Original Article
Exclusion of IL-21 in the pathogenesis of OVA-induced asthma in mice

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Abstract: Asthma is characterized by airway inflammation, mucus overproduction, and airway hyperreactivity. Cytokines, especially T helper 2-derived cytokines interleukin (IL)-4, IL-5, and IL-13, are involved in the pathogenesis of allergic diseases currently controversial. The aim of this study was to investigate the role of IL-21 in asthma airway inflammation in vivo. A murine ovalbumin (OVA)-induced allergic asthma model was used. The concentration of IL-21 in the bronchoalveolar lavage fluid (BALF) of mice was evaluated by enzyme-linked immunosorbent assay. BALF cellularity, lung histopathology, and sera IgE levels were compared between the normal control group, OVA sensitization/challenge group, and OVA sensitization/challenge plus IL-21-administered group. An OVA-induced allergic rhinitis model with IL-21 was used as a positive control and the infiltration of eosinophils in the nasal mucosa was evaluated. The concentration of IL-21 in the BALF was lower in the asthmatic group compared with the normal control group. However, no significant differences in airway eosinophilia, lung histopathology, and sera IgE levels were observed between the OVA sensitization/challenge group and OVA sensitization/challenge plus IL-21-administered group. Decreased eosinophilic infiltration of nasal mucosa was observed in the positive control allergic rhinitis model administered IL-21 during the challenge period. Exogenous administration of IL-21 alone may not alleviate allergic lung inflammation. The role of IL-21 in allergic lung inflammation needs further research.

Keywords: Allergic lung inflammation, asthma, challenge, cytokine, interleukin-21

Introduction

Asthma is a common respiratory disorder characterized by airway inflammation, bronchoconstriction, and airway hyperresponsiveness. These features clinically manifest themselves as exacerbations of wheezing and breathlessness, which have significant impact on the quality of life. Airway inflammation that occurs during atopic asthma is associated with exposure to specific allergens such as house dust mite allergens or nonspecific triggers, such as air pollution [1]. Cytokines are critical in allergic intercellular communication networks, and they contribute to disease pathology through the recruitment and activation of proinflammatory leukocytes and in the chronic phase by mediating pro-fibrotic/remodeling events. A number of studies have clearly established the importance of T helper (Th) 2-derived cytokines interleukin (IL)-4, IL-5, and IL-13 in mediating the airway inflammatory response using a murine model of allergic lung inflammation [2-5]. Th2 cytokines are produced predominantly by activated Th2 cells, which induce inflammatory cell infiltration into the airways. Recently, Th17 cells, a subset of CD4+ T cells with distinct properties from Th1, Th2, and regulatory T cells [6], are found in the lungs of patients with severe asthma [7]. In addition, the Th17 cytokine IL-17A is present in the bronchoalveolar lavage fluid (BALF) and bronchial biopsies of patients with moderate to severe asthma [8]. IL-21 is a member of the type I cytokine family with significant sequence homology to IL-2, IL-4, and IL-15 [9]. IL-21 is produced by activated CD4+ T cells, and recently it has been reported that IL-21 is produced by Th17 cells and NKT cells [10]. IL-21 biological functions are mediated by a heterodimeric receptor, formed by the
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Figure 1. Protocols for sensitization, challenge, and rmIL-21 administration.
A: Murine asthma model: mice were sensitized by i.p. immunization with OVA plus alum on days 0 and 14 followed by challenge with OVA solution by intratracheal instillation on days 24, 26, and 28. Some groups of mice were treated with rmIL-21 before challenge. B: Allergic rhinitis model: mice were sensitized by i.p. administration of OVA plus alum on days 0, 7, and 14 followed by daily injections of OVA solution into the nostrils on days 21-28 (challenge). Some groups of mice were treated with rmIL-21 administration into the nasal cavity before challenge. On day 29, mice were sacrificed and samples were collected.

Materials and methods

Mice

Female BALB/c mice (6-8 weeks old) were purchased from the Laboratory Animal Center of Hubei Province China and housed under specific pathogen-free conditions in the Laboratory Animal Center of Huazhong University of Science and Technology.

The experimental procedure was approved by the committee of Huazhong University of Science and Technology for animal research.

Animal model and recombinant mouse (rm) IL-21 administration

The allergic asthma model was induced as previously described [22]. Briefly, mice were sensitized on day 0 by intraperitoneal (i.p.) injection of 20 μg OVA (Grade V; Sigma, MO, USA) absorbed in 2 mg of alum (Pierce, Rockford, IL, USA) (200 μL/mouse). On day 14, mice were sensitized a second time with 100 μg OVA. On days 24, 26, and 28, mice were anesthetized by i.p. injection of 0.1 mL of a mixture of 10 mg/mL ketamine and 1 mg/mL xylazine diluted in sterile phosphate-buffered saline (PBS) and challenged with 200 μg OVA in 40 μL of sterile PBS by intratracheal instillation. The control group received sterile PBS with alum i.p. on days 0 and 14 and 40 μL of sterile PBS on days 24, 26, and 28 by intratracheal instillation. In the IL-21 treatment group, mice were adminis-
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A nasal allergic model was used as a positive control, and mice were treated as previously described [20]. Briefly, mice were administered 100 μg OVA and 4 mg alum in saline i.p. at a dosage of 0.2 mL/mouse. The sensitization was repeated three times at weekly intervals (days 0, 7, and 14) followed by daily injections of OVA solution (40 mg/mL in saline) into the nostrils (0.02 mL/mouse) on days 21-28 (challenge). In the IL-21 treatment group, mice were administered 20 ng of rmIL-21 intranasally 30 min before challenge (Figure 1).

Bronchoalveolar lavage and lung histopathologic examination

Bronchoalveolar lavage was performed with 0.8 mL PBS three times. The lavage fluid was centrifuged, and the supernatants were kept at -80°C until used for cytokine measurements. The cell pellets were resuspended in 0.5 mL of PBS and used for total and differential cell counts. The total number of cells in BALF was counted by hemacytometer. Eosinophils, neutrophils, and macrophages were counted in BALF using cytocentrifuge preparations stained with Diff-Quik staining. A total of 200 cells in each sample were counted. Lungs were then inflated with 0.8 mL 10% formalin, fixed in 10% formalin for at least 24 h, dehydrated through a gradient ethanol, embedded in paraffin, and the sagittal sections were cut at a thickness of 5 μm. Sections were stained with hematoxylin and eosin.

Enzyme-linked immunosorbent assay

Sera were collected from mice within 24 h after the last challenge. Concentrations of IgE were evaluated using an enzyme-linked immunosorbent assay (ELISA) kit (BioLegend, San Diego, CA). Concentrations of IL-21 in the BALF were measured by ELISA kits (R & D Systems, Minneapolis, MN).

Statistical analysis

Data were expressed as mean ± SD. Statistical significance was assessed by one-way analysis of variance followed by the Tukey’s multiple comparison test or the unpaired Student’s t-test. Statistical significance was set at p < 0.05.

Results

BALF concentration of IL-21 was decreased in asthmatic mice

The concentration of IL-21 in the BALF was determined by ELISA. IL-21 levels were decreased in asthmatic mice (40.17 ± 5.919 pg/mL) compared with those in normal mice (102.7 ± 14.44 pg/mL) (Figure 2).

Intranasal IL-21 administration alleviates infiltration of eosinophils in nasal mucosa in murine allergic rhinitis

To validate the effect of IL-21 as a positive control, we performed an experiment using an allergic rhinitis model treated with IL-21 before challenge as previously described [20]. Intranasal IL-21 administration decreased eosinophilic infiltration of the nasal mucosa in an OVA-induced murine allergic rhinitis model (Figure 3).

Administration of IL-21 did not ameliorate allergic airway inflammation

Because the concentration of IL-21 in the BALF was lower in asthmatic mice and the administration of IL-21 decreased eosinophilic infiltration of nasal mucosa in allergic rhinitis, we determined whether allergic airway inflammation could be improved by administration of IL-21 before challenge. rmIL-21 was administered intratracheally before each challenge as in Figure 1A. Inflammation around the bronchus and vessels was not affected by the
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After the last challenge, mice were sacrificed and the lungs were lavaged, cells were counted and a differential cell count was performed. There was no significant change in the number of total cells, macrophages, neutrophils, eosinophils, or lymphocytes between the asthmatic group and the IL-21 administration group (Figure 5). Similarly, there were no significant changes in sera IgE levels between the groups with and without treatment of rmIL-21 before challenge in asthmatic mice (Figure 6).

Discussion

In the present study, we examined the effect of IL-21 treatment on allergic asthmatic mice. IL-21 had no effect on allergic airway inflammation in mice. During the challenge period, intratracheal administration of IL-21 may not ameliorate airway inflammation and sera IgE levels were not affected.

Figure 3. Histological examination of nasal mucosa. A: Nasal mucosa from the normal control group. B: Nasal mucosa from the OVA-induced mice allergic rhinitis model. C: Nasal mucosa from the rmIL-21-treated group. Mice were sensitized and challenged with the allergen with or without rmIL-21 administration before challenge. In rmIL-21-treated mice, the infiltration of eosinophils in the nasal mucosa was decreased significantly compared with the OVA sensitized/challenge group.

Figure 4. Administration of IL-21 before challenge does not ameliorate OVA-induced peribronchial inflammation. Lung sections were stained with hematoxylin and eosin for the measurement of inflammatory cells around the airways. The extent of cellular infiltration of the peri-airway region at 24 h after the last OVA challenge was comparable between the OVA sensitized/challenge group (B) and the IL-21-treated group (C). (A) Histologic section of lung from the normal control group.

Conflicting data on the effects of IL-21 on allergic inflammation in animal models of various diseases have been reported. IL-21 receptor deficiency inhibited allergic cutaneous inflammation by suppressing the trafficking of cutaneous dendritic cells to draining lymph nodes in a murine model of epicutaneous sensitization using tape stripping [23]. And studies by Fröhlich A et al. [24] and Lajoie S et al. [25] found that IL-21 may play roles in enhancing allergic asthma in IL-21 receptor deficient mice using OVA-induced allergic model and HDM allergen-induced allergic mouse model respectively. In contrast, IL-21 suppressed experimental allergic rhinitis by down-regulating Th2 cytokines and preventing nasal fibroblasts from producing eotaxins [20]. To validate the effect of IL-21 as a positive control in our study, we also used an allergic rhinitis model treated with IL-21 before challenge. We found decreased eosinophilic infiltration of the nasal mucosa. These results illustrate that IL-21 levels were
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The function of IL-21 on the immune system is complex. IL-21 regulates the activation, proliferation, and survival of both CD4+ T cells and B cells, the functional activity of CD8+ T cells and NK cells, limits the differentiation of inducible regulatory T cells, and counteracts their suppressive properties on effector T cells [26, 27]. IL-21 can also negatively regulate the maturation and function of dendritic cells [28, 29]. As IgE may play a role in allergic disorders, many studies have focused on the influence of IL-21 on IgE synthesis. However, these reports also remain inconsistent. Studies by Suto et al. indicated that IL-21 down-regulated IL-4-dependent IgE production from B cells but did not affect Th2 cell differentiation [30]. Caven et al. found that IL-21 inhibited IgE production at high cell concentrations, but only a modest enhancement when using low cell input in vitro culture systems [31]. IL-21 has been reported to have pleiotropic effects on B cells in terms of cell survival, proliferation, and differentiation in a context-dependent fashion [19]. Studies also revealed that IL-21 induced B-cell apoptosis when combined with prostaglandin E2, while IL-21 alone supported the viability of cultured mouse B cells [32]. In this study, we found that IL-21 had no effect on IgE production in vivo. This may because of the differential effects of IL-21 on B cells under different inflammatory conditions.

In conclusion, this study showed that although IL-21 levels were decreased in asthmatic mice, exogenous administration of IL-21 may not alleviate allergic lung inflammation. Further research may explore the function of IL-21 on various cell types in asthmatic inflammation and its association with other cytokines.

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Disclosure of conflict of interest

None.

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