

Neurocode USA, Inc

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Plasma Neurofilament Light Chain (NfL)

Test Name	Neurofilament Light Chain
Abbreviation	Plasma NfL
CPT code	83520
Methodology	Chemiluminescent enzyme immunoassay (CLEIA)
Intended Use	Quantification of NfL in plasma to assist in the clinical assessment of neurological disorders.
Test requirements	Specimen Type: EDTA-plasma Minimum volume: 0.5 mL Preferred volume: 5 mL Rejection criteria: grossly hemolytic, icteric, or lipemic
Specimen collection	 Overnight fasting recommended 5 mL EDTA lavender-top tubes Within an hour of collection spin, aliquot, and freeze plasma. Ship frozen on dry-ice.
Specimen stability	Store frozen (-80 – -70°C preferred) Sample should be frozen until transport, avoid freeze/thaws.
Test schedule	Once a week
TAT	1 – 3 days
Reference range	Age group $20 - 29 \le 8.4 \text{ ng/L}$ $30 - 39 \le 11.4 \text{ ng/L}$ $40 - 49 \le 15.4 \text{ ng/L}$ $50 - 59 \le 20.8 \text{ ng/L}$ $60 - 69 \le 28.0 \text{ ng/L}$ $70 - 79 \le 37.9 \text{ ng/L}$ $\ge 80 \le 51.2 \text{ ng/L}$
Limitations	 This is a laboratory developed test, its performance was determined by Neurocode USA Inc. It has not been cleared or approved by the Food and Drug Administration. Elevated NfL is not a disease specific factor and may be caused by a neurodegenerative disease or traumatic brain injury. Results should be used in conjunction with clinical signs and symptoms. NfL levels measured in the evening may be more than 10% lower than those measured in the morning (Benedict et al 2020). Higher levels of NfL can occur in patients with a history of stoke, atrial fibrillation, myocardial infarction, chronic kidney disease, pregnancy, and diabetes. Lower levels may be found in person who are obese (BMI ≥ 30) (Syrjanen et al 2022).
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