



# Histamine Induced Production of Chemokine CXCL8 Through H1R/PLC and NF- $\kappa$ B Signaling Pathways in Nasal Fibroblasts

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**Background and Objectives:** Histamine has been suggested to play an important role during allergic and inflammatory reactions, affecting allergic rhinitis and chronic rhinosinusitis. CXCL8 is a pro-inflammatory chemokine and a critical factor that causes many airway inflammatory diseases including allergic rhinitis and chronic rhinosinusitis.

**Materials and Method:** Histamine cytotoxicity was measured by MTT assay. Real-time polymerase chain reaction was used to identify histamine type 1 receptor in nasal fibroblasts. The fibroblasts were then treated with histamine with or without a histamine type 1 receptor antagonist and the CXCL8 protein was assessed using an enzyme-linked immunosorbent assay (ELISA). The downstream signaling molecules, including phospholipase C and phospho-p50, were evaluated by western blot and immunofluorescent staining.

**Results:** Histamine had no significant cytotoxic effect until the concentration reached 1,000  $\mu$ M. Histamine type 1 receptor mRNA was expressed in nasal fibroblasts. CXCL8 protein expression level was significantly increased following histamine stimulation. However, the expression level of CXCL8 decreased when phospholipase C was inhibited by U73122. Histamine increased phospho-p50 expression as seen in western blot results. The BAY11-7082, NF- $\kappa$ B inhibitor significantly reduced CXCL8 production in histamine-stimulated nasal fibroblasts.

**Conclusion:** Histamine can induce the production of NF- $\kappa$ B controlled-chemokine CXCL8 by nasal fibroblasts, which supports a role for histamine in upper airway inflammatory diseases.

**KEY WORDS:** Histamine · CXCL8 · Histamine type 1 receptor · Signal pathways · Nasal fibroblasts.

## INTRODUCTION

Histamine has been recognized as a major mediator in allergic diseases such as allergic rhinitis, chronic rhinosinusitis, and nasal polyps.<sup>1)</sup> Histamine gets released by lymphocytes, neutrophils, platelets, macrophages, and eosinophils, also endothelial cells and nasal lavage fluid produces histamine released factors.<sup>2,3)</sup> This heterogeneous group of factors causing basophil degranulation may also recruit mast cell or basophil participation in late-phase allergic reactions and other chronic inflammatory conditions.<sup>4,5)</sup>

Histamine H1 receptor (H1R), originally derived from bovine, is known to be expressed in many human tissues including airway, intestinal and vascular smooth muscle and brain.<sup>6,7)</sup> Histamine-1 receptor (H1R) antagonists control clinical symptoms related to histamine expression and H1R is a primary receptor of histamine activity.<sup>8)</sup> Immunohistochemical studies have shown that strong immune responses to H1R in vascular endothelial cells increase vascular permeability, leukocyte infiltration, and edema formation.<sup>9)</sup> Nevertheless, the mechanism of this effect has not been clearly understood.

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Fibroblasts are a major structural component of tissue that provides mechanical strength by providing a fabric that supports the extracellular matrix (ECM).<sup>10</sup> In the sinonasal mucosa, fibroblasts play a major role in a structural modification in myofibroblast differentiation affecting the pathogenesis of nasal polyps.<sup>11</sup> Fibroblasts are known to produce large amounts of the ECM such as collagen and fibronectin, and they also express many receptors for cytokines, growth factors, and hormones.<sup>9</sup> They are additionally considered to be responsible for local enrollment of inflammatory cells.

CXCL8 is considered to be a major source of neutrophil chemostimulants and induces changes and directional movement, exocytosis of stored proteins, and respiratory bursts.<sup>12</sup> CXCL8 is produced by inflammatory and non-inflammatory cells such as mononuclear cells, fibroblasts and epithelial cells.<sup>13</sup> Previous studies have shown a correlation between CXCL8 and allergic diseases such as asthma.<sup>14</sup> Histamine induces CXCL8 secretion in endothelial cells and alveolar macrophages.<sup>15</sup> These data suggest that histamine may also bring on a major role in CXCL8 production in nasal fibroblasts. Nevertheless, the mechanism by which histamine induces the expression of CXCL8 in nasal fibroblasts is uncertain. To understand the pathophysiology of sinonasal diseases, it is important to understand the mechanisms influencing the nasal patency.

The aim of this study is to determine the effect of histamine on CXCL8 production and histamine receptor activity and to investigate the mechanism underlying the effect in nasal fibroblasts.

## MATERIALS AND METHODS

### Chemicals and reagents

The reagents used in this experiment included histamine, diphenhydramine (first-generation H1R antagonist), pyrillamine (first-generation H1R antagonist), fexofenadine (second-generation H1R antagonist), and U-73122 (phospholipase C inhibitor). These five reagents used were purchased from Sigma-Aldrich (St. Louis, MO, USA). BAY11-7082 (NF- $\kappa$ B inhibitor) was purchased from Calbiochem (Billerica, MA, USA). Histamine, diphenhydramine, and pyrillamine were dissolved in distilled water. Fexofenadine, BAY11-7082, and pyrillamine were used in its dimethyl sulfoxide dissolved form. The final maximum concentration of dimethyl sulfoxide was below 0.1%.

### Nasal tissues and nasal fibroblast culture

Eight normal control subjects were recruited for this experiment. All patients were recruited from the Department of Otolaryngology-Head and Neck Surgery, Korea University Medical Center. Only patients without allergies, asthma or aspirin sensitivity met inclusion criteria. Likewise, none of the participants experienced treatment with anti-allergic drugs for more than 2 months. Before obtaining nasal fibroblast cultures, informed consent was thoroughly explained and received from all participants. Normal control tissues from inferior turbinate specimens were gained by patients without any sinonasal diseases by rhinoplasty surgeries. The purity of cells was confirmed by its morphological characteristics such as spindle-shaped cell morphology and flow cytometry. Nasal fibroblasts were positive for vimentin, fibroblast markers, whereas negative for E-cadherin, epithelial cell marker. Once the cell confluency reached more than 80%, cells were enzymatically digested. The nasal fibroblasts were cultured for four passages.<sup>16</sup> The experiment was approved by the Korea University Medical Center Institutional Review Board.

### Measurement of cell viability

Nasal fibroblasts were seeded in 96-well culture plates at a density of  $10^4$  cells/well, in culture medium and incubated for 24 hours. Fibroblasts were then treated with different concentrations of histamine or H1R antagonist for another 48 hours. Then nasal fibroblasts were incubated with 3-(4, 5-dimethylthiazol-2-yl)-2, 5-diphenyltetrazolium bromide (MTT, Sigma) at a final concentration of 0.5 mg/mL. After 4 hours, MTT solution was discarded and 150  $\mu$ L dimethyl sulfoxide (Sigma-Aldrich) was added to dissolve the formazan precipitate by shaking the plates for 10 minutes at mild speed on an orbital shaker. Microplate readers (F2000, Hitachi Ltd., Tokyo, Japan) were used to read the absorbance of each well at the wavelength of 540 nm.

### Detection of histamine type 1 receptor mRNA in nasal fibroblasts

After nasal fibroblasts were stimulated by histamine (100  $\mu$ M), total RNA was extracted by a Trizol reagent (Invitrogen, Carlsbad, CA, USA). The total amount of RNA was determined by an optical system using NanoDrop 2000 (Thermo Scientific, Wilmington, DE, USA). Real-time polymerase chain reaction was performed in Quantstudio3 (Applied Biosystems, Foster City, CA, USA) using Power SYBR

Green PCR Master Mix (Applied Biosystems). Relative gene expression was calculated by evaluating real-time polymerase chain reaction data using the 2<sup>-ΔΔCt</sup> method. Experiments were repeated at least 3 times and glyceraldehyde 3-phosphate dehydrogenase (GAPDH) was used as an internal control. A primer was added to perform the polymerase chain reaction; *H1R* (sense sequence 5'-GTC TAA CAC AGG CCT GGA TT-3', antisense sequence 5'-GGA TGA AGG CTG CCA TGA TA-3') and *glyceraldehyde-3-phosphate dehydrogenase (GAPDH)* (sense sequence, 5'-GTGGATATTGTCCCAT-3' and antisense sequence, 5'-GCCCCAGCCTTCTTCATGGGG-3').

### CXCL8 measurements

The secretion levels of CXCL8 protein in nasal fibroblasts after treatment with histamine were evaluated by ELISA. Ten percentage of fetal bovine serum was used to incubate nasal fibroblasts and the supernatant media were collected. ELISA kits (R&D Systems, Minneapolis, MN, USA) were used to quantify CXCL8 expression according to manufacturer's instructions. Ninety-six-well microplates were coated with diluted capture antibody and incubated overnight at room temperature. They were blocked with 1% bovine serum albumin for 1 hour and then added collected-supernatant media and standards for 2 hours. Detection antibody was added for 2 hours after washing. Streptavidin-horseradish peroxidase and its substrate were used for the manifestation of color. Optical density was set to 450 nm. The microtiter plate reader (SpectraMax 190, Molecular Devices, Sunnyvale, CA, USA) was used to determine the CXCL8 expression and each well was examined within 30 minutes.

### Western blotting analysis

PRO-PREP™ protein extraction solution (iNtRON Biotechnology, Seongnam, Korea) was used to lyse nasal fibroblasts. The lysed materials were then placed in sodium dodecyl sulfate-polyacrylamide gel electrophoresis and were transferred to polyvinylidene difluoride membranes (Millipore Inc., Billerica, MA, USA). Membranes were blocked for an hour in a 5% non-fat dry milk and were covered overnight in phospho (p)-p50, and total-p50 with 3% bovine serum albumin at 4°C. Afterwards that, membranes were washed thoroughly for five minutes three times, and then treated with horseradish peroxidase-conjugated secondary antibodies (Vector Laboratories, Burlingame, CA, USA)

for one hour. Then, substrates from an ECL system (Pierce, Rockford, IL, USA) were added for visualization.

### Immunofluorescent staining

Nasal fibroblasts were seeded on coverslips and treated with histamine (100 μM). Nasal fibroblasts were fixed with 4% paraformaldehyde, and then permeabilized with 0.01% Triton X-100 in 1% bovine serum albumin for 10 minutes. After permeabilization, nasal fibroblast was blocked with 5% bovine serum albumin for 1 hour at room temperature. Nasal fibroblasts incubated with p-p50 (Sigma-Aldrich) overnight at 4°C and then incubated with anti-rabbit Alexa 555 secondary antibodies (Invitrogen) for 1 hour. 4'-6-diamidino-2-phenylindole was applied for counterstaining of the nucleus. An LSM700 confocal laser scanning microscope (Zeiss, Oberkochen, Germany) was used for capturing and identifying stained cells.

### Statistical data analysis

All statistical analyses were performed via SPSS Statistics 20.0.0 version (IBM Corp., Armonk, NY, USA). Statistically significant differences between groups were evaluated by one-way analysis of variation (ANOVA) for factorial comparisons and subsequently with Turkey's post-hoc trial. Outcomes are given as mean ± standard error of the mean (SEM) which were extracted from at least three independent trials. Statistical significance was defined as  $p < 0.05$ .

## RESULTS

### Effects of histamine on cytotoxicity

An MTT assay was used to examine the cytotoxicity of histamine and H1R antagonists. A cell titration curve was produced from serial dilutions of nasal fibroblasts and MTT reagent. The standard curve indicated a linear response between cell number and absorption at 570 nm. Nasal fibroblasts were examined at histamine concentrations ranging from 0 to 1,000 μM with various mixtures of histamine and H1R antagonist (fexofenadine, pyrilamine, and diphenhydramine). Cell survival was not affected by histamine until the concentration reached 1,000 μM (Fig. 1A), and it was also produced by different mixtures of histamine and each H1R antagonist for 72 hours (Fig. 1B).

### Expression of H1R mRNA and effect of histamine on CXCL8 production

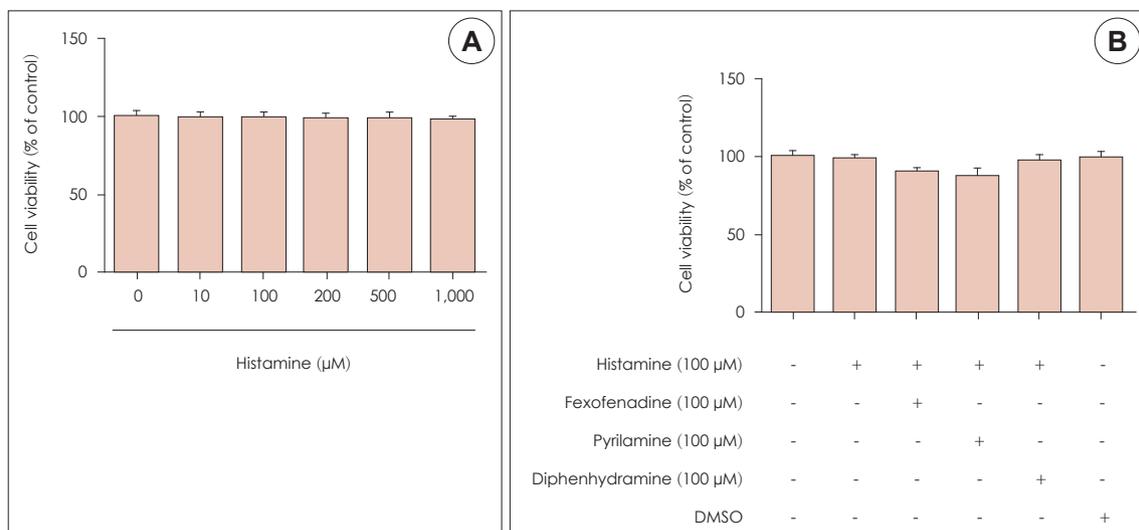
To investigate the CXCL8 expression in nasal fibroblasts, we measured CXCL8 expression using ELISA kit. H1R mRNA expression in nasal fibroblasts was identified by real-time polymerase chain reaction. H1R mRNA expression was increased significantly by histamine stimulation in a dose-dependent manner (Fig. 2A). ELISA was then used to examine whether histamine stimulated CXCL8 protein production in nasal fibroblasts. Nasal fibroblasts were treated with up to 200  $\mu$ M histamine for 24 hours. CXCL8 protein expression increased significantly in response to histamine stimulation in a dose-dependent manner (Fig. 2B).

### Effect of H1R antagonists on CXCL8 production of histamine-stimulated nasal fibroblasts

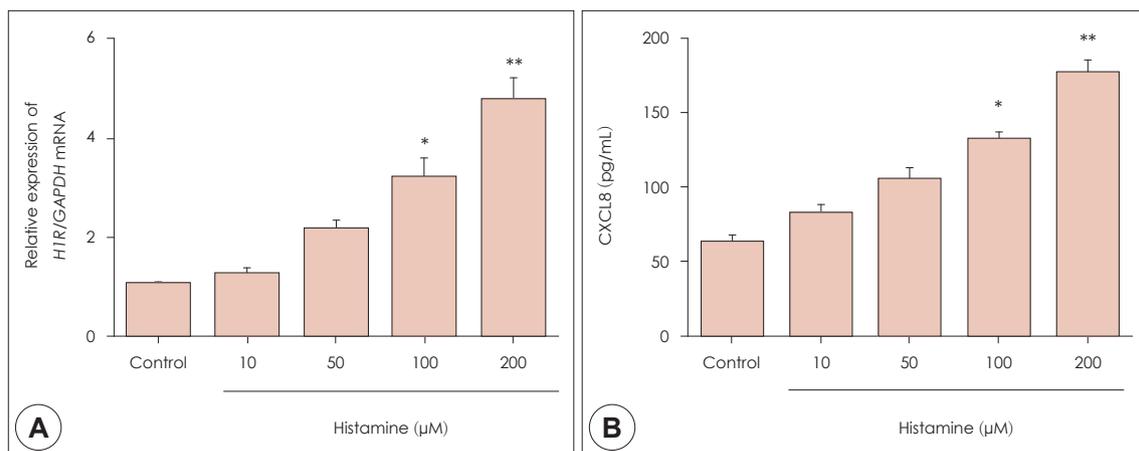
We investigated which histamine receptor was involved in the CXCL8 production. Nasal fibroblasts were pretreated with histamine receptor antagonists for H1R (fexofenadine, pyrilamine, diphenhydramine) and then stimulated by histamine for 24 hours. Fexofenadine, pyrilamine, and diphenhydramine significantly decreased CXCL8 production (Fig. 3A).

### Effect of phospholipase C on histamine-induced CXCL8 production

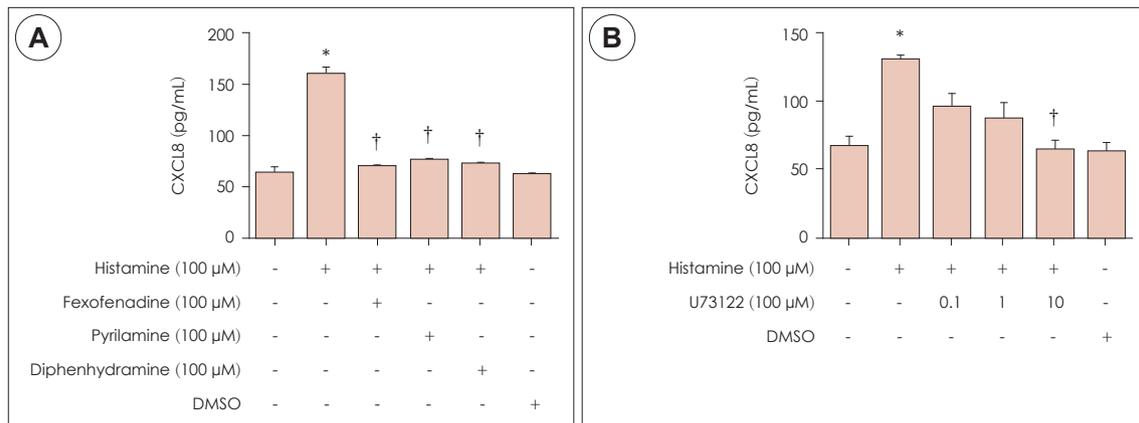
To find the signaling pathway for histamine-induced



**Fig. 1.** Effect of histamine and histamine type 1 receptor antagonists on cell viability in nasal fibroblasts. A: Cytotoxicity tests were performed using 3-(4, 5-dimethylthiazol-2-yl)-2, 5-diphenyl tetrazolium bromide (MTT) assay at various concentrations of histamine. B: MTT assays were also executed with different mixtures of histamine and each H1R antagonist (fexofenadine, pyrilamine, and diphenhydramine). DMSO: dimethyl sulfoxide.



**Fig. 2.** Histamine type 1 receptor expression and CXCL8 expression. A: The expression levels of CXCL8 protein in control and nasal polyp tissue were determined using ELISA. B: The expression level of histamine type 1 receptor mRNA was determined by real-time polymerase chain reaction. Values represent mean  $\pm$  SEM. \*:  $p < 0.05$  vs. control, \*\*:  $p < 0.01$  vs. control.



**Fig. 3.** Inhibitory effect of histamine type 1 receptor and phospholipase C antagonists on CXCL8 expression in nasal fibroblasts. A, B: The expression levels of CXCL8 protein after treatment with histamine type 1 receptor antagonists (fexofenadine, pyrilamine and diphenhydramine) and phospholipase C inhibitor (U73122) were determined via ELISA. Values represent mean  $\pm$  SEM. \*:  $p < 0.05$  vs. control, †:  $p < 0.01$  vs. histamine alone. DMSO: dimethyl sulfoxide.

CXCL8 production in nasal fibroblasts, stimulation of phospholipase C as a downstream marker of CXCL8 signaling was evaluated via ELISA. Histamine-induced CXCL8 protein production decreased in a dose-dependent manner after treatment with a phospholipase C inhibitor (U73122) (Fig. 3B). CXCL8 production was significantly decreased compared to the control after 10  $\mu$ M treatments with a phospholipase C inhibitor.

### Effect of histamine on NF- $\kappa$ B activation in histamine-induced CXCL8 production

Next, the effect of NF- $\kappa$ B activation on CXCL8 production was confirmed in histamine-stimulated nasal fibroblasts. The expression levels of p-p50 were determined by western blot in nasal fibroblasts, in the absence or presence of H1R antagonist or BAY11-7082. Histamine increased expression of p-p50 and BAY11-7082 (NF- $\kappa$ B inhibitor) suppressed p-p50 expression (Fig. 4A). In immunocytochemical staining, histamine induced phosphorylation of p50 and translocation of p-p50 into the nucleus (Fig. 4B). The expression level of CXCL8 protein was determined by ELISA. Histamine induced expression of CXCL8 protein levels. However, BAY11-7082 inhibited expression of histamine-induced CXCL8 protein levels (Fig. 4C). These results indicated that histamine-induced CXCL8 production through NF- $\kappa$ B signaling pathway.

## DISCUSSION

In the present study, we examined the effects of histamine

on CXCL8 production via phospholipase C and NF- $\kappa$ B signaling pathways in nasal fibroblasts. The results showed a clear correlation between histamine signaling pathways and CXCL8 production, which was mediated primarily through H1R. H1R antagonists inhibited CXCL8 production. The phospholipase C inhibitor, U73122, significantly reduced the effect of histamine on CXCL8 production. The NF- $\kappa$ B inhibitor (BAY11-7082) inhibited CXCL8 protein production as effectively as fexofenadine in histamine-stimulated nasal fibroblasts. These effects supported the idea that H1R inhibited the activation of phospholipase C and NF- $\kappa$ B signaling pathways. Taken together, we suggest a signal pathway of histamine-induced CXCL8 through the H1R, phospholipase C and NF- $\kappa$ B signaling pathways.

Mast cells initiate immediate hypersensitivity reactions; after an IgE-dependent stimulation, mast cells rapidly release preformed mediators such as histamine.<sup>17)</sup> Histamine plays a major role in the early clinical manifestations of allergic disorders. Mast cells reside in close proximity to potential target cells of mediator action such as fibroblasts, nerves, epithelial cells, and vascular endothelial cells. Histamine acts on target cells through distinct H1, H2, H3 and H4 receptors. Most of the effects of histamine in allergic disorders are mediated through H1 and H2 receptors.<sup>7)</sup> The production of CXCL8 is also augmented in human bronchial epithelial cells by histamine.<sup>18)</sup> As in these human epithelial cells, histamine was found to stimulate the production of IL-6 and CXCL8 in human keratinocytes in the previous study.<sup>19)</sup> Previous study reported that histamine stimulates the production of IL-6 and CXCL8, hence our



phospholipase C and NF- $\kappa$ B signaling pathways. These results suggest that antihistamines are likely to affect multiple cytokines or chemokine in addition to blocking the more common effects of histamine in allergic inflammation. To investigate the clinical use of CXCL8 pathway production, further study into other pathways and *in vivo* applications are required.

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