

MEDICAL POLICY

Medical Policy Title	Lab Testing for Alzheimer's Disease
Policy Number	2.02.16
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Next Review Date	June 2027

Our medical policies are guides to evaluate technologies or services for medical necessity. Criteria are established through the assessment of evidence based, peer-reviewed scientific literature, and national professional guidelines. Federal and state law(s), regulatory mandates and the member's subscriber contract language are considered first in the determination of a covered service.

(Link to [Product Disclaimer](#))

POLICY STATEMENT(S)

Genetic Testing

Asymptomatic Individuals

- I. Targeted genetic testing for a known familial variant in the presenilin (PSEN) genes or amyloid-beta precursor protein (APP) gene associated with autosomal dominant early-onset Alzheimer disease (EOAD) may be considered **medically appropriate** in an asymptomatic individual to determine future risks of disease when **ANY** of the following criteria are met:
 - A. The individual has a close relative (i.e., first- or second-degree relative) with a known familial variant associated with autosomal dominant early-onset Alzheimer disease;
 - B. The individual has a family history of dementia consistent with autosomal dominant Alzheimer disease, and the genetic status of the affected family members is unavailable;
 - C. Results of testing will inform reproductive decision making.

Symptomatic Individuals

- II. Targeted genetic testing for a known familial variant in the presenilin (PSEN) genes or amyloid-beta precursor protein (APP) may be considered **medically appropriate** in a symptomatic individual to determine future risk of disease when **ANY** of the following criteria are met:
 - A. An individual with early onset Alzheimer's Disease (and other etiologies have been excluded);
 - B. A family history of dementia or unknown family history;
 - C. Autosomal dominant family history of dementia with one or more cases of EOAD;
 - D. A relative with a pathological variant consistent with EOAD.
 - E. Results of testing will inform reproductive decision making.

Apolipoprotein E (APOE) Gene

- III. Genetic testing for the apolipoprotein E (APOE) gene to guide initiation or management of a U.S. Food and Drug Administration-approved amyloid-beta targeting therapy is considered **medically appropriate** in individuals with mild cognitive impairment or mild dementia associated with

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Alzheimer disease.

Fluid Biomarker Testing

- IV. CSF Biomarker testing is considered **medically appropriate** when **ALL** of the following criteria have been met:
- A. As part of the evaluation for the initiation of amyloid beta targeting therapy;
 - B. An Individual with mild cognitive impairment or mild dementia due to Alzheimer's disease;
and
 - C. Utilizing **ONE** of the following tests:
 - 1. Phospho-Tau (pTau181) and β -Amyloid (1-42) assay, (Tau181/ A β 42) (e.g., Elecsys Phospho-Tau);
 - 2. Total Tau (tTau) with A β 42 assay, (tTau/A β 42) (e.g., Elecsys Total Tau);
 - 3. β -Amyloid Ratio (1-42/1-40) Test, (A β 42/40) (e.g., Lumipulse G);
 - 4. pTau protein (e.g., ALZpath pTau217).
- V. CSF Biomarker testing, including but not limited to amyloid beta, tau, or neural thread proteins (NTP) is considered **investigational** for **EITHER** of the following indications:
- A. As an adjunct to clinical diagnosis in individuals with mild cognitive impairment;
 - B. As part of an evaluation for the continuation of amyloid beta targeting therapy.
- VI. Biomarker testing for the diagnosis or assessment of Alzheimer disease using **ANY** of the following tests is considered **investigational**, including, but not limited to:
- A. Triggering receptor expressed on myeloid cells 2 (TREM2);
 - B. MindX Memory Alzheimer test;
 - C. Neural Thread Proteins (NTP);
 - D. Urine biomarkers- (e.g., Urinary AD7c-NTP);
 - E. PrecivityAD2 test;
 - F. Elecsys Amyloid Plasma Panel Test;
 - G. CSF testing for alpha-Synuclein (e.g., SYNTap biomarker test);
 - H. Neurofilament Light Chain (NfL);
 - I. Skin biopsy biomarkers (e.g., DISCERN).

RELATED POLICIES

Corporate Medical Policy

2.02.03 Genetic Testing for Inherited Disorders

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6.01.24 Positron Emission Tomography (PET) – Head Imaging

11.01.03 Experimental or Investigational Services

Pharmacy Policy

100 Anti-Amyloid Directed Therapies

POLICY GUIDELINE(S)

- I. The Health Plan and its employees adhere to all State and Federal laws concerning the confidentiality of genetic testing and the results of genetic testing. All records, findings and results of any genetic test performed on any person shall be deemed confidential and shall not be disclosed without the written informed consent of the person to whom such genetic test relates. This information shall not be released to any person or organization not specifically authorized by the individual subject of the test or in compliance with applicable law.
- II. Genetic testing is appropriate only when performed by a qualified laboratory certified under the Clinical Laboratory Improvement Amendments of 1988 (CLIA) and offered in a setting with adequately trained health care professionals who are qualified to provide appropriate pre- and post-test counseling.
- III. Genetic testing is contract dependent. Coverage only applies to members with a valid contract; coverage is not provided for family members without a valid contract.
- IV. Supporting documentation required:
 - A. The following factors will be considered when determining the medical appropriateness of a genetic test:
 1. There must be reasonable expectations based on family history, pedigree analysis, risk factors, and/or symptomatology that a genetically inherited condition exists. Autosomal recessive disorders may be present without a family history.
 2. The genotypes to be detected by a genetic test must be shown by scientifically valid methods to be associated with the occurrence of the disease, and the analytical and clinical validity of the test must be established.
 3. The clinical utility of the test must be established (e.g., test results will influence decisions concerning disease treatment or prevention).
 4. Genetic testing should be performed for management or treatment of the patient and not only for knowledge purposes. Documentation should demonstrate how test results will impact treatment or medical management.
 5. When there is family history or phenotype suggestive of a specific syndrome, results of targeted testing for the mutation associated with the syndrome should be documented prior to any panel testing. If targeted testing has not been performed, rationale as to why panel testing is medically necessary should be documented.
- V. In the appropriate use recommendations for Lecanemab it states that either elevated amyloid on

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PET or elevated phosphorylated tau and low A β 42 level (increased p-tau/ A β 42 ratio) in the CSF is required prior to initiating treatment with Lecanemab, currently blood based biomarkers are not a reliable and fully validates and are not considered adequate to identify appropriate patients for treatment. To guide safety and risk discussions, it is recommended that APOE genotyping testing of all treatment candidates be completed before initiating therapy (Cummings et al., 2023).

In the prescribing information for LEQEMBI it suggests that testing should be performed prior to initiation of treatment to better determine the risk if ARIA. Stating that, individuals treated with LEQEMBI or other drugs in this class of medications who are ApoE ϵ 4 homozygotes (two identical pairs of genes for a specific trait) have a higher incidence of amyloid related imaging abnormalities (ARIA).

- VI. In the appropriate use criteria for Donanemab it states that biomarker confirmation of A β pathology by PET or CSF is required for patients to be eligible for treatment. To date, three CSF tests have been approved by the FDA for determination of amyloid status: A β 42/A β 40 ratio measured on the Lumipulse platform, and p-tau181/A β 42 or total tau/A β 42 ratios measured on the Elecsys platform (Rabinovici et al, 2025).

DESCRIPTION

Alzheimer's disease (AD) is the leading cause of dementia in the elderly, accounting for 50 to 75% of all cases of dementia. AD can be associated with a family history (40% of patients with AD have at least one other afflicted first-degree relative) or idiopathic. More than 90% of AD occurs after age 65 years (late-onset AD) and is characterized by gradual onset and progressive and irreversible decline in cognitive function. There is also a less common early-onset form of AD, which appears before the age of 60 and is associated with a rapid decline and severe neurochemical and neuron-pathological changes. The estimated lifetime risk of AD in the general population is about 15%. Over 100 genes, particularly on chromosomes 9, 10, and 12 have been associated with late-onset AD, while pathologic variants in chromosomes 1, 14, and 21 have been associated with early onset familial AD. Genetic testing has been investigated both in patients with probable AD and in asymptomatic family members.

Susceptibility Polymorphism at the Apolipoprotein E (APOE) Gene

The APOE lipoprotein gene is a carrier of cholesterol and is produced in the liver and brain glial cells. The APOE gene has three alleles - epsilon 2, 3, and 4 - with epsilon 3 allele being the most common. Every person carries two APOE alleles. The presence of at least one epsilon 4 allele is associated with an increased risk of AD in the range of 1.2- to 3-fold, depending on the ethnic group. For those homozygous for epsilon 4 (about 2% of the population), the risk of AD is higher than for those heterozygous for epsilon 4. The mean age of onset of AD is about 68 years for epsilon 4 homozygotes, about 77 years for heterozygotes, and about 85 years for those with no epsilon 4 allele. It should be noted that the epsilon 4 allele represents a risk factor for AD rather than a disease-causing pathologic variant.

Genetic Pathologic Variants

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Patients with early-onset AD (e.g., before age 65 but as early as 30 years) are a small subset of patients. The families of these patients may show an autosomal dominant pattern of inheritance. Three genes have been identified by linkage analysis of affected families: amyloid-beta precursor protein (APP) gene, presenilin 1 (PS1) gene, and presenilin 2 (PS2) gene. These genes have nearly 100% penetrance, absent death from other causes; however, rare cases of lack of penetrance in elderly individuals have been reported. A variety of pathologic variants within these genes have been associated with AD; pathologic variants in PS1 appear to be the most common. However, only 2% to 10% of all patients with AD have early onset AD, and genetic pathologic variants have only been identified in 30% to 50% of these patients. Therefore, overall, identifiable genetic pathologic variants are rare causes of AD.

Biomarker Tests

The Elecsys Amyloid Plasma Panel (Roche, a biotech company) measures phosphorylated Tau (pTau) 181 protein assay and apolipoprotein (APOE) E4 assay in human blood plasma. Positive results indicate the need for further confirmatory testing for AD. The panel test is intended to be used in conjunction with other clinical information in symptomatic patients who are being evaluated for AD and other causes of cognitive decline.

Roche also has Elecsys β -Amyloid (1-42) CSF and Elecsys Phospho-Tau (181P) CSF in vitro diagnostic immunoassays measuring β -Amyloid (1-42) and Phospho-Tau concentrations in CSF in adult patients with cognitive impairment who are being evaluated for Alzheimer's disease (AD) or other causes of dementia.

Lumipulse G β -Amyloid Ratio (1-42/1-40) immunoassay (Fujirebio Diagnostics, Inc.), the first in vitro diagnostic test for early detection of amyloid plaques with AD. The test uses CSF and is intended to be used in adult patients greater than or equal to 55 years, presenting with cognitive impairment who are being evaluated for AD and other causes of cognitive decline.

The Precivity mass-spec amyloid beta assay uses plasma and a proprietary mass spectrometry platform that combines quantitative measurement of amyloid beta 42 and 40 peptides in plasma along with apolipoprotein E prototype (equivalent to ApoE genotype) to calculate an individual's likelihood of amyloid plaques in the brain. The test is currently not intended to be used as a stand-alone diagnostic.

SUPPORTIVE LITERATURE

Hu et al (2025) reported the preclinical development of an immunoassay for detecting triggering receptors expressed on myeloid cells 2 (TREM2), a novel biomarker for Alzheimer's disease (AD). In this study, highly sensitive and specific TREM2 antibodies were generated using hybridoma technology, and three optimized immunoassay formats were developed. The assays demonstrated lower limits of quantitation (LLOQs) of 0.474, 0.807, and 0.415 ng/mL, respectively. All assays exhibited high sensitivity and specificity. Notably, the sandwich enzyme-linked immunosorbent assay (ELISA) showed strong potential for use in the diagnosis of Alzheimer's disease.

Pahlke et al (2025) reported the results of a systematic review and meta-analysis evaluated the diagnostic accuracy of blood-based biomarkers (BBMs) for detecting Alzheimer's disease (AD)

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pathology in cognitively impaired individuals within specialized care settings. Analyzing 49 observational studies published between 2019 and 2024, the review assessed 31 plasma-based tests measuring phosphorylated tau (p-tau217, %p-tau217, p-tau181, p-tau231) and amyloid beta (A β 42/A β 40), using cerebrospinal fluid biomarkers, amyloid PET, or neuropathology as reference standards. Diagnostic performance varied substantially across tests, with pooled sensitivities ranging from approximately 49% to 91% and specificities from 62% to 97%, leading to wide variation in post-test probabilities across different pre-test scenarios. The certainty of evidence ranged from moderate to very low, and most studies were judged to have a high risk of bias, particularly related to patient selection, index test conduct, and reference standards. Overall, the findings provide a comprehensive evidence base to inform an Alzheimer's Association–led clinical practice guideline on BBM use, while highlighting the need for ongoing evaluation as biomarker technologies and evidence continue to evolve.

Ren et al (2025) conducted a meta-analysis looking for the association of APOE ϵ 4 status in AD patients. Included in the meta-analysis were 22 eligible articles involving 571,800 cases published between 2001 and 2023. There was heterogeneity in the ability to APOE ϵ 4 status to predict AD risk, they performed a sensitivity analysis to predict whether one study impacted the pooled hazard ratios; and the results did not change when a study was eliminated. They conducted a subgroup analysis on the basis of region, study design, and cutoff method to investigate the source of the heterogeneity. There was not a reduction in the heterogeneity in the basis of region stratification and the study design. It remains unclear on of the source of heterogeneity, but this meta-analysis suggests that there is an increased risk of AD in individuals with different APOE ϵ 4 statuses; future high quality, larger sample, randomized control trials are needed.

Safransky et al (2024) conducted a retrospective study of brain donors from the national Alzheimer's Coordinating Center who had normal cognition at the time of the lumbar puncture, and CSF A β 42 and p-tau181 performed with Lumipulse assays. The primary objective of the study investigated the association between CSF biomarkers and AD and AD neuropathologic changes among those brain donors. The National Institute on Aging-Alzheimer's Association criteria was used to stage ADNC. Participants were divided into 2 categories: AD negative (no AD/low ADNC) and AD positive (intermediate /high ADNC). At autopsy confirmed AD in 20 (40.8%) individuals. The average span from LP until death was 7.76 years. CSF p-tau181/A β 42 was the optimal predictor of AD, having excellent discrimination accuracy (AUC = 0.97), followed by CSF p-tau181 (AUC = 0.92), CSF A β 42 alone (AUC = 0.92) and CSF t-tau having the lowest accuracy (AUC = 0.87). CSF p-tau181/A β 42 was associated with the Consortium to Establish a Registry for Alzheimer's (CERAD) ratings of neuritic amyloid plaque scores, and Braak staging (refers to two methods used to classify the degrees of pathology in Parkinson's and Alzheimer's disease) of the neurofibrillary tangle (NFT's).

Schraen-Maschke et al (2024) reported results from a subgroup (n=106) of the BALTAZAR study evaluating whether plasma levels of the free amyloid peptides A β 1–42 and A β 1–40 and the free plasma A β 1–42/A β 1–40 ratio is associated with the conversion of MCI to dementia over three years of follow-up total of 50 participants converted to dementia during follow-up. The risk of conversion was lower for participants in the highest quartile of free plasma A β 1–42/ A β 1–40 compared to those in the three lower quartiles. The risk of conversion in the highest quartile of total plasma A β 1–42/A β 1–40 compared to the lower quartiles was similar. The results show the relevance of the free

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plasma A β 1–42/A β 1–40 ratio for identifying MCI patients at lower risk of conversion to dementia (mainly AD) within three years. Using the threshold of 25.8% for the free plasma A β 1–42/A β 1–40 ratio, it was possible to identify MCI patients with an at least 60% lower risk of conversion to dementia. The performance of the free A β 1–42/A β 1–40 ratio in predicting conversion to dementia was similar to that of the total A β 1–42/A β 1–40 ratio. The study limitations were, the diagnosis of AD was only based on clinical and MRI; the results of the CSF biomarkers were not taken into consideration for the AD diagnosis; they excluded participants with Lewy Body, Parkinsonian, frontotemporal, or vascular dementia; and small sample size.

Bolsewig et al (2024) conducted a cross-sectional multicenter cohort study with prospective component which included individuals with dementia with Lewy bodies, Alzheimer's disease, and healthy control. 562 Participants had annual follow up for 5 years. Plasma biomarkers were measured; amyloid status was determined by CSF A β 42 concentrations and cognition evaluation by Mini-Mental State Examination (MMSE). Higher baseline glial fibrillary acidic protein (GFAP), neurofilament light (NfL), and phosphorylated-tau (P-tau) concentrations were associated with lower MMSE scores in DLB, and GFAP and NfL were associated with a faster cognitive decline. DLB participants with parkinsonism had higher concentrations of NfL ($\beta = 0.08$, 95% CI 0.02–0.14, $p = 0.006$) than those without. This study does suggest that the use of plasma biomarkers (A β 42/40, P-tau181, and P-tau231) to assess amyloid copathology in DLB, and plasma GFAP and NfL as monitoring biomarkers for cognitive symptoms in DLB.

Krishna et al (2024) reported results of a cross-sectional study of a single molecule array (Simoa) analysis of A β 1–42, total tau (t-tau), phospho-tau (p-tau 181), and neurofilament L (NfL) in the plasma samples of AD patients ($n=35$), healthy controls ($n=35$), and non-AD ($n=33$) patients from a tertiary care center in India. The non-AD dementia patients included those with frontotemporal dementia ($n=12$), vascular dementia ($n=5$), Lewy body dementia ($n=4$), and mixed dementia ($n=12$). The cutoffs used for calculating sensitivity and specificity were unclear. A model including all 4 biomarkers had sensitivity of 94% and specificity of 96% for distinguishing AD versus healthy controls. The model including all 4 biomarkers had sensitivity of 40% and specificity of 93% for distinguishing AD from non-AD dementia. Limitations included a small sample size, dissimilarity analysis between dementia types could not be assessed, supporting data from CSF measurements and neuroimaging of biomarkers regarding the diagnosis of AD or other dementias was lacking, and A β 40 in plasma was not measured.

Van Dyck et al (2023) conducted an 18-month multicenter, double blinded, phase 3 trial involving 1795 individuals with early Alzheimer's disease (mild cognitive impairment or mild dementia due to Alzheimer's disease) with evidence of amyloid on positron emission tomography (PET) or by cerebrospinal fluid (CSF) testing. Individuals were randomly assigned in a 1:1 ratio to receive intravenous lecanemab or a placebo. Primary endpoint was to see a change from the baseline at 18 months in the score on the Clinical Dementia Rating. Secondary endpoint was to see a change on the PET imaging, and changes on other dementia evaluation tools. During the trial participants had the option of participating in sub studies that evaluated the longitudinal changes in brain amyloid burden as measured by PET, brain tau pathologic features as measured by PET, and CSF biomarkers of Alzheimer's disease. Amyloid-related imaging abnormalities-edema or effusion (ARIA-E) and amyloid-related imaging abnormalities-Hemorrhage (ARIA-H) were numerically less common among

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ApoE ϵ 4 noncarriers than among carriers, with higher frequency among ApoE ϵ 4 homozygotes than among ApoE ϵ 4 heterozygotes.

Amft et al (2022) conducted a retrospective study on 103 patients with known amyloid PET status. The study aimed to evaluate the performance of the amyloid beta ($A\beta$)_{42/40} ratio for predicting amyloid positivity by PET, compared with $A\beta$ ₄₂ alone, and phosphorylated tau 181 (pTau₁₈₁)/ $A\beta$ ₄₂ and total tau (tTau)/ $A\beta$ ₄₂ ratios, using the fully automated Elecsys CSF immunoassay test by Roche Diagnostics International Ltd. in patients with a range of cognitive disorders. Results for sensitivity and specificity of CSF biomarkers and biomarker ratios versus amyloid PET were 0.93 and 0.57 for $A\beta$ ₄₂, 0.96 and 0.69 for pTau₁₈₁/ $A\beta$ ₄₂, 0.92 and 0.69 for tTau/ $A\beta$ ₄₂, and 0.94 and 0.82 for $A\beta$ _{42/40}. For area under the curve point estimates (95% confidence intervals) versus amyloid PET were 0.78 (0.68–0.88) for $A\beta$ ₄₂, 0.88 (0.81–0.95) for pTau₁₈₁/ $A\beta$ ₄₂, 0.87 (0.80–0.95) for tTau/ $A\beta$ ₄₂, and 0.90 (0.83–0.97) for $A\beta$ _{42/40}. The study proved that $A\beta$ _{42/40} ratio in detecting amyloid positivity by PET was higher than using $A\beta$ ₄₂ alone and comparable with the performance of the pTau₁₈₁/ $A\beta$ ₄₂ and tTau/ $A\beta$ ₄₂ ratios. Limitations of the study were, small sample size, use of a pre-analytical protocol not in accordance with the Elecsys CSF immunoassay method sheets, and the lack of a pre-defined cut-off for $A\beta$ _{42/40}.

Swanson et al (2021) conducted a randomized double blinded clinical trial of 854 individuals with early Alzheimer's disease (AD), mild cognitive impairment due to AD and mild dementia. The study aimed to establish the simplest dose that achieves greater or equal to 90% of the maximum treatment effect. They were randomized to receive LEQEMBI or placebo, approximately 70% were ApoE₄ carriers and 30% were ApoE₄ non-carriers. The ApoE₄ carriers were not randomized, as they were authorized to make sure that there were no ApoE₄ carriers in the 10 mg/kg group. The incidence of ARIA-H on LEQEMBI was higher in ApoE₄ carriers (13.1%) than in ApoE₄ non-carriers (4.6%)

Fink et al (2020) conducted a systematic review of biomarker accuracy for diagnosing neuropathologically defined AD in older patients with dementia. The analysis included literature published between January 2012 and November 2019, with 9 cohort studies focusing on CSF biomarkers. Overall, CSF biomarkers and ratios had moderate sensitivity (range, 62% to 83%) and specificity (range, 53% to 69%). Biomarker accuracy was higher with amyloid beta-42/pTau ratio, tTau/amyloid beta-42 ratio, and pTau compared with tTau alone. The limitations were small study size, biomarker cut points and neuropathologic AD were inconsistently defined and methods with uncertain applicability to typical clinical settings were used.

Olsson et al (2018) reported the results of a systematic review and meta-analysis that evaluated the diagnostic performance of 15 cerebrospinal fluid (CSF) and blood biomarkers for Alzheimer's disease (AD). Analyzing 231 studies involving 15,699 patients with AD and 13,018 controls, the study found that the core CSF biomarkers— $A\beta$ ₄₂, total tau (T-tau), and phosphorylated tau (P-tau)—consistently showed strong ability to distinguish AD from controls and to identify mild cognitive impairment due to AD. CSF neurofilament light (NFL) and plasma T-tau also demonstrated strong associations with AD, while several emerging CSF biomarkers (NSE, VLP-1, HFABP, and YKL-40) showed moderate diagnostic value. Other biomarkers had limited or no discriminatory power. Overall, the findings

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support the continued clinical and research use of CSF A β 42, T-tau, P-tau, and NFL as the most reliable biomarkers for Alzheimer's disease.

PROFESSIONAL GUIDELINE(S)

The Genetic Counseling and Testing for Alzheimer Disease Joint Practice Guidelines by the American College of Medical Genetics and the National Society of Genetic Counselors (Goldman et al 2011; reaffirmed in 2018) state:

- "Do not recommend pediatric testing for AD."
- "Genetic testing for AD should only occur in the context of genetic counseling (in-person or through video conference) and support by someone with expertise in this area."
- "Direct-to-consumer testing of APOE is not advised."
- "At least a three-generation family history should be obtained, with specific information regarding diagnosis of AD in affected family members, along with age of onset and age of death."
- "Testing for genes associated with early-onset autosomal dominant AD (EOAD) should be offered in the following situations:
 - o Symptomatic patients with EOAD with a family history of dementia or unknown family history.
 - o Autosomal dominant family history of dementia with one or more cases of EOAD.
 - o A relative with a pathological variant consistent with EOAD."
- "For families in which autosomal-dominant AD is unlikely:
 - o Genetic testing for susceptibility loci (e.g., APOE) is not clinically recommended due to limited clinical utility."

National Institute for Health and Care Excellence 2018 Guidelines on the Assessment, Management and Support of People Living with Dementia state:

- "APOE genotyping should not be used to diagnose Alzheimer disease."
- "If the diagnosis is uncertain and Alzheimer's disease is suspected you may consider PET, or CSF testing for either total tau or total tau and phosphorylated-tau 181 and either amyloid beta 1-42 or amyloid beta 1-42 and amyloid beta 1-40."

Jack et al (2024) reported The National Institute of Aging and Alzheimer's Association, revised 2018 criteria for the diagnosis and staging of Alzheimer's disease. The workgroup divided biomarkers into three broad categories (below only addresses CSF or plasma samples).

Core biomarkers of AD neuropathologic change (ADNPC)

Core 1

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- Aβ proteinopathy- Aβ 42
- Phosphorylated and secreted AD tau- p-tau217, p-tau181, p-tau231.

Core 2

- AD tau proteinopathy- MTBR-tau243, other phosphorylated tau forms (e.g., p-tau205), non-phosphorylated mid-region tau fragments.

Non-specific biomarkers that are important in AD pathogenesis but are also involved in other brain diseases

- Injury, dysfunction, or degeneration of neuropil- NfL
- Inflammation, Astrocytic activation- Glial Fibrillary Acidic Protein (GFAP)

Biomarkers of common non-AD copathologies

- α-synuclein- αSyn-SAA

They state:

“To date, clinical symptoms have not been used as the reference standard for regulatory approval of AD biomarkers. Our position is that AD is defined by its biology and therefore a biomarker that can accurately detect ADNPC or a validated surrogate is sufficient to establish the diagnosis of the disease.”

“The clinical use of AD biomarkers is presently intended for the evaluation of symptomatic individuals, not cognitively unimpaired individuals. We highlight the distinction between can and should. AD can be diagnosed in asymptomatic individuals, but we do not believe this should be done for clinical purposes at this time.”

“Although, the presence of abnormal Core 1 biomarkers is sufficient for confirming AD pathology in a symptomatic individual, it does not preclude the search for other contributors to the clinical symptoms, particularly other common copathologies.”

“AD biomarkers are fundamental to making an accurate diagnosis and determining likely contributions to the patient's symptoms. Although it is expected that clinicians will use AD biomarkers to determine potential eligibility for recently approved Aβ-specific therapies, clinical applications also include counseling and tailoring medications for symptomatic treatment.”

Shaw et al (2018) reported the Alzheimer's Association published appropriate use criteria for lumbar puncture and cerebrospinal fluid testing in the diagnosis of Alzheimer's disease. The recommendation is that “there should be an initial clinical diagnosis or differential diagnosis, followed by a determination of how the CSF biomarkers might contribute to the diagnosis and to the clinical decision making.” They do discuss the need for further studies that include long-term follow-up, comparing against pathological standards, effects on decision making, and elevated amyloid in cognitively normal individuals.

Hansson et al (2022) reported the Alzheimer's Association appropriate use recommendations for blood-based biomarkers in Alzheimer's disease. They state that “blood-based markers (BBM) should not yet be used as a primary end point in pivotal trials” and “recommend to cautiously start using

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BBM's in specialized memory clinics as part of the diagnostic work-up of patients with cognitive symptoms and the results should be confirmed whenever possible with CSF or PET." More data is needed before BBMs are used as a stand-alone diagnostic test for AD or use in primary care.

REGULATORY STATUS

The United States Food and Drug Administration (FDA) regulates vaccines, blood and blood products, and biologics via the Center for Biologics Evaluation and Research (CBER) which ensures the safety, efficacy, and quality of these products. Refer to the FDA vaccines/blood/biologics website. Available from: <https://www.fda.gov/vaccines-blood-biologics> [accessed 2026 May 22]

The FDA maintains information for consumers and health professionals on vaccine, blood and biologics warnings and other safety information. Available from: [Recalls \(Biologics\) | FDA](#) [accessed 2026 May 22]

In May of 2022, the U.S. Food and Drug Administration (FDA) permitted marketing for Lumipulse G β -Amyloid Ratio (1-42/1-40) immunoassay (Fujirebio Diagnostics, Inc.) The CSF fluid immunoassay was granted breakthrough device designation.

In July 2022, the FDA granted breakthrough device designation to the Elecsys Amyloid Plasma Panel (Roche). Roche has also received a Breakthrough Device Designation for the ElecsysR s-Amyloid (1-42) CSF and Elecsys Phospho-Tau (181P) CSF in vitro diagnostic immunoassays.

In November 2020, C2N Diagnostics gained CLIA certification for its Precivity mass-spec amyloid beta assay. This plasma test has received breakthrough device designation from the FDA for review as an in-vitro diagnostic.

Clinical laboratories may develop and validate tests in-house and market them as laboratory service; laboratory-developed tests must meet the general regulatory standards of the Clinical Laboratory Improvement Amendments (CLIA). Laboratories that offer laboratory-developed tests must be certified by CLIA for high-complexity testing. To date, the FDA has chosen not to require any regulatory review of these tests. AlzheimerAlert and AdMark CSF analysis are examples of tests that may be available in CLIA certified labs.

Additional diagnostic blood tests that have received FDA breakthrough device designation include AlzoSure Predict (Diadem) in January 2022 and SOBA-AD (AltPep Corporation) in March 2022.

CODE(S)

- Codes may not be covered under all circumstances.
- Code list may not be all inclusive (AMA and CMS code updates may occur more frequently than policy updates).
- (E/I) =Experimental/Investigational
- (NMN)=Not medically necessary/appropriate

CPT Codes

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Code	Description
81401	Molecular pathology procedure level 2 (e.g., 2-10 SNPs, 1 methylated variant, or 1 somatic variant [typically using nonsequencing target variant analysis], or detection of a dynamic mutation disorder/triplet repeat). Includes APOE (apolipoprotein E) (e.g., hyperlipoproteinemia type III, cardiovascular disease, Alzheimer disease), common variants (e.g., *2, *3, *4)
81405	Molecular pathology procedure level 6 (e.g., analysis of 6-10 exons by DNA sequence analysis, mutation scanning or duplication/deletion variants of 11-25 exons, regionally targeted cytogenomic array analysis). Includes PSEN1 (presenilin 1) (e.g., Alzheimer's disease), full gene sequence
81406	Molecular pathology procedure level 7 (e.g., analysis of 11-25 exons by DNA sequence analysis, mutation scanning or duplication/deletion variants of 26-50 exons). Includes PSEN2 (presenilin 2 [Alzheimer's disease 4]) (e.g., Alzheimer's disease), full gene sequence and APP (amyloid beta [A4] precursor protein) (eg, Alzheimer's disease), full gene sequence
82172	Apolipoprotein, each
82233	Beta-amyloid; 1-40 (Abeta 40)
82234	Beta-amyloid; 1-42 (Abeta 42)
83884 (E/I)	Neurofilament light chain (NfL)
84393	Tau, phosphorylated (e.g., pTau 181, pTau 217), each
84394	Tau, total (tTau)
0206U (E/I)	Neurology (Alzheimer disease); cell aggregation using morphometric imaging and protein kinase C-epsilon (PKCe) concentration in response to amylospheroid treatment by ELISA, cultured skin fibroblasts, each reported as positive or negative for Alzheimer disease (DISCERN, NeuroDiagnostics)

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Code	Description
0207U (E/I)	Neurology (Alzheimer disease); quantitative imaging of phosphorylated ERK1 and ERK2 in response to bradykinin treatment by in situ immunofluorescence, using cultured skin fibroblasts, reported as a probability index for Alzheimer disease (List separately in addition to code for primary procedure) (DISCERN, NeuroDiagnostics)
0289U (E/I)	Neurology (Alzheimer disease), mRNA, gene expression profiling by RNA sequencing of 24 genes, whole blood, algorithm reported as predictive risk score (MindX Blood Test-Memory/Alzheimer's, MindX Sciences Laboratory)
0358U	Neurology (mild cognitive impairment), analysis of B-amyloid 1-42 and 1-40, chemiluminescence enzyme immunoassay, cerebral spinal fluid, reported as positive, likely positive, or negative (Lumipulse G β -Amyloid Ratio (1-42/1-40) Test, Fujirebio Diagnostics, Inc)
0393U (E/I)	Neurology (e.g., Parkinson disease, dementia with Lewy bodies), cerebrospinal fluid (CSF), detection of misfolded α -synuclein protein by seed amplification assay, qualitative (SAAmplify α -SYNTap, Amprion Clinical Laboratory)
0412U (E/I)	Beta amyloid, A β 42/40 ratio, immunoprecipitation with quantitation by liquid chromatography with tandem mass spectrometry (LC-MS/MS) and qualitative ApoE isoform-specific proteotyping, plasma combined with age, algorithm reported as presence or absence of brain amyloid pathology. (PrecivityAD blood test)
0443U (E/I)	Neurofilament light chain (NfL), ultra-sensitive immunoassay, serum or cerebrospinal fluid (Neurofilament Light Chain (NfL), Neuromuscular Clinical Laboratory at Washington University in St. Louis School of Medicine)
0445U	B-amyloid (Abeta42) and phospho tau (181P) (pTau181), electrochemiluminescent immunoassay (ECLIA), cerebral spinal fluid, ratio reported as positive or negative for amyloid pathology (Elecsys Phospho-Tau CSF [pTau181] and β -Amyloid [1-42] CSF II [Abeta 42] Ratio, Roche Diagnostics Operations)
0459U	B-amyloid (Abeta42) and total tau (tTau), electrochemiluminescent immunoassay (ECLIA), cerebral spinal fluid, ratio reported as positive or negative for amyloid pathology

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Code	Description
	(Elecsys Total Tau CSF (tTau) and β -Amyloid CSF II Ratio, Roche Diagnostics Operations, Inc)
0479U	Tau, phosphorylated, pTau217 (ALZpath pTau217, Neurocode USA, Inc, Quanterix/ALZpath)
0503U (E/I)	Neurology (Alzheimer disease), beta amyloid (AB40, AB42, AB42/40 ratio) and tau-protein (ptau217, np-tau217, ptau217/np-tau217 ratio), blood, immunoprecipitation with quantitation by liquid chromatography with tandem mass spectrometry (LC-MS/MS), algorithm score reported as likelihood of positive or negative for amyloid plaques (PrecivityAD2, C2N Diagnostics, LLC)
0568U (E/I)	Neurology (dementia), beta amyloid (AB40, AB42, AB42/40 ratio), tau-protein phosphorylated at residue (eg, pTau217), neurofilament light chain (NfL), and glial fibrillary acidic protein (GFAP), by ultra-high sensitivity molecule array detection, plasma, algorithm reported as positive, intermediate, or negative for Alzheimer pathology (LucentAD Complete, Quanterix Corporation)
0596U (E/I)	Neurology (Alzheimer disease), plasma, 3 distinct isoform-specific peptides (APOE2, APOE3, and APOE4) by liquid chromatography with tandem mass spectrometry (LC- MS/MS), reported as an APOE prototype (effective 10/01/25) (Precivity-ApoE, C2N Diagnostics, LLC)

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HCPCS Codes

Code	Description
S3852	DNA analysis for APOE epsilon 4 allele for susceptibility to Alzheimer's disease

ICD10 Codes

Code	Description
F03.90- F03.91	Unspecified dementia, unspecified severity, without behavioral disturbance, psychotic disturbance, mood disturbance, and anxiety
G30.0-G30.9	Alzheimer's Disease (code range)
G31.1	Senile degeneration of brain, not elsewhere classified

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Code	Description
G31.84	Mild cognitive impairment of uncertain or unknown etiology
R41.81	Age-related cognitive decline

REFERENCES

Amft M, et al. The cerebrospinal fluid biomarker ratio A β 42/40 identifies amyloid positron emission tomography positivity better than A β 42 alone in a heterogeneous memory clinic cohort. *Alzheimers Res Ther*. 2022 Apr 26;14(1):60.

APA Work Group on Alzheimer's Disease and other Dementias; Rabins PV, et al. American Psychiatric Association practice guideline for the treatment of patients with Alzheimer's disease and other dementias. Second edition. *Am J Psychiatry*. 2007 Dec;164(12 Suppl):5-56.

Arvanitakis Z, et al. Diagnosis and management of dementia: review. *JAMA*. 2019 Oct 22;322(16):1589-1599.

Banning LCP, et al. Apolipoprotein E and affective symptoms in mild cognitive impairment and Alzheimer's disease dementia: a systematic review and meta-analysis. *Neurosci Biobehav Rev*. 2019 Jan;96:302-315.

Bertram L and Tanzi RE. Alzheimer's disease: one disorder, too many genes? *Hum Mol Genet*. 2004 Apr 1;13 Spec No 1:R135-41

Bertram L, et al. Systematic meta-analyses of Alzheimer disease genetic association studies: the AlzGene database. *Nat Genet*. 2007 Jan;39(1):17-23.

Bird T. [Internet]. Alzheimer disease overview: gene reviews. Initial posting 1998 Oct 23 [updated 2018 Dec 20; accessed 2026 Apr 21]. Available from: <http://www.ncbi.nlm.nih.gov/books/NBK1161/>

Bird T. [Internet]. Early-onset familial alzheimer disease. Initial posting 1999 Sept 24 [update 2012 Oct 18; accessed 2026 Apr 21]. Retired. Available from: <http://www.ncbi.nlm.nih.gov/books/NBK1236/>

Bolsewig K, European-Dementia with Lewy Bodies (E-DLB) Consortium, et al. Association of Plasma Amyloid, P-Tau, GFAP, and NfL with CSF, clinical, and cognitive features in patients with dementia with lewy bodies. *Neurology*. 2024 Jun 25;102(12):e209418.

Borenstein AR, et al. Early-life risk factors for Alzheimer disease. *Alzheimer Dis Assoc Disord*. 2006 Jan-Mar;20(1):63-72.

Brickell KL, et al. Early-onset Alzheimer disease in families with late-onset Alzheimer disease: a potential important subtype of familial Alzheimer disease. *Arch Neurol*. 2006 Sep;63(9):1307-11.

Cassidy MR, et al. Comparing test-specific distress of susceptibility versus deterministic genetic testing for Alzheimer's disease. *Alzheimer's Dement*. 2008 Nov;4(6):406-13.

Cummings J, et al. Lecanemab: appropriate use recommendations. *J Prev Alzheimers Dis*.

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2023;10(3):362-377.

Erickson P, et al. Prevalence and clinical implications of a β -amyloid-negative, tau-positive cerebrospinal fluid biomarker profile in Alzheimer Disease. *JAMA Neurol*. 2023 Jul 31;80(9):969–79.

Fink HA, et al. Accuracy of biomarker testing for neuropathologically defined alzheimer disease in older adults with dementia. *Ann Intern Med*. 2020 May 19;172(10):669-677.

Goldman JS, et al. Genetic testing and counseling for Alzheimer's disease: joint practice guidelines of the American College of Genetics and the National Society of Genetic Counselors. *Genet Med*. 2011;13(6):597-605.

Goldman JS, et al. ADDENDUM: Genetic counseling and testing for Alzheimer disease: joint practice guidelines of the American College of Medical Genetics and the National Society of Genetic Counselors. *Genet Med*. 2019 Oct;21(10):2404.

Hansson O, et al. The Alzheimer's Association appropriate use recommendations for blood biomarkers in Alzheimer's disease. *Alzheimer's Dement*. 2022 Dec;18(12):2669-2686.

Heshka JT, et al. A systematic review of perceived risks, psychological and behavioral impacts of genetic testing. *Genet Med*. 2008;10(1):19-32.

Hu J, et al. Preclinical development of an immunoassay for the detection of TREM2: a new biomarker for Alzheimer's disease. *Sci Rep*. 2025 Jul 22;15(1):26525.

Jack CR Jr, et al. Revised criteria for diagnosis and staging of Alzheimer's disease: Alzheimer's Association Workgroup. *Alzheimers Dement*. 2024 Aug;20(8):5143-5169.

Kaplow J, et al. Concordance of Lumipulse cerebrospinal fluid t-tau/A β 42 ratio with amyloid PET status. *Alzheimers Dement*. 2020 Jan;16(1):144-152.

Kim DH, et al. Genetic markers for diagnosis and pathogenesis of Alzheimers disease. *Gene*. 2014;545(2):185-93.

Knopman DS, et al. Practice parameter: diagnosis of dementia (an evidence-based review). Report of the Quality Standards Subcommittee of the American Academy of Neurology. *Neurology*. 2001 May 8;56(9):1143-53.

Krishna G, et al. Potential utility of plasma biomarker panels in differential diagnosis of alzheimer's disease. *J Alzheimers Dis Rep*. 2024 Jan 9;8(1):1-7.

Langlois CM, et al. Alzheimer's prevention initiative generation program: development of an APOE genetic counseling and disclosure process in the context of clinical trials. *Alzheimers Dement (NY)*. 2019 Nov 6;5:705-716.

Larner AJ, Doran M. Clinical phenotypic heterogeneity of Alzheimer's disease associated with mutations of the presenilin-1 gene. *J Neurol*. 2006 Feb;253(2):139-58.

Limon-Sztencel A, et al. The algorithm for Alzheimer risk assessment based on APOE promoter polymorphisms. *Alzheimers Res Ther*. 2016 May19;8(1):19.

Liu Y, et al. APOE genotype and neuroimaging markers of Alzheimer's disease: systematic review and

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meta-analysis. J Neurol Neurosurg Psychiatry. 2015;86(2):127-34.

McKhann GM, et al. The diagnosis of dementia due to Alzheimer's disease: recommendations from the National Institute on Aging-Alzheimer's Association workgroups on diagnostic guidelines for Alzheimer's disease. Alzheimers Dement. 2011 May;7(3):263-9.

Mielke MM, et al. Influence of amyloid and APOE on cognitive performance in a late middle-aged cohort. Alzheimers Dement. 2016 Mar;12(3):281-91.

National Institute for Health and Care Excellence (NICE) [Internet]. Dementia: assessment, management and support for people living with dementia and their carers [NG97]. 2018 [accessed 2026 Apr 21]. Available from: <https://www.nice.org.uk/guidance/ng97>

National Institute on Aging (NIA) [Internet]. Alzheimer's Disease Genetics Fact Sheet. Reviewed Mar 1, 2023. [accessed 2026 Apr 21]. Available from: <https://www.nia.nih.gov/health/alzheimers-disease-genetics-fact-sheet>

Olsson B, et al. CSF and blood biomarkers for the diagnosis of Alzheimer's disease: a systematic review and meta-analysis. Lancet Neurol. 2016 Jun;15(7):673-684.

Pahlke S, et al. Blood-based biomarkers for detecting Alzheimer's disease pathology in cognitively impaired individuals within specialized care settings: A systematic review and meta-analysis. Alzheimers Dement. 2025 Nov;21(11):e70828.

Palmqvist S, et al. Alzheimer's Association Clinical Practice Guideline on the use of blood-based biomarkers in the diagnostic workup of suspected Alzheimer's disease within specialized care settings. Alzheimers Dement. 2025 Jul;21(7):e70535.

Rabinovici GD, et al. Donanemab: Appropriate use recommendations. J Prev Alzheimers Dis. 2025 May;12(5):100150.

Real R, et al. Association between the LRP1B and APOE loci and the development of Parkinson's disease dementia. Brain. 2023 May 2;146(5):1873-1887.

Ren Z, et al. Association between apolipoprotein E ε4 status and the risk of Alzheimer's disease: a meta-analysis. BMC Neurosci. 2025 Jan 24;26(1):5.

Safransky M, et al. Lumipulse-measured cerebrospinal fluid biomarkers for the early detection of alzheimer disease. Neurology. 2024 Dec 10;103(11):e209866.

Schraen-Maschke S, BALTAZAR study group, et al. The free plasma amyloid Aβ₁₋₄₂/Aβ₁₋₄₀ ratio predicts conversion to dementia for subjects with mild cognitive impairment with performance equivalent to that of the total plasma Aβ₁₋₄₂/Aβ₁₋₄₀ ratio. The BALTAZAR study. Neurobiol Dis. 2024 Apr;193:106459.

Shaw LM, et al. Appropriate use criteria for lumbar puncture and cerebrospinal fluid testing in the diagnosis of Alzheimer's disease. Alzheimer's Dement. 2018 Nov;14(11):1505-1521.

Sinclair LI, et al. Possible positive effect of the APOE ε2 allele on cognition in early to mid-adult life. Neurobiol Learn Mem. 2017 Dec;146:37-46.

Swanson CJ, et al. A randomized, double-blind, phase 2b proof-of-concept clinical trial in early

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Alzheimer's disease with lecanemab, an anti-A β protofibril antibody. *Alzheimers Res Ther.* 2021 Apr 17;13(1):80.

van Dyck CH, et al. Lecanemab in Early Alzheimer's Disease. *N Engl J Med.* 2023 Jan 5;388(1):9-21.

Vilatela A, et al. Genetics of Alzheimer's disease. *Arch Med Res.* 2012 Nov;43(8):622-31.

Vöglein J, et al. Clinical, pathophysiological and genetic features of motor symptoms in autosomal dominant Alzheimer's disease. *Brain.* 2019 May 1;142(5):1429-1440.

Wishart HA, et al. Increased brain activation during working memory in cognitively intact adults with the APOE epsilon4 allele. *Am J Psychiatry.* 2006 Sep;163(9):1603-10.

Zou Z, et al. Clinical genetics of Alzheimer's disease. *Biomed Res Int.* 2014;2014:291862.

SEARCH TERMS

Not Applicable

CENTERS FOR MEDICARE AND MEDICAID SERVICES (CMS)

[Molecular Pathology Procedures \(LCD L35000\)](#) [accessed 2026 Apr 21]

[Article - Billing and Coding: Molecular Pathology Procedures \(LCA A56199\)](#) [accessed 2026 Apr 21]

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- Services are contract dependent; if a product does not cover a service, medical policy criteria do not apply.
- If a commercial product (including an Essential Plan or Child Health Plus product) covers a specific service, medical policy criteria apply to the benefit.
- If a Medicaid product covers a specific service, and there are no New York State Medicaid guidelines (eMedNY) criteria, medical policy criteria apply to the benefit.
- If a Medicare product (including Medicare HMO-Dual Special Needs Program (DSNP) product) covers a specific service, and there is no national or local Medicare coverage decision for the service, medical policy criteria apply to the benefit.
- If a Medicare HMO-Dual Special Needs Program (DSNP) product DOES NOT cover a specific service, please refer to the Medicaid Product coverage line.

POLICY HISTORY/REVISION

Committee Approval Dates

12/20/01, 09/19/02, 07/17/03, 06/17/04, 06/16/05, 04/20/06, 02/15/07, 03/20/08, 05/28/09, 05/27/10, 05/19/11, 03/15/12, 03/21/13, 03/20/14, 03/19/15, 03/17/16, 03/16/17, 04/19/18, 05/16/19, 05/21/20, 05/20/21, 05/19/22, 05/18/23, 05/16/24, 06/26/25, 06/18/26

Date

Summary of Changes

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06/18/26	<ul style="list-style-type: none">Annual Review; policy intent unchanged.
06/26/25	<ul style="list-style-type: none">Annual Review, policy statements added for biomarkers for Alzheimer's disease as investigational. Codes added to policy 82233,82234,83884, 84393, 84394, 0206U, 0207U, 0358U, 0361U, 0393U, 0443U, 0445U, 0459U, 0479U, 0503U, 0568U and Deletion of code 0346U.
01/01/25	<ul style="list-style-type: none">Summary of changes tracking implemented.
12/20/01	<ul style="list-style-type: none">Original effective date