Usher Syndrome: Deletion/Duplication Panel

Test Code: MD237
Turnaround time: 2 weeks
CPT Codes: 81228 x1, 81406 x1

Condition Description

Usher syndrome is a disorder consisting of retinitis pigmentosa (RP) and congenital hearing loss, ranging from partial or profound. Several clinical subtypes exist. Usher syndrome type I is generally characterized by profound congenital hearing loss with no vestibular function and early onset RP. Usher syndrome type II is generally characterized by mild to severe pre-lingual hearing loss with intact vestibular function and adolescent or adult onset RP. Usher syndrome type III is characterized by progressive post-lingual hearing loss, variable vestibular impairment, and late onset RP.

References:
- OMIM
- GeneReviews
- Emory and Rimoin's Principles and Practice of Medical Genetics, 5th Edition

Genes

ABHD12, ADGRV1, CDH23, CIB2, CLRN1, HARS, MYO7A, PCDH15, USH1C, USH1G, USH2A, WHRN

Indications

This test is indicated for:
- Confirmation of a clinical diagnosis of Usher syndrome.
- Carrier testing in adults with a family history of Usher syndrome.

Methodology

Deletion/Duplication Analysis: DNA isolated from peripheral blood is hybridized to a gene-targeted CGH array to detect deletions and duplications. The targeted CGH array has overlapping probes that cover the entire genomic region.

Detection

Deletion/Duplication: Detection is limited to duplications and deletions. The CGH array will not detect point or intronic pathogenic variants. 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 24 hours of collection. Do not refrigerate or freeze.

Special Instructions

Please include fundus photographs, electroretinogram (ERG) findings, visual field findings, and visual acuity, if available, for expert review and clinical correlation with test results.

Disclaimer: This information is confidential and subject to change without notice. It may not be reproduced in whole or part unless authorized in writing by an authorized EGL representative.
Related Tests

- Usher Syndrome: Sequencing Panel
- Eye Disorders: Comprehensive Sequencing
- Eye Disorders: Deletion/Duplication Panel