SYP-related XLMR: SYP Gene Deletion/Duplication

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

Condition Description

SYP-related mental retardation is an X-linked condition characterized by mild to moderate intellectual disability. No additional consistent features have been identified; however, epilepsy was noted in some of the individuals with reported SYP mutations. Female carriers had no manifestation.

Mutations in the SYP gene (Xp11.23-p11.22) cause SYP-related mental retardation. In 1122 individuals with X-linked intellectual disability, four different mutations were identified in the SYP gene. SYP encodes an integral membrane protein of small synaptic vesicles.

For patients with suspected SYPrated X-linked mental retardation, 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:

- OMIM #313475: Synaptophysin.

Genes

SYP

Indications

This test is indicated for:

- Confirmation of a clinical diagnosis of SYP-related X-linked mental retardation in an individual in whom sequence analysis was negative.
- Carrier testing in adults with a family history of SYP-related X-linked mental retardation in whom sequence analysis was negative.

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 unclootted 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 SYP gene is available and is required before deletion/duplication analysis.
- 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 only for known familial mutations to individuals who are confirmed carriers of mutations. Please contact the laboratory genetic counselor to discuss appropriate testing prior to collecting a prenatal specimen.