Inhibitory effect of soluble RAGE in disturbed flow-induced atherogenesis

CHANG HOON HA1*, SUNGHYEN KIM1*, JIHWA CHUNG1, SHUNG HYEN AN1, SUNGHA PARK3, DONGHOON CHOI3 and KIHWAN KWON1,2

1Medical Research Institute, 2Department of Internal Medicine, Cardiology Division, School of Medicine, Ewha Womans University, Yangcheon-gu, Seoul 158-710; 3Division of Cardiology, Severance Cardiovascular Hospital, Yonsei University College of Medicine, Seodaemun-gu, Seoul 120-752, Republic of Korea

Received April 2, 2013; Accepted May 15, 2013

DOI: 10.3892/ijmm.2013.1393

Abstract. Soluble receptor for advanced glycation end products RAGE (sRAGE), a secretory form of RAGE, plays an important role in suppressing RAGE signals that induce pro-inflammatory gene activation in a range of inflammatory diseases, such as Alzheimer's disease, complications of diabetes mellitus and atherosclerosis. Recent studies have suggested that fluid shear stress generated by laminar blood flow protects blood vessels from atherosclerosis, whereas low and oscillatory shear stress (OSS) generated by disturbed blood flow causes atherosclerosis. Although RAGE levels are increased in atherosclerotic plaque, the regulatory mechanisms of sRAGE in the occurrence of atherosclerotic plaque induced by disturbed blood flow remain largely unknown. This study aimed to determine the effects of sRAGE as a competitive inhibitor of RAGE in atherogenesis induced by disturbed blood flow. To determine the role of sRAGE in atherosclerosis induced by disturbed blood flow, we used a mouse model of partial carotid artery ligation and showed that low shear stress and oscillatory shear stress (OSS) cause endothelial dysfunction and accelerate atherosclerosis. They suggested that disturbed blood flow directly induces the development of atherosclerosis.

Introduction

Atherosclerosis is considered to be a chronic inflammatory disease (1). It has recently been suggested that atherosclerosis primarily proceeds from disturbed blood flow regions, such as various branching points and curvatures of the arteries (2). A previous study demonstrated that blood flow with laminar shear stress (LSS) increases the expression of specific genes and proteins and protects endothelial cells (ECs) against atherosclerosis, whereas shear stress generated by disturbed blood flow generally promotes atherogenesis. These data suggest that shear stress generated by disturbed blood flow is a main cause of atherosclerosis (3).

However, the exact regulatory mechanisms through which fluid shear stress affects atherogenesis remain unclear. Recently, Nam et al developed a mouse model of disturbed blood flow by partial carotid artery ligation and showed that low shear stress and oscillatory shear stress (OSS) cause endothelial dysfunction and accelerate atherosclerosis. They suggested that disturbed blood flow directly induces the development of atherosclerosis.

Advanced glycation end products (AGEs) and their cell surface receptors have been implicated in the pathogenesis of diabetic complications. The receptor for advanced glycation end products (RAGE) is a transmembrane receptor for AGEs and other ligands, such as high mobility group box 1 (HMGB1). The binding of RAGE and these ligands activate ECs and monocytes, leading to inflammation (5). A number of studies have demonstrated that the interaction between RAGE and its ligands leads to the development and progression of atherosclerotic disease. In animal studies, a correlation...
between RAGE and atherosclerosis has been demonstrated (6). Apolipoprotein E (apoE) and RAGE (apoE<sup>-/-</sup>/RAGE<sup>−/−</sup>) double knockout mice have been shown to develop a significantly lower number of atherosclerotic lesions compared with apoE<sup>−/−</sup> mice with an intact expression of RAGE (7). Although RAGE expression is increased in atherosclerotic plaque, it is unknown whether arterial hemodynamics regulate RAGE expression and activation (5).

Soluble RAGE (sRAGE) is the soluble receptor form of RAGE lacking the intracellular domain through the proteolytic cleavage of RAGE or alternative RNA splicing in humans (8). Typically, sRAGE functions as a competitive inhibitor of RAGE by interacting with AGEs and other ligands, such as HMGB1, resulting in the inhibition of RAGE-induced cellular signaling, tissue damage and dysfunction (9). The administration of sRAGE to diabetic ApoE<sup>−/−</sup> mice has been shown to repress atherogenesis, as well as to stabilize established atherosclerosis (6,10,11). In a model of carotid arterial injury, the interruption of RAGE-ligand interaction by treatment with sRAGE has been shown to result in the reduced proliferation of smooth muscle cells and neointimal formation in both diabetic and non-diabetic rats (12).

The regulatory mechanisms of sRAGE in the occurrence of atherosclerotic plaque induced by disturbed blood flow remain largely unknown. In this study, we demonstrate that RAGE expression is directly modulated by fluid shear stress and that sRAGE inhibits RAGE-induced pro-inflammatory responses induced by OSS. Therefore, sRAGE, as a competent inhibitor of RAGE plays a protective role during the development of atherosclerosis originating from disturbed blood flow.

Materials and methods

**Cell culture and blood flow experiments.** Human umbilical vein endothelial cells (HUVECs; Gibco, Paisley, Scotland, UK) were grown in Medium 200 with 5% fetal bovine serum and low-serum growth supplement (LSGS; Cascade Biologics Inc., Portland, OR, USA) (13). U937 cells were purchased from the Korean Cell Line Bank (KCLB). Cells were cultured in RPMI-1640 (Gibco, Scotland, UK) with 10% fetal bovine serum (Gibco) and 1% penicillin-streptomycin (Corning Cellgro, Manassas, VA, USA) at 37°C and 5% CO<sub>2</sub>.

Confluent HUVECs cultured in 60-mm dishes were exposed to fluid shear stress. Cells were exposed to flow in a cone and plate viscometer. We exposed a unidirectional steady flow (shear stress of 15 dyne/cm<sup>2</sup>) for LSS and a bidirectional disturbed flow (shear stress of ±5 dyne/cm<sup>²</sup>) for OSS as previously described (14).

**Model of partial carotid artery ligation.** This animal study was performed in accordance with the Guidelines for Animal Experiments of the Animal Experimentation Ethics Committee of Ewha Womans University, Seoul, Korea. To investigate the function of sRAGE in blood vessels under disturbed flow conditions, we generated a mouse model of disturbed flow by partial ligation of the carotid artery as previously described (4). Male mice were ligated at 6 weeks of age. ApoE<sup>−/−</sup> and C57BL/6 mice were purchased from the Central Animal Laboratory, Inc., Seoul, Korea. Partial carotid artery ligated mice are viable and show acutely induced disturbed blood flow with low shear stress and OSS, resulting in endothelial dysfunction and atherosclerosis. All mice were fed a chow diet and provided with water ad libitum until partial ligation. Partial ligation of the left carotid artery (LCA) was carried out as described in a previous study (4).

ApoE<sup>−/−</sup> mice were treated with sRAGE (80 µg/kg; A&RT Co., Korea) or PBS through intraperitoneal injection daily for 2 weeks and subsequently sacrificed. C57BL/6 mice were sacrificed at 1, 3 and 7 days following partial carotid artery ligation.

**Immunoﬂuorescence staining.** The mice were sacrificed and perfused with saline containing heparin (10 U/ml) through the left ventricle. The LCA and right carotid artery (RCA) were collected en bloc with the trachea and esophagus. For the cryosections, tissue was embedded in Tissue-Tek optimum cutting temperature (OCT) medium, frozen on dry ice and stored at -80°C until use (15). Cross cryosections (4 µm) were air-dried for 1 h at room temperature. The tissue sections were then placed in 10% formalin for 10 min at room temperature for fixation. They sections were then washed in PBS for 5 min to remove the OCT compound and then blocked with 10% normal donkey serum (Santa Cruz Biotechnology, Inc., Santa Cruz, CA, USA) in PBS for 2 h. The tissue sections were then stained with rabbit anti-RAGE (1:50; Abcam, Cambridge, MA, USA) or rabbit anti-CD31 (1:50; Abcam) overnight at 4°C. The samples were then incubated for 2 h in the dark with anti-rabbit secondary antibody (1:200; Santa Cruz Biotechnology, Inc.). Nuclei were counterstained with DAPI (100 ng/ml; Santa Cruz Biotechnology, Inc.) for 8 min in the dark. The tissue sections were then routinely stained with hematoxylin and eosin (H&E) (16). The slides were viewed on an Olympus BX51 microscope (Olympus America Inc., Melville, NY, USA).

**Immunohistochemical staining.** Cryosections were fixed in 10% formalin for 10 min and blocked with peroxide block (Cell Marque, Rocklin, CA, USA) for 10 min at room temperature. Samples were incubated with rabbit anti-RAGE antibodies (1:50, 1:100; Abcam) or rabbit anti-CD31 (1:50; Abcam) for 1 h at room temperature. To visualize primary antibodies, the Polink-2 Plus HRP Detection kit (Golden Bridge International, Inc., Mukilteo, WA, USA) was used. Nuclei were counterstained with Mayer's hematoxylin (Merck KGaA, Darmstadt, Germany) (6). Oil red O staining was carried out with using frozen sections as previously described (17).

**Monocyte adhesion assay.** HUVECs were treated with sRAGE (1 µg/ml) followed by exposure to static, laminar (15 dyne/cm<sup>²</sup>) and oscillatory flow (±5 dyne/cm<sup>²</sup>) for 24 h. For the positive controls, cells were treated with 10 ng/ml tumor necrosis factor-α (TNF-α, 10 ng/ml) for 6 h. The medium was then removed, and U937 cells were added to the dishes and incubated for 30 min at 37°C. The unbound cells in the dishes were then removed by washing 3 times with serum-free medium. The adherent cells were counted in 5 randomly selected optical fields in each well. Phase-contrast microphotographs of the cells in the plates were taken using an Olympus CKX41 inverted microscope (Olympus America Inc.).

**RNA isolation, reverse-transcription polymerase chain reaction (PCR).** Total RNA was isolated from the cultured
HUVECs by the use of a Total RNA Isolation kit (Qiagen Inc., Hilden, Germany) (13). First-strand cDNA was synthesized using the Super Script III first-strand synthesis system (Invitrogen, Carlsbad, CA, USA). cDNA was amplified by PCR for 30 cycles (Eppendorf AG, Hamburg, Germany). The following oligonucleotide primers were used in this study: human RAGE sense, 5’-AGCGGCTGAATGGAAACTGAAACA-3’ and antisense, 5’-GAAGGGCAAGGGCACAACCATC-3’; human HMGB1 sense, 5’-GCGACTCTGTGCGTCGCTGA-3’ and antisense, 5’-ACGGGCCTTGTCCGCTTTCGTC-3’; human VCAM-1 sense, 5’-GGCCTCAGTCGTGGTGA-3’ and antisense, 5’-AACCCCATTCAGCGCTCA-3’. The experiments were repeated 3 times.

Statistical analysis. All data are expressed as the means ± SEM from least 3 independent experiments with different sample sizes. A paired t-test was used to assess the significance of the results between 2 groups. Differences between 3 or more groups were analyzed by contrast analysis, using Super ANOVA. A p-value <0.05 was considered to indicate a statistically significant difference.

Results

**sRAGE represses RAGE-dependent inflammatory effects under OSS conditions.** To determine whether sRAGE is involved in the RAGE-dependent pathway in ECs, we treated HUVECs with sRAGE (1 µg/ml of EC medium) and subsequently exposed the cells to LSS (fluid shear stress, 15 dyne/cm²) or OSS (shea stress, ±5 dyne/cm²) for 24 h.

The cell lysates were collected, and the mRNA expression of RAGE was analyzed by RT-PCR. As shown in Fig. 1A, sRAGE significantly inhibited the OSS-induced expression of RAGE in the HUVECs. We also observed that sRAGE reduced the mRNA expression level of HMGB1, a ligand of RAGE. These results suggest that sRAGE plays an important role in the RAGE-dependent pathway as a competent inhibitor of RAGE under OSS conditions.

LSS plays a critical role in enhancing the survival of ECs and regulating vascular tone; it also exerts anti-inflammatory and anti-atherosclerotic effects (18-22). Previous studies have shown that OSS generally upregulates genes with pro-inflammatory and atherogenic properties (22-25). These effects occur through the LSS-dependent induction of certain genes, such as Krüppel-like factor -2 (KLF2) (26-29). KLF-2 is induced by LSS, which in turn regulates a number of flow-responsive genes (26,27,30). Furthermore, KLF2 has been shown to regulate leukocyte adhesion to the endothelium by downregulating the expression of adhesion molecules that recruit leukocytes, such as vascular cell adhesion molecule-1 (VCAM-1) (26,31,32). We used RT-PCR to analyze the expression of KLF2 and VCAM-1. The mRNA expression level of KLF2 was increased by LSS; however, it was significantly reduced by OSS. However, sRAGE significantly upregulated KLF2 expression in HUVECs under OSS (Fig. 1B). By contrast, when the cells were treated with sRAGE, the OSS-induced VCAM-1 expression was markedly attenuated (Fig. 1B). Takeb together, these data demonstrate that sRAGE inhibits RAGE-dependent inflammation induced by OSS.
expression of VCAM-1, which regulates the recruitment of monocytes to the ECs (26). To determine the potential role of sRAGE in repressing the OSS-induced pro-inflammatory effects, we examined monocyte adhesion to HUVECs under static, LSS and OSS conditions with or without treatment with sRAGE (Fig. 2). ECs challenged with the pro-inflammatory cytokine, TNF-α, were used as the positive controls. As shown in Fig. 2, the TNF-α and OSS-stimulated cells exhibited augmented monocyte adhesion. By contrast, the sRAGE-treated ECs displayed significantly less monocyte adhesion even under static conditions with TNF-α treatment and OSS. Taken together, these results indicate that sRAGE regulates monocyte-EC adhesion, stimulated by OSS.

Partial carotid artery ligated mice have enhanced RAGE expression in regions of disturbed blood flow. To directly examine variations in RAGE expression under OSS conditions in vivo, we used a mouse model of disturbed blood flow with low OSS induced by partial carotid artery ligation (4).

As shown in Fig. 3, RAGE expression was markedly increased in the partially ligated LCA, whereas there was a significantly lower expression level of RAGE in the non-ligated RCA (Fig. 3A). To confirm whether shear stress modulates the expression of RAGE in ECs, we performed immunohistochemical staining on tissues from partially ligated C57BL/6 mice (Fig. 3B). The endothelial layer was stained with CD31 as an EC marker. The levels of RAGE expression were significantly enhanced in the partially ligated LCA compared with the non-ligated RCA. This suggests that shear stress modulates the expression of RAGE in the vasculature. These findings are consistent with those from previous studies, showing that RAGE transcript levels are highly expressed at the site of disturbed flow compared to the site of normal blood flow in the arteries of swine (33). In our study, the expression of RAGE was increased in the LCA even as early as 1 day following ligation (Fig. 3C). On day 7, RAGE expression was markedly upregulated in the LCA compared with the RCA.

sRAGE reduces atherosclerotic plaque formation in vivo. To explore the role of sRAGE in disturbed flow-induced atherosclerotic plaque formation in vivo, we used a mouse model of partial carotid artery ligation using ApoE⁻/⁻ mice. The mice were fed a high-fat diet for 2 weeks following surgery. sRAGE (80 µg/kg) or PBS was injected into the intraperitoneal space, daily for 2 weeks. As shown in Fig. 4, the LCA of the control group showed severe atherosclerotic plaque formation as determined by Oil red O staining (Fig. 4A). However, the LCA of the sRAGE-treated group showed significantly reduced plaque formation (Fig. 4B). To better understand the effect of sRAGE
in disturbed flow-induced atherosclerotic plaque formation, the severity of stenosis (Fig. 4C) and plaque area (Fig. 4D) were compared between the control and sRAGE-treated group. The non-ligated RCA in both groups did not show any evidence of luminal stenosis and atherosclerotic plaque formation. In contrast to the RCA, the ligated LCA of the control group
showed severe stenosis and a significantly increased plaque area, while the sRAGE-treated group showed significantly reduced luminal stenosis and plaque area. These results suggest that sRAGE inhibits the development of atherosclerotic plaque induced by disturbed blood flow.

**Discussion**

The novel finding of the present study is that sRAGE plays a critical role as an inhibitor of atheromatous plaque formation induced by disturbed blood flow. First, sRAGE repressed the pro-inflammatory response induced by OSS through the downregulation of RAGE, HMGB1 and VCAM-1 expression. According to a previous study, HMGB1, the specific ligand for RAGE, is upregulated at sites of activated ECs during vascular inflammation (9). In the current study, the expression of HMGB1 under OSS conditions was significantly inhibited following treatment with sRAGE. Monocyte binding was also inhibited following treatment with sRAGE through the decreased expression of VCAM-1. RAGE expression itself was decreased by sRAGE under OSS conditions. Therefore, these findings suggest that sRAGE may be an important inhibitor of vascular inflammatory responses generated by disturbed blood flow.

Furthermore, the expression of KLF2, an anti-inflammatory factor, was induced following treatment with sRAGE under OSS conditions. This result indicates that sRAGE induces the expression of atheroprotective factors which protect cells in a pro-inflammatory environment, such as during the early atherosclerotic process.

Second, we demonstrated that sRAGE suppresses OSS-induced inflammatory responses, such as monocyte-EC adhesion in *in vitro* experiments, as well as the development of atherosclerotic plaque in partially ligated carotid arteries of ApoE−/− mouse. We showed that RAGE expression was enhanced in the LCA, particularly in the endothelium of the lesion of disturbed blood flow, and that the administration of sRAGE suppressed atheromatous plaque formation in the partially ligated carotid arteries of ApoE−/− mice. These findings are consistent with those from previous studies, showing that the treatment of ApoE−/− mice with sRAGE reduced the development and progression of atherosclerosis in a dose-dependent manner and that the progression of atherosclerosis was halted by the administration of sRAGE in diabetic ApoE−/− mice (5,6). However, these previous studies did not determine the correlation between disturbed blood flow, RAGE expression, sRAGE and atheromatous plaque formation. By comparison, in our study, RAGE expression was increased by disturbed blood flow and it was suppressed following treatment with sRAGE. Furthermore, we showed that sRAGE suppressed atheromatous plaque formation induced by disturbed blood flow. Blood vessels, particularly ECs are consistently exposed to fluid shear stress, the dragging force generated by blood flow. Certain studies have suggested...
that fluid shear stress modulates vascular homeostasis and the focal distribution of atherosclerosis by regulating endothelial gene expression (19,34). Investigating the regulation of blood flow-mediated genes is therefore mandatory in order to gain a better understanding of atheroprotection.

To our knowledge, this is the first study demonstrating that sRAGE regulates the process of atherogenesis induced by disturbed blood flow. Previous studies using static-cultured ECs have demonstrated that inflammatory signaling is down-regulated by sRAGE through the blockade of RAGE (35,36). However, in this study, we demonstrate that fluid shear stress itself regulates the expression of RAGE and its ligand, HMGB1. Another major finding of this study was that sRAGE plays a crucial role in the regulation of the RAGE-dependent pathway involving the expression of pro-inflammatory and anti-inflammatory genes (VCAM-1 and KLF2) even in the absence of additional RAGE ligands, such as HMGB1 and certain S100 family members. This study unveils a novel role of sRAGE in regulating atherogenesis in response to hemodynamic forces, and thus provides the foundation for novel approaches for the treatment of atherosclerosis.

Acknowledgements

The present study was supported by the Bio and Medical Technology Development Program of the National Research Foundation (NRF) funded by the Korean government (MEST) (no. 2011-0019695).

References