Effects of microRNA-21 on the interleukin 12/signal transducer and activator of transcription 4 signaling pathway in asthmatic mice

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Abstract

Objective: To study the effect of microRNA-21 (miRNA-21) on the regulation of the interleukin 12 (IL-12)/signal transducer and activator of transcription 4 (STAT4) pathway in the lung tissue of asthmatic mice.

Material and methods: Forty five male C57BL/6 mice were randomly divided into three groups of 15 mice each: normal control, asthmatic model, and dexamethasone. Our mouse model of allergic asthma was established using OVA sensitization and challenge. Hematoxylin and eosin staining was performed to observe the pathological changes in lung tissue morphology. Both the total cell number and the amount of eosinophils (EOS) in the bronchoalveolar lavage fluid (BALF) were manually counted. The expression of miRNA-21 was detected by real time quantitative PCR. The expression levels of IL-12 and STAT4 in lung tissue were assayed via western blot, and immunohistochemistry was used to observe the distribution of their expression.

Results: The expression levels of miRNA-21 as well as the total number of BALF cells and EOS were significantly higher in the asthmatic model group than in the control or dexamethasone groups, with significantly higher amounts found in the dexamethasone group than in the control group. The expression levels of IL-12 and STAT4 proteins were lower in the asthmatic model group than in the control and dexamethasone groups, with a significantly lower expression of IL-12 and STAT4 in the dexamethasone group than in the control group.

Conclusions: The expression level of miRNA-21 was significantly increased and the expression level of IL-12 and STAT4 proteins was significantly decreased in allergic asthmatic mice compared with normal control mice. These findings suggest a role for miRNA-21 and the IL-12/STAT4 pathway in the development of allergic asthma.

Key words: allergic asthma, miRNA-21, IL-12, signal transducer and activator of transcription 4 (STAT4).
Material and methods

Mice

We purchased 45 specific pathogen-free (SPF) male mice with a C57BL/6 background (20-22 g) from Shanghai SLAC Laboratory Animal Co., Ltd. (China). The mice used in the studies were sacrificed when they were 6-8-weeks-old.

Laboratory equipment

Real-time polymerase chain reaction (qPCR) thermal cycler (Agilent-Stratagenegene, USA); electrophoresis apparatus (BioRad, USA); western blot transfer unit (BioRad, USA); electric atomizer (GSK, UK); microplate reader (AI, USA); optical microscope (Olympus, Japan).

Establishment of asthmatic mouse model

Mice were housed under SPF conditions at 24 ±2°C and 55 ±5% relative humidity with a 12-h light-dark cycle. A commercial diet and potable water were available ad libitum. All experiments were approved by the Animal Research Ethics Board of Lishui University (Lishui, Zhejiang Province, China). Mice were randomly divided into three groups (n = 15 mice per group): the model group; the dexamethasone treatment group; and the control group. On days 0 and 8, mice were administered ovalbumin (OVA; Sigma, USA) dissolved in 1.0 ml of aluminum hydroxide gel (60 μg of OVA and 2.25 mg of aluminum hydroxide gel) by intraperitoneal (i.p.) injection. Commencing on day 15, mice in the model group received an ultrasonic nebulization of 2% OVA in a 5 ml solution for 30 min every other day over 7 days. Mice in the dexamethasone treatment group received an i.p. injection of 0.3 mg/kg dexamethasone (Sigma, USA) prior to OVA nebulization. Mice in the control group received saline instead of OVA.

Hematoxylin/eosin (HE) staining of lung tissue

A portion of the right upper lung tissue was obtained, fixed with 4% paraformaldehyde, dehydrated and embedded in paraffin, then sectioned (5-μm thickness). Sections were baked for 1 h at 60°C, hydrated through a graded ethanol series, and stained for 10 min with hematoxylin, then stained for 3 min with eosin. After washing under running tap water, sections were dehydrated through a graded alcohol series, treated with xylene, and mounted with neutral gum. Pathological changes in sections were observed using an optical microscope.

Cell count within bronchoalveolar lavage fluid (BALF)

Mice in all groups were anaesthetized through an i.p. injection of 5 ml/kg 20% urethane solution after the last OVA nebulization event. Serum and left lung tissue was collected from each mouse; bronchoalveolar lavage was conducted and BALF collected. The total cell number and number of eosinophils (EOS) were counted using a microscope and a high power field-of-view.

Real-time polymerase chain reaction analysis of microRNA-21 expression in lung tissue

A portion of the right lung from each mouse was obtained, and total RNA extracted using Trizol (Invitrogen, USA). We used electrophoresis to determine the quality of the extracted RNA by observing bands corresponding to 28S, 18S and 5S RNA. The purity and quantity of RNA samples was determined by measuring the absorbance at 260 (A260) and 280 nm (A280) and calculating the A260/280 ratio. Only samples with an A260/280 ratio greater than 1.8 were used. We used 1 μg of total RNA for reverse transcription reactions. The generated cDNA (1 ml) was used as a template in qPCR assays which comprised 25 ml. A melt curve for each amplicon was created to determine the specificity of the amplification reaction. We used U6 SnRNA as an internal control.

Western blotting analysis of interleukin 12 and signal transducer and activator of transcription 4 in lung tissue

Proteins were extracted from the lower right section of lung tissues following homogenization of the tissue and the application of a lysis buffer. The concentration of proteins in each sample was determined, and then samples were subjected to polyacrylamide gel electrophoresis (PAGE) for 2 h at 100 V. Proteins were transferred to membranes (0.5 A, 60 min) and probed with goat anti-mouse IL-12 (1 : 100 dilution; Santa Cruz Biotechnology, USA) and goat anti-mouse STAT4 (1 : 100; Santa Cruz Biotechnology, USA) antibodies at 4°C overnight. Samples were repeatedly washed with phosphate-buffered saline (PBS) for 5 min, then incubated with a horseradish peroxidase (HRP)-conjugated rabbit anti-goat antibody (1 : 4000; Sigma, USA) at room temperature for 1 h. Positive signals were detected using chemiluminescence techniques. We used β-tubulin as an internal control.

Immunohistochemical analysis of interleukin 12 and signal transducer and activator of transcription 4 in lung tissues

Formalin-fixed paraffin-embedded lung tissue sections were de-waxed, then rehydrated through a graded alcohol series. Sections were treated with 3% (v/v) H2O2 to inhibit endogenous peroxidase activity. The antigen was retrieved by microwaving, and sections blocked using normal sheep serum. Sections were then probed with IL-12- and STAT4-specific primary (1 : 100; Santa Cruz Biotechnology, USA) and secondary (1 : 4000; Sigma, USA) antibodies. We added SP (Jiangsu Beyotime, China) reagent and 3,3’-diaminobenzidine (DAB; Beyotime, Jiangsu, China)
to visualize positive brown signals. Nuclear staining was conducted using hematoxylin. Sections were dehydrated and mounted with gum, then examined using a microscope. Differences in protein expression levels among different groups were compared by calculating the average gray value.

**Statistical analysis**

Data were analyzed using SPSS18.8 (SPSS Inc., USA). The values we have presented are the mean ± standard deviation (X ± s). Single factor analysis of variance (ANOVA) was used for data comparison among groups. A Bonferroni test was used to determine homogeneity of variance, and Dunnett’s T3 analysis was used when variance was missing. A P-value of less than 0.05 was considered statistically significant.

**Results**

**Pathological changes in lung tissues**

Alveolar wall tissues of mice in the control group had an integral structure. The alveolar cavity exhibited no exudation, and the alveolar wall had not thickened (Fig. 1A). The alveolar wall structure in the model group was damaged, with the alveolar wall thickened, and a large number of inflammatory cells had infiltrated around the bronchus (Fig. 1B). Lung tissue alveolar wall damage and inflammatory cell infiltration in the dexamethasone group were decreased compared with that in the model group (Fig. 1C).

**Bronchoalveolar lavage fluid results**

The number of cells in BALF samples from the control, model, and dexamethasone groups is presented in Table 1. The total cell and EOS numbers in BALF samples in the model group were significantly higher than those in mice of the control group. These values were significantly reduced (p < 0.05) following dexamethasone intervention, but still higher than those in normal mice.

**Expression profile of microRNA-21 in lung tissues**

Expression levels of miRNA-21 mRNA in the control, model, and dexamethasone groups are presented in Table 2. The miRNA-21 expression levels in asthmatic mice from the model group were significantly higher (p < 0.05) than those in normal mice from the control group. Expression levels of miRNA-21 were significantly lower (p < 0.05) in mice treated with dexamethasone, but still higher than those in normal mice.

**Expression of interleukin 12 and signal transducer and activator of transcription 4 in lung tissues**

Lung IL-12 and STAT4 protein expression levels are presented in Fig. 2, and integral optical density ratios compared

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**Table 1. Comparison of bronchoalveolar lavage fluid (BALF) cell counts among the model, dexamethasone, and control groups**

| Group          | Total cells in BALF (×10⁴) | No. of EOS in BALF (×400 photo) |
|----------------|----------------------------|---------------------------------|
| model group    | 86.39 ±22.53*              | 241.78 ±49.23*                  |
| dexamethasone group | 28.13 ±7.39**             | 78.63 ±22.51**                  |
| control group  | 9.46 ±3.07                 | 23.56 ±5.97                    |
| F              | 126.75                     | 195.40                          |
| P              | < 0.05                     | < 0.05                          |

*compared with the control group, p < 0.05; **compared with the model group, p < 0.05

**Table 2. Expression levels of miRNA-21 in mice from the model, dexamethasone, and control groups**

| Item          | MicroRNA-21 levels |
|---------------|--------------------|
| model group   | 4.95 ±1.21*        |
| dexamethasone group | 2.21 ±0.67*       |
| control group | 1.02 ±0.23         |
| F             | 92.967             |
| P             | < 0.05             |

*compared with the control group, p < 0.05; *compared with the model group, p < 0.05
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in Table 3. Interleukin 12 and STAT4 levels in asthmatic mice of the model group were significantly reduced \((p < 0.05)\) compared with control mice. Interleukin 12 and STAT4 protein expression levels were significantly improved \((p < 0.05)\) following glucocorticoid intervention; however, these levels were still lower than those in control mice.

**Immunohistochemical detection of interleukin 12 and signal transducer and activator of transcription 4 in lung tissues**

The expression of IL-12 and STAT4 in the lung tissues of asthmatic mice from the model group were significantly decreased \((p < 0.05)\) compared with those in the control group. Expression levels were significantly enhanced \((p < 0.05)\) following dexamethasone intervention. Interleukin 12 was expressed in the cytoplasm, and STAT4 was expressed mainly in the nucleus with some expression seen in the cytoplasm (Figs. 3 and 4).

**Discussion**

Research has shown that a variety of miRNAs can affect the occurrence and development of asthma [10, 11]. Wang *et al.* [12] showed that miRNAs can lead to degradation of mRNA and inhibit protein translation. These miRNAs bind to corresponding target sites on the 3'-untranslated region (UTR) of the mRNA, thereby achieving post-translational regulation of gene transcription. Changes in the expression levels of miRNA133a [13], miRNA26a [14] and miR-

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**Table 3. Relative integral optical density values for interleukin 12 (IL-12) and signal transducer and activator of transcription 4 (STAT4) in mice from the model, dexamethasone, and control groups**

| Item          | IL-12         | STAT4         |
|---------------|---------------|---------------|
| model group   | 1.32 ±0.25*   | 1.69 ±0.31*   |
| dexamethasone | 2.98 ±0.78*#  | 3.48 ±0.87*#  |
| control group | 4.26 ±0.97    | 5.34 ±1.02    |
| F             | 60.67         | 79.17         |
| P             | < 0.05        | < 0.05        |

*compared with the control group, \(p < 0.05\); #compared with the model group, \(p < 0.05\).
NA155 [15, 16] can lead to changes in the airway, such as remodeling, thereby aggravating asthma symptoms.

In asthmatic mice, the alveolar wall is damaged by increased secretions. The number of inflammatory cells around the airway is significantly increased. However, injuries are improved after dexamethasone intervention. The total cell number, along with the number of EOS in BALF samples in asthmatic mice are significantly higher \( (p < 0.05) \) than those in normal mice; however, these numbers are significantly decreased \( (p < 0.05) \) following dexamethasone intervention, but still higher than those in normal mice. We found that miRNA-21 expression levels were significantly higher \( (p < 0.05) \) in asthmatic mice compared with those in normal mice. These miRNA levels were significantly decreased \( (p < 0.05) \) following glucocorticoid intervention, but were still lower than those in control mice.

Recent studies have shown that miRNA-21 target sites are located within IL-12 mRNA, therefore miRNA-21 can significantly inhibit the translation process after IL-12 transcription [17]. Our findings indicate that miRNA-21 can regulate IL-12 and STAT4 translation post-transcriptionally, resulting in less IL-12 and STAT4 proteins. Our results reflect those of previous studies. Interleukin 12 is an important cell factor in maintaining the Th1/Th2 balance in the body [18-20]. The receptors are dimerized by combining with IL-12 receptors at the T0 cell surface, with JAK kinases activated by the dimerized receptor. This triggers phosphorylation of intracellular receptors that area tyrosine residues. Phosphorylated intracellular receptors begin to raise STAT4 levels and transfer it to the nucleus, where STAT4 is combined with the target gene promoter sequence to promote transcription of the target gene so that T0 cells are differentiated into Th1 cell subsets. The Th1 cell subsets can mediate the cellular immune response and the occurrence of delayed type hypersensitivity reactions. These results indicate that IL-12 can regulate the expression of STAT4, and that IL-12/STAT4 signaling channels can inhibit asthma and other type I allergic reactions [21-24].

We have shown that miRNA-21 expression levels in asthmatic mice are significantly increased, and that miRNA-21 is specific to IL-12 mRNA allowing for inhibition of IL-12 translation. This miRNA also results in reduced STAT4 expression levels, which can lead to Th1/Th2 balance disorders. Type I allergic reactions can occur more easily under the stimulation of external allergens, thereby promoting the occurrence and development of asthma. The expression levels of miRNA-21 are decreased following glucocorticoid intervention, such that IL-12 post-transcriptional translation is increased. When IL-12 expression levels are significantly increased, the expression level of STAT4 is increased, thereby promoting re-establishment of the Th1/Th2 balance and reducing the incidence of asthma.

The authors declare no conflict of interest.

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