Emery-Dreifuss Muscular Dystrophy, X-linked: EMD Gene Deletion/Duplication

Test Code: DEMDX
Turnaround time: 2 weeks
CPT Codes: 81404 x1

Condition Description

The clinical diagnosis of Emery-Dreifuss muscular dystrophy (EDMD) is based on the presence of the following triad:

- Joint contractures that begin in early childhood: contractures of the elbow flexors, Achilles tendons (heels), and neck extensors resulting in limitation of neck flexion, followed by limitation of extension of the entire spine
- Slowly progressive wasting and weakness of the humero-peroneal/scapulo-peroneal muscles in the early stages that later extends to the scapular and pelvic girdle muscles
- Cardiac disease with conduction defects and arrhythmias: atrial fibrillation, supraventricular and ventricular arrhythmias, atrio-ventricular and bundle-branch blocks, dilated cardiomyopathy

Age of onset, severity, and progression of muscle and cardiac involvement demonstrate both inter- and intrafamilial variability. Clinical variability ranges from early onset with severe presentation in childhood to late onset with slow progression in adulthood. In general, joint contractures appear during the first two decades, followed by muscle weakness and wasting. Cardiac involvement usually occurs after the second decade.

The two genes known to be associated with EDMD are EMD, encoding emerin and causing X-linked EDMD (XL-EDMD), and LMNA, encoding lamins A and C and causing autosomal dominant EDMD (AD-EDMD) and autosomal recessive EDMD (AR-EDMD). The diagnosis of X-linked EDMD is based on immunodetection of emerin in various tissues and molecular genetic testing of EMD. The diagnosis of AD-EDMD and AR-EDMD is based on clinical findings, family history, and molecular genetic testing of LMNA. About 45% of individuals with EDMD who have emerin detected on immunocytochemistry and/or immunoblotting have no mutation identified in EMD or LMNA, suggesting that these individuals are either misdiagnosed or that other as yet unidentified genes are involved in EDMD.

This testing is for mutations in the EMD gene (Xq28) only. For testing of the LMNA gene, please see the test description for Limb-Girdle Muscular Dystrophy Type 1B: LMNA Full Gene Sequencing (test code SLMNA).

Sequencing of the EMD gene detects an EMD mutation in more than 99% of individuals with established X-linked inheritance and/or with no emerin detected by immunodetection methods.

For patients with suspected X-linked EDMD, sequence analysis is recommended as the first step in mutation identification. For patients in whom mutations are not identified by full gene sequencing, deletion/duplication analysis is appropriate.

References:
- GeneReviews: Emery-Dreifuss Muscular Dystrophy
- OMIM #310300 Emery-Dreifuss Muscular Dystrophy 1

Genes

EMD

Indications

This test is indicated for:

- Confirmation of a clinical diagnosis of X-linked Emery-Dreifuss muscular dystrophy in individuals who have tested negative for sequence analysis
- Carrier testing in adult females with a family history of X-linked Emery-Dreifuss muscular dystrophy who have tested negative for sequence analysis

Methodology

DNA isolated from peripheral blood is hybridized to a CGH array to detect deletions and duplications. The targeted CGH array has overlapping probes which cover the entire genomic region.

Detection

Detection is limited to duplications and deletions. The CGH array will not detect point or intronic mutations. Results of molecular analysis must be interpreted in the context of the patient's clinical and/or biochemical phenotype.

Specimen Requirements

Submit only 1 of the following specimen types
Type: DNA, Isolated

Specimen Requirements:
- Microtainer
- 3µg

Isolation using the Perkin Elmer™Chemagen™ Chemagen™ Automated Extraction method or Qiagen™ Puregene kit for DNA extraction is recommended.

Specimen Collection and Shipping:
Refrigerate until time of shipment in 100 ng/µL in TE buffer. Ship sample at room temperature with overnight delivery.

Type: Whole Blood (EDTA)

Specimen Requirements:
- EDTA (Purple Top)
  - Infants and Young Children (2 years of age to 10 years old): 3-5 ml
  - Older Children & Adults: 5-10 ml
  - Autopsy: 2-3 ml unclotted cord or cardiac blood

Specimen Collection and Shipping:
Ship sample at room temperature for receipt at EGL within 72 hours of collection. Do not freeze.

Special Instructions

Submit copies of diagnostic biochemical test results with the sample, if appropriate. 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 EGL Genetics, please submit a copy of the sequencing report with the test requisition.

Related Tests

- Sequence analysis of the **EMD** gene is available and is required before deletion/duplication analysis.
- Analysis of the **LMNA** gene is also available for autosomal recessive and autosomal dominant Emery-Dreifuss MD.
- Custom diagnostic mutation analysis (KM) is available to family members if mutations are identified by targeted mutation testing or sequencing analysis.
- Prenatal testing is available for known familial mutations only. Please contact the laboratory genetic counselor to discuss appropriate testing prior to collecting a prenatal specimen.