Lab Dept: Anatomic Pathology

Test Name: COMPREHENSIVE EPILEPSY PANEL

**General Information** 

Lab Order Codes: CEPL

Synonyms: N/A

**CPT Codes:** 81479 – Unlisted molecular pathology procedure

**Test Includes:** Sequence Analysis and Exon-level Deletion/Duplication testing. The panel

analyzes 187 genes that are associated with both syndromic and non-

syndromic causes of epilepsy.

Logistics

**Test Indications:** Given the clinical overlap between epilepsy disorders, comprehensive

testing allows for more efficient evaluation of multiple conditions based on a single indication for testing. Identification of the molecular basis of the disease in an affected individual may confirm diagnosis, predict prognosis, and encourage testing of additional family members to inform reproductive risk. Note: This assay includes sequencing and deletion analysis of

UBE3A, but does not detect uniparental disomy or imprinting center defects.

**Lab Testing Sections:** Anatomic Pathology - Sendouts

**Referred to:** Invitae (INVT test: 03401)

Phone Numbers: MIN Lab: 612-813-6280

STP Lab: 651-220-6550

**Test Availability:** 24 hours, daily

**Turnaround Time:** 10 – 21 days

**Special Instructions:** N/A

Specimen

Specimen Type: Whole blood

Container: Lavender (EDTA) top tube

Alternate: Buccal Swab

**Draw Volume:** 3 mL (Minimum: 1.5 mL) whole blood

**Processed Volume:** Do Not process or centrifuge whole blood.

Submit EDTA whole blood in original container.

**Collection:** Routine blood collection

**Special Processing:** Lab Staff: Store unprocessed blood at room temperature. Ship at room

temperature via overnight courier, using a cool pack in hot weather. If shipment is delayed, the specimen should be refrigerated for up to 14 days.

Patient Preparation: None

Sample Rejection: Mislabeled or unlabeled specimen

Interpretive

**Reference Range:** An interpretive report will be provided

Critical Values: N/A

**Limitations:** The clinical sensitivity of this test is dependent on the patient's underlying

genetic condition.

**Methodology:** Next-generation Sequencing (NGS)

**References:** <u>Invitae</u> January 2019

**Updates:** 8/25/2015:CPT update.

4/16/2018: CPT and gene list update.

1/16/2019: Testing moved from GeneDx to Invitae

8/17/22: Added Alt specimen