

Glial Fibrillary Acidic Protein, Plasma

Patient ID C7028846-0006523	Patient Name VALIDATIONSOFT, TESTINGBGFAP		Birth Date 1991-11-25	Sex F	Age 33
Order Number X100507025	Client Order Number C70288460006523	Ordering Physician CLIENT, CILENT	Report Notes		
Account Information C7028846 DLMP Rochester		Collected 19 Aug 2025 07:00			

## **Glial Fibrillary Acidic Protein, P**

1 SDL

**Reference Value** ≤ 32.6 pg/mL

High The testing method is a chemiluminescent enzyme immunoassay manufactured by Fujirebio, Inc. and performed on the Lumipulse analyzer. Values obtained with different assay methods or kits may be different and cannot be used interchangeably.

This test is not intended as a screening or standalone diagnostic assay; correlation with clinical findings is recommended.

**Received:** 19 Aug 2025 13:41 **Reported:** 19 Aug 2025 13:41

## **Laboratory Notes**

1 This test was developed and its performance characteristics determined by Mayo Clinic in a manner consistent with CLIA requirements. It has not been cleared or approved by the US Food and Drug Administration.

## **Performing Site Legend**

Code	Laboratory	Address	Lab Director	<b>CLIA Certificate</b>
SDL	Mayo Clinic Laboratories - Rochester Superior Drive	3050 Superior Drive NW, Rochester MN 55905	Nikola Baumann Ph.D.	24D1040592