Plasma β-Amyloid 1-40 Is Associated With the Diffuse Small Vessel Disease Subtype

Merixtell Gomis, MD; Tomás Sobrino, PhD; Angel Ois, MD; Mònica Millán, MD; Ana Rodríguez-Campello, MD; Natalia Pérez de la Ossa, MD; Raquel Rodríguez-González, BSc; Jordi Jiménez-Conde, MD, PhD; Elisa Cuadrado-Godia, MD; Jaume Roquer, MD, PhD; Antoni Dávalos, MD, PhD

Background and Purpose—The underlying mechanisms of small vessel disease (SVD) subtypes are diffuse arteriopathy (diffuse-SVD) or microatheroma (focal-SVD). Endothelial dysfunction by β-amyloid peptide (Aβ) deposition has been associated with lacunar infarcts and leukoaraiosis, but its specific relationship with SVD subtypes is unknown. We hypothesized that plasma Aβ levels can play a different role in SVD subtypes in patients with acute lacunar stroke.

Methods—We studied 149 patients with acute ischemic stroke of SVD etiology according to Trial Of Org 10172 In Acute Stroke Treatment criteria and 25 age-matched control subjects. Patients were classified into focal-SVD: 39 patients with isolated lacunar infarct without leukoaraiosis and diffuse-SVD: 110 patients with an isolated lacunar infarct with leukoaraiosis or with multiple lacunar infarcts with or without leukoaraiosis. Baseline data included vascular risk factors and extensive laboratory tests, including plasma Aβ levels.

Results—Median [quartiles] Aβ1-40 levels (40.4 [35.1, 50.5] versus 55.1 [42.3, 69.6] pg/mL), but not Aβ1-42 levels, were significantly higher in the diffuse-SVD group than in focal-SVD group (P<0.001) and control subjects (P<0.001). No differences in Aβ1-40 levels were found between focal-SVD and control subjects. Logistic regression analysis showed that age (OR, 1.06; 95% CI, 1.01 to 1.12), history of hypertension (OR, 3.5; 95% CI, 1.3 to 9.2), and plasma Aβ1-40 levels over the median value (OR, 17.3; 95% CI, 3.0 to 99 for the third quartile and OR, 6.0; 95% CI, 1.6 to 23 for the fourth quartile) were independently associated with the diffuse-SVD subtype.

Conclusions—Plasma β-amyloid1-40 levels are independently associated with the diffuse-SVD subtype. These results are consistent with the pathophysiological role of fraction Aβ1-40 in disrupting endothelial vascular function. (Stroke. 2009;40:3197-3201.)

Key Words: acute stroke ■ beta-amyloid protein ■ leukoaraiosis ■ small vessel disease

The pathogenesis of cerebral small vessel disease (SVD) is incompletely understood. Pathological studies have suggested that there may be 2 types of SVD that can be differentiated on brain imaging.1 A diffuse arteriopathy of the perforating arteries with hyaline deposition: a pattern referred to as lipohyalinosis (diffuse-SVD) and localized small vessel microatheroma at the origin of the deep perforating arteries (focal-SVD). Diffuse-SVD is associated with multiple small lacunar infarcts with leukoaraiosis and focal-SVD with single large lacunar infarcts without leukoaraiosis.2,3 Endothelial dysfunction may play an important role in the pathogenesis of the diffuse-SVD subtype.4–6 Plasma β-amyloid peptide (Aβ) is a peptide consisting of either 42 (Aβ42) or 40 (Aβ40) amino acids derived from a proteolytic processing of the amyloid precursor protein.7 Insoluble Aβ fibrils are the predominant constituents of senile plaques, one of the pathological hallmarks of Alzheimer disease,8,9 and of cerebrovascular amyloid in the related condition of cerebral amyloid angiopathy.10 Plaque amyloid is primarily comprised of Aβ42, whereas vascular amyloid is formed by the Aβ40 species. Recent evidence suggests that amyloid precursor protein overexpression and Aβ accumulation impair cerebral circulation.11,12 In vitro studies have suggested direct physiological or toxic effects of Aβ1-40 in the regulation of cerebral circulation by endothelial cells.13 In humans, recent studies have demonstrated an association among plasma Aβ concentrations, leukoaraiosis, and lacunar infarcts.14,15 However, the specific relationship between Aβ and SVD subtypes is unknown. We hypothesized that plasma Aβ levels can play a different role in SVD subtypes in patients with acute lacunar stroke.

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From the Stroke Unit (M.G., M.M., N.P.d.l.O., A.D.), Department of Neurosciences, Hospital Universitari Germans Trias i Pujol (Badalona), Department of Medicine of the Universitat Autònoma de Barcelona, Barcelona, Spain; the Clinical Neurosciences Research Laboratory (T.S., R.R.-G.), Hospital Clínico Universitario, University of Santiago de Compostela La Coruña, Spain; and Unitat d’Ictus (A.O., A.R.-C., J.J.-C., E.C.-G., J.R.), Servei de Neurologia, Hospital del Mar, Departament de Medicina de la Universitat Autònoma de Barcelona, IMIM-Hospital del Mar, Barcelona, Spain. Correspondence to Merixtell Gomis, MD, Stroke Unit, Neurosciences Department, Hospital Universitari Germans Trias i Pujol, Carretera de Canyet s/n Badalona, Barcelona, Spain. E-mail mgomis.germanstrias@gencat.cat © 2009 American Heart Association, Inc.
Subjects and Methods

Study Population
BasicMar16 is a stroke register designed as a tool to study epidemiologic data in a hospital-based population of a single center in Barcelona serving a population of approximately 300,000.

From January 2005 to November 2007, 1106 consecutive patients with acute stroke were admitted. Of them, 194 were diagnosed as SVD according to Trial Of Org 10172 In Acute Stroke Treatment criteria14 at hospital discharge. We excluded 27 patients without stored blood samples because they did not give their consent to participate in BasicMar register. We also excluded 8 patients whose Trial Of Org 10172 In Acute Stroke Treatment classification changed at the final visit and 10 patients who we lost contact with and we were unable to follow-up at 3 months. Finally, 149 patients were eligible for this study; all survived the 3-month follow-up period.

Demographic data, stroke risk factors, acute phase clinical and biological data, stroke severity at hospital admission, and functional outcome at 3 months after stroke were prospectively recorded. SVD subtype was classified according to the Trial Of Org 10172 In Acute Stroke Treatment criteria15 using in-hospital and outpatient data and defined as a clinical lacunar syndrome with a compatible acute lesion on MRI or CT. Exclusion criteria included the presence of subcortical infarction >15 mm in diameter or cortical infarction of any size; carotid or vertebral artery stenosis >50%; and potential cardiac sources of embolism constituting high or moderate risk under Trial Of Org 10172 In Acute Stroke Treatment criteria.15 Patients were re-evaluated at 3 months after stroke onset to assess morbidity and mortality rates and confirm the initial Trial Of Org 10172 In Acute Stroke Treatment classification.17 We also included 25 age-matched control subjects without SVD. Control subjects were subjects with stroke mimics events that were included in the BasicMar register (seizures=5, hypoglycemia=4, conversion disorder=8, migraine with aura=2, peripheral neuropathy=3, and dizziness=3). None of them had a final diagnosis of stroke and all these patients had a normal neuroimaging study. Informed consent was obtained from the patients or relatives, and the study was approved by the Ethics Committee of Hospital del Mar, Barcelona.

Data Collection
Patients completed the extensive BasicMar16 clinical protocol, including demographic data (age and sex) and determination of the following vascular risk factors: arterial hypertension (defined as evidence of at least 2 blood pressure measurements >140/90 mm Hg recorded on different days before stroke onset, a physician’s diagnosis, or use of antihypertensive medication), diabetes (fasting serum glucose level >7.0 mmol/L, a physician’s diagnosis, or use of diabetic medication), hyperlipidemia (serum cholesterol concentration >12.2 mmol/L or serum triglyceride concentration >1.1 mmol/L, a physician’s diagnosis, or use of medication), alcohol overuse (>60 g/day), and current smoking status. We also recorded weight, height, waist circumference, systolic and diastolic blood pressure, axillary temperature, severity of symptoms based on National Institutes of Health Stroke Scale score17 at admission, and prestroke antplatelet and statins use. Biological determinations included leukocyte count and plasma creatinine levels. All patients had brain imaging, electrocardiogram, and Duplex or Doppler ultrasound study of the intra- and extracranial vessels. When clinical suspicion was high for a cardioembolic source of stroke, echocardiography was performed. The control group (stroke mimics) was evaluated by using the same clinical protocol, but ultrasound studies and echocardiography were performed only in doubtful cases.

Plasma Amyloid β Levels
Blood samples were collected on admission in glass test tubes, centrifuged at 3000 g for 10 minutes, and immediately frozen and stored at -80°C. Serum Aβ42 levels were measured with commercially available quantitative enzyme-linked immuno-nsorbent assay kits obtained from Biosource Europe SA, Belgium. Determinations were performed in an independent laboratory blinded to clinical and neuroimaging data.

Subtyping of Lacunar Stroke
Patients were retrospectively classified into 2 groups based on neuroimaging findings: focal-SVD group, including 39 patients with isolated lacunar infarct without leukoaraiosis, and diffuse-SVD group, including 110 patients with an isolated lacunar infarct with leukoaraiosis or with multiple lacunar infarcts (≥3) with or without leukoaraiosis.

Lacunar Brain Infaracts and Leukoaraiosis
We obtained axial T1-, T2-, and proton density-weighted scans on 1.5-T MRI scanners (GE Signa) using 5-mm slice thickness. Slice thicknesses for CT scans varied from 5 to 8 mm.

Leukoaraiosis was defined as ill-defined hyperintensities ≥5 mm on both T2 and proton density/fluid-attenuated inversion recovery MRI images without prominent hyperintensities on T1-weighted MRI scans and as ill-defined and moderately hypodense areas of ≥5 mm on CT. Leukoaraiosis was quantified on MRI by using the Fazekas scale.19 This method yields 2 separate scores for subcortical and deep white matter lesions and periventricular lesions. The 4-point Fazekas scale of increasing severity was used to classify each score with 0 indicating a patient without leukoaraiosis.

Lacunes were defined as well-defined hypodense areas of >2 mm and <15 mm on CT and as hyperintensities >2 mm and <15 mm on both T2 and proton density with hypointensities on T1-weighted and fluid-attenuated inversion recovery MRI images. If lesions with these characteristics were ≤2 mm, they were considered perivascular spaces, except around the anterior commissure, where perivascular spaces can be larger. Patients with both CT and MRI studies were classified in the diffuse or focal SVD group according to the MRI assessment.

Statistical Analysis
Quantitative variables are presented as mean±SD or median [interquartile range], as appropriate, and qualitative variables are presented as percentages. The χ2 test or Mann-Whitney U test was used to assess differences in continuous variables, and the χ2 test was used to compare proportions. The association between Aβ1-40 levels and the variables listed in Table 1 was analyzed by using the Mann-Whitney U test and Spearman correlation.

The relationship between Aβ fractions and SVD subtype was analyzed by logistic regression analysis (enter method). Due to a lack of linearity of the ORs, Aβ1-40 was entered in the logistic model classified in quartiles according to these 3 cutoff values, 39, 53, and 68 pg/mL. It has been suggested that the ratio of Aβ1-42 to Aβ1-40 may be more important than separate levels, at least for Alzheimer disease.20,21 Therefore, we also analyzed the association between the ratio of plasma Aβ1-42/Aβ1-40 and the diffuse SVD subtype. Potential confounders of the SVD subtype were included in the logistic models; we selected those factors that have been associated with SVD in previous reports or those associated with Aβ1-40 in the present work in bivariate analyses with a probability value <0.05. Statistical analyses were performed with the SPSS 13.0 software package.

Results
SVD subtype was diagnosed according to findings on acute MRI (n=104 [69.8%]) or CT alone (n=45 [30.2%]). A total of 39 patients were classified in the focal-SVD group (26 by MRI); none of them had leuкоaraiosis. Two patients in the focal-SVD group had 2 lacunar infarcts in the MRI, whereas the other 37 had only one. The 110 participants classified in the diffuse-SVD group (78 by MRI) included patients with an isolated lacunar infarct with leuкоaraiosis (n=41), with
Table 1. Medical History and Baseline Characteristics by SVD Subtype Groups

<table>
<thead>
<tr>
<th></th>
<th>Focal-SVD (n=39)</th>
<th>Diffuse-SVD (n=110)</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean age, years</td>
<td>66.3±10.8</td>
<td>73.6±9.8</td>
<td>0.0001</td>
</tr>
<tr>
<td>Males</td>
<td>69.2</td>
<td>63.6</td>
<td>0.33</td>
</tr>
<tr>
<td>Weight, kg (n=143)</td>
<td>73.2±13.6</td>
<td>74.3±15.3</td>
<td>0.68</td>
</tr>
<tr>
<td>Height, cm (n=143)</td>
<td>163.7±9.2</td>
<td>162.4±9.8</td>
<td>0.47</td>
</tr>
<tr>
<td>Waist circumference (n=117)</td>
<td>100.2±16.1</td>
<td>100.8±17.3</td>
<td>0.86</td>
</tr>
<tr>
<td>History of arterial hypertension</td>
<td>46.2</td>
<td>73.6</td>
<td>0.002</td>
</tr>
<tr>
<td>Diabetes mellitus</td>
<td>25.6</td>
<td>40.9</td>
<td>0.06</td>
</tr>
<tr>
<td>Hyperlipidemia</td>
<td>33.3</td>
<td>33.6</td>
<td>0.56</td>
</tr>
<tr>
<td>Peripheral arterial disease</td>
<td>7.7</td>
<td>6.4</td>
<td>0.80</td>
</tr>
<tr>
<td>Current smoking</td>
<td>53.8</td>
<td>32.4</td>
<td>0.016</td>
</tr>
<tr>
<td>Alcohol overuse</td>
<td>35.9</td>
<td>40.0</td>
<td>0.40</td>
</tr>
<tr>
<td>Serum glucose, mg/dL</td>
<td>145.7±88.2</td>
<td>136.4±58.5</td>
<td>0.46</td>
</tr>
<tr>
<td>Leukocyte count, ×10⁹</td>
<td>7942.2±2438.7</td>
<td>7912.3±2475</td>
<td>0.94</td>
</tr>
<tr>
<td>Cholesterol, mg/dL (n=118)</td>
<td>215.6±43.3</td>
<td>195.2±49.2</td>
<td>0.047</td>
</tr>
<tr>
<td>Low-density lipoprotein, mg/dL (n=94)</td>
<td>153.5±33.9</td>
<td>143.9±44</td>
<td>0.32</td>
</tr>
<tr>
<td>High-density lipoprotein, mg/dL (n=94)</td>
<td>52.8±16.3</td>
<td>47.8±16.8</td>
<td>0.19</td>
</tr>
<tr>
<td>Creatinine, mg/dL</td>
<td>0.8 [0.7–1]</td>
<td>0.9 [0.7–1]</td>
<td>0.31</td>
</tr>
<tr>
<td>Triglycerides, mg/dL (n=118)</td>
<td>156.6±107.1</td>
<td>134.6±62.1</td>
<td>0.17</td>
</tr>
<tr>
<td>Aβ₁₋₄₀ pg/mL</td>
<td>60.6 [35.6, 76.7]</td>
<td>61.3 [37.7, 64.8]</td>
<td>0.83</td>
</tr>
<tr>
<td>Aβ₁₋₄₂ pg/mL</td>
<td>40.4 [35.1, 50.5]</td>
<td>55.1 [42.3, 69.6]</td>
<td>0.0001</td>
</tr>
<tr>
<td>Aβ₁₋₄₀/Aβ₁₋₄₂ pg/mL</td>
<td>1.29 [0.89, 1.79]</td>
<td>0.92 [0.74, 1.27]</td>
<td>0.0001</td>
</tr>
<tr>
<td>Systolic blood pressure, mm Hg</td>
<td>156.7±26.2</td>
<td>159.8±27.5</td>
<td>0.55</td>
</tr>
<tr>
<td>Diastolic blood pressure, mm Hg</td>
<td>83.2±17.2</td>
<td>83.2±14.9</td>
<td>0.97</td>
</tr>
<tr>
<td>Temperature, °C (n=124)</td>
<td>36.2±0.2</td>
<td>36.1±0.5</td>
<td>0.16</td>
</tr>
<tr>
<td>Premorbid modified Rankin Scale score ≥2</td>
<td>2.6</td>
<td>3.6</td>
<td>0.043</td>
</tr>
<tr>
<td>National Institutes of Health Stroke Scale</td>
<td>3 [1, 4]</td>
<td>3 [2, 4]</td>
<td>0.86</td>
</tr>
<tr>
<td>Antiplatelet pretreatment</td>
<td>20.5%</td>
<td>34.5%</td>
<td>0.07</td>
</tr>
<tr>
<td>Statins pretreatment; (n=143)</td>
<td>16.2%</td>
<td>23.8%</td>
<td>0.23</td>
</tr>
</tbody>
</table>

Values are percentages, mean (SD) or median [quartiles] (n) no. of patients with available data.

Discussion

This study demonstrates that plasma Aβ₁₋₄₀ levels, but not Aβ₁₋₄₂, are strongly associated with the diffuse SVD subtype. The effect remained after controlling for relevant factors associated with either Aβ₁₋₄₀ or the diffuse SVD subtype such as age,²² renal function,²³ and history of arterial hypertension.²⁴
Two types of cerebral SVD have been proposed based on neuropathological studies: the first subtype shows single large lacunar infarcts caused by atherosclerosis in the larger perforating arteries, and the second is defined by multiple smaller lacunar infarcts resulting from a diffuse arteriopathy affecting the smaller perforating vessels in which the underlying pathology is lipohyalinosis usually due to hypertension. Distinct pathology of the subtypes is supported by MRI studies, which have shown that the subtype with multiple lacunar infarcts is usually associated with leukoaraiosis. This disease is characterized by white matter abnormalities that correlate with cognitive impairment. The classification used in this study is justified by reported clinical and pathological findings.

Considerable evidence suggests that endothelial dysfunction may play an important role in the diffuse SVD subtype. Two major mechanisms have been proposed: chronic hypoperfusion and increased blood–brain barrier permeability with leakage of plasma components into the vessel wall and surrounding brain parenchyma. Although we cannot rule out a confounding effect by yet unknown factors, the present results are consistent with the pathophysiological role of Aβ fraction in disrupting endothelial vascular function.

Endothelial dysfunction by Aβ deposition has been related to lacunar infarcts and leukoaraiosis. Amyloid precursor protein is cleaved by secretases to produce several peptides. Aβ(1-40) is a normal component of blood and cerebrospinal fluid, and in vitro studies have suggested direct physiological or toxic effects of Aβ(1-40) on the blood vessel wall. Interestingly, these studies in cell cultures and transgenic mouse models provide evidence that Aβ(1-40), but not Aβ(1-42), causes reactive oxygen species-mediated cerebrovascular dysfunction.

Plasma Aβ(1-40) levels were found to be markedly elevated in patients with ischemic stroke compared with age-matched control subjects in one study. Patients with cardiovascular and large artery atherosclerotic infarcts had higher Aβ(1-40) levels than patients with SVD infarctions, and levels of Aβ(1-20) correlated positively with infarct size and stroke severity. However, this study included only 12 patients classified as SVD and the presence of leukoaraiosis was not analyzed. Elevated circulating Aβ(1-40) in patients with ischemic stroke may derive from the brain as a consequence of the acute ischemic insult, but it might also be the result of chronic vascular insufficiency. In an experimental model in rodents, chronic cerebral hypoperfusion elicited the cleavage of the amyloid precursor protein into Aβ-sized fragments. Our results may reasonably rule out a secondary increase of plasma Aβ(1-40) levels after an acute lacunar stroke because levels were similar in control and focal-SVD groups. In contrast, our results support the idea that previous Aβ(1-40) levels increase as a result of diffuse-SVD disease. This, in turn, supports the pathophysiological role of fraction Aβ(1-40) in disrupting endothelial vascular function.

This study has some strengths that should be acknowledged: Stroke-mimic patients with comparable vascular risk factors but without any SVD were studied as control subjects; Aβ plasma levels were measured in an independent laboratory without knowledge of the clinical factors and neuroimaging findings; and neuroimages were also evaluated blinded to all other data. We recognize also some limitations. Aggregation and clearance of Aβ may be influenced by the apolipoprotein E polymorphism, but that was not determined in this study. However, 2 studies have reported different results. In the Rotterdam Scan Study, increased plasma Aβ levels were positively associated with lacunar infarcts and leukoaraiosis in apolipoprotein E e4 carriers. This association was not found in the Gurol et al study performed on subjects with Alzheimer disease, mild cognitive impairment, and cerebral amyloid angiopathy. Another limitation is the relatively small sample size of the focal-SVD group.

The observation that plasma Aβ(1-40) levels are strongly associated with diffuse SVD subtype is novel. Prospective studies are needed to confirm our results.

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None.

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