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### Article

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## ARTICLE OPEN



# Plasma Amyloid- $\beta$ dynamics in late-life major depression: a longitudinal study

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Depressed individuals are twice as likely to develop Alzheimer's disease (AD) as compared to controls. Brain amyloid- $\beta$  (A $\beta$ ) deposition is believed to have a major role in AD pathogenesis but studies also suggest associations of A $\beta$  dynamics and depression. The aim of this study was to test if plasma A $\beta$  levels are longitudinally associated to late-life depression. We measured plasma levels of amyloid- $\beta_{1-40}$  (A $\beta$ 40) and amyloid- $\beta_{1-42}$  (A $\beta$ 42) peptides longitudinally for three consecutive years in 48 cognitively intact elderly subjects with late-life major depressive disorder (LLMD) and 45 age-matched cognitively healthy controls. We found that the A $\beta$ 42/A $\beta$ 40 plasma ratio was significantly and steadily lower in depressed subjects compared to controls ( $p < 0.001$ ). At screening, A $\beta$ 42/A $\beta$ 40 plasma did not correlate with depression severity (as measured with Hamilton Depression Scale) or cognitive performance (as measured with Mini-Mental State Examination) but was associated to depression severity at 3 years after adjustment for age, education, cognitive performance, and antidepressants use. This study showed that reduced plasma A $\beta$ 42/A $\beta$ 40 ratio is consistently associated with LLMD diagnosis and that increased severity of depression at baseline predicted low A $\beta$ 42/A $\beta$ 40 ratio at 3 years. Future studies are needed to confirm these findings and examine if the consistently lower plasma A $\beta$ 42/A $\beta$ 40 ratio in LLMD reflects increased brain amyloid deposition, as observed in AD subjects, and an increased risk for progressive cognitive decline and AD.

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## INTRODUCTION

Major depressive disorder (MDD) is a well-established risk factor for Alzheimer's disease (AD) [1, 2]. Depression is also a part of the clinical syndrome in the early stages of AD and can be a prodromal manifestation [3, 4]. A meta-analysis of studies of depression and dementia [5] showed that depressed individuals are nearly twice as likely to develop dementia as compared to controls [6]. A 7-year longitudinal study [7] showed that for each point increase in the 10-item Center for Epidemiologic Studies Depression Scale (CES-D), risk of developing AD increased by an average of 19%. Lifetime depression has been associated with increases in AD-related brain pathology, including elevated density of neuritic plaques and neurofibrillary tangles [8, 9]. However, not all studies have found a link between depression and AD [10–12], and depression may be part of the dementia prodrome thus highlighting the complexity of the relationship between these two conditions.

In a bid to elucidate the relationship between depression and AD, several reports, including from our group [13], have suggested that disturbances in amyloid- $\beta$  (A $\beta$ ) production or aggregation

status may play a role in the disease onset, alone or in combination with other factors (e.g., inflammation). A $\beta$  deposition in the brain is a hallmark of AD, but abnormal A $\beta$  levels have also been observed in individuals with MDD, both in the cerebrospinal fluid [13], and in the brain [14, 15]. In AD, numerous studies have shown a reduction in both the plasma and CSF A $\beta$ 42/A $\beta$ 40 ratio which has been associated with increases in amyloid PET uptake [16, 17]. Studies of PET amyloid scans in MDD have provided conflicting results with some reports of increased amyloid build up [14, 18–21], or no change [22, 23], and most surprisingly even reductions in amyloid compared to controls as well [24]. The basis for these conflicting PET findings may be due to methodological differences across studies. These include heterogeneity of depression with respect to age of first depressive episode, degree of response to treatment, magnitude of HPA axis and microglia activation accompanying each depressive episode as well as in the frequency of the e4 allele which have all been implicated in A $\beta$  dynamics.

A reduction in plasma A $\beta$ 42/A $\beta$ 40 ratio has also been observed in the very early phases of AD and was found to be associated

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with future decline in cognition [25, 26]. Cross-sectional studies in depressed elderly adults have also reported decreased plasma A $\beta$ 42/A $\beta$ 40 ratios compared to age-matched controls [27]. However, to our knowledge, no longitudinal studies have evaluated plasma A $\beta$  levels in subjects with late-life major depression (LLMD).

In this study, we tested the hypothesis that plasma A $\beta$  levels differ between elderly individuals with LLMD and age-matched controls and that disease severity at baseline predicts long-term plasma A $\beta$  status. We explored this hypothesis in a group of 48 cognitively intact depressed participants and 45 controls who were followed up longitudinally for 3 years. In addition, we examined whether plasma A $\beta$  levels were accompanied by changes in cognitive performance and severity of depressive symptoms over time. Since brain derived neurotrophic factor (BDNF), a neurotrophin that plays an important role in cognition and depression [28, 29] is also known to regulate A $\beta$  production and deposition [30, 31], we also examined the correlations between plasma BDNF and A $\beta$  at Baseline.

## METHODS

### Subjects and study design

The study was approved by the Nathan Kline Institute for Psychiatric Research and the New York University School of Medicine Institutional Review Boards. Participants were recruited via advertisements in local newspapers and flyers, or via the Memory Education and Research Initiative. Participants were prescreened over the phone during which a description of the study and participation requirements were described in detail before coming to the clinic for screening. Participants provided written informed consent prior to the study at the screening visit and were compensated up to \$450 for their participation. One-hundred and thirty-three total participants were enrolled in the study. Thirty subjects did not complete the study: 3 died, 15 were lost to follow-up or did not return calls to schedule, 4 moved to a different location, and 8 withdrew from participation. Ten subjects completed all study visits but had unusable plasma samples and were excluded from the primary analysis. Of the 133 subjects entering the study, 93 completed all four study visits (baseline and three follow-up visits), had available plasma samples and were included in the primary analysis. Of these 93 subjects, 48 had a diagnosis at baseline of LLMD, and 45 were aged-matched controls.

In order to be enrolled in the study participants had to be 60 years, have a normal MRI scan (Fazeka score < 3), absence of unstable medical illness, and clinically normal lab values. To be enrolled into the LLMD group, participants had to meet the criteria for Major Depressive Disorder (MDD) based on the Structured Clinical Interview for the Diagnostic and Statistical Manual of Mental Disorders (DSM)-IV (SCID) conducted by a board-certified psychiatrist. Antidepressant use was allowed in the LLMD Group. To be enrolled in the control group, participants could not have a history of MDD or any other major psychiatric disorder. Additionally, all participants, regardless of LLMD or control group, had to be cognitively normal at screening. Normal cognition was defined as: Global Clinical Dementia Rating Scale score of 0 and a Mini-Mental State Exam score > 25.

### Study procedure

During the screening phase of the study, participants underwent testing at the Nathan Kline Institute or at the New York University Medical Center, over three successive visits, one week apart. During the first Screening Visit, participants provided informed consent, blood for APOE genotype was collected, and the Hamilton Depression Rating Scale (HAM-D) was administered. During the second Screening Visit, a general medical history was taken, the Mini-Mental State Examination (MMSE), the Geriatric Depression Scale (GDS) and Hamilton Anxiety Scale were also administered. During the third Screening Visit, participants received an MRI scan of the head and blood sample for routine laboratory tests was collected. After completing the screening phase and qualifying for the study, participants completed the Baseline Visit. During the Baseline Visit, participants underwent a full neuropsychological assessment, a psychiatric interview, a physical and neurological exam, and repeated mood scales (HAM-D, GDS, etc.). In general, the time between Screening and Baseline was approximately two weeks, but in some cases, there was a longer interval (i.e., up to a month). The three follow-up visits (Follow-up 1–3 Visits) took place in the same locations

and were spaced approximately one-year apart. During the follow-up visits, participants completed the same comprehensive neuropsychological evaluation that they completed at the Baseline Visit. A clinical evaluation to assess for change in mood, a physical and neurological exam, the mood rating scales and routine laboratory tests were completed. Blood draws for A $\beta$ 40 and A $\beta$ 42 measurements were taken at Baseline and at each of the three follow-up visits.

### Measurements of depression status and cognitive performance

Depression status and severity were measured with the 21-item Hamilton Depression Rating Scale (HAM-D) which has a range of 0–55. Two measures of depression severity are captured from this measure, the HAM-D-17, which sums the total of the first 17 items and the HAM-D-21, which sums all items and has additional questions about depressive subtypes. The HAM-D-17 is used to determine depression severity with a cut-off of 17 for moderate depression [32]. Global cognitive performance was assessed with the Mini-Mental State Examination (MMSE) which has a range of 0–30 and a score of 25 as education-adjusted cut-off for normality. Verbal learning and memory were measured, with the Buschke Selective Reminding Test Total score and 20-min Delayed Recall Score, which were selected from the larger neuropsychological test battery. Tests were administered at Baseline and at the 3 yearly follow-ups.

### Plasma A $\beta$ measurement

Blood was collected in standard 10 ml EDTA tubes (BD Vacutainer). Immediately after blood draw, tubes were inverted gently 10–12 times for mixing and centrifuged at 3,000 rpm for 15 min at room temperature. Plasma aliquots were transferred to polypropylene low binding tubes and immediately stored at  $-80^{\circ}\text{C}$ . Plasma A $\beta$ 42 and A $\beta$ 40 concentrations were determined using biochemical the INNO-BIA plasma A $\beta$  forms assay, format B (Fujirebio, Ghent, Belgium), a multiplex microsphere-based Luminex xMAP technique for simultaneous detection of A $\beta$  X-42 and A $\beta$ X-40 as previously described [33]. The plasma A $\beta$ 42/A $\beta$ 40 ratio was calculated by dividing the plasma A $\beta$ 42 by A $\beta$ 40 concentrations at each corresponding visit.

### Plasma BDNF measurement

BDNF protein levels in plasma samples were measured using an enzyme-linked immunosorbent assay (ELISA) method (BDNF Emax Immunoassay System, Promega, USA), according to the manufacturer's instructions as described previously (see Pillai, Bruno [29]). Briefly, 96-well flat bottom immunoplates were incubated with an anti-BDNF monoclonal antibody at  $4^{\circ}\text{C}$  overnight followed by incubation with standards and samples for 2 h at room temperature. Following washing with TBST wash buffer, plates were incubated for 2 h with Anti-Human BDNF polyclonal antibody. Subsequently, Anti-immunoglobulin Y-horse-radish peroxidase conjugate was added followed by the addition of TMB One solution to develop the color. The reaction was stopped by the addition of HCl 1 N and the absorbance was read at 450 nm on a microplate reader. BDNF concentrations were determined using the BDNF standard curve values (ranging from 7.8 to 500 pg/ml purified BDNF). All the samples were analyzed in duplicate in one session by an investigator blind to experimental set up.

### Statistical Analysis

Comparisons of variables measured longitudinally [Hamilton Depression Rating Scale Score (HAM-D), cognitive tests (MMSE, Buschke Selective Reminding Test Total and Delay Recall), A $\beta$ 42 and A $\beta$ 40 plasma levels and their ratio] between depressed and control subjects were performed with repeated-measures analysis of variance (RMANOVA). The potential effects of a number of variables at screening (MMSE score, age, education, diagnosis, HAM-D, antidepressant use, APOE genotype) on A $\beta$  plasma levels at Year 3 were evaluated with stepwise linear regression and partial correlations. The main analyses were performed with all subjects completing the study ( $n = 93$ ). Power calculations were determined before and after the study with the G\*Power 3.1 software. For this sample size, the post-power analysis confirms that we had had a power  $(1-\beta) > 0.85$  in detecting significant ( $\alpha = 0.05$ ) differences equal or superior to 20% in plasma A $\beta$ 42, A $\beta$ 40 or A $\beta$ 42/A $\beta$ 40 between depressed and control groups. Sensitivity analyses were carried out excluding those subjects with abnormal MMSE score or MRI findings at Baseline ( $n = 6$ ) and produced results very similar to those of the main analyses, so the reported findings include all subjects completing the study. Statistical analysis was run in

**Table 1.** Baseline demographic and clinical characteristics (standard deviations in parentheses) of the subjects completing the study.

Baseline scores	LLMD (n = 48)		Controls (n = 45)		P-value
Age (years)	67.9 (5.8)		68.4 (6.3)		0.740
Education (years)	16.5 (2.6)		16.5 (2.6)		0.856
MMSE Score	29.6 (0.9)		29.6 (0.9)		0.723
Hamilton depression rating scale score (HAM-D)	18.6(9.3)		1.5 (2.8)		<0.001
Baseline frequencies	LLMD		Controls		P-value
	N	%	N	%	
Sex (Female)	24	50	24	53	0.748
Family history of Alzheimer's disease	5	10	10	22	0.122
Apolipoprotein ε genotype (presence of at least one allele)					
ε2	10	21	12	27	0.508
ε3	45	94	43	96	0.700
ε4	15	31	7	16	0.075
Number of subjects taking antidepressants at Baseline	29	60	0	0	0.004
Number of subjects taking SSRIs at Baseline	17	35	0	0	<0.001

Continuous variables were compared with Student *t*-test, while frequencies were compared with Chi Square test.

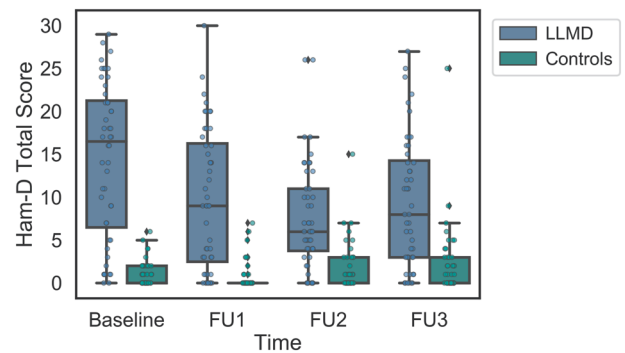
SPSS v24.0. Alpha of  $p < 0.05$  was used to indicate statistical significance. Figures were generated using Seaborn 0.10.0.

## RESULTS

Table 1 summarizes the demographic and clinical characteristics. Antidepressant use is described in Supplemental Table S1. A total of 48 elderly LLMD (mean age at Baseline = 67.9 years) and 45 control elderly subjects (mean age at Baseline = 68.4 years) were followed over a 3-year period. Both the LLMD and the control groups differed only on the HAM-D total score ( $p < 0.001$ ), with the LLMD group significantly more depressed than controls. Both groups were cognitively intact as measured by the MMSE at Baseline (LLMD: Mean score = 29.6; Controls: mean score = 29.6). We found no group differences in the proportion of individuals with factors known to influence Aβ dynamics, including in the number of ε4 or ε2 allele carriers and those with a family history of AD. Supplemental Table S2 shows demographic and clinical characteristics of the 40 subjects who were not included in the primary analysis. Their baseline characteristics were very similar to those of the 93 subjects included in the primary analysis. An analysis of cognition over time is found in the Supplemental Results. In summary, there were no groups differences in any of the cognitive indices at baseline and at the end of 3 years.

### Severity of depression

Figure 1 shows mean total HAM-D scores of the LLMD and control groups at Baseline and at the 3 follow-up visits. Two-way RMANOVA indicates that mean HAM-D scores of the LLMD group were significantly different than those of the control group at all time points ( $p < 0.001$ ). There was a significant interaction between the fixed factor time and diagnosis ( $p < 0.001$ ) indicating that mean HAM-D scores in depressed patients significantly decreased over time compared to Baseline. Although, the LLMD groups' depressive symptoms improved relative to Baseline, across the 3-year period the LLMD group remained significantly more depressed than the control group.

**Fig. 1** Boxplots of HAM-D Score. Boxplots with individual datapoints superimposed of total HAM-D score of the LLMD (blue) and control (green) groups and baseline and each yearly follow-up visits. \* $p < 0.001$  between LLMD and controls.**Table 2.** Means and standard deviations (SD) of plasma levels of Aβ40, Aβ42, and Aβ42/Aβ40 plasma ratio for cognitively intact participants with major depressive disorder (LLMD) and controls at Baseline and Follow-Up Visits 1, 2 and 3.

Plasma Aβ	Time	LLMD (n = 48)	Controls (n = 45)	t-test (p-value)
Aβ42 (pg/mL)	Baseline	17.9 (6.6)	17.9 (5.5)	0.942
	FU1	19.0 (6.8)	21.5 (5.6)	0.052
	FU2	19.4 (6.4)	21.2 (6.1)	0.170
	FU3	19.8 (6.8)	20.7 (5.4)	0.497
Aβ40 (pg/mL)	Baseline	143.3 (39.6)	122.6 (39.2)	0.013*
	FU1	158.8 (41.4)	148.3 (40.9)	0.222
	FU2	151.7 (31.7)	148.1 (42.4)	0.645
	FU3	155.7 (40.1)	145.3 (39.8)	0.213
Aβ42/Aβ40	Baseline	0.13 (0.1)	0.15 (0.5)	0.005*
	FU1	0.12 (0.4)	0.15 (0.5)	0.001*
	FU2	0.13 (0.4)	0.15 (0.5)	0.031*
	FU3	0.13 (0.2)	0.15 (0.4)	0.082

Univariate comparisons using independent samples *t*-test with the corresponding *p*-value for each timepoint are listed. \*Significant difference at  $p < 0.05$  level between LLMD and controls.

### Aβ plasma levels

Table 2 reports the means and standard deviations of plasma concentrations of Aβ42, Aβ40, and Aβ42/Aβ40 ratio across visits in the LLMD and controls groups.

Although small changes were observed in Aβ42 and Aβ40 over time (Table 3), the variations were similar in depressed and control groups and no significant variations of the plasma concentrations were observed for the two peptides over time. Figure 2 shows boxplots of plasma Aβ42, Aβ40 and Aβ42/Aβ40 ratio of the LLMD and controls groups at Baseline and at the follow-up visits. Results of two-way mixed RMANOVA did not reveal significant differences between LLMD and control groups in either Aβ42 or Aβ40 plasma levels. However, the mean Aβ42/Aβ40 plasma ratios of the LLMD group were significantly lower than those of the control group at all time points ( $p = 0.005$ ) (Fig. 2c). There was no significant interaction between time and diagnosis indicating that the temporal trends of the mean Aβ42/Aβ40 ratio of the two groups were parallel. We did not find significant differences in mean



**Table 3.** Repeated Measure ANOVA F-Test results for testing the interaction between time and groups (depressed and controls) for HAM-D, Cognitive Tests, A $\beta$ 40, A $\beta$ 42, and A $\beta$ 42/A $\beta$ 40 ratio.

Outcome variable	F test (within-subject)		P-value	Effect size ( $\eta^2$ )
	df **	F statistic		
HAM-D	2.64, 240.38	15.18	$p < 0.001^*$	0.14
MMSE	2.43, 221.26	0.53	0.627	0.006
Total recall	2.87, 261.11	3.63	0.015*	0.04
Delay recall	2.93, 266.71	2.47	0.064	0.03
A $\beta$ 40	2.85, 259.26	1.51	0.214	0.02
A $\beta$ 42	2.79, 254.01	1.69	0.174	0.02
A $\beta$ 42/A $\beta$ 40	2.63, 239.09	1.66	0.184	0.02
Outcome variable	F test (between-subject)		P-value	Effect size ( $\eta^2$ )
	df	F statistic		
A $\beta$ 42/A $\beta$ 40	1.91	8.205	0.005*	0.08

\*Significant interaction of time and diagnosis. \*\*degrees of freedom adjusted for violations of Mauchly's test using the Huynh-Feldt correction.

A $\beta$ 42/A $\beta$ 40 plasma ratio at Baseline ( $p = 0.93$ ) between LLMD patients taking antidepressants and those not treated (mean = 0.128 (SD = 0.046), and 0.129 (SD = 0.034), respectively).

#### Association of depressive symptoms and future plasma A $\beta$ levels at follow up

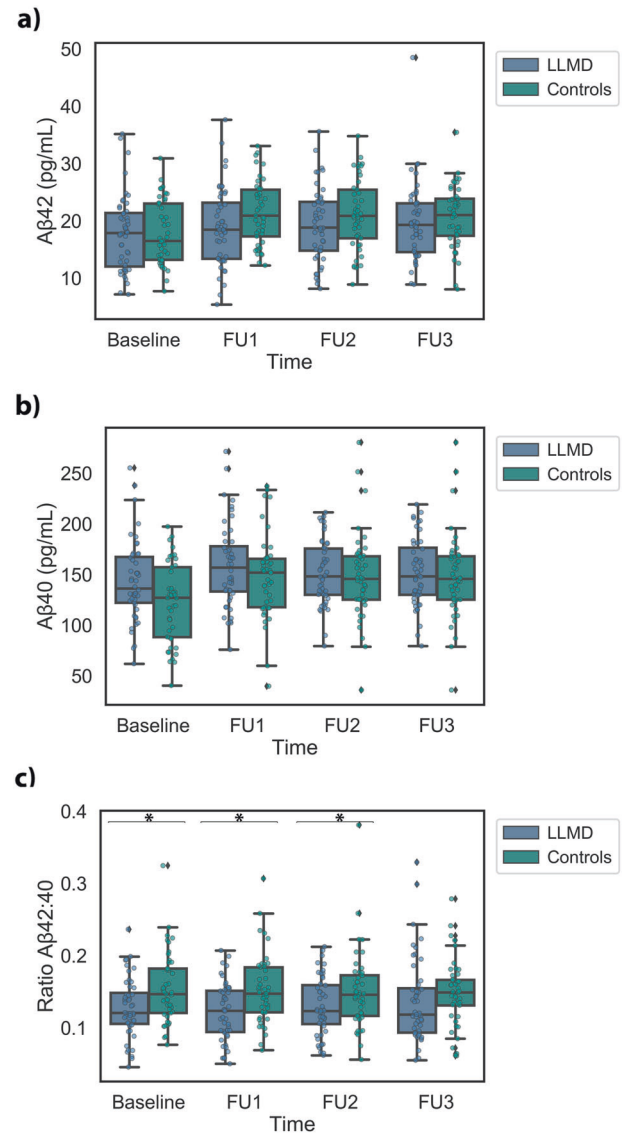
We used a stepwise linear regression to model the associations of depressive symptoms at screening and the A $\beta$ 42/A $\beta$ 40 plasma ratios at the different visits (Table 3). The predictors included MMSE score, age, education, HAM-D, APOE status and antidepressant use at screening. Since there was a significant reduction in the HAM-D total score from the screening to the baseline visit we tested both the Screening and the Baseline HAM-D score in predicting the A $\beta$ 42/A $\beta$ 40 plasma ratio. We found that HAM-D scores collected at screening significantly predicted A $\beta$ 42/A $\beta$ 40 ratio at year 1 ( $\beta = -0.33$ ,  $p = 0.001$ ) and year 3 ( $\beta = -0.24$ ,  $p = 0.018$ ). Similarly, the HAM-D score measured at baseline also significantly predicted A $\beta$ 42/A $\beta$ 40 ratio at year 1 ( $p = .003$ ) and year 3 ( $p = 0.040$ ). We also examined possible differences in subjects with early- (<60 years,  $n = 29$ ) vs late- ( $\geq 60$  years,  $n = 19$ ) onset of LLMD. Neither age of first depressive episode (mean  $\pm$  SD = 53.5  $\pm$  12.0 years) nor first onset at age 60 or older was significantly correlated with the plasma A $\beta$ 42/A $\beta$ 40 ratio, in LLMD, at any timepoint.

#### Relationship between plasma A $\beta$ and BDNF

The relationship between plasma A $\beta$  and BDNF was explored using Pearson correlations. There were no significant correlations between plasma A $\beta$ 42, A $\beta$ 40 or A $\beta$ 42/A $\beta$ 40 ratio and BDNF for the whole group. There were also no significant correlations for LLMD or controls, or for males and females, when run separately.

#### DISCUSSION

We found that the plasma A $\beta$ 42/A $\beta$ 40 ratio was consistently lower in depressed subjects compared to controls over a period of 3 years. This finding is consistent with previous cross-sectional studies indicating a reduced plasma A $\beta$ 42/A $\beta$ 40 ratio in older



**Fig. 2** Boxplots of Plasma A $\beta$  Variables. Boxplots with individual datapoints superimposed of plasma A $\beta$ 42 (panel a), A $\beta$ 40 (panel b) and A $\beta$ 42/A $\beta$ 40 ratio (panel c) of the LLMD and controls groups at Baseline and at the follow-up visits (FU). \* $p < 0.01$  between LLMD and controls.

adults with late-life depression [27]. Since no significant differences were observed in cognitive performance between depressed and control groups, or as a function of plasma A $\beta$  levels, it is unlikely that greater cognitive decline contributed to the lower plasma A $\beta$ 42/A $\beta$ 40 ratio in the LLMD group. Our study also identified an association of depressive symptoms at baseline and the plasma A $\beta$ 42/A $\beta$ 40 ratios in subsequent visits. This finding supports the hypothesis that depressive symptoms may drive plasma A $\beta$  dynamics, which should be explored in further studies.

The lower plasma A $\beta$ 42/A $\beta$ 40 ratio we observed in elderly depressed subjects was associated with higher plasma A $\beta$ 40 with no significant group difference in plasma A $\beta$ 42 levels. Thus, future studies should determine if this peptide alone contributes to the significantly lower A $\beta$ 42/A $\beta$ 40 ratio found in LLMD. It also is possible that the plasma A $\beta$ 42/A $\beta$ 40 ratio is a more sensitive biomarker of A $\beta$  dynamics since it represents the equilibrium of the two, rather than a single peptide. This conclusion agrees with the generally accepted notion that A $\beta$ 40 serves to control for inter-individual differences in total A $\beta$  production, as well as

possible pre-analytical confounders [34]. Although the precise origin of circulating blood A $\beta$  is not known [35], numerous studies in AD support that a reduction in the plasma A $\beta$ 42/A $\beta$ 40 ratio is associated with brain amyloidosis [36, 37]. Therefore, future studies, using amyloid PET, should validate whether the decreased plasma A $\beta$ 42/A $\beta$ 40 ratio which we observed in elderly depressives reflects increased brain amyloid burden or changes in soluble A $\beta$  levels, especially circulating A $\beta$ 40, which may be independent of brain amyloid load.

Previous studies have examined brain and CSF amyloid indices in elderly depressives; however, the number of studies is limited and the results conflicting. In a subgroup of the cohort described in the current report, mean CSF A $\beta$ 42 levels at baseline were significantly lower in depressed subjects compared to controls but then increased over time and were no longer significantly different from controls after 3 years [38]. The increase in CSF A $\beta$ 42 levels in depressed subjects coincided with improvement in depressive symptoms. These findings were consistent with state-dependent effects of depressive symptoms on CSF A $\beta$ 42 rather than with the prevailing opinion that this disorder might reflect prodromal AD. This interpretation is also supported by our current findings that no notable differences in cognitive ability were observed across groups, or as a function of A $\beta$  levels, although we cannot rule out that these differences might yet emerge later. Furthermore, the plasma A $\beta$ 42/A $\beta$ 40 ratio at 3 years was predicted by baseline severity of depressive symptoms, and not by cognitive indices.

Statistical analyses indicated that group differences in plasma A $\beta$ 42/A $\beta$ 40 ratio that we observed cannot be ascribed to differences in the frequency of the  $\epsilon$ 4 allele or to a family history of AD. In addition, individuals with a first episode of depression earlier in life (EOMD) showed a reduction in the plasma A $\beta$ 42/A $\beta$ 40 ratio that was comparable to that observed in individuals with late onset major depression (LOMD). This suggests that the reduction found in plasma A $\beta$ 42/A $\beta$ 40 levels in EOMD is unlikely to reflect preclinical AD and is consistent with epidemiological data that depression early in life may be a risk factor for AD [39]. Furthermore, these findings raise the possibility that in cognitively intact depressed elderly, factors intrinsic to depression such as increased severity of symptoms might influence A $\beta$  metabolism independently of preclinical or prodromal AD, or alternatively that brain A $\beta$  deposition that does not lead to overt cognitive symptoms (i.e., in the absence of neurofibrillary tangles or neuronal death) promotes depressive symptoms in the elderly. Several lines of evidence from the preclinical and clinical literature have presented compelling evidence that the applications of various types of stress, sleep deprivation, administration or elevations in corticosterone or corticotropin-releasing hormone, which have all been implicated in depression, can result in significant elevation in soluble and aggregated forms of A $\beta$  [40, 41] as well as elevation in total and phosphorylated tau [42]. These effects may be in part mediated by brain-region-specific increases in neuronal activity which has been reported in depression [43]. Additionally, a subtle loss of hippocampal integrity associated with depression, might result in a reduction of inhibitory input to the hypothalamic-pituitary-adrenal axis and cortisol elevations which have been implicated in altered A $\beta$  metabolism [44].

The reduced plasma A $\beta$ 42/A $\beta$ 40 ratio which we observed in LLMD subjects may involve both central and peripheral mechanisms. A decrease in plasma A $\beta$ 42/A $\beta$ 40 ratio is observed also in both AD subjects and in those with early or prodromal AD stages [45, 46] and has been associated to accumulation of A $\beta$ 42 in brain deposits with consequent sequestration of the A $\beta$ 42 peptide from CSF and plasma. A similar mechanism must be excluded in LLMD because it has been demonstrated that the longitudinal pattern of CSF A $\beta$ 42 concentrations does not depend on cognitive status but reflects depression severity with lower levels of CSF A $\beta$ 42 associated with more severe depressive symptoms [38]. Increased

platelet activation associated with depression [47] may have also influenced plasma A $\beta$ 42/A $\beta$ 40 ratio levels since platelets are known to produce A $\beta$ , particularly A $\beta$ 40, regardless of the increase in brain A $\beta$  deposition [48]. The increase in circulating levels of A $\beta$ 40 has also been implicated in the increase in A $\beta$  plaques formation through a number of mechanisms, including NF- $\kappa$ B-dependent endothelial cell activation, neuroinflammation, increase in APP metabolism and A $\beta$  production [49, 50]. Therefore, future studies should determine whether the increased platelet activation associated with depression causes an increase in circulating A $\beta$ 40 levels which in turn increases brain A $\beta$  deposition and consequently the risk of AD onset.

Although previous studies have shown an association between plasma BDNF levels and amyloid burden, our findings did not show significant association between BDNF and A $\beta$  indices. The lack of correlation between plasma BDNF and A $\beta$ 42, or A $\beta$ 40 in the plasma samples in our study could be due the differences in the methodology used, tissue samples examined (plasma vs. brain) and/or the subject population (AD vs late-life depression). In addition, the ELISA kit used in our study does not differentiate proBDNF and mature BDNF levels. ProBDNF is known to enhance A $\beta$  levels and accelerate its deposition in the brain [51]. Future studies using isoform specific BDNF antibodies are warranted to examine the relative roles of mature and proBDNF on A $\beta$  burden.

The main limitation of this study is the small sample size. Additionally, the results of this study were focused on plasma indices. Although we collected CSF A $\beta$  measures, it was only in a small subset. Therefore, we were not able to formally compare the plasma results with the corresponding CSF measures as there was an insufficient number of subjects with both plasma and CSF A $\beta$  determinations. In addition, our study did not examine the relationship between lower plasma A $\beta$ 42/A $\beta$ 40 ratio and indices of the HPA axis and microglial activation as well as whole brain and region-specific PET amyloid binding [52].

Future studies of LLMD should consider collecting CSF, plasma, and brain indices to further elucidate the relationship between these measures. Additionally, future studies should explore plasma A $\beta$  indices using mass spectrometry or ultrasensitive immunoassays (Simoa) with higher sensitivity and specificity for reliably measuring A $\beta$  and tau levels in blood [53] in this population to determine their relationship to depression diagnosis and to its aforementioned clinical features. If confirmed, more effective treatment of depressive symptoms, insomnia, and platelet activation in patients with depression may lead to a lowering of AD biomarkers and the associated risk of AD. Additionally, an important clinical question is whether we can delay the onset of AD in some individuals by treating their depressive symptoms.

In conclusion, our study demonstrates that the association of depression and plasma A $\beta$  status is robust. Future studies should determine if successful treatment of depressive symptoms in cognitively normal depressed subjects will impact the longitudinal plasma A $\beta$ 42/A $\beta$ 40 ratio and the potential long-term AD risk.

## DATA AVAILABILITY

The data generated during the current study are not publicly available because the approved IRB protocol does not include a provision for deposit in a public repository. The data may be made available from the corresponding author on reasonable request.

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## AUTHOR CONTRIBUTIONS

N.P. conceived, designed, and supervised the whole work, was the PI of the funded R01, was responsible for evaluation of study subjects, for collection of data and storage of samples. N.P. prepared the first draft of the manuscript. H.Z. and K.B.'s labs determined plasma Aβ levels. They also provided feedback on the statistical analysis and assisted in the interpretation of the results and preparation of the manuscript. A.P.'s lab determined plasma BDNF levels. He also helped with the interpretation of the results and writing of the manuscript. C.R.P. and D.B. performed the statistical analyses, assisted in the interpretation of the results and writing of the manuscript. C.R.P. generated the tables and figures. B.P.I., J.R.C., and R.S.O. assisted with the interpretation of the results and with manuscript preparation. All authors reviewed the manuscript and provided critical comments.

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## COMPETING INTERESTS

H.Z. has served at scientific advisory boards for Alector, Eisai, Denali, Roche Diagnostics, Wave, Samumed, Siemens Healthineers, Pinteon Therapeutics, Nervgen, AZTherapies and CogRx, has given lectures in symposia sponsored by Cellectricon, Fujirebio, Alzecure and Biogen, and is a co-founder of Brain Biomarker Solutions in Gothenburg AB (BBS), which is a part of the GU Ventures Incubator Program. KB has served as a consultant, at advisory boards, or at data monitoring committees for Abcam, Axon, Biogen, JOMDD/Shimadzu, Julius Clinical, Lilly, MagQu, Novartis, Prothena, Roche Diagnostics, and Siemens Healthineers, and is a co-founder of Brain Biomarker Solutions in Gothenburg AB (BBS), which is a part of the GU Ventures Incubator Program. All unrelated to the work presented in this paper. R.S.O. is a member of 'The Alliance for Sleep' which is sponsored by Idorsia U.S.

## ADDITIONAL INFORMATION

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