Medium Chain Acyl Co-A Dehydrogenase (MCADD): ACADM Gene Sequencing

Test Code: MV
Turnaround time: 4 weeks
CPT Codes: 81479 x1

Condition Description

Medium chain acyl-CoA dehydrogenase deficiency (MCADD) is an autosomal recessive disorder of fatty acid oxidation, the process by which the body metabolizes fats for energy in the absence of glucose. MCADD deficiency results in the inability to break down medium sized fatty acids (6-12 carbon atoms in length). As a result, these fatty acids accumulate mainly in the liver and to a lesser extent in the heart and kidneys.

MCADD generally presents between two months and two years of life, but can present as early as two days of life and as late as adulthood. Affected children are healthy and usually asymptomatic until symptoms are triggered by prolonged fasting or an illness that causes a decreased caloric intake, like the flu, a cold, or an ear infection. The inability to convert fats to energy can lead to:

- hypoglycemia
- vomiting
- lethargy
- coma
- apnea
- cardiac arrest
- sudden unexplained death

About 20-25% of MCADD patients die from their first symptomatic episode. MCADD is believed to account for up to 2.5% of sudden infant death syndrome (SIDS) cases.

MCADD results from mutations in the acyl-CoA dehydrogenase, medium-chain (ACADM) gene located on chromosome 1p31. The MCAD protein functions within the mitochondria at the first step in beta-oxidation of medium chain fatty acids. Sequencing of ACADM is recommended for patients with a biochemical diagnosis of MCADD as a complementary method to confirm the presence of mutations in a proband, to identify carriers among the proband's relatives, and to provide prenatal diagnosis in families with known mutations. Sequencing is recommended only after mutation analysis for the K304E and Y42H mutations has been performed. For patients with mutations not identified by full gene sequencing, a separate deletion/duplication assay is available using a targeted CGH array.

Reference:

Genes

ACADM

Indications

This test is indicated for:

- Patients who are found to have symptoms of MCADD (including elevated urine dicarboxylic acids and/or elevated medium chain acylcarnitines).
- Infants who are hypoglycemic, have unexplained seizures, or have a family history of SIDS (Sudden Infant Death Syndrome).
- Patients with symptoms of MCADD with only one/no mutations identified through the common mutation panel.

Methodology

The 12 coding exons and immediate flanking regions of ACADM are amplified by PCR and sequenced in both forward and reverse directions. The patient’s gene sequences are then compared to a normal reference sequence. Sequence variations are classified as previously described mutations, novel mutations, or variations of unknown significance. This analysis may detect novel variants of unclear effect that require further studies.

Detection

This assay will detect over 97% of the sequence variants in the coding region and splice junctions. Mutations in the promoter region, some mutations in the introns and other regulatory elements, large deletions, and insertion mutations will not be detected by this assay.

Results of molecular analysis should be interpreted in the context of the patient's biochemical phenotype.

Specimen Requirements

Submit only 1 of the following specimen types

* Preferred specimen type: Whole Blood

Type: Whole Blood

Specimen Requirements:
In EDTA (purple top) or ACD (yellow top) tube:
Infants (2 years): 3-5 ml
Older Children & Adults: 5-10 ml

Specimen Collection and Shipping: Refrigerate until time of shipment. Ship sample within 5 days of collection at room temperature with overnight delivery.

**Type: Saliva**

Specimen Requirements:

Oragene™ Saliva Collection kit (available through EGL) used according to manufacturer instructions.

Specimen Collection and Shipping: Store sample at room temperature. Ship sample within 5 days of collection at room temperature with overnight delivery.

**Special Instructions**

Please submit copies of diagnostic biochemical test results along with the sample. Contact the laboratory if further information is needed. Sequence analysis is required before deletion/duplication analysis by targeted CGH array. If sequencing is performed outside of Emory Genetics Laboratory, please submit a copy of the sequencing report with the test requisition.

**Related Tests**

- Acylcarnitine Analysis - Plasma (PA) and Organic Acid Analysis (OA) are used in the diagnosis and evaluation of patients with metabolic conditions, such as MCADD.
- MCADD Common Mutation Analysis (MC) is available for the K304E and Y42H mutations, prior to sequencing.
- Known Mutation Analysis (KM) is available to family members if mutations are identified by sequencing.
- Deletion/Duplication Assay is available separately for individuals where mutations are not identified by sequence analysis. Refer to the test requisition or contact the laboratory for more information.
- Prenatal testing is available to couples who are confirmed carriers of mutations. Please contact the laboratory genetic counselor to discuss appropriate testing prior to collecting a prenatal specimen.