Association of methylation and expression of IL-4 and IFN-γ in peripheral blood of asthmatic patients.

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Abstract

This study aims to compare the expressions and methylation of interleukin-4 (IL-4) and interferon-γ (IFN-γ) between Han and Uighur populations as well as their relationships with asthma. A total of 108 asthmatic patients (including 73 Han patients and 35 Uyghur patients) were included into this study, together with 150 healthy populations (including 120 Han patients and 30 Uyghur patients) as the control. The levels of IL-4 and IFN-γ were measured by Polymerase Chain Reaction (PCR), their protein expressions were detected by Enzyme-Linked Immuno Sorbent Assay (ELISA), and their methylation degrees were detected by SNaPshot. Compared with the control group, the expressions of IL-4 and IFN-γ genes and proteins were increased in the asthma group (P<0.05); the methylation degree of IL4-1SF locus was higher (P<0.05); the analysis toward the asthma group showed that the expressions of IL-4 and IFN-γ genes in the Uyghur patients were slightly higher while their protein expressions were significantly decreased; however, there was no significant difference in the methylation level of IL-4 and IFN-γ among different loci between the Han and Uyghur patients (P>0.05). IL-4 and IFN-γ are overexpressed in asthmatic patients, especially in Han patients. The methylation degree of IL4-1SF locus in asthmatic patients is increased, but there is no difference between Han and Uyghur populations.

Keywords: Asthma, Epigenetics, Gene expression, Methylation.
Materials and Methods

Subjects

The asthmatic patients treated in the First Affiliated Hospital of Xinjiang Medical University and the People’s Hospital of Xinjiang Uygur Autonomous Region from Dec 2014 to Jul 2015 was enrolled. Inclusion criteria: (1) Aging 18-60 y old; (2) With a disease history ≥ 1 y; (3) Had lived locally in Xinjiang for at least 5 y and had not long-term lived in outer regions; (4) Met with the diagnostic criteria of bronchial asthma prevention and treatment guidance (2008): recurrent wheezing, shortness of breath, chest tightness or cough, which was more related to the exposure to allergens, cold air, physical or chemical stimuli, viral upper respiratory tract infection, or exercise; scattered or diffuse expiratory phase-based wheezing can be heard in the lungs at the onset; the expiratory phase was prolonged; the patients with atypical clinical symptoms should have at least one positive result of the following tests: bronchial provocation test or exercise test, bronchodilation test (FEV1 was increased by more than 15%, and the absolute value of FEV1 was increased by >200 ml), intraday variability or circadian volatility of PEF ≥ 20%; 5) signed the informed consent. Exclusion criteria: (1) with other airway infectious diseases; (2) With other allergic diseases; (3) With hypertension, diabetes, or other metabolic disorders; (4) Administrated immunosuppressive agents within 1 week. A total of 150 asthmatic patients that met with the above criteria were enrolled in this study, including 120 Han patients and 30 Uygur patients, with a mean age as (35.81 ± 11.37) y. A total of 108 healthy population with normal physical examination results were included as the control group, with the mean age as (37.21 ± 10.62) y. This study was conducted in accordance with the declaration of Helsinki. This study was conducted with approval from the Ethics Committee of Xinjiang Medical University. Written informed consent was obtained from all participants.

Polymerase chain reaction (PCR)

All patients were sampled with 5 ml blood, and the CD4+ lymphocytes were collected by magnetic activated cell sorting (MACS) from 4 ml, then IL-4 mRNA and IFN-γ mRNA were extracted by Trizol and the expression level of both were detected after PCR amplification [14].

Enzyme-linked immuno sorbent assay (ELISA)

The protein expression of IL-4 and IFN-γ were detected by ELISA (Xin Le Biological Engineering Co. Ltd., Shanghai, China) based on the plasma used for PCR detection [15].

SNaPshot

The remain blood (1 ml) was used to extract DNA after cell lysis, and the methylation were detected by EZ DNA methylation modification Kit (ZYMO RESEARCH Biotechnology company, Orange County, California, USA) [16].

Statistical analysis

SPSS17.0 was used for the statistical analysis; the mRNA expression levels (IFN-γ, IL-4 and GAPDH) were expressed as mean ± standard deviation (x ± s); the computed tomography (CT) values and protein expression levels were analysed using the t-test of two independent samples, with P<0.05 considered as statistical significance. As for the methylation data, the raw data collected from the ABI3730XL sequencer were analysed by GeneMapper (Applied Bio systems Co. Ltd., USA) and SPSS17.0; the methylation levels of IFN-γ and IL-4 were expressed as mean ± standard deviation (x ± s)%; the comparison of the methylation degree between two genes was compared using the t-test of two independent samples, with P<0.05 considered as statistical significance.

Results

Expressions of IL-4 and IFN-γ

Compared with the control group, the levels of IL-4 and IFN-γ mRNA in the asthmatic patients were slightly increased, but the differences were statistically significant (P<0.05). The protein expressions of IL-4 and IFN-γ were similar to those of the mRNAs (P<0.05) (Table 1).

Table 1. IL-4/IFN-γ expression in asthma group and control group (x̄ ± s).

<table>
<thead>
<tr>
<th>Items</th>
<th>Asthma group (n=150)</th>
<th>Control group (n=108)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>IL-4 mRNA</td>
<td>10.54 ± 2.41</td>
<td>9.95 ± 1.91</td>
<td>0.047</td>
</tr>
<tr>
<td>IFN-γ mRNA</td>
<td>8.33 ± 2.97</td>
<td>6.86 ± 1.80</td>
<td>0</td>
</tr>
<tr>
<td>IL-4 protein</td>
<td>2107.74 ± 611.75</td>
<td>444.19 ± 138.87</td>
<td>0</td>
</tr>
<tr>
<td>IFN-γ protein</td>
<td>394.59 ± 134.40</td>
<td>319.08 ± 110.30</td>
<td>0</td>
</tr>
</tbody>
</table>

Compared with the Han asthmatic patients, the mRNA levels of IL-4 and IFN-γ in Uygur asthmatic patients were statistically increased in the Uygur asthmatic patients (P<0.05), but the protein and mRNA expression levels were lower; there was no significant difference in the IL-4 protein between the two groups (P=0.219), but the expression of IFN-γ protein exhibited statistical difference (P=0.004) (Table 2).

Table 2. IL-4/IFN-γ expression in asthma patients in different races (x̄ ± s).

<table>
<thead>
<tr>
<th>Items</th>
<th>Uygur race (n=30)</th>
<th>Han race (n=120)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>IL-4 mRNA</td>
<td>10.69 ± 2.99</td>
<td>9.74 ± 1.88</td>
<td>0.038</td>
</tr>
<tr>
<td>IFN-γ mRNA</td>
<td>9.62 ± 3.57</td>
<td>8.14 ± 2.75</td>
<td>0.029</td>
</tr>
<tr>
<td>IL-4 protein</td>
<td>1961.78 ± 530.88</td>
<td>2121.92 ± 627.40</td>
<td>0.219</td>
</tr>
<tr>
<td>IFN-γ protein</td>
<td>264.28 ± 97.23</td>
<td>331.69 ± 111.96</td>
<td>0.004</td>
</tr>
</tbody>
</table>
**Methylation levels of IL-4 and IFN-γ**

Compared with the control group, the methylation degree of IL-4-1SF locus in the asthmatic patients was significantly increased (P=0.040), but other loci, namely IFN-γ1SF, IFN-γ2SR, IL-4-2SR, and IL-44SR showed no significant difference in the methylation degree (P>0.05) (Table 3).

Table 3. Methylation sights of IL4-1SF, IL4-2SR, IL-44SR, IFN-γ1SF and IFN-γ2SR in asthma group and control group (x̄ ± s,%).

<table>
<thead>
<tr>
<th>Items</th>
<th>Asthma group (n=150)</th>
<th>Control group (n=108)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>IL-41SF</td>
<td>97.04 ± 5.13</td>
<td>92.30 ± 15.88</td>
<td>0.04</td>
</tr>
<tr>
<td>IL-42SR</td>
<td>91.05 ± 1.78</td>
<td>90.75 ± 2.02</td>
<td>0.256</td>
</tr>
<tr>
<td>IL-44SR</td>
<td>93.58 ± 1.54</td>
<td>93.44 ± 1.56</td>
<td>0.515</td>
</tr>
<tr>
<td>IFN-γ1SF</td>
<td>94.87 ± 1.17</td>
<td>95.14 ± 9.02</td>
<td>0.066</td>
</tr>
<tr>
<td>IFN-γ2SR</td>
<td>96.53 ± 6.53</td>
<td>96.59 ± 7.28</td>
<td>0.517</td>
</tr>
</tbody>
</table>

Compared with the Han asthmatic patients, the methylation levels of IL-4 and IFN-γ loci in the Uighur asthmatic patients were similar, showing no statistical significance (P>0.05) (Table 4).

Table 4. Methylation sights of IL4-1SF, IL4-2SR, IL-44SR, IFN-γ1SF and IFN-γ2SR in Han and Uygur patients (x̄ ± s,%).

<table>
<thead>
<tr>
<th>Items</th>
<th>Uygur race (n=30)</th>
<th>Han race (n=120)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>IL-41SF</td>
<td>96.32 ± 7.43</td>
<td>97.29 ± 4.04</td>
<td>0.392</td>
</tr>
<tr>
<td>IL-42SR</td>
<td>90.99 ± 2.02</td>
<td>91.07 ± 1.71</td>
<td>0.826</td>
</tr>
<tr>
<td>IL-44SR</td>
<td>93.79 ± 1.75</td>
<td>93.50 ± 1.46</td>
<td>0.389</td>
</tr>
<tr>
<td>IFN-γ1SF</td>
<td>95.12 ± 0.78</td>
<td>95.13 ± 0.93</td>
<td>0.905</td>
</tr>
<tr>
<td>IFN-γ2SR</td>
<td>96.48 ± 0.89</td>
<td>96.55 ± 0.54</td>
<td>0.621</td>
</tr>
</tbody>
</table>

**Discussion**

One study of Taiwan reveals that the prevalence of asthma has increased from 12.99%-16.86%, and environmental changes are suspected to be the major driver of asthma with air pollution identified as an important exposure, but it is still a complex genetic trait meaning that the phenotype is the result of the interaction of multiple genes [17-20].

This study revealed that the mRNA and protein levels of IL-4 in the asthma group were significantly higher than the healthy control, confirming that asthma is mainly characterized by Th2 hyperthyroidism-based immune responses, suggesting that IL-4 antagonists can be used clinically as targeted therapy against asthma, which can interrupt the IL-4 signal pathway, thereby reducing the number of pro-inflammatory factors in the airway and improving patients’ conditions [21]. IL-4 can promote the secretion of Th2 cells by reducing the secretion of IFN-γ, which is the main process that mediates the pathological and physiological changes of asthma. Compared with the Han asthmatic patients, the IL-4 mRNA expression was slightly elevated in the Uighur asthmatic patients, but there was no difference in the IL-4 protein expression, suggesting that IL-4 may not be the reason that causes the different incidences of asthma in Han and Uighur populations.

IFN-γ is an important Th1 cytokines, can inhibit the release of Th2 cytokines, and can also regulate the immune function [22]. Studies have shown that one of the important reasons for allergic asthma in children is the reduction of Th1-type cytokines [23]. IFN-γ can inhibit the expression of IL-4 gene and the synthesis of IgE by IL-4, thus participating in the occurrence and development of inflammatory processes in asthma [24], and it can also affect the proliferation and differentiation of Th2 and Th1 as well as antagonize the role of IL-4. Nowadays, current studies have confirmed that the Th1 cells can produce not only IL-2 but also IFN-γ [25]. Compared with IL-4, the Th1 cytokine, IFN-γ, was also highly expressed in the asthma group than the control group, not consistent with the recognized Th1 immune response reduction-based pathogenesis of asthma, and it may be caused the secretion differences of IFN-γ at the transcription level and the translation level.

However, in recent years, Th1 and Th2 have been found to be able to induce, instead of mutually antagonizing, the activity of bronchial asthma, respectively, which means the existence of mutual promotion between these two [26]. The fact that the IFN-γ gene and protein expressions were increased in the asthma group can be interpreted as the results of the mutual promotion between Th1 and Th2 rather than mutual antagonism.

However, further studies are still needed to verify these different results. Further analysis targeting the Han and Uygur asthmatic patients showed that although the IFN-γ mRNA level in the Uygur asthmatic patients was slightly higher than the Han asthmatic patients, the protein expression level was significantly reduced, suggesting that the difference in the plasma concentration of this factor may be the reason leading to the difference in the pathogenesis of asthma between Han and Uygur populations.

The results of SNPhot typing showed that the methylation degree of IL-41SF locus in the asthma group was significantly higher than the control group. It is generally believed that the regulatory role of DNA hyper methylation is associated with gene silencing, suggesting that the hyper methylation of IL-41SF is closely related to the occurrence of asthma, so it can be used to find new treatment directions against asthma. However, because there was no significant difference in the methylation degrees of IL-4 and IFN-γ between the Uygur and Han asthmatic patients, it suggests that the methylation degree may not be the cause resulting in the different prevalence between Uygur and Han patients; or it may be caused by too small Uygur patients, which needs to further expand the sample size so as to verify the results.
Acknowledgement

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Conflicts of Interest

The authors declare no conflict of interest.

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