Alzheimer’s disease normative cerebrospinal fluid biomarkers validated in PET amyloid-β characterized subjects from the Australian Imaging, Biomarkers and Lifestyle (AIBL) study

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**Affiliations:**[^a] Florey Institute of Neuroscience and Mental Health, The University of Melbourne, VIC, Australia |[^b] Department of Nuclear Medicine and Centre for PET, Austin Health, Heidelberg, VIC, Australia |[^c] CSIRO Digital Productivity/Australian e-Health Research Centre and Cooperative Research Centre
Background: The cerebrospinal fluid (CSF) amyloid-\(\text{\textregistered}\) (A\(\text{\textregistered}\)1-42, total-tau (T-tau), and phosphorylated-tau (P-tau181P) profile has been established as a valuable biomarker for Alzheimer’s disease (AD). Objective: The current study aimed to determine CSF biomarker cut-points using positron emission tomography (PET) A\(\text{\textregistered}\) imaging screened subjects from the Australian Imaging, Biomarkers and Lifestyle (AIBL) study of aging, as well as correlate CSF analyte cut-points across a range of PET A\(\text{\textregistered}\) amyloid ligands. Methods: A\(\text{\textregistered}\) pathology was determined by PET imaging, utilizing 11C-Pittsburgh Compound B, 18F-flutemetamol, or 18F-florbetapir, in 157 AIBL participants who also underwent CSF collection. Using an INNOTEST assay, cut-points were established (A1-42 >544 ng/L, T-tau <407 ng/L, and P-tau181P <78 ng/L) employing a rank based method to define a “positive” CSF in the sub-cohort of amyloid-PET negative healthy participants (n=97), and compared with the presence of PET demonstrated AD pathology. Results: CSF A\(\text{\textregistered}\)1-42 was the strongest individual biomarker, detecting cognitively impaired PET positive mild cognitive impairment (MCI)/AD with 85% sensitivity and 91% specificity. The ratio of P-tau181P or T-tau to A\(\text{\textregistered}\)1-42 provided greater accuracy, predicting MCI/AD with 92% sensitivity and specificity. Cross-validated accuracy, using all three biomarkers or the ratio of P-tau or T-tau to A\(\text{\textregistered}\)1-42 to predict MCI/AD, reached \(\text{\textregistered}\)92% sensitivity and specificity. Conclusions: CSF A\(\text{\textregistered}\)1-42 levels and analyte combination ratios demonstrated very high correlation with PET A\(\text{\textregistered}\) imaging. Our study offers additional support for CSF biomarkers in the early and accurate detection of AD pathology, including enrichment of patient cohorts for treatment trials even at the pre-symptomatic stage.

Keywords: Alzheimer’s disease, amyloid-\(\text{\textregistered}\), cerebrospinal fluid biomarkers, positron emission tomography A\(\text{\textregistered}\) imaging, tau

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