Stickler Syndrome: Deletion/Duplication Panel

Test Code: MD236
Turnaround time: 2 weeks
CPT Codes: 81228 x1

Condition Description

Stickler syndrome is a genetically heterogeneous connective tissue disorder that typically results in abnormalities of the ocular, auditory, and skeletal systems. Individuals can have a characteristic flat facial appearance that results from underdeveloped bones in the midface. Pierre Robin sequence, hearing impairment/loss and joint hypermobility are common. Eye manifestations include high myopia, cataract, retinal detachment, and vitreous abnormalities. While the disorder is completely penetrant, much phenotypic variability exists. Stickler syndrome can be inherited in an autosomal dominant (COL2A1, COL11A1 and COL11A2 genes) or autosomal recessive (COL9A1 and COL9A2 genes) manner.

References:
- OMIM
- GeneReviews

Genes

COL11A1, COL11A2, COL2A1, COL9A1, COL9A2

Indications

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

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.

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

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