



## **Test Update 827**

**Posted Date** 12/21/2022

**Effective Date** 01/16/2023

**Test Name** [Eosinophil Derived Neurotoxin](#)

**Update Type** [New Tests](#)

### **NEW TEST**

#### **Eosinophil Derived Neurotoxin, Serum**

Order Code: EDN

Fee Code: pending (CPT 83520)

Reference Laboratory: Mayo EDN

Mayo Clinic Laboratories Eosinophil Derived Neurotoxin, Serum assay will be available effective January 16, 2023. This test is used for evaluating patients suspected to have a condition associated with eosinophilia or hypereosinophilia, evaluating patients with elevated peripheral blood eosinophil counts, and managing patients with elevated eosinophil-derived neurotoxin in the context of eosinophil-associated diseases. This test replaces Mayo Clinic Laboratories test ID FECP (order code MMLR) Eosinophil Cationic Protein referred to Quest Diagnostics.

Collection Instructions: Collect specimen in an SST (preferred) or red top tube. Centrifuge, aliquot 0.5 mL (minimum 0.3 mL) of serum into a plastic vial within 12 hours of collection and refrigerate.

Reference Range: <70 mcg/L Normal; 70 - 99 mcg/L Borderline; > or = 100 mcg/L Elevated.

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