

ORIGINAL ARTICLE

INCREASES IN PLASMA A β 40 LEVELS AND THE A β 40/42 RATIO IN PATIENTS WITH DIABETES MELLITUS AND DEMENTIA

Yaeko Ueno¹⁾, Motohiro Nagaoka²⁾, Takumi Nakamura³⁾, Naoko Abe³⁾, Takashi Nakata³⁾, Yasuhito Wakasaya³⁾, Etsuro Matsubara⁴⁾, Takeshi Kawarabayashi³⁾ and Mikio Shoji³⁾

Abstract Aim: The plasma A β 40/42 ratio is a possible biomarker for the onset of Alzheimer's disease (AD). We here measured plasma A β 40 and A β 42 levels in patients with diabetes mellitus (DM) and dementia in order to clarify the relationship between DM and AD.

Methods: Fifty-three patients, including 33 patients with DM and 25 patients with dementia, were assessed using the Mini-mental state examination (MMSE) and brain MRI, plasma A β 40 and A β 42, blood sugar levels, and HbA1c % were measured, and the genotype of apolipoprotein E was determined. Plasma A β levels and blood sugar levels were measured in 16 out of 53 patients, following fasting and 2 hours after breakfast.

Results: Plasma A β 40 levels and the A β 40/42 ratio were increased in the DM with dementia group ($p < 0.01$, $p < 0.001$), while HbA1c % correlated with the A β 40/42 ratio in the non-dementia group. MMSE scores were also associated with the plasma A β 40/42 ratio and A β 40 levels in the non-dementia group, independent of the presence of ApoE ϵ 4. We did not observed significant direct responses of plasma A β proteins to an increase in blood sugar levels.

Conclusion: These results suggest that plasma A β metabolism are closely related chronic hyperglycemia before the onset of dementia.

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Key words: Dementia; Diabetes mellitus; HbA1c; Plasma A β 40; Plasma A β 40/42ratio.

Introduction

Recent studies identified diabetes mellitus (DM) as a risk factor for not only vascular dementia (VaD), but also Alzheimer's disease (AD)¹⁻³⁾. The Rotterdam Study and Hisayama Study also reported that DM was a significant risk factor for all-cause dementia and AD^{4, 5)}. Uncontrolled DM has been shown to increase the risk of AD and VaD⁶⁾. Furthermore, the severity of DM has been associated with cognitive function and a greater decline in the elderly⁷⁾. The Alzheimer's Disease Neuroimaging Initiative (ADNI) study found that subjects

with baseline normal blood sugar levels exhibited less of a decline in cognition and whole-brain volume as well as lower conversion rates from MCI to AD⁸⁾. However, 2 major prospective studies showed that glucose intolerance and insulin resistance were not associated with AD pathology⁹⁾ and that the rate of cognitive decline was slower in AD patients with DM¹⁰⁾.

The deposition of senile plaque amyloids consisting of aggregated A β , the appearance of neurofibrillary tangles, and neuronal cell loss are hallmark characteristics of AD pathology. Two major species of A β , A β 40 and A β 42, are

¹⁾ Department of Internal Medicine, ²⁾ Department of Psychiatry, Ube Kyoritsu Hospital, 16-23, Gojumezama, Ube, 755-0005, Japan

³⁾ Department of Neurology, Hirosaki University Graduate School of Medicine, Hirosaki, Japan

⁴⁾ Department of Neurology, Oita University Faculty of Medicine 1-1 Idaigaoka, Hasama, Yufu, Oita 879-5593, Japan

Correspondence: Y. Ueno

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processed from amyloid β precursor protein (APP), with A β 42 containing more hydrophobic amino acids in the C-terminus than A β 40. A β 42 is the early and causal component of amyloid deposits as diffuse plaques in the AD brain, followed by A β 40 deposits in core plaques and cerebral blood vessels. Recent studies confirmed that cerebrospinal fluid A β 42 and A β 40 were the most sensitive biomarkers for predicting and diagnosing AD¹¹⁻¹³). Plasma A β 42 and A β 40 can be assayed and the plasma A β 40/A β 42 ratio has recently been suggested as a biomarker of and major risk factor for AD¹²⁻¹⁵). A relationship has also been reported between cognitive function and the plasma A β 40/A β 42 ratio¹⁶⁻¹⁹).

To clarify the role of DM in the onset of dementia in AD, we here measured plasma A β 40 and A β 42 levels in 53 patients in our hospital and revealed that long-term hyperglycemia increased the plasma A β 40/A β 42 ratio, leading to clinical dementia.

Materials and Methods

Subjects consisted of 53 patients, including 35 females and 18 males, 33 of whom had DM and 25 had dementia. Thirty out of the 33 DM patients had type 2 DM while 3 had type 1 DM. Of the 25 dementia patients, 18 had AD, 5 had AD with cerebrovascular disease, and 2 had dementia with Lewy bodies. Regarding the clinical backgrounds of the 33 patients with DM, the mean duration of the disease was 16.1 years. Fourteen out of 33 patients were treated with insulin therapy. Half of these patients had diabetic microangiopathic complications such as retinopathy or nephropathy.

Neurological examinations, the Mini-mental state examination (MMSE), brain MRI, and a volumetric analysis using the Voxel-based Specific Regional analysis system for AD (VSRAD plus)²⁰) were carried out. The

diagnosis of DM depended on the diagnostic criteria of the Japanese Diabetes Society in 2010. Borderline glucose intolerance cases were excluded from DM patients in order to examine its effects on cognition under chronic hyperglycemia. The diagnosis of dementia was dependent on ICD10 and core clinical criteria by the National Institute on Aging and Alzheimer's disease association workgroups (AAN/AA). Neuropsychiatric examinations using HDS-R (Hasegawa dementia rating scale-revised), clock-drawing, cube copy, kana pick-up test, Kohs block design test, test your memory (TYM) Scale were carried out. Brain MRI (Hitachi APERTO INSPIRE 0.4 tesla) was conducted on all patients. Hippocampal and whole brain atrophy was analyzed by VSRAD plus.²⁰

Plasma A β 40 and A β 42 levels, the A β 40/42 ratio, the genotype of apolipoprotein E, blood glucose levels and serum C-peptide immunoreactivity (CPR), HbA1c %, and body mass index (BMI) were examined after informed consent and the agreement of patients and their families had been obtained. Blood samples were collected casual time in outpatient, and plasma was separated by centrifugation at 3,000 rpm for 10 minutes and stored at -45°C before the measurement of A β 40 and A β 42 using WAKO ELISA kits.¹³) Plasma A β 40 and A β 42 levels and blood glucose levels and serum CPR were measured in 14 out of 33 DM patients following fasting time and 2 hours after breakfast in hospitalized patient.

Statistical analysis

Statistical analyses were performed using SPSS ver11.01 (SPSS Inc.) and ver 22 and Graph Pad prism 6 (Graphpad Software, Inc.) using the Mann-Whitney U test, a one-way ANOVA, student t test, and paired *t*-test.

Table 1 Basic Patient Characteristics

	Total	Diabetes Mellitus (-)		Diabetes Mellitus (+)	
		Dementia (-) (DM-DEM-)	Dementia (+) (DM-DEM+)	Dementia (-) (DM+DEM-)	Dementia (+) (DM+DEM+)
Number	53	10	10	18	15
Mean age(years)	76.2 ± 9.3	73.3 ± 11.6	82.6 ± 5.4	70.9 ± 8.5	80.2 ± 6.3
Female/male	35/18	9/1	8/2	10/8	8/7
MMSE score	23.5 ± 4.2	27.2 ± 2.0	21.8 ± 1.9	26.2 ± 2.5	19.3 ± 3.8
HSD-R	22.9 ± 4.7	25.9 ± 2.7	20.1 ± 4.1	25.9 ± 3.3	19.1 ± 3.7
BS mmol/L	7.9 ± 3.2	5.9 ± 1.1	5.8 ± 1.4	9.2 ± 3.7	8.9 ± 3.0
HbA1c %	6.8 ± 1.5	5.7 ± 0.3	5.6 ± 0.4	7.5 ± 1.5	7.4 ± 1.6
BMI	22.8 ± 3.4	22.4 ± 4	21 ± 2.5	23.5 ± 4.1	23.5 ± 2
ApoEε4 n[%]	13 (25%)	2 (20%)	2 (20%)	5 (28%)	4 (27%)
Aβ40 fmol/ml	107.7 ± 45.6	83.7 ± 20.3	105.3 ± 33.6	102.7 ± 50.1	131.2 ± 51.5**
Aβ42 fmol/ml	11 ± 3.8	11.0 ± 2.8	11.9 ± 6.4	10.6 ± 3.2	11.2 ± 2.9
Aβ40/42	9.8 ± 2.7	7.7 ± 0.8	9.5 ± 2.4	9.6 ± 2.8	11.7 ± 2.7***
VSRAD plus					
Hippocampus score	1.6 ± 1.1	1.0 ± 0.6	2.2 ± 1.3*	1.1 ± 0.7	2.4 ± 1.1***
Whole Brain %	11.4 ± 4.4	7.8 ± 4.1	13.0 ± 4.1**	9.8 ± 3.4	14.8 ± 2.8***

DM: Diabetes Mellitus; DEM: dementia; MMSE: Mini-mental state examination score; HSD-R: Hasegawa dementia rating scale-revised; BS: casual blood sugar level, mmol/ml; BMI: Body mass index; VSRAD plus: Voxel-based Specific Regional analysis system for Alzheimer's Disease plus version, Hippocampus score: a Z-score in the medial temporal area including the entorhinal cortex, and whole brain % means the percentage of the voxel area showing atrophy (> Z-score 2) relative to the whole brain normal control database.

Results

The basic profiles of our subjects were described in Table 1. In the 20 subjects without DM (DM-), 10 patients did not have dementia (DM-DEM- group) while the other 10 did (DM-DEM+ group). Among the 33 subjects with DM (DM+), 18 did not have dementia (DM+DEM- group) while 15 did (DM+DEM+ group). No significant differences were observed in the mean age with or without DM; however, it was approximately 10 years higher in the groups with dementia. Mean MMSE scores were 27.2 ± 2.0 in the DM-DEM- group, 21.8 ± 1.9 in the DM-DEM+ group, 26.2 ± 2.5 in the DM+DEM- group, and 19.3 ± 3.8 in the DM+DEM+ group. No significant difference was observed in the frequency of ApoEε4 among these groups. Mean casual blood glucose levels were 5.9 ± 1.1 mmol/ml in DM-DEM-, 5.8 ± 1.4 mmol/ml in DM-DEM+, 9.2 ± 3.7 mmol/ml in DM+DEM- and 8.9 ± 3.0 mmol/ml in DM+DEM+. High blood glucose levels were

observed in the DM (+) groups. HbA1c % were 5.7 ± 0.3 %, 5.6 ± 0.4 %, 7.5 ± 1.5 %, and 7.4 ± 1.6 % in DM-DEM-, DM-DEM+, DM+DEM-, and DM+DEM+, respectively.

Aβ40 levels were 83.7 ± 20.3 fmol/ml in DM-DEM-, 105.3 ± 33.6 fmol/ml in DM-DEM+, 102.7 ± 50.1 fmol/ml in DM+DEM-, and 131.2 ± 51.5 fmol/ml in DM+DEM+. Aβ42 levels were 11.0 ± 2.8 fmol/ml, 11.9 ± 6.4 fmol/ml, 10.6 ± 3.2 fmol/ml, and 11.2 ± 2.9 fmol/ml in DM-DEM-, DM-DEM+, DM+DEM-, and DM+DEM+, respectively. Aβ40/42 ratios were 7.7 ± 0.8, 9.5 ± 2.4, 9.6 ± 2.8, and 11.7 ± 2.7 in DM-DEM-, DM-DEM+, DM+DEM-, and DM+DEM+, respectively. Aβ40 levels were significantly higher in DM+DEM+ (P<0.01) than in DM-DEM-, DM-DEM+, and DM+DEM-. However, no significant differences were observed in Aβ42 levels between the groups. The mean Aβ40/42 ratio was also significantly higher in DM+DEM+ (P<0.001) (Fig. 1). These results suggested that DM or dementia elevated Aβ40 levels and the Aβ40/42 ratio in plasma.

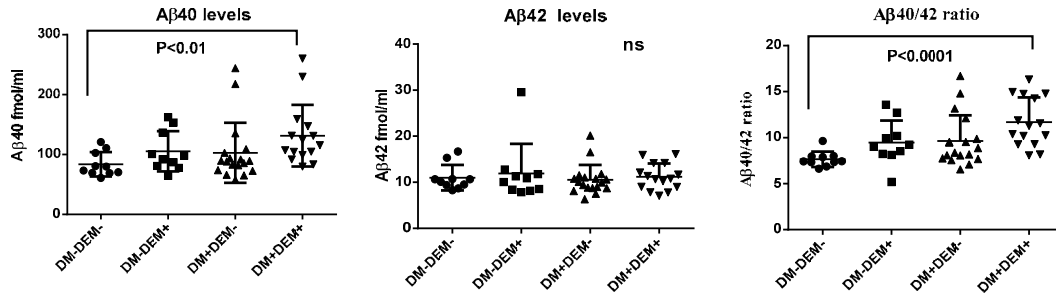


Fig.1 Plasma A β 40 and A β 42 levels and the A β ratio in four group. A β 40 levels and A β ratio were significantly higher in the DM+DEM+ group ($P < 0.01$; $P < 0.0001$, respectively).

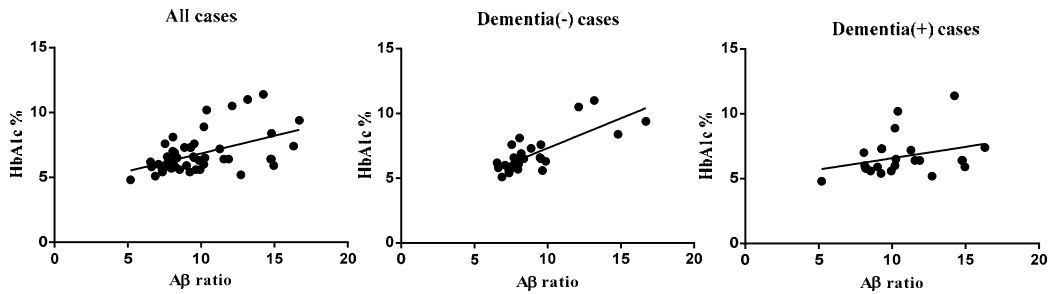


Fig.2 The relationship between the A β ratio and HbA1c. Correlations were observed between HbA1c % and the A β 40/42 ratio in all subjects ($Y = 0.2729X + 4.11$; $P < 0.0002$) and in Dementia (-) cases ($Y = 0.4613X + 2.709$; $P < 0.0001$).

We analyzed the relationship between HbA1c % and the A β 40/42 ratio, and found correlations in all subjects ($Y = 0.2729X + 4.11$; $P < 0.0002$) and the DEM- groups ($Y = 0.4613X + 2.709$; $P < 0.0001$) (Fig. 2). To examine this relationship in more detail, we separated all cases into 4 subgroups according to HbA1c % (7 %) and the presence of dementia. The A β 40/42 ratio was elevated in patients with HbA1c % less than 7 % and with dementia ($P < 0.001$; Fig. 3). Moreover diabetics belong to the group of more than 7 % had high level of the A β ratio with or without dementia. These results demonstrated that long-term hyperglycemia road influenced the 40/42 ratio, suggesting that the metabolism of A β is altered in DM patients without dementia.

To analyze the relationship between cognitive function and plasma A β levels, a linear regression analysis was performed between

MMSE scores and the A β 40/42 ratio and A β 40 and A β 42 levels. MMSE scores correlated with the A β 40/42 ratio ($Y = -0.2485X + 14.68$; $P = 0.0045$) and A β 40 levels ($Y = -3.482X + 189.8$; $P = 0.081$). However, a relationship was not observed between MMSE scores and A β 42 levels (Fig. 4). No significant differences were noted in A β 40 levels (109.8 ± 49.6 SE vs 101.2 ± 31), A β 42 levels (10.92 ± 2.9 vs 11.56 ± 5.9), or the A β 40/42 ratio (9.97 ± 2.9 vs 9.36 ± 2.28) between groups without or with ApoE ϵ 4 (Fig. 5). These results suggested that the plasma A β 40/42 ratio and A β 40 levels were closely related to cognitive function independent of the presence of ApoE ϵ 4.

To clarify the direct relationship between blood glucose and plasma A β levels, blood samples from 14 inpatients were examined for blood glucose control before breakfast and 2

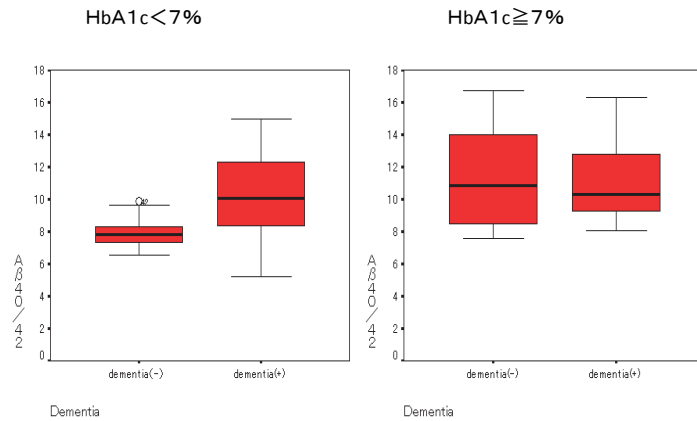


Fig.3 The relation of between the Aβ40/42 ratio and HbA1c in two group according to the blood sugar control.

The Aβ ratio was elevated significantly in patients with HbA1c less than 7% and with dementia ($P < 0.001$), but in patients more than 7% not differed with or without dementia.

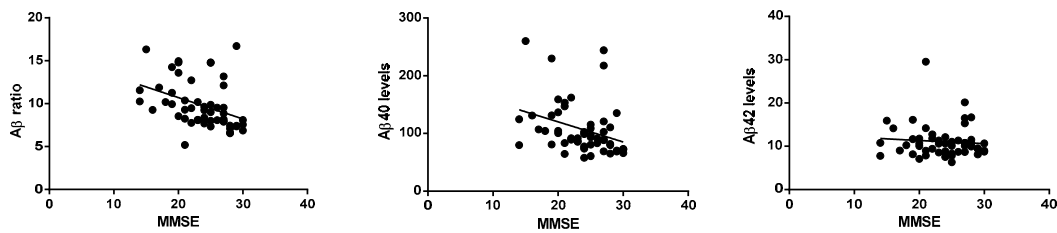


Fig.4 Aβ levels and Cognitive impairment.

MMSE scores correlated with the Aβ ratio ($Y = -0.2485X + 14.68$; $P = 0.0045$) and Aβ40 levels ($Y = -3.482X + 189.8$; $P = 0.081$). However, no relationship was observed between MMSE scores and Aβ42 levels.

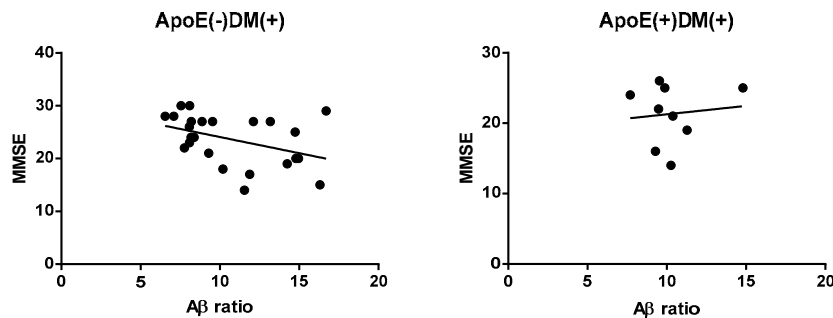


Fig.5 Relationship between the Aβ ratio and ApoEε4 in dementia patients.

A correlation between decreased MMSE scores and a higher Aβ ratio was observed in patients with DM without ApoEε4. However, no significant relationship was found in patients with DM and ApoEε4.

hours after. Figure 6 shows serial alterations in these 4 markers. Blood sugar levels significantly increased from 7.32 mmol/L to 9.50 mmol/L ($P < 0.002$). However, no significant changes were observed in Aβ40 levels (from 148 fmol/ml

to 151 fmol/ml), Aβ42 levels (from 11.87 fmol/ml to 12.14 fmol/ml), or the Aβ ratio (from 12.58 to 12.49), suggesting that blood glucose levels did not directly affect plasma Aβ levels or their ratios, daily fluctuations in blood glucose levels

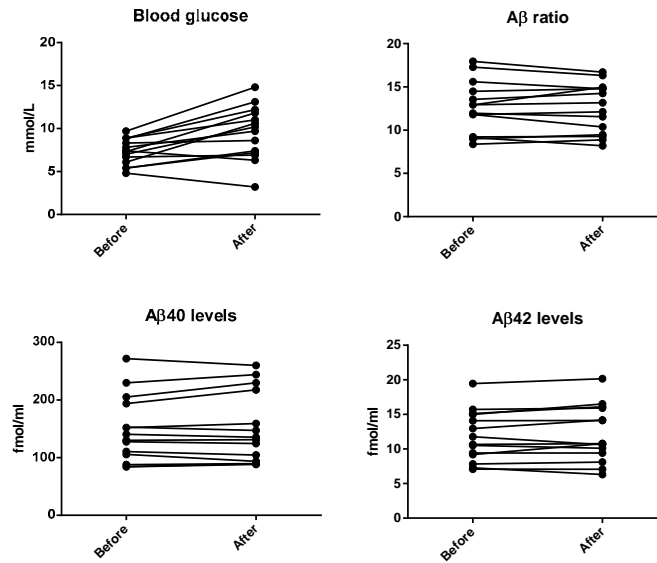


Fig.6 Direct association between blood sugar and A β levels. Blood sugar levels were significantly increased ($P < 0.002$). However, A β 40 and A β 42 levels and the A β ratio were not significantly affected.

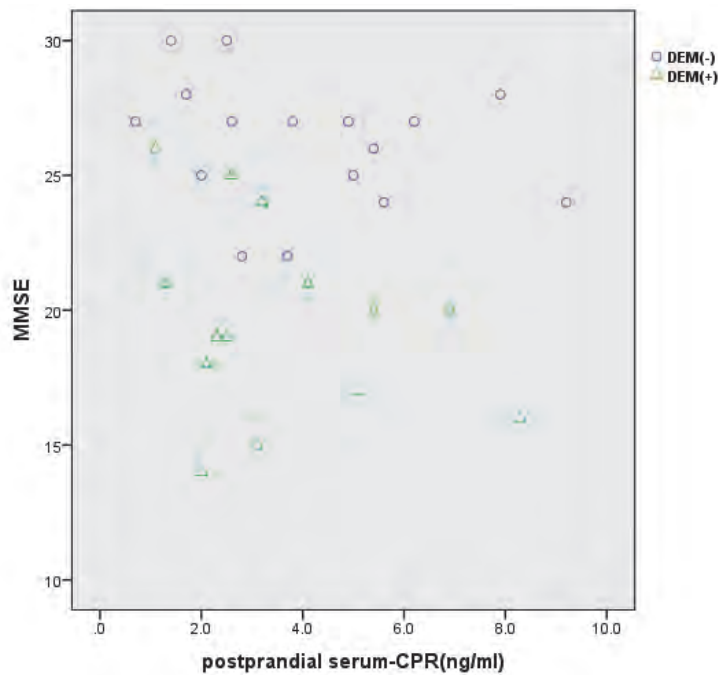


Fig.7 Presence of dementia and the level of postprandial serum CPR in type2 diabetes (n=30). Patients with dementia showed by open triangles, have any level of serum CPR that indicate endogenous insulin secretion ability.

did not change the A β ratio or A β 40 levels, and long-term disturbances in glucose metabolism affected plasma A β metabolism in DM patients. In this series we measured serum CPR that was an index to express endogenous insulin

production instead of serum insulin levels because insulin therapy had already done in many patients. The result showed that patients with various insulin production abilities could suffer from cognitive impairment (Fig 7).

Discussion

We herein revealed that plasma A β 40 levels and the A β 40/42 ratio were increased in the DM with dementia group and that HbA1c % correlated with the A β 40/42 ratio, especially in the non-dementia group. A decline in cognitive function was associated with the plasma A β ratio and A β 40 levels in the non-dementia group, independent of the presence of ApoE ϵ 4. However, no significant changes were observed in plasma A β 42 levels in any factors, and we did not detect any direct response of plasma A β 40 levels or the A β 40/42 ratio to increased blood sugar levels. These results demonstrated that plasma A β 40 levels and the A β 40/42 ratio are closely related with long-term hyperglycemia before the onset of dementia.

A systematic review of 14 eligible longitudinal population-based studies revealed the high risk of VaD and AD in individuals with DM and suggested that vascular disease and alterations in the metabolism of glucose, insulin, and A β underlie its pathophysiology.¹⁾ Although the risk of AD is less than that with ApoE ϵ 4, DM is a significant independent risk.²⁾ Hyperinsulinemia disrupts the clearance of A β from the brain by competing with insulin-degrading enzymes, and directly affects the accumulation of A β in the AD brain.^{3, 22)} Insulin resistance in the brain also induces central inflammation, cerebrovascular inflammation, oxidative stress, and mitochondrial dysfunction, and advanced glycation end products appear to be important mediators that facilitate the pathology of AD.²³⁻²⁵⁾

A prospective cohort study conducted by Graff-Radford *et al.* showed that the plasma A β 42/40 ratio may be a useful biomarker for predicting the development of mild cognitive impairment (MCI) or AD.¹⁶⁾ The Rotterdam Study demonstrated that a high concentration of A β 40, but not A β 42 at baseline was associated

with an increased risk of dementia.¹⁷⁾ Low or decreasing plasma A β 42 levels during a 3- to 6-year follow-up were associated with cognitive decline in MCI and a cognitively intact control cohort.¹⁸⁾ A low plasma A β 42/40 ratio has also been associated with a greater decline among the elderly without dementia over 8 years.¹⁹⁾ The Australian Imaging Biomarker and Lifestyle (AIBL) study of aging revealed a decrease in the A β 42/40 ratio in patients with AD that was also inversely correlated with the neocortical amyloid burden. Over 18 months, plasma A β 42 levels decreased in subjects with MCI and in those transitioning from a healthy condition to MCI.²¹⁾ This finding supports our result in which the A β 40/42 ratio was related to cognitive impairment before the onset of dementia. Our results also indicate that long-term hyperglycemia is a causal factor that increases plasma A β 40 levels and the A β 40/42 ratio, leading to cognitive impairment before the onset of dementia.

The relationship between HbA1c % and plasma A β levels has not yet been elucidated in detail.²⁶⁾ Our results clearly show that the A β 40/42 ratio correlated with HbA1c %. When subjects were limited to patients without dementia, the A β 40/42 ratio strongly correlated with higher HbA1c % (≥ 7 %, $p < 0.0001$). These results also suggest that patients with chronic hyperglycemia had impaired A β metabolism before the onset of dementia. And another aspect in this study the plasma A β 40/42 ratio was significantly high in a group with dementia when HbA1c % was less than 7 %. It is thought that the plasma A β 40/42 ratio is possible biomarker of dementia in without diabetes or in controlled diabetes, but not in poorly controlled diabetes.

In the present analysis of 53 subjects, a negative correlation was observed between plasma A β 40/42 ratios and MMSE scores. The increase observed in the plasma A β 40/42

ratio was mainly attributed to elevated A β 40 levels rather than A β 42 levels. Van Oijen *et al.* previously showed that increased plasma A β 40 levels were the most important risk factor for the development of dementia, and that this risk was further increased by a decrease in plasma A β 42 levels. They also reported that increased A β 40 levels were often observed in patients with white matter lesions in the brain.¹⁷⁾ A β 40 has been implicated in vascular toxicity, which may impair the regenerative function of vascular endothelial cells.²⁹⁾ In addition, the deposition of A β 40 in the walls of blood vessels has been observed with amyloid angiopathy in the elderly and AD patients. Therefore, the condition of increased A β 40 levels, such as in hyperglycemia, may facilitate the progression of microangiopathy, leading to cerebral ischemia, which is consistent with the results of the present study.

By a large-scaled study a history of severe hypoglycemic episodes has been associated with a greater risk of dementia, and severely diabetic patients with increased HbA1c % levels are more likely to develop hypoglycemia.²⁷⁾ An impaired acute insulin response at midlife was associated with an increased risk of AD up to 35 years later.²⁸⁾ We recognize the fact that dementia could be develop in various insulin secretion ability in type 2 diabetes (Figure7). It seemed dementia in type 2 diabetes have heterogeneity, and the lower insulin secretion group have more frequency of both hypoglycemia and sugar toxic state. On the other hand it is said type 2 diabetes caused dementia from hyperinsulinemia, we suggest the possibility that not severe diabetes with higher insulin secretion ability have more typical pathologies of alzheimer's dementia in comparison with so-called diabetic dementia.

In the present study, no significant differences were observed in the prevalence of ApoE ϵ 4 in each group or between those with and without

DM, and it reflects that reported in the healthy Japanese population. A negative correlation was noted between MMSE scores and A β 40/42 ratios in the patient group without ApoE ϵ 4, but not in another group with ApoE ϵ 4. These results suggest that the plasma A β 40/42 ratio is related to cognitive impairments independent of ApoE ϵ 4 levels.

A previous study reported that A β 40 levels significantly increased after a glucose load.²⁵⁾ However, we did not find any significant changes in A β 40 levels, A β 42 levels, or the A β 40/42 ratio after a food load in spite of a significant increase in the mean blood glucose level. Thus, there was no significant short-term change in A β levels in response to hyperglycemia after a meal. This may be partly attributed to lower blood sugar levels after a meal than those before receiving anti-diabetic drugs, and also to smaller daily fluctuations in plasma A β levels than those in cerebrospinal fluid A β levels.

Conclusion

Plasma A β 40 levels and the A β 40/42 ratio were increased in DM patients with dementia and HbA1c % correlated with the A β 40/42 ratio. A cognitive impairment was also related to the plasma A β 40/42 ratio and A β 40 levels in DM patients without dementia independent of the presence of ApoE ϵ 4. We did not find any direct response of plasma A β 40 levels or the A β 40/42 ratio to increased blood sugar levels. These results indicate that plasma A β 40 levels and the A β 40/42 ratio are closely related to long-term hyperglycemia before the onset of dementia. Moreover in controlled or not-diabetics the A β 40/42 ratio may possibly be biomarker of dementia.

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Disclosure statement

The authors declare no conflict of interest.

References

- 1) Biessels GJ, Staekenborg S, Brunner E, Brayne C, Scheltens P. Risk of dementia in diabetes mellitus: a systematic review. *Lancet Neurol.* 2006 Jan;5(1):64-74.
- 2) Profenno LA, Porsteinsson AP, Faraone SV. Meta-analysis of Alzheimer's disease risk with obesity, diabetes, and related disorders. *Biol Psychiatry.* 2010 Mar 15;67(6): 505-12.
- 3) Mayeux R, Stern Y. Epidemiology of Alzheimer disease. *Cold Spring Harb Perspect Med.* 2012 Aug 1;2(8).
- 4) Ott A, Stolk RP, van Harskamp F, Pols HA, Hofman A, Breteler MM. Diabetes mellitus and the risk of dementia: The Rotterdam Study. *Neurology.* 1999 Dec 10;53(9):1937-42.
- 5) Ohara T, Doi Y, Ninomiya T, Hirakawa Y, Hata J, Iwaki T, et al. Glucose tolerance status and risk of dementia in the community: the Hisayama study. *Neurology.* 2011 Sep 20;77(12):1126-34.
- 6) Xu WL, von Strauss E, Qiu CX, Winblad B, Fratiglioni L. Uncontrolled diabetes increases the risk of Alzheimer's disease: a population-based cohort study. *Diabetologia.* 2009 Jun;52(6):1031-9.
- 7) Yaffe K, Falvey C, Hamilton N, Schwartz AV, Simonsick EM, Satterfield S, et al. Diabetes, glucose control, and 9-year cognitive decline among older adults without dementia. *Arch Neurol.* 2012 Sep;69(9):1170-5.
- 8) Morris JK, Vidoni ED, Honea RA, Burns JM; Alzheimer's Disease Neuroimaging Initiative. Impaired glycemia increases disease progression in mild cognitive impairment. *Neurobiol Aging.* 2014 Mar;35(3):585-9.
- 9) Sanz C, Andrieu S, Sinclair A, Hanaire H, Vellas B; REALFR Study Group. Diabetes is associated with a slower rate of cognitive decline in Alzheimer disease. *Neurology.* 2009 Oct 27;73(17):1359-66.
- 10) Thambisetty M, Beason-Held LL, An Y, Kraut M, Metter J, Egan J, Ferrucci L, O'Brien R, Resnick SM. Impaired glucose tolerance in midlife and longitudinal changes in brain function during aging. *Neurobiol Aging.* 2013 Oct;34(10):2271-6.
- 11) Kanai M, Matsubara E, Isoe K, Urakami K, Nakashima K, Arai H, et al. Longitudinal study of cerebrospinal fluid levels of tau, A β 1-40, and A β 1-42(43) in Alzheimer's disease: a study in Japan. *Ann Neurol.* 1998 Jul;44(1): 17-26.
- 12) Fagan AM, Xiong C, Jasielec MS, Bateman RJ, Goate AM, Benzinger TL, et al; Dominantly Inherited Alzheimer Network. Longitudinal change in CSF biomarkers in autosomal-dominant Alzheimer's disease. *Sci Transl Med.* 2014 Mar 5;6(226):226ra30.
- 13) Xu W, Kawarabayashi T, Matsubara E, Deguchi K, Murakami T, Harigaya Y, et al. Plasma antibodies to Abeta40 and Abeta42 in patients with Alzheimer's disease and normal controls. *Brain Res.* 2008 Jul 11;1219:169-79.
- 14) Shoji M. Biomarkers of the dementia. *Int J Alzheimers Dis.* 2011;2011:564321
- 15) Schupf N, Zigman WB, Tang MX, Pang D, Mayeux R, Mehta P, Silverman W. Change in plasma A β peptides and onset of dementia in adults with Down syndrome. *Neurology.* 2010 Nov 2;75(18):1639-44.
- 16) Graff-Radford NR, Crook JE, Lucas J, Boeve BF, Knopman DS, Ivnik RJ, et al. Association of low plasma Abeta42/Abeta40 ratios with increased imminent risk for mild cognitive impairment and Alzheimer disease. *Arch Neurol.* 2007 Mar;64(3): 354-62.
- 17) van Oijen M, Hofman A, Soares HD, Koudstaal PJ, Breteler MM. Plasma Abeta(1-40) and Abeta(1-42) and the risk of dementia: a prospective case-cohort study. *Lancet Neurol.* 2006 Aug;5(8):655-60.
- 18) Seppälä TT, Herukka SK, Hänninen T, Tervo S, Hallikainen M, Soininen H, Pirttilä T. Plasma Abeta42 and Abeta40 as markers of cognitive change in follow-up: a prospective, longitudinal, population-based cohort study. *J Neurol Neurosurg Psychiatry.* 2010 Oct;81(10):1123-7.

- 19) Yaffe K, Weston A, Graff-Radford NR, Satterfield S, Simonsick EM, Younkin SG, et al. Association of plasma beta-amyloid level and cognitive reserve with subsequent cognitive decline. *JAMA*. 2011 Jan 19;305(3):261-6.
- 20) Nakatsuka T, Imabayashi E, Matsuda H, Sakakibara R, Inaoka T, Terada H. Discrimination of dementia with Lewy bodies from Alzheimer's disease using voxel-based morphometry of white matter by statistical parametric mapping 8 plus diffeomorphic anatomic registration through exponentiated Lie algebra. *Neuroradiology*. 2013 May;55(5):559-66.
- 21) Rembach A, Faux NG, Watt AD, Pertile KK, Rumble RL, Trounson BO, et al. AIBL research group. Changes in plasma amyloid beta in a longitudinal study of aging and Alzheimer's disease. *Alzheimers Dement*. 2014 Jan;10(1):53-61.
- 22) Qiu WQ, Folstein MF. Insulin, insulin-degrading enzyme and amyloid-beta peptide in Alzheimer's disease: review and hypothesis. *Neurobiol Aging*. 2006 Feb;27(2):190-8.
- 23) Yang Y, Song W. Molecular links between Alzheimer's disease and diabetes mellitus. *Neuroscience*. 2013 Oct 10;250:140-50.
- 24) Fishel MA, Watson GS, Montine TJ, Wang Q, Green PS, Kulstad JJ, et al. Hyperinsulinemia provokes synchronous increases in central inflammation and beta-amyloid in normal adults. *Arch Neurol*. 2005 Oct;62(10):1539-44.
- 25) Takeda S, Sato N, Uchio-Yamada K, Sawada K, Kunieda T, Takeuchi D, et al. Diabetes-accelerated memory dysfunction via cerebrovascular inflammation and Abeta deposition in an Alzheimer mouse model with diabetes. *Proc Natl Acad Sci U S A*. 2010 Apr 13;107(15):7036-41.
- 26) Yaffe K, Blackwell T, Whitmer RA, Krueger K, Barrett Connor E. Glycosylated hemoglobin level and development of mild cognitive impairment or dementia in older women. *J Nutr Health Aging*. 2006 Jul-Aug;10(4):293-5.
- 27) Whitmer RA, Karter AJ, Yaffe K, Quesenberry CP Jr, Selby JV. Hypoglycemic episodes and risk of dementia in older patients with type 2 diabetes mellitus. *JAMA*. 2009;301:1565-1572.
- 28) Rönnekaa E, Zethelius B, Sundelöf J, Sundström J, Degerman-Gunnarsson M, Berne C, et al. Impaired insulin secretion increases the risk of Alzheimer disease. *Neurology*. 2008 Sep 30;71(14):1065-71.
- 29) Hayashi S, Sato N, Yamamoto A, Ikegame Y, Nakashima S, Ogihara T, Morishita R. Alzheimer disease-associated peptide, amyloid beta40, inhibits vascular regeneration with induction of endothelial autophagy. *Arterioscler Thromb Vasc Biol*. 2009 Nov;29(11):1909-15.