

RESEARCH

Open Access



Plasma amyloid- β levels, cerebral atrophy and risk of dementia: a population-based study

Saima Hilal^{1,2}, Frank J. Wolters², Marcel M. Verbeek³, Hugo Vanderstichele⁴, M. Kamran Ikram^{2,5}, Erik Stoops⁴, M. Arfan Ikram^{1,2} and Meike W. Vernooij^{1,2*}

Abstract

Background: Plasma amyloid- β (A β) levels are increasingly studied as a potential accessible marker of cognitive impairment and dementia. However, it remains underexplored whether plasma A β levels including the novel A β peptide 1–38 (A β _{1–38}) relate to preclinical markers of neurodegeneration and risk of dementia. We investigated the association of plasma A β _{1–38}, A β _{1–40}, and A β _{1–42} levels with imaging markers of neurodegeneration and risk of dementia in a prospective population-based study.

Methods: We analyzed plasma A β levels in 458 individuals from the Rotterdam Study. Brain volumes, including gray matter, white matter, and hippocampus, were computed on the basis of 1.5-T magnetic resonance imaging (MRI). Dementia and its subtypes were defined on the basis of internationally accepted criteria.

Results: A total of 458 individuals (mean age, 67.8 \pm 7.7 yr; 232 [50.7%] women) with baseline MRI scans and incident dementia were included. The mean \pm SD values of A β _{1–38}, A β _{1–40}, and A β _{1–42} (in pg/ml) were 19.4 \pm 4.3, 186.1 \pm 35.9, and 56.3 \pm 6.2, respectively, at baseline. Lower plasma A β _{1–42} levels were associated with smaller hippocampal volume (mean difference in hippocampal volume per SD decrease in A β _{1–42} levels, –0.13; 95% CI, –0.23 to –0.04; p = 0.007). After a mean follow-up of 14.8 years (SD, 4.9; range, 4.1–23.5 yr), 79 persons developed dementia, 64 of whom were diagnosed with Alzheimer's disease (AD). Lower levels of A β _{1–38} and A β _{1–42} were associated with increased risk of dementia, specifically AD (HR for AD per SD decrease in A β _{1–38} levels, 1.39; 95% CI, 1.00–2.16; HR for AD per SD decrease in A β _{1–42} levels, 1.35; 95% CI, 1.05–1.75) after adjustment for age, sex, education, cardiovascular risk factors, apolipoprotein E ϵ 4 allele carrier status, and other A β isoforms.

Conclusions: Our results show that lower plasma A β levels were associated with risk of dementia and incident AD. Moreover, lower plasma A β _{1–42} levels were related to smaller hippocampal volume. These results suggest that plasma A β _{1–38} and A β _{1–42} maybe useful biomarkers for identification of individuals at risk of dementia.

Keywords: Plasma amyloid- β levels, Magnetic resonance imaging, Atrophy, Dementia, Population-based

Background

Dementia due to Alzheimer's disease (AD), constituting up to 70% of all dementias, is characterized by deposition of amyloid plaques and neurofibrillary tangles in the brain parenchyma [1]. Amyloid- β (A β) 1–40 (A β _{1–40}) and A β _{1–42}

peptides derived from amyloid precursor protein are present in these parenchymal plaques, and it is hypothesized that A β peptide abnormalities begin early in the neurodegenerative pathological cascade [2, 3]. According to this amyloid cascade hypothesis, an imbalance between the production and clearance of the A β peptides, especially the A β _{1–42} isoform, leads to their aggregation in the cortical tissue and vessels [3]. This A β deposition in the brain subsequently affects plasma concentrations, thus suggesting that circulating levels of A β could potentially be used as markers of disease risk [4].

* Correspondence: m.vernooij@erasmusmc.nl

¹Department of Radiology and Nuclear Medicine, Erasmus University Medical Center, Office no. 2505, Wytemaweg 80, 3015, CN, Rotterdam, The Netherlands

²Department of Epidemiology, Erasmus University Medical Center, Rotterdam, The Netherlands

Full list of author information is available at the end of the article



$A\beta_{1-40}$ and $A\beta_{1-42}$ peptides are the two most studied AD biomarkers, which can be measured in plasma through minimally invasive techniques and thus can feasibly be determined in large samples. Several large community-based studies have reported on the association of high plasma $A\beta_{1-40}$ and $A\beta_{1-42}$ levels and lower $A\beta_{1-42}/A\beta_{1-40}$ ratios at baseline with risk of dementia and reduced cognitive measure scores [5–7]. Two studies have shown a link between higher levels of $A\beta_{1-42}$ but not $A\beta_{1-40}$ with risk of AD [8, 9]. Conversely, some studies reported an association of increased concentrations of $A\beta_{1-42}$ and $A\beta_{1-40}$ levels and higher $A\beta_{1-42}/A\beta_{1-40}$ ratio with reduced risk of dementia [10, 11]. Longitudinal studies with repeated measurements of $A\beta$ have shown that individuals in preclinical stages of the disease [12] and persons with newly diagnosed AD [6] showed significant reductions in plasma $A\beta_{1-42}$ concentrations over time. Though most of the study results are inconsistent, they nevertheless do suggest that plasma $A\beta$ levels may be differentially associated with the risk for AD, possibly reflective of the stage of the disease and the complex pattern of production and clearance from the brain [4]. Limited data also suggest that increased circulating levels of $A\beta_{1-40}$ and $A\beta_{1-42}$ may induce cerebral atrophy, detected on magnetic resonance imaging (MRI) studies as cortical thinning in asymptomatic elderly persons through mechanisms such as synaptic deficits and neuronal loss [13–15]. More studies of a general population are needed to investigate the relationship between $A\beta$ and markers of neurodegeneration as well as risk of dementia to further comprehend the underlying mechanisms in a subclinical phase.

Besides $A\beta_{1-40}$ and $A\beta_{1-42}$ peptides, which have been investigated extensively, another isoform— $A\beta_{1-38}$ —is of interest because it is reported to be elevated in the cerebrospinal fluid (CSF) of patients with sporadic AD [16]. Moreover, it is suggested to be a sensitive and specific marker for diagnosing AD over the range of other non-AD dementias, and subsequent studies confirmed the suitability of $A\beta_{1-38}$ as an additional biomarker for differential diagnosis of dementia [17, 18]. Despite this ongoing research, little is known regarding whether plasma $A\beta$ levels, including the novel $A\beta_{1-38}$, are associated with preclinical markers of neurodegeneration, such as gray matter, white matter, and hippocampal atrophy, and the risk of dementia in a large population-based setting. We examined the association of plasma $A\beta_{1-38}$, $A\beta_{1-40}$, and $A\beta_{1-42}$ levels with neurodegenerative markers and risk of dementia in a subsample of the Rotterdam Study.

Methods

Study population

The Rotterdam Study is a population-based prospective cohort study of middle-aged and elderly persons living in the Ommoord district in the city of Rotterdam, the

Netherlands. All participants in this study undergo reexamination every 3–4 years and are being followed continuously for events, including occurrence of dementia. This study is embedded within the first cohort, which was initiated in 1990 with 7983 participants at baseline (aged ≥ 55 yr). Persons were randomly selected ($n = 563$) during the second visit of the first cohort (1995–1996) and were invited to undergo neuroimaging [19]. Blood samples of the same individuals were drawn in the 1998 and 1999. After individuals with insufficient plasma ($n = 22$), no MRI scans ($n = 73$), and prevalent dementia ($n = 10$) were removed, the final sample size consisted of 458 persons [20].

Plasma assessment

Blood samples were drawn into ethylenediaminetetraacetic acid (EDTA) tubes for plasma collection. After centrifugation ($2500 \times g$, $+4^\circ\text{C}$ for 20 min), plasma samples were stored at -80°C within 60 minutes of collection. Plasma levels of $A\beta_{1-38}$, $A\beta_{1-40}$, and $A\beta_{1-42}$ were quantified by EUROIMMUN β -Amyloid 1–38, 1–42, and 1–40 plasma enzyme-linked immunosorbent assays (EUROIMMUN, Lübeck, Germany), which have been validated and described in more detail previously [21, 22]. For quality control (QC) purposes, QC samples were produced by pooling of EDTA plasma samples from individual participants. After aliquoting, samples were stored at -80°C . The samples were coded QC1, QC2, QC8, QC9, and QC10 and were used in the three plasma amyloid assays. QC in the three assays was ± 2 SD of each amyloid concentration (in pg/ml) across study participants, with the ranges of concentration values detectable being 5.9–18.7 pg/ml for $A\beta_{1-38}$, 67.6–161.8 pg/ml for $A\beta_{1-40}$, and 46.6–55.6 pg/ml for $A\beta_{1-42}$. The average coefficients of variation of measurement of $A\beta_{1-38}$, $A\beta_{1-40}$, and $A\beta_{1-42}$ in QC plasma samples during the study were 11.04%, 5.72%, and 8.70%, respectively [22].

Brain imaging

Brain MRI was performed using a 1.5-T MRI system (VISION MR; Siemens AG, Erlangen, Germany) to obtain T1-weighted, proton density, T2-weighted, and high-resolution inversion recovery double-contrast three-dimensional half-Fourier-acquired single-shot turbo spin echo (HASTE) sequences [23]. Image preprocessing and the tissue classification algorithm have been described elsewhere [23]. Briefly, the k -nearest neighbors brain tissue classifier technique was used to classify voxels into CSF, gray matter, normal white matter, and white matter hyperintensities. Intracranial volume was the sum of the CSF, gray matter, normal white matter, and white matter hyperintensities. We used a validated nonrigid registration algorithm to map brain regions to the template scan. Hippocampal volumes were manually outlined on coronal

HASTE sequences perpendicular to the long axis of the hippocampus [24].

Assessment of dementia

Participants were screened for dementia at baseline and at follow-up examinations [20]. Screening was performed using the Mini Mental State Examination (MMSE) and the Geriatric Mental State Schedule (GMS) organic level. Individuals with screen-positive results (MMSE < 26 or GMS organic level > 0) subsequently underwent an examination and informant interview with the Cambridge Examination for Mental Disorders of the Elderly. Additionally, the whole cohort was continuously monitored for dementia through computerized linkage of the study database and digitized medical records of general practitioners and the Regional Institute for Outpatient Mental Health Care. When required and available, neuroimaging was used to facilitate dementia diagnosis. A consensus panel led by a consultant neurologist established the final diagnosis according to standard criteria for dementia (*Diagnostic and Statistical Manual of Mental Disorders, Third Edition-Revised*). The diagnosis of AD was made using the National Institute of Neurological and Communicative Disorders and Stroke-Alzheimer's Disease and Related Disorders Association [25] and National Institute of Neurological Disorders and Stroke-Association Internationale pour la Recherche et l'Enseignement en Neurosciences criteria for vascular dementia [26]. The first cohort was followed for dementia until 15 years (based on maximum follow-up) after baseline examination (i.e., January 2015). Follow-up for dementia was complete for 99.5% of potential person-years in this cohort.

Covariate assessment

Data on demographics and medical history were recorded on the same day of dementia screening. Blood pressure was measured in two readings using a random zero sphygmomanometer in a sitting position, and the mean of both measurements was calculated. Mean arterial blood pressure was calculated as two-thirds of the diastolic blood pressure plus one-third of the systolic blood pressure. Serum total cholesterol levels were measured using an automated enzymatic procedure. Diabetes mellitus was defined as fasting blood glucose ≥ 7 mmol/L or receiving treatment for diabetes. Smoking was categorized into ever versus never smokers. Education was treated as the number of years of formal education. Apolipoprotein E (APOE) genotype was determined using PCRs on coded DNA samples. Distribution of APOE genotype and allele frequencies were in Hardy-Weinberg equilibrium. APOE- $\epsilon 4$ carrier status was defined by the presence of at least one $\epsilon 4$ allele.

Statistical analysis

Plasma A β levels and brain tissue volumes were standardized (by subtracting each variable by population mean divided by SD). Plasma A β levels were expressed as per-SD decrease. The A β_{1-40} /A β_{1-42} ratio was calculated using raw values of A β_{1-40} and A β_{1-42} levels. We first performed linear regression models to determine the association between A β levels and brain tissue volumes (total brain volumes, white matter volume, gray matter volume, and hippocampal volume). The models were adjusted for age, sex, intracranial volume, mean arterial blood pressure, total cholesterol, diabetes, smoking, and APOE- $\epsilon 4$ carrier status. We tested the independent effects of A β levels with brain tissue volumes by adding all three isoforms together in the regression models.

Using Cox proportional hazards models, we calculated HRs with corresponding 95% CIs for dementia and its subtypes with per-SD decrease in A β level. Participants were censored within the follow-up period at date of event diagnosis, death, or loss to follow-up, whichever came first. The proportional hazards assumption was tested by adding the interaction terms of A β_{1-38} , A β_{1-40} , A β_{1-42} , and A β_{1-40} /A β_{1-42} ratio with follow-up time in different models. All Cox proportional hazards models were initially adjusted for age, sex, and education and subsequently for mean arterial blood pressure, total cholesterol, diabetes, smoking, APOE- $\epsilon 4$ carrier status, and the other A β isoforms using a similar approach to the one described above. We also investigated whether the association between plasma A β and dementia was different in carriers and noncarriers of the APOE- $\epsilon 4$ allele. The level of significance was set to 5%, and all tests were two-sided. Statistical analyses were performed using IBM SPSS Statistics version 24 software (IBM, Armonk, NY, USA).

Results

Baseline characteristics of the participants are shown in Table 1. The mean age of the participants was 67.8 ± 7.7 years, and 232 (50.7%) were women. Diabetes was present in 55 (12.1%) of the study population, whereas the frequency of ever smokers was 317 (69.7%). Almost 30% ($n = 137$) persons were APOE- $\epsilon 4$ carriers. The mean \pm SD values of A β_{1-38} , A β_{1-40} , and A β_{1-42} were 19.4 ± 4.3 , 186.1 ± 35.9 , and 56.3 ± 6.2 , whereas for white matter volume, gray matter volume, and hippocampal volume, the respective values were 354.1 ± 85.4 , 522.7 ± 55.8 , and 6.4 ± 0.9 . The correlation between A β_{1-38} and A β_{1-40} was 0.81 (Pearson's correlation coefficient, R); between A β_{1-38} and A β_{1-42} , $R = 0.24$, and between A β_{1-40} and A β_{1-42} , $R = 0.25$.

Table 2 shows the cross-sectional analysis between A β levels and brain tissue volumes adjusted for age, sex, vascular risk factors, APOE- $\epsilon 4$ carrier status, intracranial volume, and other A β isoforms, when appropriate. Lower A β_{1-38} , A β_{1-40} , and A β_{1-42} levels and lower A β_{1-40} /A β_{1-42}

Table 1 Baseline characteristics of the study population

| Variables | Subsample of Rotterdam Study (n = 458) |
|--|---|
| Demographic and vascular risk factors | |
| Age, yr, mean (SD) | 67.8 (7.7) |
| Women, n (%) | 232 (50.7) |
| Education, yr, mean (SD) | 10.7 (3.4) |
| Mean arterial blood pressure, mmHg, mean (SD) | 96.5 (12.7) |
| Total cholesterol, mmol/L, mean (SD) | 5.7 (0.93) |
| Diabetes mellitus, n (%) | 55 (12.1) |
| Smoker, ever, n (%) | 317 (69.7) |
| APOE-ε4 carriers, n (%) | 137 (30) |
| Plasma levels of Aβ isoforms, pg/ml, mean (SD) | |
| Aβ ₁₋₃₈ | 19.4 (4.3) |
| Aβ ₁₋₄₀ | 186.1 (35.9) |
| Aβ ₁₋₄₂ | 56.3 (6.2) |
| MRI markers, ml, mean (SD) | |
| White matter volume | 354.1 (85.4) |
| Gray matter volume | 522.7 (55.8) |
| Intracranial volume | 1126.2 (113.9) |
| Hippocampus volume | 6.4 (0.9) |

Abbreviations: , APOE Apolipoprotein, Aβ Amyloid-β, MRI Magnetic resonance imaging

ratio were not associated with total brain, gray, and white matter volumes. A significant association was observed between lower plasma Aβ₁₋₄₂ levels and smaller hippocampal volume (mean difference in hippocampal volume per SD decrease in Aβ₁₋₄₂ levels, -0.13; 95% CI, -0.23 to -0.04; $p = 0.007$).

During a mean follow-up of 14.8 years (SD, 4.9; range, 4.1–23.5 yr), 241 individuals died. In the same follow-up, 79 persons developed dementia. Of these 79 individuals with dementia, 64 were diagnosed with AD and 15 with vascular dementia. The interaction terms of Aβ levels with follow-up time in all models were nonsignificant (p value for interaction between Aβ₁₋₃₈ and time, $p = 0.832$; between Aβ₁₋₄₀ and time, $p = 0.820$; between Aβ₁₋₄₂ and

time, $p = 0.998$; and between Aβ₁₋₄₀/Aβ₁₋₄₂ ratio and time, $p = 0.984$). In the multivariable analysis, lower levels of Aβ₁₋₃₈ and Aβ₁₋₄₂ were associated with increased risk of dementia (HR for dementia per SD decrease in Aβ₁₋₃₈, 1.33; 95% CI, 1.01–1.89; HR for dementia per SD decrease in Aβ₁₋₄₂, 1.27; 95% CI, 1.02–1.58) (Table 3). No association was observed between Aβ₁₋₄₀ and Aβ₁₋₄₀/Aβ₁₋₄₂ ratio and dementia. When the analysis was performed separately for dementia subtypes, lower levels of Aβ₁₋₃₈ and Aβ₁₋₄₂ were associated with increased risk of AD (HR for AD per SD decrease in Aβ₁₋₃₈, 1.39; 95% CI, 1.00–2.16; HR for AD per SD decrease in Aβ₁₋₄₂, 1.35; 95% CI, 1.05–1.75). The HRs for AD and vascular dementia were closely similar but

Table 2 Association of amyloid-β levels with neurodegenerative markers

| Plasma Aβ levels (per SD decrease) | Total brain volume, mean difference (95% CI), ^a p value | Total gray matter volume, mean difference (95% CI), ^a p value | Total white matter volume, mean difference (95% CI), ^a p value | Total hippocampal volume, mean difference (95% CI), ^a p value |
|--|--|--|---|--|
| Aβ ₁₋₃₈ | 0.03 (-0.03; 0.08), $p = 0.362$ | 0.02 (-0.10; 0.15), $p = 0.715$ | 0.01 (-0.09; 0.12), $p = 0.819$ | 0.04 (-0.11; 0.19), $p = 0.582$ |
| Aβ ₁₋₄₀ | -0.04 (-0.09; 0.02), $p = 0.171$ | -0.08 (-0.21; 0.06), $p = 0.248$ | 0.01 (-0.11; 0.12), $p = 0.912$ | -0.01 (-0.17; 0.14), $p = 0.869$ |
| Aβ ₁₋₄₂ | 0.00 (-0.03; 0.04), $p = 0.812$ | -0.06 (-0.14; 0.03), $p = 0.174$ | 0.04 (-0.03; 0.12), $p = 0.251$ | -0.13 (-0.23; -0.04), $p = 0.007$ |
| Aβ ₁₋₄₀ /Aβ ₁₋₄₂ ratio | 0.03 (-0.02; 0.09) $p = 0.231$ | 0.06 (-0.07; 0.19), $p = 0.380$ | -0.00 (-0.11; 0.11) $p = 0.998$ | -0.10 (-0.25; 0.05), $p = 0.194$ |

Abbreviations: Aβ amyloid-β, SD standard deviation, CI confidence interval

^a Adjusted for age, sex, mean arterial blood pressure, total cholesterol, diabetes, apolipoprotein ε4 carrier status, smoking, intracranial volume and other Aβ levels

Table 3 Association of amyloid- β levels with incident dementia and its subtypes

| Plasma A β levels (per SD decrease) | Incident dementia (n = 79) HR (95% CI) ^a | Alzheimer's dementia (n = 64) HR (95% CI) ^a | Vascular dementia (n = 15) HR (95% CI) ^a |
|--|--|---|--|
| A β ₁₋₃₈ | 1.33 (1.01–1.89) ^b | 1.39 (1.00–2.16) ^b | 1.20 (0.49–2.96) |
| A β ₁₋₄₀ | 0.99 (0.69–1.43) | 0.95 (0.60–1.49) | 0.74 (0.31–1.79) |
| A β ₁₋₄₂ | 1.27 (1.02–1.58) ^b | 1.35 (1.05–1.75) ^b | 1.05 (0.61–1.79) |
| A β ₁₋₄₀ /A β ₁₋₄₂ ratio | 0.92 (0.64–1.32) | 0.97 (0.64–1.49) | 1.30 (0.54–3.11) |

Abbreviations: A β amyloid- β , SD standard deviation, HR hazard ratio, CI confidence interval

^aAdjusted for age, sex, education, mean arterial blood pressure, total cholesterol, diabetes, apolipoprotein ϵ 4 carrier status, smoking and other A β levels

^bSignificant at $p < 0.05$

insignificant in cases of vascular dementia (Table 3). The association between lower levels of A β and risk of dementia among APOE- ϵ 4 carriers and noncarriers appeared similar, such that lower levels of A β ₁₋₃₈, A β ₁₋₄₀, and A β ₁₋₄₂ were associated with increased risk of dementia in both strata. A stronger association was observed for A β ₁₋₃₈ with increased risk of dementia among APOE- ϵ 4 carriers (HR for dementia per SD decrease in A β ₁₋₃₈, 1.58; 95% CI, 1.01–2.89), whereas for A β ₁₋₄₂, this association was observed only among APOE- ϵ 4 noncarriers (HR for dementia per SD decrease in A β ₁₋₄₂, 1.47; 95% CI, 1.09–1.99) (Table 4).

Discussion

In this study, we showed that lower levels of plasma A β were not associated with preclinical markers of neurodegeneration (i.e., total gray matter and white matter volumes), except for A β ₁₋₄₂, which was associated with smaller hippocampal volume in elderly individuals. Individuals with lower levels of A β ₁₋₃₈ and A β ₁₋₄₂ had an independent increased risk of dementia, specifically AD. These findings suggest that A β ₁₋₃₈ and A β ₁₋₄₂ may be involved in different pathways leading to dementia.

Previous research has shown that higher plasma A β levels (1–40 and 1–42 peptides) at baseline were associated with cognitive dysfunction and faster cognitive decline, regardless of dementia status at follow-up, supporting the notion that plasma A β may induce a variety of brain pathologies (including cortical atrophy) earlier in life [27]. A recent study consisting of 100 participants also reported that higher plasma A β ₁₋₄₂ levels were associated with

thinner temporal cortex in cognitively normal elderly persons [13]. However, apart from the latter study, no studies have yet combined the structural brain changes with plasma A β levels to assess the differences in cognitively normal elderly individuals. Moreover, the effects of novel A β ₁₋₃₈ in addition to A β ₁₋₄₀ and 1–42 on cerebral atrophy and whether it is an important fluid biomarker for neurodegeneration and dementia has not been explored previously. Contrary to what we expected, our findings showed that lower plasma A β levels (including A β ₁₋₃₈) were not related to brain tissue volumes, except for hippocampal volume, where a significant association was observed between lower A β ₁₋₄₂ levels and smaller hippocampal volume in the elderly. The possible reason for the lack of association between A β ₁₋₃₈ and brain atrophy could be that plasma A β ₁₋₃₈ levels may reflect vascular disease in the brain rather than neurodegeneration. On one hand, this is supported by our previous study in which we showed that A β ₁₋₃₈ reflected microvascular damage in the brain and possibly induced adverse changes by inflammation, imbalance of oxygen free radicals, and apoptosis [22]. On the other hand, the association between A β ₁₋₄₂ and the hippocampus in the present study supports the notion that as A β ₁₋₄₂ starts to deposit in the brain after the age of 60 years, lower plasma A β levels that follow this deposition relate to hippocampal atrophy. A recent study has shown that accumulation of A β and tau pathologies in the brain were related to a decrease in hippocampal volume, including its critical subcompartments (i.e., CA1 and subiculum) in the earliest stages of AD prodromes [28]. Moreover, another study with healthy control individuals

Table 4 Association of amyloid- β levels with incident dementia in carriers and non-carriers of apolipoprotein ϵ 4 allele

| Plasma A β levels (per SD increase) | Incident dementia (n = 79) HR (95% CI) ^a | |
|--|--|--|
| | APOE- ϵ 4 carriers (n = 35) | APOE- ϵ 4 non-carriers (n = 44) |
| A β ₁₋₃₈ | 1.58 (1.01–2.89) ^b | 1.10 (0.66–1.83) |
| A β ₁₋₄₀ | 1.03 (0.58–1.83) | 1.04 (0.64–1.71) |
| A β ₁₋₄₂ | 1.07 (0.76–1.52) | 1.47 (1.09–1.99) ^b |
| A β ₁₋₄₀ /A β ₁₋₄₂ ratio | 0.72 (0.41–1.26) | 0.75 (0.45–1.16) |

Abbreviations: A β amyloid- β , SD standard deviation, HR hazard ratio, CI confidence interval, APOE Apolipoprotein

^a Adjusted for age, sex, education, mean arterial blood pressure, total cholesterol, diabetes, smoking and other A β levels

^bSignificant at $p < 0.05$

and persons with subjective cognitive complaints reported greater cortical thickness at intermediate levels of A β pathology (measured using CSF) [29]. By contrast, our data suggest that lower plasma A β levels (which are thought to indirectly reflect accumulation of amyloid in brain) do not necessarily correlate with universal volumetric decline in all structures (gray matter and white matter volumes), which is supported by some recent data [30].

Several lines of evidence suggest that A β levels in the CSF and plasma are in dynamic equilibrium with each other and that increased A β production in the brain gives rise to raised levels in the plasma [31]. It is further reported that as A β starts to deposit in the brain in the form of plaques, this in turn leads to lower plasma A β levels, which has been related to a higher risk of dementia [32]. Previous cross-sectional studies examining the association between A β levels and dementia have been mainly inconsistent [33–36]. It has been further reported that because plasma A β levels tend to change over the course of the dementia process, longitudinal studies are more useful in assessing the link between A β levels and risk of dementia in asymptomatic individuals [7]. Thus far, limited data have shown that plasma A β_{1-42} levels significantly decline in concentration in persons with newly diagnosed AD compared with individuals with prevalent AD and control subjects [12]. It is also suggested that plasma A β_{1-42} levels decline at an average rate of 12% per year among individuals with mild cognitive impairment [6]. Our results add to the previous reports by showing that lower levels of baseline plasma A β_{1-38} , in addition to A β_{1-40} and A β_{1-42} , were associated with a reduced risk of dementia. The association of A β_{1-38} and A β_{1-42} with dementia persisted when all the isoforms were added together in the model, indicating an independent link with increased dementia risk. These observations further support the findings that A β_{1-38} and A β_{1-42} may be generated independently by γ -secretase and that the production of these peptides is not coordinately regulated [16]. This was further confirmed in *in vitro* experiments involving γ -secretase modulators (sulindac sulfide), where A β_{1-38} levels were increased upon treatment with sulindac sulfide with no concurrent effect on A β_{1-42} levels, thus arguing against a precursor–product relationship [37, 38]. Though no association was observed between A β_{1-38} levels and neurodegenerative markers in this study, a link was still observed between lower levels of A β_{1-38} and increased risk of dementia. As mentioned before, the A β_{1-38} isoform is a marker of vascular pathology, and a higher level may indicate the activation of different inflammatory cascades (cytokines, cluster of differentiation 40 ligand, and tumor necrosis factor α) [22], which may increase the vulnerability to dementia, but such a relationship requires further clarification. Moreover, the relationship of plasma A β levels and brain amyloid deposition is suggested to be further

complicated by dynamics of the blood-brain barrier and other possible sources of A β materials outside the central nervous system, including platelets and skeletal muscle cells [32].

With respect to the subtypes of dementia, similar associations were observed between plasma A β_{1-38} and A β_{1-42} levels with AD and vascular dementia. Though the significant results were observed only in cases of AD, this finding might be related to the fact that most of the individuals diagnosed with incident dementia had AD (72%). Also, in an elderly population, mixed pathology is commonly observed, which might explain the similar estimates for AD and vascular dementia. Moreover, the association of lower levels of A β_{1-38} with increased risk of dementia was more significant in carriers of the APOE- ϵ 4 allele than in noncarriers. Though an opposite link was observed between A β_{1-42} and increased dementia risk in APOE- ϵ 4 noncarriers, the similarity in the direction of effect estimates further suggests a role of mixed pathology in the development of dementia, regardless of APOE carrier status.

Strengths of our study include its prospective design, the population-based setting, volumetric quantification of the brain tissues, and virtually complete follow-up for dementia in the older cohort. There are some potential limitations of our study. First, because plasma samples were collected after MRI acquisition with a 3-year time window, the effect estimates calculated in this study may represent over- or underestimations. Second, plasma A β levels provide an indirect measure of brain-specific A β pathology, and an *in vivo* analysis of brain-specific A β burden (such as with amyloid positron emission tomography) could allow more accurate measures of A β burden. Finally, we lacked repeated measurement of plasma A β concentrations, which limits its ability to better estimate the trajectory of plasma levels over time in relation to risk of dementia. However, this could be a subject for further studies.

Conclusions

This study provides evidence that lower plasma A β levels, specifically A β_{1-38} and A β_{1-42} , are associated with increased risk of dementia, specifically AD. Future studies should examine whether inclusion of novel plasma A β_{1-38} levels as an additional biomarker can provide further information on risk of developing dementia and AD dementia.

Abbreviations

AD: Alzheimer's disease; APOE: Apolipoprotein E; A β : Amyloid- β ; CSF: Cerebrospinal fluid; EDTA: Ethylenediaminetetraacetic acid; GMS: Geriatric Mental State Schedule; HASTE: Half-Fourier-acquired single-shot turbo spin echo; MMSE: Mini Mental State Examination; MRI: Magnetic resonance imaging; QC: Quality control

Acknowledgements

We thank Andrea Vermeij and Jeannette Vergeer for plasma analysis in this study.

Funding

This study is part of the Cerebral Amyloid Angiopathy Vascular Imaging and fluid markers of Amyloid deposition (CAVIA) project (no. 733050202), which has been made possible by ZonMW. The CAVIA project is part of 'Memorabel', the research and innovation programme for dementia, as part of the Dutch national 'Deltaplan for Dementia' (zonmw.nl/dementia research). The CAVIA project is a consortium of Radboudumc, Leiden University Medical Center (LUMC), Erasmus Medical Center (EMC), Vrije Universiteit Medical Center (VUMc), ADX Neurosciences, Philips Healthcare, Stony Brook University and Massachusetts General Hospital. The sponsor had no role in study design; the collection, analysis and interpretation of data; or in the writing of the manuscript.

Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Authors' contributions

SH participated in data acquisition and performed statistical analysis as well as drafting and revision of the manuscript. FJW, MMV, HV, MKI and ES participated in data acquisition and revised the manuscript for intellectual content. MAI and MWV were responsible for the study concept and design, obtaining funding, and drafting and revising the manuscript. All authors read and approved the final manuscript.

Ethics approval and consent to participate

The Rotterdam Study was approved by the medical ethics committee according to the Population Study Act Rotterdam Study, as well as by the Ministry of Health, Welfare and Sports of the Netherlands (PG/OGZ 3098756). Written informed consent was obtained from all participants prior to study recruitment.

Consent for publication

Not applicable

Competing interests

SH, FJW, MMV, HV, MKI, MAI and MWV report no competing interests. ES is a shareholder of ADx NeuroSciences NV, Gent, Belgium, which provided the kits for plasma assessment of amyloid- β levels.

Publisher's Note

Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Author details

¹Department of Radiology and Nuclear Medicine, Erasmus University Medical Center, Office no. 2505, Wytemaweg 80, 3015, CN, Rotterdam, The Netherlands. ²Department of Epidemiology, Erasmus University Medical Center, Rotterdam, The Netherlands. ³Department of Neurology and Laboratory Medicine, Donders Institute for Brain, Cognition and Behavior, Radboud Alzheimer Centre, Radboud University Medical Center, Nijmegen, The Netherlands. ⁴ADx NeuroSciences, Ghent, Belgium. ⁵Department of Neurology, Erasmus University Medical Center, Rotterdam, The Netherlands.

Received: 14 March 2018 Accepted: 30 May 2018

Published online: 30 June 2018

References

- Selkoe DJ. Alzheimer's disease: genes, proteins, and therapy. *Physiol Rev*. 2001;81:741–6.
- Jack CR Jr, Knopman DS, Jagust WJ, Shaw LM, Aisen PS, Weiner MW, et al. Hypothetical model of dynamic biomarkers of the Alzheimer's pathological cascade. *Lancet Neurol*. 2010;9:119–28.
- Hardy J, Selkoe DJ. The amyloid hypothesis of Alzheimer's disease: progress and problems on the road to therapeutics. *Science*. 2002;297:353–6.
- Shah NS, Vidal JS, Masaki K, Petrovitch H, Ross GW, Tilley C, et al. Midlife blood pressure, plasma β -amyloid, and the risk for Alzheimer disease: the Honolulu Asia Aging Study. *Hypertension*. 2012;59:780–6.
- Graff-Radford NR, Crook JE, Lucas J, Boeve BF, Knopman DS, Ivnik RJ, et al. Association of low plasma $A\beta_{42}/A\beta_{40}$ ratios with increased imminent risk for mild cognitive impairment and Alzheimer disease. *Arch Neurol*. 2007;64:354–62.
- Mayeux R, Honig LS, Tang MX, Manly J, Stern Y, Schupf N, et al. Plasma $A\beta_{40}$ and $A\beta_{42}$ and Alzheimer's disease: relation to age, mortality, and risk. *Neurology*. 2003;61:1185–90.
- van Oijen M, Hofman A, Soares HD, Koudstaal PJ, Breteler MM. Plasma $A\beta_{1-40}$ and $A\beta_{1-42}$ and the risk of dementia: a prospective case-cohort study. *Lancet Neurol*. 2006;5:655–60.
- Lopez OL, Kuller LH, Mehta PD, Becker JT, Gach HM, Sweet RA, et al. Plasma amyloid levels and the risk of AD in normal subjects in the Cardiovascular Health Study. *Neurology*. 2008;70:1664–71.
- Schupf N, Tang MX, Fukuyama H, Manly J, Andrews H, Mehta P, et al. Peripheral $A\beta$ subspecies as risk biomarkers of Alzheimer's disease. *Proc Natl Acad Sci U S A*. 2008;105:14052–7.
- Lambert JC, Schraen-Maschke S, Richard F, Fievet N, Rouaud O, Berr C, et al. Association of plasma amyloid- β with risk of dementia: the prospective Three-City Study. *Neurology*. 2009;73:847–53.
- Chouraki V, Beiser A, Yonkin L, Preis SR, Weinstein G, Hansson O, et al. Plasma amyloid- β and risk of Alzheimer's disease in the Framingham Heart Study. *Alzheimers Dement*. 2015;11:249–57.
- Pomara N, Willoughby LM, Sidtis JJ, Mehta PD. Selective reductions in plasma $A\beta$ 1–42 in healthy elderly subjects during longitudinal follow-up: a preliminary report. *Am J Geriatr Psychiatry*. 2005;13:914–7.
- Llado-Saz S, Aizenstein M, Cantero JL. Increased levels of plasma amyloid- β are related to cortical thinning and cognitive decline in cognitively normal elderly subjects. *Neurobiol Aging*. 2015;36:2791–7.
- Zahs KR, Ashe KH. β -Amyloid oligomers in aging and Alzheimer's disease. *Front Aging Neurosci*. 2013;5:28.
- Selkoe DJ. Soluble oligomers of the amyloid- β -protein impair synaptic plasticity and behavior. *Behav Brain Res*. 2008;192:106–13.
- Reinert J, Martens H, Huettenrauch M, Kolbow T, Lannfelt L, Ingelsson M, et al. $A\beta_{38}$ in the brains of patients with sporadic and familial Alzheimer's disease and transgenic mouse models. *J Alzheimers Dis*. 2014;39:871–81.
- Welge V, Fiege O, Lewczuk P, Mollenhauer B, Esselmann H, Klafki HW, et al. Combined CSF tau, p-tau181 and amyloid- β 38/40/42 for diagnosing Alzheimer's disease. *J Neural Transm*. 2009;116:203–12.
- Wiltfang J, Esselmann H, Bibl M, Smirnov A, Otto M, Paul S, et al. Highly conserved and disease-specific patterns of carboxyterminally truncated $A\beta$ peptides 1–37/38/39 in addition to 1–40/42 in Alzheimer's disease and in patients with chronic neuroinflammation. *J Neurochem*. 2002;81:481–96.
- Ikram MA, Vrooman HA, Vernooij MW, van der Lijn F, Hofman A, van der Lugt A, et al. Brain tissue volumes in the general elderly population. The Rotterdam Scan Study. *Neurobiol Aging*. 2008;29:882–90.
- de Bruijn RF, Bos MJ, Portegies ML, Hofman A, Franco OH, Koudstaal PJ, et al. The potential for prevention of dementia across two decades: the prospective, population-based Rotterdam Study. *BMC Med*. 2015;13:132.
- Lerche S, Brockmann K, Pilotto A, Wurster I, Sunkel U, Hoberg MA, et al. Prospective longitudinal course of cognition in older subjects with mild parkinsonian signs. *Alzheimers Res Ther*. 2016;8:42.
- Hilal S, Akoudad S, van Duijn CM, Niessen WJ, Verbeek MM, Vanderstichele H, et al. Plasma amyloid- β levels, cerebral small vessel disease, and cognition: the Rotterdam Study. *J Alzheimers Dis*. 2017;60:977–87.
- Ikram MA, van der Lugt A, Niessen WJ, Koudstaal PJ, Krestin GP, Hofman A, et al. The Rotterdam Scan Study: design update 2016 and main findings. *Eur J Epidemiol*. 2015;30:1299–315.
- den Heijer T, Geerlings MI, Hoebeek FE, Hofman A, Koudstaal PJ, Breteler MM. Use of hippocampal and amygdalar volumes on magnetic resonance imaging to predict dementia in cognitively intact elderly people. *Arch Gen Psychiatry*. 2006;63:57–62.
- McKhann GM, Knopman DS, Chertkow H, Hyman BT, Jack CR Jr, Kawas CH, 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;7:263–9.
- Roman GC, Tatemichi TK, Erkinjuntti T, Cummings JL, Masdeu JC, Garcia JH, et al. Vascular dementia: diagnostic criteria for research studies: report of the NINDS-AIREN international workshop. *Neurology*. 1993;43:250–60.
- Cosentino SA, Stern Y, Sokolov E, Scarmeas N, Manly JJ, Tang MX, et al. Plasma β -amyloid and cognitive decline. *Arch Neurol*. 2010;67:1485–90.
- Tardif CL, Devenyi GA, Amaral RSC, Pelleieux S, Poirier J, Rosa-Neto P, et al. Regionally specific changes in the hippocampal circuitry accompany progression of cerebrospinal fluid biomarkers in preclinical Alzheimer's disease. *Hum Brain Mapp*. 2018;39:971–84.

29. Fortea J, Sala-Llonch R, Bartres-Faz D, Llado A, Sole-Padullés C, Bosch B, et al. Cognitively preserved subjects with transitional cerebrospinal fluid β -amyloid 1–42 values have thicker cortex in Alzheimer's disease vulnerable areas. *Biol Psychiatry*. 2011;70:183–90.
30. Jack CR Jr, Knopman DS, Jagust WJ, Petersen RC, Weiner MW, Aisen PS, et al. Tracking pathophysiological processes in Alzheimer's disease: an updated hypothetical model of dynamic biomarkers. *Lancet Neurol*. 2013;12:207–16.
31. Ghersi-Egea JF, Gorevic PD, Ghiso J, Frangione B, Patlak CS, Fenstermacher JD. Fate of cerebrospinal fluid-borne amyloid- β -peptide: rapid clearance into blood and appreciable accumulation by cerebral arteries. *J Neurochem*. 1996;67:880–3.
32. Kawarabayashi T, Younkin LH, Saido TC, Shoji M, Ashe KH, Younkin SG. Age-dependent changes in brain, CSF, and plasma amyloid β protein in the Tg2576 transgenic mouse model of Alzheimer's disease. *J Neurosci*. 2001;21:372–81.
33. Mehta PD, Pirttilä T, Mehta SP, Sersen EA, Aisen PS, Wisniewski HM. Plasma and cerebrospinal fluid levels of amyloid- β proteins 1–40 and 1–42 in Alzheimer disease. *Arch Neurol*. 2000;57:100–5.
34. Iwatsubo T. Amyloid- β protein in plasma as a diagnostic marker for Alzheimer's disease. *Neurobiol Aging*. 1998;19:161–3.
35. Blennow K, Mattsson N, Schöll M, Hansson O, Zetterberg H. Amyloid biomarkers in Alzheimer's disease. *Trends Pharmacol Sci*. 2015;36:297–309.
36. Blennow K, Hampel H, Weiner M, Zetterberg H. Cerebrospinal fluid and plasma biomarkers in Alzheimer disease. *Nat Rev Neurol*. 2010;6:131–44.
37. Page RM, Baumann K, Tomioka M, Perez-Revuelta BI, Fukumori A, Jacobsen H, et al. Generation of $A\beta_{38}$ and $A\beta_{42}$ is independently and differentially affected by familial Alzheimer disease-associated presenilin mutations and γ -secretase modulation. *J Biol Chem*. 2008;283:677–83.
38. Czirr E, Cottrell BA, Leuchtenberger S, Kukar T, Ladd TB, Esselmann H, et al. Independent generation of $A\beta_{42}$ and $A\beta_{38}$ peptide species by γ -secretase. *J Biol Chem*. 2008;283:17049–54.

Ready to submit your research? Choose BMC and benefit from:

- fast, convenient online submission
- thorough peer review by experienced researchers in your field
- rapid publication on acceptance
- support for research data, including large and complex data types
- gold Open Access which fosters wider collaboration and increased citations
- maximum visibility for your research: over 100M website views per year

At BMC, research is always in progress.

Learn more biomedcentral.com/submissions

