Lab Dept: Anatomic Pathology

Test Name: COMPARATIVE GENOMIC HYBRIDIZATION (CGH)

WITH LIMITED GBAND STUDY

General Information

Lab Order Codes: CGLBN

Synonyms: Comparative Genomic Hybridization, Microarray; aCGH with Limited G

Band

CPT Codes: 81228 – Cytogenomic constitutional microarray analysis

88230 - Tissue culture for non-neoplastic disorders; lymphocyte

88261 - Chromosome analysis; count 5 cells, 1 karyotype with banding

Test Includes: Microarray with chromosome analysis by G-banding.

Logistics

Test Indications: For detection of small duplications and deletions in patients with normal G-

banded karyotypes. For characterization of abnormalities detected by G-

banding.

Lab Testing Sections: Anatomic Pathology - Sendouts

Referred to: MHealth Fairview - University of Minnesota Medical Center – Cytogenetics

FV Test code: LAB5988

Phone Numbers: MIN Lab: 612-813-6280

STP Lab: 651-220-6550

Test Availability: Daily, 24 hours

Turnaround Time: Performed Monday-Friday. Results are reported within 7-21 days.

Special Instructions: Please see Container and Draw Volume

Specimen

Specimen Type: Whole Blood or cord blood

Container: Green (sodium heparin, no gel) AND Lavender (EDTA) top tubes

Both tubes are required.

Note: A Yellow (ACD) tube can be substituted for the Lavender top tube.

Draw Volume: 6-10 mL (3-5 mL per tube) (Minimum: 2 mL [1 mL per tube]) blood

Note: 1-3 mL per tube minimum for infants and children, with 3 mL strongly

preferred

Processed Volume: Same as Draw Volume

Collection: Routine Venipuncture. Heelstick or fingerstick is not acceptable.

Special Processing: Lab Staff: Do Not Centrifuge. Specimen should remain in the original

collection container. Store and ship at room temperature. Do not freeze or

refrigerate. Must arrive within 24 hours of collection.

Specimens must be received in the Cytogenetics Laboratory Mon-Fri by 5:30 pm; weekends and holidays by 4:30 pm. Specimens received after

these cut-offs will be processed the following day.

Patient Preparation: None

Sample Rejection: Clotted or frozen sample; mislabeled or unlabeled specimens; incorrect

anticoagulant.

Interpretive

Reference Range: Interpretive report

Critical Values: N/A

Limitations: Post-natal studies only.

Methodology: Copy number chromosomal microarray (CMA) with limited chromosome

analysis by G-banding. Congenital.

References: MHealth Fairview Reference Laboratories September 2023

Updates: 9/5/2023: Updated reference lab test codes, added specimen stability

notes, corrected methodology, updated blood optimal and minimal volumes,

added blood collection by venipuncture is required.