Elevated Plasma Level of Soluble Form of RAGE in Ischemic Stroke Patients with Dementia

Sung-Chun Tang1 · Kai-Chien Yang2 · Chaur-Jong Hu3 · Hung-Yi Chiou4 · Chau Chung Wu2 · Jiann-Shing Jeng1

Received: 5 September 2017 / Accepted: 26 October 2017 / Published online: 2 November 2017
© The Author(s) 2017. This article is an open access publication

The receptor for advanced glycation end products (RAGE) and its downstream pathways are involved in various inflammatory and immune responses. Importantly, there is soluble RAGE (sRAGE) that forms either by alternative splicing of RAGE messenger ribonucleic acid as the endogenous soluble form of RAGE (esRAGE) or by proteolytic cleavage of full-length RAGE protein. This study aimed to investigate the associations of the plasma levels of sRAGE and esRAGE in ischemic stroke (IS) patients with and without dementia. This cross-sectional study recruited patients with IS at a university medical center. Vascular dementia was defined as the scale of Clinical Dementia Ranking (CDR) ≥ 1. Standard enzyme-linked immunoabsorbent assay was used to measure the plasma concentration of sRAGE and esRAGE. From November 2014 to October 2015, a total of 172 IS patients (mean age: 72.1 ± 7.5 years, 64.5% male) were recruited, including 73 with CDR = 0, 63 with CDR = 0.5, and 36 with CDR ≥ 1. In univariate analysis, IS patients with dementia were older and had more diabetes mellitus, less atrial fibrillation, and higher post-stroke modified Rankin Scale scores than those without dementia. Plasma levels of sRAGE and esRAGE were significantly higher in IS patients with than those without dementia (1.44 ± 1.29 vs. 1.03 ± 0.48 and 0.39 ± 0.40 vs. 0.24 ± 0.13 ng/mL, both p < 0.01). Importantly, both parameters remained independent after adjustment for clinical variables (OR 2.683, p = 0.013 and OR 39.192, p = 0.006, respectively). In summary, plasma sRAGE and esRAGE were elevated in those with dementia compared with those without dementia among IS patients.

Keywords Ischemic stroke · Vascular dementia · sRAGE · esRAGE

Abbreviations
VD Vascular dementia (VD)
RAGE Receptor for advanced glycation end products
sRAGE Soluble form of RAGE
esRAGE Endogenous soluble form of RAGE
AD Alzheimer’s disease
IS Ischemic stroke
CDR Clinical Dementia Ranking
ELISA Enzyme-linked immunoassay
CAD Coronary artery disease
AF Atrial fibrillation
MRS Modified Rankin Scale
MMSE Mini–Mental State Examination
MoCA Montreal Cognitive Assessment

Introduction

Stroke is the third leading cause of death worldwide and the most frequent cause of permanent disabilities (Bhatnagar et al. 2010). Although stroke mortality has declined in recent years, the increased survival rates may translate into an increased number of cognitively impaired stroke patients (Pendlebury and Rothwell 2009). Vascular dementia (VD) is
one of the most common causes of dementia. Around 10% of stroke survivors may develop dementia within 3 months after stroke, and an additional 20% develop dementia during the subsequent 3 years (O’Brien and Thomas 2015). However, whether there are certain post-stroke pathophysiological mechanisms specifically involving the development of VD remains controversial (Thiel et al. 2014; Pugazhenthi et al. 2017).

The receptor for advanced glycation end products (RAGE) is located on the plasma membrane of many types of cells including neurons (Alexiou et al. 2010). A variety of ligands, including the advanced glycation end products (AGE), high-mobility group box 1 (HMGB1), binds to RAGE. Importantly, there is soluble RAGE (sRAGE) that forms either by alternative splicing of RAGE messenger ribonucleic acid as the endogenous soluble form of RAGE (esRAGE) or by proteolytic cleavage of full-length RAGE protein (Kim et al. 2005). Biological studies have demonstrated that sRAGE may have diverse roles reflecting the activation of cellular RAGE signaling and also functioning as a decoy to compete with membrane RAGE for ligand binding (Lotze and Tracey 2005).

Recently, RAGE has been shown to be involved in the neurodegenerative pathways in Alzheimer’s disease (AD) (Pugazhenthi et al. 2017; Gasiorowski et al. 2017). A few studies have found decreased levels of plasma sRAGE in patients with AD compared to those with VD or non-demented controls (Qian et al. 2012; Xu et al. 2017; Liang et al. 2013; Emanuele et al. 2005). However, none of these studies directly compared the difference between stroke patients with or without dementia. In our study, we measured plasma levels of sRAGE and esRAGE in patients with ischemic stroke (IS) and correlated the results with neurocognitive status.

Materials and Methods

Study Design and Population

This prospective case series cross-sectional study was conducted at the National Taiwan University Hospital (NTUH), with the approval of the Research Ethics Committee of NTUH. Written informed consent was obtained from the patients or from their next of kin without a significant cognitive impairment.

Patients with a history of IS who had been receiving regular follow-up at the Neurology Outpatient Clinic from November 2014 to October 2015 were recruited. The diagnosis of IS was confirmed and characterized by head MRI or CT examinations (performed at least 24 h after stroke onset to clearly demonstrate the infarct region). Patients with a known active infection, cancer, renal disease (creatinine > 2.0 mg/dL), autoimmune disorder, current steroid treatment, or poor diabetes control (hemoglobin A1c > 8.0%) were excluded. A detailed history of stroke, vascular risk factors, and comorbidity was obtained from each patient. Body mass index (BMI) was calculated as weight divided by the square of height.

Dementia Diagnosis and Severity Rating

Clinical diagnoses of VD were made by neurologists in accordance with internationally accepted criteria for dementia (National Institute of Neurological Disorders and Stroke and Association Internationale pour la Recherche et l’Enseignement en Neurosciences [NINDS-AIREN]) (Tierney et al. 1989; Roman et al. 1993). Dementia severity was measured by the Clinical Dementia Rating (CDR) scale. The Mini–Mental State Examination (MMSE) and Montreal Cognitive Assessment (MoCA) testing was performed as part of the clinical evaluation. Post-stroke outcome was defined by a modified Rankin Scale (mRS) score.

Blood Sampling

Human Plasma Collection and Measurements

Blood samples were drawn at the time of recruitment. A 10-mL sample of blood was drawn, then centrifuged at 300 g for 15 min, and subsequently aliquoted into 1.5-mL tubes and stored at −80 °C until ready for use. The plasma levels of total sRAGE and esRAGE were determined using a commercially available enzyme-linked immunoassay kit (RAGE: R&D Systems, Minneapolis, MN, esRAGE Human ELISA Kit, B-Bridge International, Cupertino, CA) according to the manufacturer’s protocol. Measurements were taken in duplicate, and the results were averaged. Samples with obvious hemolysis, which was visually detected by a pink to red tinge inside, were not used for measurements.

Statistical Analyses

Statistical analyses were performed using SPSS version 17.0 software (Chicago, IL). The clinical characteristics of VD and non-VD patients were compared using the Fisher’s exact test, Chi-squared test, Student’s t test, and Mann–Whitney U test with relevant variables as indicated. Multivariable logistic regression analysis models were used with VD as the dependent variable. The independent variables in the analyses were age, gender, diabetes mellitus, hypertension, atrial fibrillation, and post-stroke mRS. Figures were made using GraphPad Prism 4.0 software, and data were expressed as mean ± SEM. Comparisons between subgroups were made with Student’s t test, logistic regression, and ANOVA followed by Newman–Keuls post hoc analysis.
Results

Study Subject Demographics

During the study period, consecutive patients who were regularly followed up at the Neurology Outpatient Clinic of NTUH and fulfilled the study criteria were recruited. A total of 172 IS patients (64.5% male) were included in our study. The average time interval between stroke onset and recruitment into the study was 85.0 ± 54.0 months. The clinical information for normal and mild cognitive impairment (CDR ≤ 0.5, n = 136) and VD patients (CDR ≥ 1, n = 36) is listed in Table 1. Compared to IS patients without dementia, those with dementia were older, had more diabetes mellitus and less atrial fibrillation, lower mean scores on the MMSE and MoCA, a lower rate of functional independence, and a longer time interval between stroke onset to study recruitment (all p < 0.05).

RAGE Signaling-Related Plasma Biomarkers in Study Subjects

Figure 1 shows plasma levels of sRAGE and esRAGE in IS patients without normal cognitive function, mild cognitive impairment, and dementia. In univariate analysis, plasma levels of sRAGE and esRAGE were higher in IS patients with dementia compared to those without dementia (both p < 0.001). After adjustment for clinical parameters including age, gender, diabetes mellitus, hypertension, atrial fibrillation, interval between stroke onset to study recruitment, and post-stroke mRS, plasma levels of sRAGE and esRAGE remained significantly higher in stroke patients with dementia compared with those without dementia [odds ratio 2.65 (1.21–5.79) and 35.90 (2.65–485.53), p = 0.014 and 0.007, respectively].

Discussion

In our study, we aimed to clarify the difference between IS patients with and without dementia. We recruited patients with a history of IS, collected blood samples to measure the levels of sRAGE and esRAGE, and evaluated cognitive functional status at the same time. Our results suggested that IS patients with dementia had significantly higher levels of sRAGE and esRAGE compared to those without dementia. These findings may indicate the direct involvement of intracellular RAGE signaling in the development of VD.

A review of the previous literature revealed that levels of serum or plasma sRAGE in patients with VD had been

![Fig. 1](image-url)
investigated in only a few studies. In 2005, Emanuele et al. reported that levels of sRAGE in patients with AD (n = 152) were significantly lower than those with VD (n = 91) and controls (n = 161) (Emanuele et al. 2005). In 2012, Qian et al. recruited patients with acute stroke (n = 152) (Qian et al. 2012). Blood sample was collected at the time of admission, and cognitive function was evaluated at 2 weeks after stroke. The results showed that serum levels of sRAGE at the time of stroke onset were lower in patients with poststroke cognitive impairment than in those without poststroke cognitive impairment. In 2013, Ling et al. measured the plasma levels of sRAGE in patients with AD (n = 126), VD (n = 98), and cognitively normal controls (n = 98) (Liang et al. 2013). The results showed that levels of sRAGE were significantly lower in the group of AD compared with the group of VD patients and the controls. There was no difference in sRAGE levels between patients with VD and the controls. In 2016, Xu et al. measured sRAGE concentration in 36 patients with AD, 12 with VD, and 35 with cognitively normal controls, and the data showed that levels of sRAGE were not statistically different among these three groups (Xu et al. 2017). Overall, serum/plasma levels of sRAGE might be lower in patients with AD compared to other types of dementia and normally cognitive controls. Whether levels of sRAGE in patients with VD are lower than in non-AD subjects is uncertain. In addition, none of the studies directly compared the levels of sRAGE between stroke patients with and without dementia. Therefore, we recruited only stroke patients with and without dementia, and thus, the data of clinical and biological parameters should be able to reflect the factor of dementia after statistical analysis. It is important to mention that this had not been addressed previously in other studies. In our study, although there were some differences in clinical parameters between the IS patients with and without dementia, levels of sRAGE and esRAGE remained statistically significant after adjustment for clinical parameters.

In our previous studies, we measured the levels of sRAGE in patients with acute stroke within 48, 48–72 h, and 5–7 days after stroke onset (Tang et al. 2013, 2016). The data suggested that the level of sRAGE has a dynamic pattern after the onset of stroke from initially high to lower through the acute to subacute phases of stroke. In addition, higher levels of sRAGE in stroke within 48 h after onset are significantly associated with unfavorable functional outcome. Therefore, we proposed that circulating sRAGE in acute stroke may mainly reflect the activation of cellular RAGE rather than function as a decoy receptor. Similarly, our results of higher levels of plasma sRAGE in IS patients with dementia than in those without dementia may indicate the activation of intracellular RAGE signaling-mediated inflammatory response in the development of dementia after stroke.

Importantly, not only sRAGE levels, but also levels of esRAGE were elevated in our IS patients with dementia. The significant difference between IS patients with and without dementia was even greater than for sRAGE after multivariate analysis. Since esRAGE is produced via the alternative splicing of RAGE messenger ribonucleic acid, this finding not only supports the finding that sRAGE increased in IS patients with dementia, but also further strengthens the importance of endogenous RAGE signaling activation in the pathophysiological mechanism of VD.

Our study has several limitations. First, this is a single hospital study, and the possibility of case selection bias cannot be ignored. Second, this is a case series cross-sectional study, so only association but not causal relation between sRAGE and VD can be established from our study result. Besides, it is also possible that our recruited cases may have mixed dementia or already have cognitive impairment before the onset of stroke. Further study with a prospective cohort study design may help to clarify this issue. Third, our study is a purely clinical observation and can only present the data based on the measured values. Whether the levels of sRAGE and esRAGE are reflecting the activity of intracellular RAGE signaling, as we proposed, requires further investigation. Theoretically, it would have been informative to investigate the expression of RAGE protein in the brain between IS patients with or without dementia and correlate the results with plasma levels of sRAGE. Nevertheless, our study did demonstrate a novel phenomenon that plasma levels of sRAGE and esRAGE increased in stroke patients with VD compared to those without VD.

In conclusion, among IS patients, plasma levels of sRAGE and esRAGE were elevated in those with dementia.

Acknowledgements We thank the patients and their families for their helpful collaboration to the success of our research.

Funding No other funding sources, except Taiwan Ministry of Science and Technology Grant (105-2314-B-002 -001), involved in the study design, data collection, analysis and interpretation, or writing of this report.

Compliance with Ethical Standards

Conflict of interest The authors declare that they have no conflict of interests.

Authors’ Contributions SCT, KCY, CJH, and CCW collected patient samples. SCT, KCY, HYC, and JSJ did the data analysis. SCT and JSJ wrote the manuscript. CJH, CCW, and JSJ revised the manuscript. All authors read and approved the final manuscript.

Ethics Approval and Consent to Participate This prospective case series cross-sectional study was approved by the Research Ethics Committee of NTUH. Written informed consent was obtained from the patients or from their next of kin without a significant cognitive impairment.
Open Access  This article is distributed under the terms of the Creative Commons Attribution 4.0 International License (http://creativecommons.org/licenses/by/4.0/), which permits unrestricted use, distribution, and reproduction in any medium, provided you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made.

References

Alexiou, P., Chatzopoulou, M., Pegklidou, K., & Demopoulos, V. J. (2010). RAGE: A multi-ligand receptor unveiling novel insights in health and disease. *Current Medicinal Chemistry, 17*(21), 2232–2252.

Bhatnagar, P., Scarborough, P., Smeeton, N. C., & Allender, S. (2010). The incidence of all stroke and stroke subtype in the United Kingdom, 1985–2008: A systematic review. *BMC Public Health, 10*, 539. https://doi.org/10.1186/1471-2458-10-539.

Emanuele, E., D’Angelo, A., Tomaino, C., Binetti, G., Ghidoni, R., Politi, P., et al. (2005). Circulating levels of soluble receptor for advanced glycation end products in Alzheimer disease and vascular dementia. *Archives of Neurology, 62*(11), 1734–1736.

Gasiorowski, K., Brokos, B., Echeverria, V., Barreto, G. E., & Leszek, J. (2017). RAGE-TLR crosstalk sustains chronic inflammation in neurodegeneration. *Molecular Neurobiology*. https://doi.org/10.1007/s12035-017-0419-4.

Kim, W., Hudson, B. I., Moser, B., Guo, J., Rong, L. L., Lu, Y., et al. (2005). Receptor for advanced glycation end products and its ligands: A journey from the complications of diabetes to its pathogenesis. *Annals of the New York Academy of Sciences, 1043*, 553–561.

Liang, F., Jia, J., Wang, S., Qin, W., & Liu, G. (2013). Decreased plasma levels of soluble low density lipoprotein receptor-related protein-1 (sLRP) and the soluble form of the receptor for advanced glycation end products (sRAGE) in the clinical diagnosis of Alzheimer’s disease. *Journal of Clinical Neuroscience, 20*(3), 357–361.

Lotze, M. T., & Tracey, K. J. (2005). High-mobility group box 1 protein (HMGB1): Nuclear weapon in the immune arsenal. *Nature Reviews Immunology, 5*(4), 331–342.

O’Brien, J. T., & Thomas, A. (2015). Vascular dementia. *Lancet, 386*(10004), 1698–1706.

Pendlebury, S. T., & Rothwell, P. M. (2009). Prevalence, incidence, and factors associated with pre-stroke and post-stroke dementia: A systematic review and meta-analysis. *Lancet Neurology, 8*(11), 1006–1018.

Pugazhenthi, S., Qin, L., & Reddy, P. H. (2017). Common neurodegenerative pathways in obesity, diabetes, and Alzheimer’s disease. *Biochimica et Biophysica Acta*. 1863(5), 1037–1045.

Qian, L., Ding, L., Cheng, L., Zhu, X., Zhao, H., Jin, J., et al. (2012). Early biomarkers for post-stroke cognitive impairment. *Journal of Neurology, 259*(10), 2111–2118.

Roman, G. C., Tatamichi, T. K., Erkinjuntti, T., Cummings, J. L., Masdeu, J. C., Garcia, J. H., et al. (1993). Vascular dementia: diagnostic criteria for research studies: Report of the NINDS-AIREN International Workshop. *Neurology, 43*(2), 250–260.

Tang, S. C., Wang, Y. C., Li, Y. I., Lin, H. C., Manzanero, S., Hsieh, Y. H., et al. (2013). Functional role of soluble receptor for advanced glycation end products in stroke. *Arteriosclerosis, Thrombosis, and Vascular Biology, 33*(3), 585–594.

Tang, S. C., Yeh, S. J., Tsai, L. K., Hu, C. J., Lien, L. M., Peng, G. S., et al. (2016). Cleaved but not endogenous secretory RAGE is associated with outcome in acute ischemic stroke. *Neurology, 86*(3), 270–276.

Thiel, A., Cechetto, D. F., Heiss, W. D., Hachinski, V., & Whitehead, S. N. (2014). Amyloid burden, neuroinflammation, and links to cognitive decline after ischemic stroke. *Stroke, 45*(9), 2825–2829.

Tierney, M. C., Snow, W. G., & Fisher, R. H. (1989). Dementia diagnosis. *Neurology, 39*(11), 1559–1560.

Xu, X. Y., Deng, C. Q., Wang, J., Deng, X. J., Xiao, Q., Li, Y., et al. (2017). Plasma levels of soluble receptor for advanced glycation end products in Alzheimer’s disease. *International Journal of Neuroscience, 127*(5), 454–458.