Inhibitory Effect of S100A8 on Neutrophil Apoptosis by Cytokine Release of Normal and Allergic Monocytes

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S100A8 functions as an essential factor in inflammatory response. Cytokine release of monocytes and regulation of neutrophil apoptosis are important steps in pathogenesis of allergy. This study aims to examine the relation between cytokine release of monocytes due to S100A8 and neutrophil apoptosis. S100A8 enhanced the release of IL-6 and IL-8 in monocytes of normal and allergic subjects. Treatment of supernatants of normal and allergic monocytes with S100A8 blocked neutrophil apoptosis by inhibition of caspase 9 and caspase 3 activation. The secretion signal induced by S100A8 is involved in TLR4, Src family protein, PKC δ , ERK1/2, p38 MAPK, JNK, and NF- κ B. These findings may contribute to understanding the complex pathogenesis of allergic diseases by determining inflammatory responses associated with S100A8, monocytes, and neutrophils.

Key Words: Allergy, Monocytes, Neutrophils apoptosis, S100

S100A8, known as MRP8, is included in the S100 protein family and expressed in various cell types, including neutrophils, monocytes, and myeloid dendritic cells (Goyette and Geczy, 2011). Intracellular S100A8 functions as a scavenger of reactive oxygen species (ROS) and telomerase activity after intracellular calcium binding. Extracellular S100A8 regulates β 2 integrin-dependent neutrophil chemotaxis and expression of IL-1 β and TNF- α in Toll-like receptor (TLR4)-dependency (Donato et al., 2013). In monocytes, S100A8 induces cell migration by microtubule reassembly (Vogl et al., 2004). A recent study reported on the activation of monocytes due to S100A8 in TLR4-dependency using transcriptome evaluation (Fassl et al., 2015). Pathogenesis of allergic diseases is elicited by a variety of causes including gene mutation, toxic environment, and immune dysregulation (Holgate, 2008). Cytokine secretion and inhibition of neutro-

phil apoptosis increase inflammatory responses, resulting in exacerbation of allergy, and association of S100A8 and allergic diseases such as asthma and atopic dermatitis has been reported (Kim et al., 2014). We have, for the first time, demonstrated that S100A8 secreted by house dust mite directly inhibits neutrophil apoptosis in normal and asthmatic subjects (Kim et al., 2013).

A total of 10 allergy patients including 5 allergic asthma and 5 allergic rhinitis subjects (average = 31.6 years) were recruited from Eulji University Hospital. Allergic status was based on the presence of positive results of a skin prick test ($\geq 2+$), multiple allergen simultaneous test (MAST) ($\geq$ clas2), or measurement of specific HDM IgE using the Pharmacia Unicap 100 system to common allergens. Levels of total IgE of normal and allergic subjects using an ADVIA Centaur immunoassay (Siemens Medical Solutions Diag-

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Fig. 1. S100A8 induces IL-6, and IL-8 and cytokine secretion suppresses neutrophil apoptosis via inhibition of the caspase 9/3 pathway. (A) Monocytes isolated from normal and allergic subjects were incubated in the absence or presence of 1 μ g/mL S100A8 for the indicated time. (B) Monocytes were incubated with and without 1 μ g/mL of S100A8 for 48 h. The supernatant (Sup) was collected and added to the fresh neutrophils isolated from normal and allergic subjects. Neutrophils apoptosis was analyzed by measuring the binding of annexin V-FITC and PI. Data are presented relative to the control, which was set at 100% as the means $\pm$ S.E.M. ** P <0.01 indicates a significant difference between the control and S100A8-treated groups or between the control supernatant and supernatant-treated groups. (C) Normal neutrophils were incubated with the supernatant (Sup) or the S100A8-treated supernatant of normal monocytes. Caspase 9 and caspase 3 were detected by Western blotting. ERK2 expression was used as an internal control.

nostics, Erfurt, Germany) were 68.5 IU/mL and 565.3 IU/mL, respectively. Additionally, 8 normal subjects between 18 and 30 years of age (average = 23.0 years) were recruited as controls. This study was approved by the Institutional Review Board of Eulji University for normal volunteers allergic subjects (EU14-33). All participants in this study gave their written informed consent. Human monocytes and neutrophils were isolated from the heparinized peripheral blood of healthy persons and allergic subjects using Ficoll-Hypaque gradient centrifugation (Kim et al., 2014). Both a CD16 microbeads magnetic cell sorting kit and monocyte isolation kit II (Miltenyi Biotec, Bergisch Gladbach, Germany) were used for isolation of neutrophil and monocyte, respectively. The cells were washed after hypotonic lysis to remove erythrocytes. This method routinely yielded greater than 97% purity of neutrophil and monocyte, respectively. An annexin V-fluorescein isothiocyanate (FITC) apoptosis detection kit (BD Biosciences, San Diego, CA, USA) was used for detection of neutrophil apoptosis. Activation of caspase 9 and 3, and were confirmed by Western blotting. The concentrations of IL-6 and IL-8 in a cell supernatant after treatment with S100A8 for 48 h were measured with a sandwich enzyme-linked immunosorbent assay (ELISA) using OptEIA™ Set

human IL-6 and IL-8 (BD Biosciences, San Diego, CA, USA). The DNA-binding activity of NF- κ B was evaluated using EZ-Detect™ transcription factor kits for NF- κ B p65 (PIERCE, Rockford, IL) as described in the manufacturer's instructions. DNA binding specificity was assessed using wild type or mutant NF- κ B oligonucleotides. Chemiluminescent detection was performed using a luminometer. Here, we hypothesized that S100A8 induces the secretion of cytokines associated with neutrophil survival such as IL-6 and IL-8 in monocytes, and the cytokine release regulates constitutive apoptosis of neutrophils. To determine whether or not the effect of S100A8 on cytokine release occurs in allergic diseases, we evaluated the change of secretion of IL-6 and IL-8 due to S100A8 in monocytes separated from normal and allergic subjects. S100A8 induced the elevation of IL-6 and IL-8 in normal and allergic monocytes compared with control (Fig. 1A). Supernatant treated with S100A8 inhibited apoptosis of neutrophils isolated from normal peripheral blood, while control supernatant without S100A8 treatment did not affect neutrophil apoptosis (Fig. 1B). Supernatant treated with S100A8 suppressed allergic neutrophil apoptosis. Control supernatant of allergic monocytes without S100A8 treatment also inhibited the apoptosis of allergic

neutrophils in contrast to the normal result. Activation of caspase 9 and caspase 3 was inhibited by S100A8 in a time-dependent manner (Fig. 1C). TLR4i, an inhibitor of TLR4, PP2, an inhibitor of Src family protein, rottlerin, an inhibitor of PKC δ , PD98059, an inhibitor of ERK, SB202190, an inhibitor of p38 MAPK, SP600125, an inhibitor of JNK and BAY-11-7085, an inhibitor of NF- κ B significantly blocked the increased expression of IL-6 and IL-8 ($P<0.05$) (Fig. 2A). Activation of NF- κ B induced by S100A8 was significantly

inhibited by TLR4i, PP2, rottlerin, PD98059, SB202190, and SP600125 ($P<0.05$) (Fig. 2B). S100A8 functions as a pathological factor in induction of allergic diseases, autoimmune diseases, and malignant tumors (Jin et al., 2014; Kang et al., 2015). However, a few research groups have demonstrated that S100A8 plays a role as a protector in pathogenesis of allergy. S100A8 reduces mast cell degranulation and eosinophil chemotaxis in ovalbumin-sensitized mice, and the S100A8 concentration of induced sputum in

Fig. 2. Cytokine secretion enhanced by S100A8 is involved in TLR4, Src family protein, PKC δ , MAPKs, and NF- κ B in normal and allergic monocytes. (A) Normal (upper panel) and allergic monocytes (lower panel) were pre-treated for 1 h with and without 2 μ M TLR4i, 20 μ M PP2, 5 μ M rottlerin, 20 μ M PD98059 (PD), 20 μ M SB202190 (SB), 20 μ M SP600125 (SP) and 10 μ M BAY-11-7085 (BAY), after which the cells were incubated for 48 h in the absence and presence of S100A8 (1 μ g/mL). The supernatant was collected and analyzed by ELISA. (B) Normal monocytes were pre-treated for 1 h with and without 2 μ M TLR4i, 20 μ M PP2, 5 μ M rottlerin, 20 μ M PD98059 (PD), 20 μ M SB202190 (SB) and 20 μ M SP600125 (SP), after which the cells were incubated with S100A8 (1 μ g/mL) for 1 h. After harvested cells were lysed, NF- κ B in the lysates was detected by luciferase assay. Data are expressed as the means $\pm$ S.E.M. * $P<0.05$ and ** $P<0.01$ indicate a significant difference between the control and S100A8-treated groups or between the S100A8-treated and inhibitor-treated groups.

asthma patients was lower than in normal subjects (Zhao et al., 2011). Our results showed that S100A8 induced the expression of IL-6 and IL-8 in normal and allergic monocytes (Fig. 1A). Cytokines secreted by S100A8 have an anti-apoptotic effect on neutrophil apoptosis, which is mediated by inhibition of caspase 9 and caspase 3 (Figs. 2A and B). IL-6 and IL-8 are neutrophil survival factors and pleiotropic cytokines, which affect transition of allergic phase, activation of stromal cells, and chemotaxis (Kim and Lee, 2017). Our study supports that S100A8 promotes a pro-inflammatory effect including cytokine release and inhibition of neutrophil apoptosis in allergic diseases. These results match with our previous data related to S100A9, which binds to TLR4 (Lee et al., 2016). Apoptosis of allergic neutrophils is inhibited by control supernatant, but not that of normal neutrophils. These results indicate that response of neutrophils to factors secreted from monocytes differs between normal and allergic monocytes. Because the number of normal and allergic subjects is limited in this study, we will proceed in our study with an increased number of experimental subjects and also examine the effects of S100A8 depending on allergen-specific subjects such as existence of total IgE and house dust mite-specific IgE.

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CONFLICT OF INTEREST

The authors have no conflicts of interest, financial or otherwise, to declare.

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