Interleukin-12 Peripheral Blood Levels in Asthmatic Children

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Interleukin-12 (IL-12) was measured in 45 asthmatic children aged 3 to 16 years. The assessments were performed on 20 children during an episode of acute exacerbation and on 25 children during remission. There was no significant difference between the mean IL-12 level during exacerbation (1.63 ± 2.08 pg/mL) and during remission (0.88 ± 0.56 pg/mL) (p = .83). A positive, but insignificant, correlation was found between forced expiratory volume in 1 second and IL-12 (p = .634). IL-12 levels were significantly lower in children with a positive family history of asthma (1.13 ± 1.78 pg/mL) compared with those without (1.31 ± 1.06 pg/mL) (p < .012), supporting the theory that the gene–environment interactions affect the immune responses. IL-12 peripheral blood levels had no detectable impact on the course of established asthma in the study population.

Key words: *asthma, family history, IL-12*

The prevalence of atopic asthma, a T helper (Th)2-dependent disease, is reaching epidemic proportions, possibly owing to improved hygiene in industrialized countries.[1] The chronic inflammation of the airways in asthma is characterized by the presence of Th2 cells in sputum, bronchoalveolar lavage, and mucosal biopsy specimens,[2] and the dominance of Th2 cells is responsible for the pathogenesis of allergic diseases.[3–6] The priming of T cells requires the activation of dendritic cells (DCs),[7] which are generally considered to be the principal antigen-presenting cells (APCs) involved in the generation of polarized effector cells.[4] DCs have a major influence on the pattern of Th1/Th2 polarization by releasing cytokines, especially interleukin-12 (IL-12).[5] The route of the antigen encounter with the DC and the subtype of the APC can profoundly influence Th differentiation.[7] IL-12, the critical Th1-polarizing cytokine[8] produced by DCs after stimulation by various antigens, including the endotoxin lipopolysaccharide (LPS), which is derived from the cell walls of gram-negative bacteria[9] and it is ubiquitous in the domestic environment.[10] The functional active form of IL-12 is a heterodimer composed of two disulphide-linked chains, p35 and p40, secreted by APCs[11] and by activated Th1 cells; this heterodimer downregulates Th2 responses.[12]

Studies in asthma models concluded that the administration of IL-12 before and during the period of allergen challenge prevented allergen-induced airway eosinophilia, airway hyperresponsiveness, the production of Th2 cytokines, and the production of allergen-specific serum immunoglobulin E,[13] whereas it increased the production of interferon-γ and enhanced apoptosis of CD4$^+$ T cells in allergic airway infiltrates.[14]

The main aim of the current study was to assess the relationship of IL-12 peripheral blood levels and the two states of asthma, acute exacerbation and remission, as demonstrated by the kinetics of DC activation by environmental stimuli and, consequently, Th1/Th2 polarization.

Patients and Methods

Patients

The study was conducted between April 2004 and September 2005 after having been approved by the local institutional review board (Helsinki Committee). The parents of all of the participants provided written informed consent.

The study population included 45 asthmatic children, aged 3 to 16 years (median age 9.5 ± 3.4 years), who were

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recruited from the pediatric emergency department and the pediatric pulmonology outpatient clinic. The inclusion criteria were a diagnosis of asthma prior to study entry confirmed by a pediatric pulmonologist and being treated by inhaled corticosteroids for at least the two previous months. The exclusion criteria were evidence of pneumonia on the chest radiograph, fever, already being treated by rescue medication (β_2 agonists and systemic steroids), and having a chronic or acute illness other than asthma.

The participants were divided into two groups. Group A consisted of 20 children who were recruited when they arrived to the emergency department or to the outpatient clinic during an acute exacerbation of asthma. Group B included 25 asthmatic children who visited the outpatient clinic for a regular follow-up evaluation and were in remission.

The demographic information on the participants is presented in Table 1.

Respiratory Symptoms and Asthma

In the study children $\geq$ 6 years of age, diagnosis of asthma prior to recruitment was by clinical parameters and by demonstrating bronchial hyperreactivity in pulmonary function tests with a provocation by either exercise or the adenosine test. Children younger than 6 years were diagnosed by clinical parameters, including a history of recurrent episodes of coughing, wheezing, and breathlessness that were relieved by bronchodilators and steroids.

The following data were collected for each patient: respiratory symptoms throughout the 2 months prior to the current presentation, the regular treatment regimen during the past few months, a family history of asthma and other diseases, smoking habits of family members, and the presence of pets at home. A comprehensive physical examination included the measurement and recording of weight, height, respiratory rate, heart rate, and oxygen saturation. Respiratory symptoms were evaluated according to respiratory rate, respiratory chest recession, auscultatory breath sounds, and general condition. Acute exacerbation was defined by the presence of coughing, tachypnea according to age,[15] respiratory muscle retractions, and auscultatory findings of wheezing with prolonged expiration. Remission was defined when the child was free of cough for at least 2 weeks, the respiratory rate was $\leq$ 20 breaths/minute, there were no respiratory muscle retractions, and normal breath sounds were heard on auscultation.

Lung Function Tests

Spirometry was performed by a Vitalograph compact II spirometer (Vitalograph Ltd., UK) in patients older than 6 years. The predicted results were analyzed according to Polgar normative data.[16]

IL-12 Measurements

IL-12 levels in serum samples were analyzed in duplicate using a commercial kit: Quantikine high sensitivity human IL-12 immunoassay (catalogue number HS120, R&D Systems Inc., Minneapolis, MN). The Quantikine high-sensitivity immunoassay kit uses an amplification system in which the alkaline phosphatase reaction provides a cofactor that activates a redox cycle leading to the formation of a coloured product. The minimum detectable dose of human IL-12 is typically less than 0.5 pg/mL. This assay employs the quantitative sandwich enzyme immunoassay technique. A monoclonal antibody specific for IL-12 has been precoated onto a microplate. Standards and samples are pipetted into the wells, and any IL-12 present is bound by the immobilized antibody. After washing away any unbound substances, an enzyme-linked polyclonal antibody specific for IL-12 is added to the wells. Following

Table 1. Demographic Data on the Two Groups of Study Children

Demographic Data	Group A (n = 20)	Group B (n = 25)	p Value (Test)
Males/females	13/7	19/6	.42 (chi-square)
Median current age (yr)	8.2 ± 3.1	10.5 ± 3.3	.05 (Mann-Whitney)
Family history of asthma	12 (60%)	12 (48%)	.42 (two-way ANOVA)
Passive smoking	8 (40%)	7 (28%)	.4 (chi-square)
Pets at home	6 (30%)	8 (32%)	.89 (chi-square)
Inhaled corticosteroids			.52 (Fisher exact test)
Budesonide	10 (45%)	12 (48%)	
Fluticasone	11 (55%)	13 (52%)	

ANOVA = analysis of variance.

a wash to remove any unbound antibody-enzyme reagent, a substrate solution is added to the wells. After an incubation period, an amplifier solution is added to the wells and colour develops in production to the amount of IL-12 bound in the initial step. The colour development is stopped, and the intensity of the colour is measured.

Statistical Methods

The assumption of normal distribution of continuous parameters was examined using the Kolmogorov-Smirnov and Wilk-Shapiro tests. Since the IL-12 levels were not distributed normally, we also analyzed the reciprocal transformation of this parameter to confirm the results. Comparisons between the two groups of patients for demographic parameters (gender, age, passive smoking, family history of asthma, and pet at home) and other factors (IL-12, forced expiratory volume in 1 second [FEV_1] measures) were performed using the t-test for independent samples, the Mann-Whitney nonparametric test, and the chi-square test, as applicable. In addition, a two-way analysis of variance (ANOVA) was performed to examine the association between each group's family history of asthma and IL-12 levels. Spearman nonparametric correlation coefficients were calculated to study the relationship between all continuous parameters and IL-12. Significance was set at $p = .05$, and *SPSS for Windows* software version 13.0 (SPSS Inc, Chicago, IL) was used for the analysis.

Results

The patient population consisted of 45 children clinically diagnosed as being asthmatic: 20 were studied during an acute episode of asthma (group A) and 25 were studied during a remission (group B). There were no significant differences between the two study groups in terms of the male to female ratio, pets at home, passive smoking, and family history of asthma (see Table 1).

The median age of the children in group A (8.21 ± 3.16 years) was significantly lower than the median age of the children in group B (10.52 ± 3.33, $p < .025$), but this difference had no impact on the IL-12 analysis between two groups.

The mean IL-12 level was 1.63 ± 2.08 pg/mL (range 0.3–8.3 pg/mL) in group A and 0.88 ± 0.56 pg/mL (range 0.4–3.2 pg/mL) in group B; the difference between the two groups was not significant (t-test $p = 1.0$ and Mann-Whitney test $p = .83$). The IL-12 levels in both groups were significantly lower in children with a positive family

history of asthma compared with children without. The mean blood level of IL-12 was 1.13 ± 1.78 pg/mL in children with a family history of asthma and 1.31 ± 1.06 pg/mL in children without a family history of asthma (two-way ANOVA test $p < .012$) (Figure 1).

Plotting the related percent predicted values of FEV_1 to IL-12 levels demonstrated a positive correlation between FEV_1 and IL-12: higher IL-12 levels were correlated to the higher percent predicted values of FEV_1 (Figure 2). These positive correlations did not, however, reach a level of significance (Spearman correlation coefficient, $p = .634$).

Discussion

The findings of this study demonstrated that there were no significant differences between the mean peripheral blood levels of IL-12 during asthma exacerbation and those during remission. Thus, IL-12 peripheral blood levels do not reflect the state of asthma. Although the children who were tested during an acute attack were significantly younger than the children who were tested while in remission, Tomita and colleagues reported that age plays no role in the production of IL-12.[17] We cannot provide an explanation for the unexpected results, and further clinical studies are needed to elucidate them.

IL-12 levels and family Hx of asthma
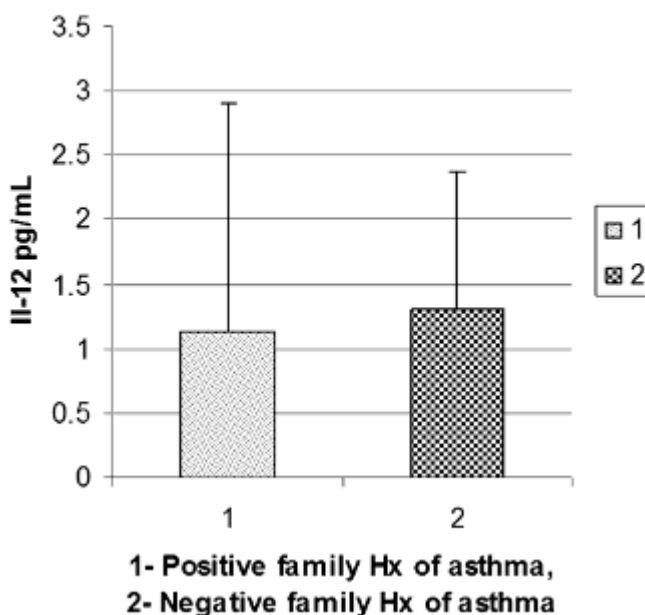
Figure 1. Correlation of interleukin-12 levels to family history (Hx) of asthma. $p < .012$ (two-way ANOVA test).

IL-12 vs. FEV1%
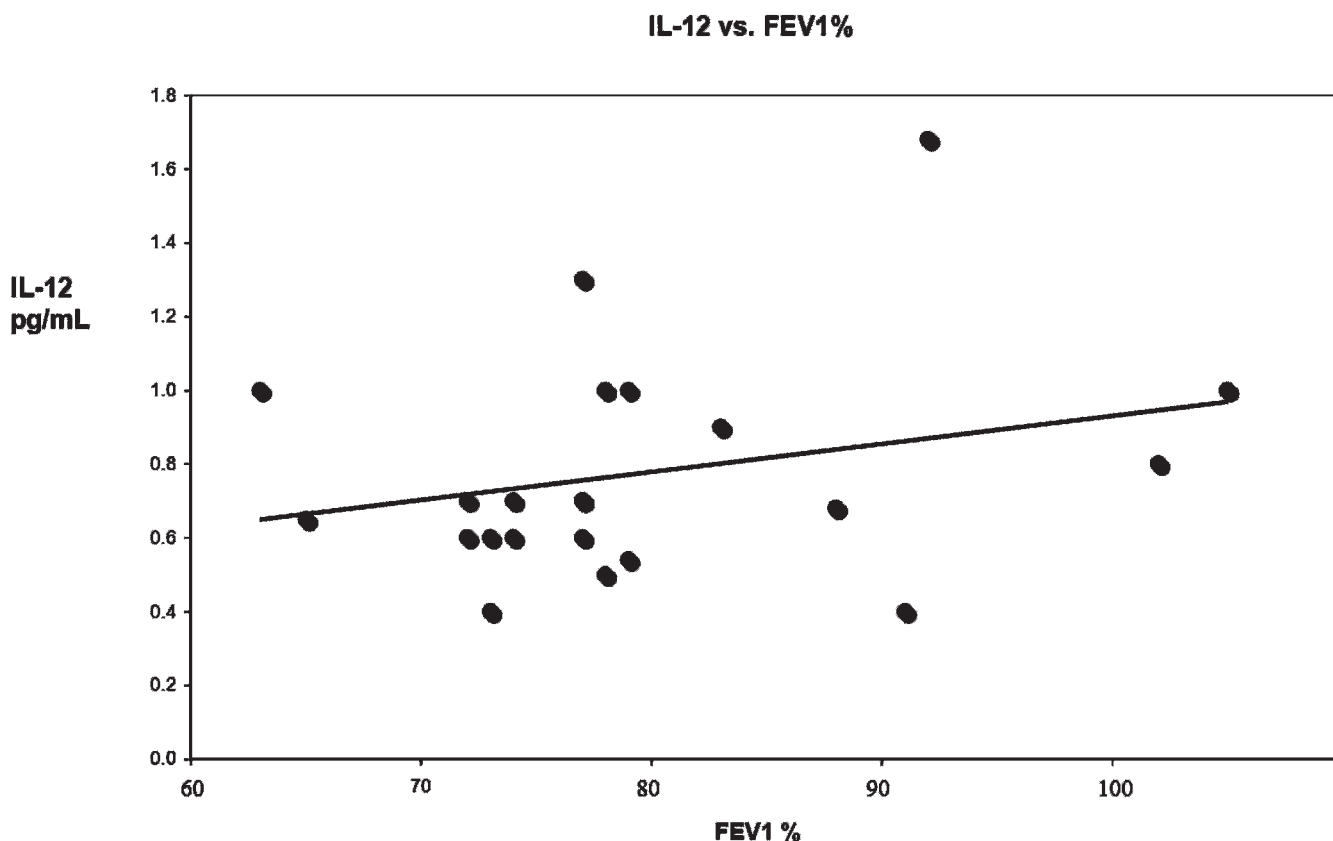
Figure 2. Interleukin-12 (IL-12) peripheral blood levels versus forced expiratory volume in 1 second percent (FEV$_1$%) predicted values. There were positive correlations between FEV$_1$% predicted values and IL-12 peripheral blood levels, but they did not reach a level of significance (p = .6).

All of the participants were treated by inhaled corticosteroids. Since there are no publications of a direct influence of inhaled corticosteroids on IL-12 blood levels, we presume that the inhaled corticosteroids were not an influencing factor on these results.

IL-12, a critical Th1 polarizing cytokine[8] and a heterodimer[18] produced mainly by DCs and Th1 cells, has been shown to have potent immune activity by inducing Th1 responses, suppressing Th2 responses, and enhancing apoptosis of CD4$^+$ T cells in allergic airways.[14] An exogenous form of IL-12 given to a mouse model reduced established airway responses, such as eosinophilic infiltration and airway hyperresponsiveness.[19] Endotoxins, which are inflammatory LPS molecules derived from a gram-negative bacteria wall, are ubiquitous in the indoor environment,[20] and significantly associated with pets at home.[21] A bimodal, dose-dependent relationship between environmental exposure to LPS and the presence of immune responses supports the hygiene theory by demonstrating a preponderance of Th2 at low LPS exposure and a preponderance of Th1 at high LPS exposure.[22] Gereda and colleagues suggested that indoor

endotoxin exposure early in life may protect against allergen sensitization.[23] Inhaled endotoxins trigger macrophages and other myeloid cells, including myeloid DCs, through CD14, a LPS-binding protein, to release cytokines, including IL-12.[24] IL-12 redirects the Th cells responses toward the Th1 immune response.[25] Asthmatic patients with severe airflow obstruction showed an impairment of IL-12 production.[26] A study of peripheral blood mononuclear cells showed that severe asthmatics had significantly less positive staining for IL-12 after stimulation with LPS compared with mild asthmatics and controls.[17] The genetic predisposition that determines the effect of environmental endotoxins and allergic reactions and the production of IL-12 was influenced by the polymorphism in the CD14 gene.[27] In opposition to all of the above-cited reports, the study of the national survey of endotoxins in US housing demonstrated that household endotoxin exposure is a significant risk factor for increased asthma symptoms and wheezing.[28] Braun-Fahrlander and colleagues claimed that exposure to endotoxins at school age was an increased risk factor for asthma exacerbations.[29]

Blockade of IL-12 was recently demonstrated in an animal model to have differential effects on allergic airway inflammation, depending on the timing of the blockade. Blocking IL-12 during the sensitization process aggravated the subsequent development of allergic airway inflammation, whereas neutralization of IL-12 during the challenge phase in previously sensitized mice abolished eosinophilic airway inflammation.[30]

The significantly lower mean level of IL-12 in children with a positive family history of asthma compared with the higher mean level in children without a positive family history supports the theory that the gene–environment interactions affect the immune responses.[31] Moreover, higher IL-12 levels correlated with higher FEV_1 levels, although not significantly so. In conclusion, we found no relationship between IL-12 peripheral blood levels and the course of established asthma in asthmatic children aged 3 to 16 years. The small cohort of children precludes our arriving at any firm conclusions, so we recommend larger clinical studies to validate our findings.

Acknowledgements

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