S100A8 Induces Secretion of MCP-1, IL-6, and IL-8 via TLR4 in Jurkat T Cells

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In the pathogenesis of inflammatory diseases such as allergies, S100A8 acts as an important molecule and T lymphocytes are essential cytokine-releasing cells. In this study, we investigated the effect of S100A8 on release of cytokines, specifically MCP-1, IL-6, and IL-8 in T cells, and its associated signaling mechanism. S100A8 increased secretion of MCP-1, IL-6, and IL-8 in a time- and dose-dependent manner. Elevated secretion of MCP-1, IL-6, and IL-8 due to S100A8 was inhibited by the TLR4 inhibitor TLR4i, the PI3K inhibitor LY294002, the PKCδ inhibitor rottlerin, the ERK inhibitor PD98059, the p38 MAPK inhibitor SB202190, the JNK inhibitor SP600125, and the NF-κB inhibitor BAY-11-7085. S100A8 induced phosphorylation of ERK, p38 MAPK, and JNK in a time-dependent manner, and activation was suppressed by TLR4i, LY294002, and rottlerin. S100A8 induced NF-κB activation by Iκ-Bα degradation, and NF-κB activity was suppressed by PD98059, SB202190, and SP600125. These results indicate that S100A8 induces cytokine release via TLR4. Study of PI3K, PKCδ, MAPKs, and NF-κB will contribute to elucidation of the S100A8-involved mechanism.

Key Words: S100A8, Lymphocytes, Cytokine, TLR4

INTRODUCTION

S100A8, a calcium-binding protein, is included in the S100 protein family and induces a variety of physiological and pathological responses (Donato et al., 2013; Kim et al., 2013). S100A8 is important in embryo development and is associated with myeloid differentiation. S100A8 also inhibits production of reactive oxygen species (ROS) and stabilizes generation of nitric oxide. Dysregulation of S100A8 is an essential step in the outbreak and aggravation of allergic diseases, autoimmune diseases, and malignant tumors (Jin et al., 2014; Kang et al., 2015; Zheng et al., 2015). Receptors binding to S100A8 are known as both TLR4 and RAGE. In particular, TLR4 functions as an essential receptor in S100A8-induced inflammation (Pruenster et al., 2015). T Lymphocytes regulate both innate and adaptive immune responses by secreting a variety of cytokines, including IL-6,
IL-8/CXCL8, and MCP-1/CCL2 (Tsai et al., 2011; Kang et al., 2014; Kim et al., 2015). Overexpression of these cytokines induces excess migration, proliferation, and differentiation, resulting in inflammation (Kang et al., 2015). We have recently reported that the house dust mite increases expression of IL-6, IL-8, and MCP-1 in normal and allergic lymphocytes, which results in inhibition of neutrophil apoptosis (Lee et al., 2016).

MATERIALS AND METHODS

Reagents

RPMI 1640 and fetal bovine serum (FBS) were purchased from Life Technologies Inc. (Gaithersburg, MD). CLI-095, an inhibitor of Toll-like receptor (TLR) 4 (TLR4i), was purchased from Invivogen (San Diego, CA, USA). PI3K inhibitor (Ly294002), PKCδ inhibitor (rotterlin), MEK inhibitor (PD98059), p38 MAPK inhibitor (SB202190), JNK inhibitor (SP600125), and NF-κB inhibitor (BAY-11-7085) were purchased from Calbiochem (San Diego, CA, USA). Antibodies against phospho-ERK1/2, phospho-p38 MAPK, phospho-JNK, lκ-Bα, phospho-NF-κB, ERK1/2, p38 MAPK, JNK, NF-κB, β-actin were purchased from Santa Cruz Biotechnology (Santa Cruz, CA, USA).

Cell culture

Jurkat T cells were obtained from the American Type Culture Collection (Manassas, VA, USA) and were maintained in RPMI 1640 medium with 10% heat-inactivated fetal bovine serum (FBS), penicillin (100 U/ml), and strepto-
mycin (100 μg/ml).

**Enzyme-linked immunosorbent assay (ELISA)**

The concentrations of IL-6, IL-8, and MCP-1 in a cell supernatant were measured with a sandwich enzyme-linked immunosorbent assay (ELISA) using OptEIATM Set human IL-6, IL-8, and MCP-1 (BD Biosciences, San Diego, CA, USA) according to the manufacturer's instructions.

**Western blotting**

After being treated with arazyme, neutrophils were harvested and then lysed in 50 μL lysis buffer (20 mM HEPES, 400 mM NaCl, 1 mM EDTA, 1 mM EGTA, 25% glycerol, 1 mM dithiothreitol, 0.1 mM Na3VO4, and protease inhibitors). They were centrifuged at 12,000 g for 15 min at 4°C. The protein samples (50 μg/lane) were separated by 10% sodium dodecyl sulfate-polyacrylamide gel electrophoresis and the proteins were transferred to nitrocellulose filters. The blots were incubated with primary and secondary antibodies, and then developed using the enhanced chemiluminescence detection system (Amersham Pharmacia Biotech).

**NF-κB p65 transcription factor assay**

The DNA-binding activity of NF-κB was assessed using EZ-DetectTM transcription factor kits for NF-κB p65 (Pierce, Rockford, IL) according to the manufacturer's instructions. DNA binding specificity was assessed using wild type or mutant NF-κB oligonucleotides. Chemiluminescence detection was performed using a luminometer.

**Statistical analysis**

Data were expressed as the means ± SD. Statistical differences were analyzed using a paired *t*-test for two-group comparisons and one-way ANOVA for comparison of more than two groups. All analyses were conducted using the SPSS statistical software package (Version 10.0, Chicago, IL), and a *P* value < 0.05 was considered to indicate statistical significance.

**RESULTS AND DISCUSSION**

**S100A8 increases secretion of MCP-1, IL-6, and IL-8 in Jurkat T lymphocytes**

In this study, we used Jurkat cells as an *in vitro* model for investigating the effect of S100A8 on cytokine release in T lymphocytes. As shown in Fig. 1, S100A8 significantly increased production of MCP-1, IL-6, and IL-8 in a time and dose-dependent manner (*P < 0.05*). Both MCP-1 and IL-8 trigger inflammatory responses by inducing migration, activation, and survival of monocytes and neutrophils (Kim...
et al., 2015). IL-6 is a pleiotropic cytokine, which is involved in inflammation after stimulation with an infectious antigen and allergen (van der Poll et al., 1997; Kim et al., 2014). These results indicate that S100A8 is associated with the inflammatory response mediated by T cells.

Fig. 3. S100A8 induces MAPKs and NF-κB activation via TLR4, PI3K, PKCδ in Jurkat T cells. (A) Jurkat T cells were incubated with S100A8 (5 μg/mL) for the indicated time. (B) Jurkat T cells were pre-treated for 1 h with and without TLR4i, 10 μM Ly294002 (LY), and 5 μM rottlerin, after which the cells were incubated for 30 min in the absence and presence of S100A8 (5 μg/mL). The harvested cells were lysed and phosphorylation of ERK, p38 MAPK, and JNK in the lysates was detected by Western blotting. (C and D) Jurkat T cells were incubated with S100A8 (5 μg/mL) for the indicated time after which harvested cells were lysed, and expression of IκBα and phosphorylation of NF-κB in the lysates was detected by Western blotting. (E) Jurkat T cells were pre-treated for 1 h with and without 10 μM PD98059 (PD), 10 μM SB202190 (SB), and 10 μM SP600125 (SP), after which the cells were incubated for 2 h in the absence and presence of S100A8 (5 μg/mL). After harvested cells were lysed, NF-κB in the lysates was detected by luciferase assay. Data are presented relative to the control, which was set at 100%. *P < 0.05 indicates a significant difference between the control and S100A8-treated groups or the S100A8-treated and inhibitor-treated groups.
DP induces release of MCP-1, IL-6, and IL-8 via activation of TLR4, PI3K, PKCδ, MAPKs, and NF-κB in Jurkat T lymphocytes

Since S100A8 induces cytokine secretion, we examined the associated signaling mechanism by using various signaling inhibitors. TLR4i, LY294002, rottlerin, PD98059, SB202190, SP600125, and BAY-11-7085 significantly suppressed secretion of MCP-1, IL-6, and IL-8 induced by S100A8 despite different degrees of inhibition (\( P < 0.05 \)) (Fig. 2). These results show that the S100A8-mediated mechanism involves TLR4, PI3K, PKCδ, MAPKs, and NF-κB. As shown in Fig. 3A and B, S100A8 induced phosphorylation of ERK, p38 MAPK, and JNK, whereas activation was blocked by TLR4i, LY294002, and rottlerin, indicating that TLR4, PI3K, and PKCδ are upstream molecules of MAPK activation. S100A8 also activated NF-κB by Iκ-Bα degradation and NF-κB phosphorylation in a time-dependent manner (Fig. 3C and D). NF-κB activity due to S100A8 was inhibited by MAPKs such as PD98059, SB202190, and SP600125 (Fig. 3E). We previously reported that the house dust mite induces secretion of IL-6, IL-8, and MCP-1 via the PAR2/PI3K/Akt/ERK/NF-κB pathway (Lee et al., 2016). However, this study shows that TLR4 is associated with cytokine expression due to S100A8 in Jurkat T cells. These differences may have various causes, including separation of B and T cells and the action of different stimulators. These findings shed new light on the complex mechanism induced by S100A8.

Conflict of interest

The authors have no financial conflicts of interest.

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