Client: Example Client ABC123
123 Test Drive
Salt Lake City, UT 84108
UNITED STATES

Physician: Doctor, Example

Patient: Patient, Example

DOB	[redacted]
Gender:	Male
Patient Identifiers:	01234567890ABCD, 012345
Visit Number (FIN):	01234567890ABCD
Collection Date:	00/00/0000 00:00

Medium Chain Acyl-CoA Dehydrogenase (ACADM) 2 Mutations

ARUP test code 0051205

MCADPCR Specimen	Whole Blood
MCAD Mutation A985G	Negative
MCAD Mutation T199C	Negative
Medium Chain Acyl-CoA Interpretation	See Note

Indication for testing: Carrier screening or diagnostic testing for MCAD deficiency.

Result
A985G: Negative
T199C: Negative

This sample is negative for the ACADM gene mutations, c.985A>G and c.199T>C. Persons affected with medium-chain acyl-CoA dehydrogenase (MCAD) deficiency may have rare mutations not detected by this assay. If the patient has biochemical and/or clinical evidence of MCAD deficiency, plasma acylcarnitine profile testing and/or ACADM gene sequencing is recommended.

This result has been reviewed and approved by [redacted], M.D., Ph.D.

H=High, L=Low, *=Abnormal, C=Critical

BACKGROUND INFORMATION: Medium Chain Acyl-CoA Dehydrogenase (ACADM) 2 Mutations

CHARACTERISTICS: Limited mitochondrial fatty acid beta-oxidation leading to hypoglycemia, lethargy, seizures, hypoketotic aciduria, vomiting, hepatomegaly, hepatic failure, encephalopathy, and sudden death. Manifestations often triggered by prolonged fasting or other metabolic stressors.
INCIDENCE: 1 in 15,000
INHERITANCE: Autosomal recessive.
CAUSE: Deleterious ACADM gene mutations.
CLINICAL SENSITIVITY: 75 percent for MCAD deficiency.
MUTATIONS TESTED: ACADM mutations c.985A>G (p.K329E, also known as K304E) and c.199T>C (p.Y67H, also known as Y42H).
METHODOLOGY: Polymerase chain reaction and fluorescence monitoring.
ANALYTICAL SENSITIVITY AND SPECIFICITY: 99 percent.
LIMITATIONS: Diagnostic errors can occur due to rare sequence variations. ACADM mutations other than c.985A>G and c.199T>C will not be detected.

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

VERIFIED/REPORTED DATES

Procedure	Accession	Collected	Received	Verified/Reported
MCADPCR Specimen	21-166-402415	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
MCAD Mutation A985G	21-166-402415	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
MCAD Mutation T199C	21-166-402415	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Medium Chain Acyl-CoA Interpretation	21-166-402415	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00

END OF CHART

H=High, L=Low, *=Abnormal, C=Critical