
Lab Dept: **Anatomic Pathology**

Test Name: **TGFBR1 DELETION/DUPLICATION HDT ARRAY**

General Information

Lab Order Codes: R1DD

Synonyms: Aortic aneurysm, familial thoracic 5 (AAT5) TGFBR1;Loeys-Dietz syndrome, type 1A (LDS1A) TGFBR1

CPT Codes: 81479 – Molecular Pathology, Unlisted Procedure

Test Includes: Deletion/Duplication HDT Array uses 60mer oligonucleotide sequences designed to selectively complement target areas with an extremely high degree of specificity. High resolution and sensitivity are achieved by assigning a number of probes to each exonic region to allow for a minimum of 3 probes within each 300-500 base section of the genomic DNA sequence. Non-coding intervening sequences are targeted at a minimum with approximately half of the density used for exonic regions. Many intervening sequences are more densely covered. The minimum CNV size detected by this high-density array is 300-500 nucleotides, which is the technical limit of the assay using these stringent parameters.

Logistics

Test Indications: For confirmation of symptoms and the clinical diagnosis related to Aortic aneurysm, familial thoracic 5 (AAT5) TGFBR1 or Loeys-Dietz syndrome, type 1A (LDS1A) TGFBR1.

AAT5 and AAT3 have been linked to mutations in the transforming growth factor Beta receptor type I and II genes (TGFBR1 and TGFBR2). Patients may have aneurysms of the aorta and other arteries. TGFBR2 mutations are currently estimated to be responsible for 5% of familial thoracic aortic aneurysms and dissections (TAAD).

Loeys-Dietz syndrome (LDS) is a recently described syndrome caused by mutations in one of two genes, TGFBR1 (Type 1A) or TGFBR2 (Type 1B). The reported phenotype is highly variable and overlaps considerably with Ehlers-Danlos syndrome IV or Marfan syndrome. A high percentage of patients have aortic root aneurysm or other arterial aneurysms. Some have cardiac abnormalities including patent ductus or atrial septal defects. Many display arterial tortuosity. Skeletal defects may include hypertelorism, pectus defects, joint laxity, craniosynostosis, arachnodactyly, scoliosis, talipes equinovarus, camptodactyly and malar hypoplasia. Some have dural ectasia. Additional features may include uterine, spleen or bowel rupture, thin, translucent, hyperextensible or velvety skin with atrophic scars and easy bruising. Blue sclera and a bifid uvula have also been observed.

Lab Testing Sections: Anatomic Pathology - Sendouts

Referred to: Connective Tissue Gene Tests (CTGT)

Phone Numbers: MIN Lab: 612-813-6280
STP Lab: 651-220-6550

Test Availability: Daily, 24 hours

Turnaround Time: 1 – 2 weeks

Special Instructions: Please include a completed CTGT [Request form](#) with the patient or specimen to the laboratory.

Specimen

Specimen Type: Whole blood

Container: Lavender top (EDTA) tube

Draw Volume: 6 mL (Minimum: 3 mL) blood

Processed Volume: Same as Draw Volume

Collection: Routine venipuncture

Special Processing: Lab Staff: **Do Not** centrifuge. Specimen should be sent in original collection container. Send via overnight shipping with to reach CTGT Monday through Friday. If weekend or holiday when drawn, store at refrigerated temperatures. Please include a [CTGT Shipment Packing Slip](#) with the shipment.

Patient Preparation: None

Sample Rejection: Mislabeled or unlabeled specimen

Interpretive

Reference Range: Interpretive report

Critical Values: N/A

Limitations:

Published estimates of test sensitivity for genes linked to certain disorders can be very inaccurate, and that it is difficult to predict the probability of detecting a mutation in any single gene for one individual. The following factors contribute to this challenge: Many disorders have overlapping phenotypes; some disorders are linked to mutations in more than one gene; in some instances genes remain to be linked to specific disorders; for most disorders, proper diagnosis requires that clinical findings are considered along with genetic findings.

Methodology:

High-Density Targeted Array

References:

[Connective Tissue Gene Tests](#) October 2010
(484) 224-2900 Fax (484) 244-2904

Updates:

2/11/2013: CPT update
7/10/2013: CPT update CTGT, previously listed as 81404