Elevation of specific immunoglobulin A antibodies to both allergen and bacterial antigen in induced sputum from asthmatics

D-H. Nahm, H-Y. Kim, H-S. Park


ABSTRACT: The antigenic specificity and pathogenetic significance of immunoglobulins in airway secretion from asthmatic patients have not been established. Elevated levels of B-cells and immunoglobulin (Ig)A antibodies have been reported in sputum of asthmatics and these levels correlate with the eosinophil counts and levels of degranulated cytotoxic proteins from eosinophils. This study aimed to investigate the antigen specificity and possible pathogenetic significance of antibodies in airway secretion from asthmatic patients.

Specific IgA and IgG antibodies to both allergen (Dermatophagoides farinae) and bacterial antigen (capsular polysaccharide antigen from Streptococcus pneumoniae) were measured in bronchial secretion from 16 atop ic asthmatic patients sensitized to D. farinae and 12 nonatopic, nonasthmatic controls by enzyme-linked immunosorbent assay. Sputum was induced by inhalation of hypertonic saline. Eosinophil cationic protein (ECP) levels in sputum from asthmatic patients were measured by the Pharmacia CAP system.

Levels of IgA to both D. farinae and S. pneumoniae and IgG to D. farinae in the sputum from asthmatic patients were significantly higher than those from controls (p<0.005). No significant difference was found in the levels of IgG to S. pneumoniae between the two groups. In asthmatic patients, there were significant correlations between IgA to D. farinae and S. pneumoniae (r=0.76, p=0.003). Sputum ECP levels correlated significantly with IgA to D. farinae (r=0.55, p=0.03) and S. pneumoniae (r=0.56, p=0.03) and IgG to D. farinae (r=0.52, p=0.04), but not with IgG to S. pneumoniae in asthmatic patients.

In conclusion, specific immunoglobulin A antibodies to both allergen and bacterial antigen were elevated in induced sputum from atopic asthmatics and their possible involvement in eosinophil degranulation was suggested.

The respiratory mucosa responds to various bacterial organisms and inhaled airborne allergens. Patients with chronic airway disease showed elevated levels of IgA antibody against bacterial antigen in their sputum [12], whereas, patients with bronchial asthma showed elevated levels of IgA antibody to inhalant allergen in sputum [13]. However, antigenic specificity of IgA antibodies in airway secretion from asthmatic patients has not been fully determined.

To investigate the antigen specificity and possible pathogenetic significance of IgA and IgG antibodies in the airway secretion from patients with asthma, IgA and IgG antibodies to allergen and bacterial antigen were measured in induced sputum from patients with atopic asthma.

**Methods**

**Subjects**

Induced sputum samples were collected from 16 atopic asthmatic patients who showed a positive response to house dust mite (*Dermatophagoides farinae*, Bencard Co., Brentford, UK) on a skin-prick test (table 1). All patients had a typical clinical history of asthma and documented reversibility of forced expiratory volume in one second (FEV1) >15% of predicted value. All patients were non-smokers and had a mild degree of asthma. None had received inhaled or oral corticosteroids for at least 3 months before the study. Subjects who had suffered a recent respiratory tract infection or exacerbation of asthma were excluded. Twelve healthy, nonatopic, nonsmoking and non-asthmatic controls were recruited from the medical students and staff of Ajou University Hospital. Some of the asthmatic subjects and all of the controls were participants in the previous study [10]. All subjects underwent a skin-prick test with common aeroallergens (house dust mite, grass pollens, tree pollens, weed pollens, cat, dog and moulds; Bencard Co.). A positive skin test was defined when the mean wheal diameter of the allergen was 3 mm greater than that of the negative control (normal saline). This study was approved by the Ethics Committee of Ajou University Hospital and all subjects gave their written informed consent. The study was conducted from July to August 1995, during the season when the environmental house dust mite allergen reaches peak levels in Korea [14].

**Sputum and saliva sampling**

Sputum was induced according to the method previously described [15]. Immediately before sputum induction, each subject was pretreated with 200 µg salbutamol, administered by means of a metered-dose inhaler. The subjects then inhaled nebulized sterile 3% saline solution for 20 min through an ultrasonic nebulizer (Omron Co., Tokyo, Japan). Subjects were encouraged to cough throughout the procedure and they regularly stopped to expectorate sputum. Before sputum expectoration, each subject was instructed to rinse their mouth with water and blow their nose, then to cough and expectorate sputum into a clean plastic container. Saliva samples were taken before induction of sputum in 12 asthmatic patients.

**Processing of sputum and saliva**

Collected sputum and saliva were processed immediately as in the method previously described [10]. The volumes of saliva and induced sputum were determined and an equal volume of phosphate-buffered saline (PBS) was added. The samples were then mixed using a vortex mixer and centrifuged for 20 min at 1,500g. The supernatants were aspirated and frozen at -20°C. The pellets were further treated with dithiothreitol (Sputolysin; Calbiochem Biochemicals, San Diego, CA, USA) as described previously [15], resuspended in saline, cytocentrifuged (Model 7 cytospin; Shadan Scientific Products, Miami, FL, USA) and stained by the Wright-Giemsa method. Eosinophils were counted from at least 200 nonsquamous cells on each slide.

**Measurement of total immunoglobulin levels**

Total IgA concentrations in induced sputum were measured by nephelometry with a Beckman Array System (Beckman Instruments, Brea, CA, USA). Total IgG concentrations in induced sputum were measured by a sandwich enzyme-linked immunosorbent assay (ELISA) method using affinity-purified polyclonal goat antibodies to human IgG (Sigma Chemical Co., St Louis, MO, USA) as described previously [4].

**ECP measurement**

The ECP level in induced sputum was measured using the Pharmacia CAP system (Pharmacia Diagnostics, Uppsala, Sweden) in 16 asthmatic patients. The lowest detection limit for ECP measurement was 2 ng·mL⁻¹. When ECP levels were below the lowest detection limit, they were treated as zero for statistical analysis.

**Measurement of specific antibodies to house dust mite (Dermatophagoides farinae) and Streptococcus pneumoniae antigens**

Specific antibodies to *D. farinae* and *Streptococcus pneumoniae* antigens were measured by ELISA according to the method reported previously [12, 13, 16]. Microtitre plates (Immulon 2; Dynatech Laboratories, Chantilly, VA, USA) were coated with either *D. farinae* antigen (kindly provided by J. Weiss at Diagnostic Products Corporation, Los Angeles, CA, USA) at a concentration of 2 µg·well⁻¹ in 0.1 M carbonate buffer (pH 9.6) and pneumococcal polysaccharide antigens (mixed antigens from 23 serotypes, Pasteur Méieux Sérums et Vaccins, Lyon, France) at a concentration of 0.1 µg·well⁻¹ in PBS, overnight at
4°C. Plates were washed and blocked with 3% bovine serum albumin in PBS with 0.05% Tween-20 and then incubated with duplicated samples (50 µL) of induced sputum and saliva at 1:50 dilutions overnight at 4°C. After washing, plates were incubated with peroxidase-conjugated affinity-purified goat anti-IgA or anti-IgG antibodies (Sigma) at 1:50 dilutions overnight at 4°C. After washing, the substrate solution consisting of 0.04% (w/v) orthophenylenediamine (OPD) dissolved in 24.3 mM citric acid, 51.4 mM NaH2PO4 (pH 5.0) and 0.03% H2O2 was added. After 15 min, the reaction was stopped by adding 2.5 N H2SO4. Colour development was measured with an ELISA reader (Molecular Devices, Menlo Park, CA, USA) at 490 nm. Amounts of specific antibodies in the samples were calculated from control curves made by optical densities from serial dilutions of positive control samples which showed high titre antibodies using a spline fit program (SoftMax, Molecular Devices). The specific antibody contents in the reference samples were designed as 100 arbitrary units (AU)·mL⁻¹. In each assay system intra-assay and interassay variations were below 15%. To assess the parallelism between standard and samples, interdilutional variations of each assay were calculated and the values were below 15%.

**ELISA inhibition**

ELISA inhibition was performed using sputum samples with high titre specific antibodies. Serially 10-fold diluted *D. farinae* and *S. pneumoniae* polysaccharide antigens starting from 1 mg·mL⁻¹ were added to diluted positive sputum samples for 16 h at 4°C and the specific antibodies were measured by ELISA as described above. The results of ELISA inhibition were calculated as the per cent inhibition according to the following formula: Inhibition (%) = ((Absorbance by addition of buffer only - Absorbance by addition of antigen)/Absorbance by addition of buffer only) × 100.

**Statistics**

Data were expressed as the mean and standard error of the mean (±SEM). The Mann-Whitney U-test was used to assess differences between groups. A Spearman's rank correlation was calculated to assess correlations between the data. The differences between induced sputum and saliva samples from the same subjects were compared by a Wilcoxon signed-rank test. A p-value <0.05 was regarded as statistically significant.

**Results**

**Levels of specific antibodies to Dermatophagoides farinae and Streptococcus pneumoniae in induced sputum**

Levels of IgA to both *D. farinae* and *S. pneumoniae* and IgG to *D. farinae* in sputum from asthmatic patients were significantly higher than those from controls (p<0.005) (fig. 1). No significant difference was found in the levels of IgG to *S. pneumoniae* between the two groups (p>0.05).

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**Fig. 1.** Levels of specific immunoglobulin (Ig)A (a, b) and specific IgG antibodies (c, d) to *Dermatophagoides farinae* and *Streptococcus pneumoniae* in induced sputum from controls and asthmatics. *: p>0.05; +: p<0.005, significant difference between groups.
ELISA inhibition

Dermatophagoides farinae and S. pneumoniae antigen inhibited specific antibody bindings over 80% in each specific IgG and IgA assay in a dose-dependent manner and there was no evidence of cross-antigenicity between the two antigens (data not shown).

Table 2. – Correlations between immunoglobulin levels and eosinophil counts (%) or eosinophil cationic protein (ECP) levels in induced sputum from asthmatic patients (n=16)

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<thead>
<tr>
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<th>Sputum eosinophil count</th>
<th>ECP</th>
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<tbody>
<tr>
<td></td>
<td>r</td>
<td>p-value</td>
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<tr>
<td>Total IgA</td>
<td>0.50</td>
<td>0.053</td>
</tr>
<tr>
<td>Total IgG</td>
<td>0.45</td>
<td>0.21</td>
</tr>
<tr>
<td>Specific IgA to D. farinae</td>
<td>0.31</td>
<td>0.26</td>
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<tr>
<td>Specific IgG to D. farinae</td>
<td>0.33</td>
<td>0.20</td>
</tr>
<tr>
<td>Specific IgA to S. pneumoniae</td>
<td>0.32</td>
<td>0.21</td>
</tr>
<tr>
<td>Specific IgG to S. pneumoniae</td>
<td>0.33</td>
<td>0.20</td>
</tr>
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</table>

Ig: immunoglobulin; D. farinae: Dermatophagoides farinae; S. pneumoniae: Streptococcus pneumoniae.

Table 3. – Fluid phase measurements in induced sputum and saliva from 12 asthmatic patients

<table>
<thead>
<tr>
<th></th>
<th>Induced sputum</th>
<th>Saliva</th>
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<tbody>
<tr>
<td>Total IgA µg·mL⁻¹</td>
<td>337±74</td>
<td>120±23*</td>
</tr>
<tr>
<td>Total IgG µg·mL⁻¹</td>
<td>240.7±105.9</td>
<td>14.3±3.2***</td>
</tr>
<tr>
<td>Specific IgA to D. farinae AU·mL⁻¹</td>
<td>29.4±8.7</td>
<td>4.8±1.2***</td>
</tr>
<tr>
<td>Specific IgG to D. farinae AU·mL⁻¹</td>
<td>12.7±8.1</td>
<td>0.28±0.13***</td>
</tr>
<tr>
<td>Specific IgA to S. pneumoniae AU·mL⁻¹</td>
<td>38.4±9.5</td>
<td>4.6±1.6***</td>
</tr>
<tr>
<td>Specific IgG to S. pneumoniae AU·mL⁻¹</td>
<td>20.4±8.5</td>
<td>0.38±0.10***</td>
</tr>
</tbody>
</table>

Values are presented as mean±SEM. Ig: immunoglobulin; D. farinae: Dermatophagoides farinae; S. pneumoniae: Streptococcus pneumoniae. *: p<0.05; ***: p<0.001, significant difference between groups.

Correlations between fluid-phase measurements of immunoglobulins in induced sputum from asthmatic patients

There were significant correlations between IgA to D. farinae and IgA to S. pneumoniae (r=0.76, p=0.003). However, there was a weaker borderline significant correlation between IgG to D. farinae and IgG to S. pneumoniae (r=0.49, p=0.06) (fig. 2).

Correlations between immunoglobulin levels and eosinophil counts or ECP levels in induced sputum from asthmatic patients

There was no significant correlation between eosinophil counts (%) and immunoglobulin levels in induced sputum from asthmatic patients (p>0.05) (table 2). In asthmatic patients, levels of sputum ECP correlated significantly with those of total IgA (r=0.82, p=0.002), IgA to D. farinae (r=0.55, p=0.03), IgA to S. pneumoniae (r=0.56, p=0.03) and IgG to D. farinae (r=0.52, p=0.04), but not with total IgG (r=0.42, p=0.1) or IgG to S. pneumoniae (r=0.30, p=0.25) (table 2).

Levels of specific antibodies to S. pneumoniae in induced sputum and saliva from 12 asthmatic patients

All fluid-phase measurements including total IgA, total IgG, specific IgA and IgG antibodies to both S. pneumoniae and D. farinae were significantly higher in induced sputum than in saliva from 12 asthmatic patients (total IgA, p<0.05; others, p<0.001) (table 3).

Discussion

This study demonstrated that IgA antibodies to both allergen and bacterial antigen are elevated in induced sputum samples from atopic asthmatic patients. Elevations of B-cells [17] and IgA antibodies [11] in induced sputum of asthmatics have been reported. Although increased levels of allergen-specific IgE, IgG and IgA antibodies in nasal
secrections from patients with allergic rhinitis [18] and house dust mite-specific IgA antibodies in sputum from mite-sensitive asthmatic patients have been reported [13], information on the antigen specificity of these antibodies in airway secretion from asthmatic patients has been limited. In this study, it was demonstrated that IgA antibodies to both house dust mite antigen and pneumococcal polysaccharide antigen are increased in induced sputum from mite-sensitive asthmatic patients.

In a previous study, we demonstrated that the majority of IgA antibodies in induced sputum from atopic asthmatics was not diffused out passively from the systemic circulation but produced locally [10]. There are several possible explanations for the elevation of IgA antibodies to pneumococcal polysaccharide antigen in induced sputum from mite-sensitive asthmatic patients.

The first explanation is infection or colonization of S. pneumoniae in the airway mucosa and stimulation of IgA antibody response in asthmatic patients. It is known that S. pneumoniae inhabit the upper respiratory tract of healthy adults and patients with chronic obstructive pulmonary disease (COPD) [19]. Only a small proportion of patients with COPD (~7%) was found to have colonization of the lower respiratory tract by S. pneumoniae [20, 21]. As the patients included in this study did not have any clinical symptoms or signs suggesting respiratory infection, the possibility of pneumococcal infection in the majority of these patients seems to be very low.

Another possibility is nonspecific expansion of pre-existing IgA-producing B cells in the airway mucosa due to increased levels of IL-5 in the airway secretion of asthmatic patients [22]. In in vitro conditions, IL-5 acts on B cells to enhance the production of IgA antibodies [23] and the application of recombinant IL-5 on to human nasal mucosa increases the production of IgA antibodies in nasal secretion [24]. Inhalation of allergen could increase the concentration of IL-5 in airway secretions from atopic asthmatics [25]. Capsular polysaccharide antigen of S. pneumoniae is one of the thymus-independent antigens and a humoral immune response against this antigen does not depend on the presence of antigen-specific T cells, although T-cell factors are required [26]. It may be speculated that activation of house dust mite-specific T-lymphocytes with natural allergen exposure results in overproduction of IL-5, stimulating both house dust mite-specific B-cells and S. pneumoniae-specific B-cells in the bronchial mucosa to produce specific IgA antibodies. The significant close quantitative correlations between specific IgA antibodies to house dust mite antigen and pneumococcal polysaccharide antigen in this study also support this hypothesis.

A significant correlation between sputum eosinophilia and B-cell counts in asthmatic patients has been reported [17] and the levels of soluble marker for eosinophil activation (ECP) correlated significantly with total IgA levels in induced sputum of asthmatic patients [10, 11]. These results suggest a possible contribution of B-cell and IgA antibodies to the pathogenesis of eosinophilic inflammation of the asthmatic airway; however, their exact role in the pathogenesis of asthma is still undefined. Several in vitro data have suggested that immunoglobulins in respiratory secretions may participate in eosinophil degranulation in respiratory allergic diseases [8], although only a few in vivo data have supported this hypothesis [10, 11]. In in vitro conditions, immobilization of immunoglobulins on a solid phase is essential in immunoglobulin-induced eosinophil degranulation and plastic beads have been used for this purpose [8]. However, solid-phase matrices to immobilize immunoglobulins for eosinophil degranulation were not defined in the asthmatic airway.

Allergens present in the airway mucosa could be a potential candidate. The allergen coated on a microtitre plate can degranulate eosinophils in the presence of sera from patients sensitized to the same allergen [27]. Moreover, the amounts of eosinophil degranulation correlated with the level of allergen-specific antibodies in the serum and depletion of allergen-specific antibodies abolished the eosinophil degranulation [27]. Recently, the same group reported that bronchoalveolar lavage (BAL) fluid obtained from ragweed-sensitive asthma after segmental allergen challenge could induce eosinophil degranulation on allergen-coated plastic surfaces and they also revealed significant correlations between amounts of eosinophil degranulation and specific IgG and IgA antibodies to ragweed in BAL fluid [28]. In this study, significant correlations were found between the levels of ECP, and IgA and IgG antibodies to house dust mite in induced sputum from mite-sensitive asthmatic patients. The patients may have been in a state of continuous low-dose allergen challenge, because this study was undertaken during the period when environmental house dust mite allergen reaches peak levels in Korea [14]. These findings suggest that allergen-specific IgG and IgA antibodies in the airway secretion may contribute to allergic inflammation by stimulating eosinophil degranulation.

Sputum induction is a useful, reliable and noninvasive method for obtaining airway secretion from asthmatic patients [29–31]. Most previous studies on bronchial asthma using induced sputum have focused on inflammatory cells and mediators [15, 29–31] and only rarely have studies dealt with allergen-specific antibodies. One of the disadvantages of studying proteins from induced sputum is the contamination of oropharyngeal secretions including saliva. To overcome this problem, two sampling methods were tried. Firstly, mucus plugs were extracted from sputum samples to minimize salivary contamination [29]. Secondly, the entire induced sputum sample was analysed and compared with the results from the saliva [15]. In this study, the latter method was used because selection of a mucus plug provides more viscous samples as well as smaller amounts of specimen for fluid-phase analysis. The levels of total IgA, total IgG, antigen-specific IgA and IgG antibodies in induced sputum samples from asthmatic patients were significantly higher than those in salivary samples. These results suggest that the majority of IgA and IgG antibodies in induced sputum may originate from subglottic airways and sputum induction is a useful noninvasive tool for studying allergen-specific antibodies in airway secretions from asthmatic patients.

In conclusion, specific immunoglobulin A antibodies to both allergen and bacterial antigen were elevated in induced sputum from atopic asthmatics and their possible involvement in eosinophil degranulation was suggested. Further study may be essential to clarify the pathogenetic role of immunoglobulins in asthmatic airways.

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References