

88235-Amniotic Fluid culture (if appropriate)

88240-Cryopreservation (if appropriate)

### **Reflex Tests:**

Test Id	Reporting Name	CPT Units	CPT Code	Always Performed	Available Separately
CULFB	Fibroblast Culture for Genetic Test	1	88233	No	Yes
CULAF	Amniotic Fluid Culture/Genetic Test	1	88235	No	Yes
MATCC	Maternal Cell Contamination, B	1	81265	No	Yes
_STR1	Comp Analysis using STR (Bill only)	1	81265	No	No, (Bill only)
_STR2	Add'l comp analysis w/STR (Bill Only)	1	81266	No	No, (Bill only)

#### **Result Codes for Reflex Tests:**



Congenital Heart Disease Gene Panel, Varies

Test ID	Result ID	Reporting Name	Туре	Unit	LOINC®
CULAF	52304	Result Summary	Alphanumeric		50397-9
CULAF	52306	Interpretation	Alphanumeric		69965-2
CULAF	52305	Result	Alphanumeric		82939-0
CULAF	CG767	Reason for Referral	Alphanumeric		42349-1
CULAF	52307	Specimen	Alphanumeric		31208-2
CULAF	52308	Source	Alphanumeric		31208-2
CULAF	52309	Method	Alphanumeric		85069-3
CULAF	54641	Additional Information	Alphanumeric		48767-8
CULAF	52310	Released By	Alphanumeric		18771-6
CULFB	52327	Result Summary	Alphanumeric		50397-9
CULFB	52329	Interpretation	Alphanumeric		69965-2
CULFB	52328	Result	Alphanumeric		82939-0
CULFB	CG770	Reason for Referral	Alphanumeric		42349-1
CULFB	CG899	Specimen	Alphanumeric		31208-2
CULFB	52331	Source	Alphanumeric		31208-2
CULFB	52332	Method	Alphanumeric		85069-3
CULFB	54625	Additional Information	Alphanumeric		48767-8
CULFB	52333	Released By	Alphanumeric		18771-6
MATCC	53285	Result Summary	Alphanumeric	phanumeric	
MATCC	53286	Result	Alphanumeric	phanumeric	
MATCC	53287	Interpretation	Alphanumeric		69047-9
MATCC	53288	Reason for referral	Alphanumeric	ımeric	
MATCC	53289	Specimen	Alphanumeric	Alphanumeric	
MATCC	53290	Source	Alphanumeric	Alphanumeric 31208	
MATCC	53291	Released By	Alphanumeric	Alphanumeric 18771-6	
MATCC	55150	Method	Alphanumeric		85069-3

### **Reference Values:**

An interpretive report will be provided