

Beta-Amyloid 1-42 and 1-40 with Ratio, BioPharma Diagnostics, Spinal Fluid

Patient ID	Patient Name		Birth Date	Sex	Age
SA00166208	SAMPLE REPORT, AMYRB N		1961-02-25	F	62
Order Number	Client Order Number	Ordering Physician	Report Notes		
SA00166208	SA00166208	CLIENT, CLIENT			
Account Information		Collected			
C7028846 DLMP Rochester		01 Feb 2024 08:00			

SDL

SDI

Beta-Amyloid Panel, BioPharma, CSF

Abeta40, CSF

158 pg/mL

ADDITIONAL INFORMATION

For research use only. This test was developed and its performance characteristics determined by Mayo Clinic in a manner consistent with CLIA requirements. This test has not been cleared or approved by the U.S. Food and Drug Administration.

Abeta42, CSF

38 pg/mL

ADDITIONAL INFORMATION

For research use only. This test was developed and its performance characteristics determined by Mayo Clinic in a manner consistent with CLIA requirements. This test has not been cleared or approved by the U.S. Food and Drug Administration.

Beta-Amyloid Ratio

0.241 ratio

SDL

Reference Value ≥ 0.073

Beta-Amyloid Ratio Interpretation

SDL

1-800-533-1710

AMYRB

A normal beta-Amyloid ratio (1-42/1-40) of ≥ 0.073 is consistent with a negative (normal) amyloid positron tomography (PET) scan result. This result indicates a reduced likelihood that a patient's cognitive impairment is due to Alzheimer's disease.

ADDITIONAL INFORMATION

The testing method is a chemiluminescent enzyme immunoassay manufactured by Fujirebio, Inc. and performed on the Lumipulse analyzer. The beta-Amyloid ratio is calculated by using individual measurements of the beta-Amyloid 1–42 and beta-Amyloid 1–40.

The Lumipulse beta-Amyloid ratio (1–42/1–40) results must be interpreted in conjunction with other patient clinical information. This test is not intended as a screening or stand-alone diagnostic assay.

Values obtained with different assay methods or kits may be different and cannot be used interchangeably.

Received: 02 Feb 2024 14:49

Reported: 02 Feb 2024 14:51

Performing Site Legend

Code	Laboratory	Address	Lab Director	CLIA Certificate
SDL	Mayo Clinic Laboratories - Rochester Superior Drive	3050 Superior Drive NW, Rochester MN 55905	William G. Morice M.D. Ph.D.	24D1040592