The effect of montelukast on eosinophil apoptosis: induced sputum findings of patients with mild persistent asthma

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ABSTRACT

Background: Apoptosis may be important in limiting airway eosinophilia. Treatment with leukotriene antagonists decreases the number of eosinophils in both peripheral blood and sputum.

Aim: To assess the effect of montelukast on eosinophil apoptosis in a group of patients with mild persistent asthma (MPA) and to compare this effect with the apoptotic effect of fluticasone propionate (FP).

Methods: Randomly selected patients with MPA (n = 22) who had not taken anti-inflammatory therapy within the preceding 12 months were included in the study. The sputum induction procedure was performed and the patients were divided into two groups: group 1 (n = 10) received FP 250 µg/day and group 2 (n = 22) received montelukast 10 mg/day orally for 4 weeks. Sputum induction was repeated after the treatment period. The resulting cytospin slides were stained by Wright’s stain and morphologic changes in apoptotic eosinophils were assessed by the use of light microscopy by two blinded expert pathologists. Serum soluble Fas ligand (sFasL) concentrations were measured by an ELISA method at baseline and after treatment in both groups, as well as in a group of healthy subjects.

Results: In within-group comparisons, the apoptotic ratio (AR) increased at the end of the study period in group 1 (p = 0.05). In the group treated with FP the ratio of sputum eosinophils significantly decreased (p = 0.02), and the AR significantly increased (p < 0.005). No differences were found in the two study groups in serum sFasL levels at the end of the treatment period compared with baseline values (p > 0.05).

Conclusion: Our findings demonstrate that 4 weeks’ treatment with a CysLT receptor antagonist (montelukast) resulted in an increase in eosinophil apoptosis comparable to that produced by FP, suggesting that induction of apoptosis may be a potential mechanism for the mode of action of CysLT receptor antagonists in asthma.

Key words: Mild persistent asthma. Eosinophil. Apoptosis. Induced sputum. Montelukast. Fluticasone propionate. Light microscopy. Soluble Fas ligand.

RESUMEN

Contexto: La apoptosis puede ser importante para limitar los eosinófilos de las vías respiratorias. El tratamiento con antagonistas de los leucotrienos reduce el número de eosinófilos tanto en la sangre periférica como en el esputo.

Objetivo: Evaluar el efecto del montelukast (M) sobre la apoptosis eosinofílica en un grupo de pacientes con asma leve persistente (ALP) para compararlo con el efecto apoptótico del propionato de fluticasona (PF).

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Métodos: Para el estudio se seleccionaron aleatoriamente pacientes con ALP (n = 22) que no habían recibido terapia antiinflamatoria durante los 12 meses previos. Tras proceder a la inducción del esputo, los pacientes se dividieron en dos grupos: uno recibió 250 μg/día de propionato de fluticasona (grupo 1, n = 10) y el otro 10 mg/día de M por vía oral (grupo 2, n = 12) durante 4 semanas. Tras el período de tratamiento se repitió la inducción del esputo. Los preparados resultantes obtenidos por citocentrifugación se tiñeron con colorante de Wright y dos patólogos expertos que desconocían el origen de las muestras las observaron al microscopio de luz para evaluar los cambios morfológicos de los eosinófilos apoptóticos. Se midieron las concentraciones séricas del ligando Fas soluble (sFasL) mediante el método ELISA antes y después del tratamiento en ambos grupos, midiéndose simultáneamente en un grupo de sujetos sanos.

Resultados: Comparando los resultados en un mismo grupo, el índice apoptótico (IA) aumentó al final del período de estudio en el grupo 1 (p = 0,05). En el grupo tratado con PF, el índice de eosinófilos en el esputo disminuyó significativamente (p = 0,02), mientras que hubo un aumento significativo del IA (p < 0,005). Los niveles séricos de sFasL no variaron al final del período de tratamiento en comparación con los niveles iniciales en ambos grupos de estudio (p > 0,05).

Conclusión: Nuestros hallazgos demuestran que un tratamiento de cuatro semanas con un antagonista de los receptores de Cys-LT, el montelukast, produce un aumento de la apoptosis eosinofilica comparable al que produce el propionato de fluticasona, lo que sugiere que la inducción de la apoptosis puede ser un mecanismo de acción de los antagonistas de los receptores de Cys-LT en el asma.


MÉTODOS

Patient selection

Randomly selected mild persistent asthmatic patients who admitted to our outpatient clinic between January 2001 and January 2002 were included in the study. Asthma diagnosis was based on a history of recurrent symptoms of wheezing, shortness of breath, cough and demonstration of objective signs of reversible airway obstruction by means of at least 12 % increase in FEV1 after 15 minutes with an in-halation of 250 μg salbutamol or a PC20 methacholine < 8mg/ml as stated by the American Thoracic Society and International Asthma Guidelines[5]. Asthma severity was determined by the frequency of asthma symptoms, pulmonary function tests, and
medication requirements according to the international guidelines by a chest physician/allergolog. Selected patients had not been taking inhaled corticosteroids, leukotriene receptor antagonists, theophylline and long acting beta2 agonists within the preceding 12 months of the study. All subjects had been free of airway infection for at least 6 weeks before investigation. The study protocol was approved by local ethical committee and all participants gave written informed consent.

Study design

Before randomization subjects’ characteristics were documented by a questionnaire. Allergy prick tests with common allergen extracts were performed at baseline. Following sputum induction procedure patients were randomly divided into two groups whether to take montelukast 10 mg/day orally (Singulair™, Merck Sharp&Dohme) or inhaled fluticasone propionate 250 microg/day (Flixotide™, GlaxoSmithKlein) for 4 weeks. Sputum induction and evaluation was made again after this treatment period. At baseline and in the end of 4 weeks spirometry was made. Following sputum induction and evaluation was made again after this treatment period. At baseline and in the end of 4 weeks spirometry was performed. Serum was collected for soluble Fas ligand assay at baseline and following treatment.

Sputum induction

Sputum induction was performed between 9.00 and 10.00 AM according to the method proposed by Pin and coworkers and slightly modified by Pizzichini and coworkers and adapted to Pizzichini and coworkers’ method. Before inhalation of hypertonic saline solution, all patients inhaled two puffs of salbutamol (200 microg) and 10 minutes later underwent spirometry. Then an aerosol of sterile 3 % saline solution was generated by an ultrasonic nebuliser (particle diameter 0.5-6 microns, 3cc, minimum spray time: 3 dk, oscillation frequency: 1.63 MHz) and inhaled for 7 minutes through a mouthpiece without a valve or nose clip. If the patient could not expectorate at this step the concentration of saline was increased. If the FEV1 fell by 20 % or if troublesome symptoms occurred nebulisation was discontinued. After each period of inhalation subjects were asked to rinse their mouth and throat carefully, swallow the water and blow the nose to minimize contamination with saliva and postnasal drip. Then they were instructed to try to cough sputum into a container.

Sputum examination

Sputum was processed as soon as possible within 15 minutes according to the method proposed by Pizzichini and coworkers. It was pooled into a Petri dish and all portions that macroscopically appeared free of salivary contamination were placed in a 15 ml polystyrene tube using a straight blunted 115 mm forceps. The mixture was vortexed for 15 seconds and gently aspirated in and out of a Pasteur pipette to ensure mixing. To stop the effect of DDT on the cell suspension a further 4 volumes of Dulbecco phosphate-buffered saline (D-PBS) was added. The suspension was filtered through 48 mm nylon gauze to remove cell debris and mucus. The resulting clear suspension was centrifugated at 790 X g for 10 minutes and the supernatant was aspirated and stored in Eppendorf tubes at −70°C for later assay. The pellet was resuspended in a volume of D-PBS, 200-600 microl. Seventy-five microliters of the cell suspension adjusted to 1.0 x 10⁷/ml were placed into caps of Shandon III cytocentrifuge and two coded cytospins were prepared at 450 rpm for 6 minutes. The slides were air-dried and stained by Wright’s stain. Morphologic changes of apoptotic eosinophils were assessed by the use of light microscopy. Presence of cell shrinkage and nuclear coalescence ie, shift from bilobed to monolobed nucleus were used to identify apoptotic eosinophils in cytospin preparations from sputum samples. The features of a normal and apoptotic eosinophil in sputum from an asthmatic study patient are shown in figure 1A and 1B.

Apoptotic eosinophils were estimated by counting 300 cells in each slide and expressed as the apoptotic cell percentage (apoptotic eosinophils/normal + normal, bilobed eosinophils). Slides were coded and cell counts were performed by two expert pathologists who had no knowledge about the clinical characteristics of patients.

Serum soluble Fas ligand assay

Serum soluble Fas ligand concentrations were measured by the use of Enzyme-linked immunosorbent assay (ELISA) (Bender MedSystems, humans-
Fas ligand BMS260/2, Vienna, Austria). Soluble Fas ligand levels of the patients were compared with healthy controls.

Statistical analysis

Data were analysed using the SPSS-PC package, Windows 7.5 version. Results are presented as mean ± SD. Mann-Whitney U test used to study differences between the two independent groups. Comparisons between data from two dependent groups were analysed using Wilcoxon signed-rank test. Values of p < 0.05 were considered statistically significant.

RESULTS

Study groups

A total of 24 patients were included in the study. Following the baseline sputum induction procedure 2 patients were excluded as they couldn’t successfully produce adequate samples. Remaining 22 patients were randomly divided into two groups: Group 1 (n = 10) was treated with fluticasone dipropionate and Group 2 (n = 12) was treated with montelukast. Demographic characteristics of the study groups are shown in table I.

Safety and success of sputum induction

Sputum induction was successfully performed in the study groups without any complication including bronchoconstruction. Nineteen subjects produced adequate samples at the first attempt. The rest with inadequate samples at the first occasion produced better quality sputum at the following second attempt.

Comparison of two groups by means of asthma history and FEV1 values

There was no difference in FEV1 measurements between groups both at baseline and at the end of
4 weeks treatment (table I). Duration of asthma and rate of atopy also did not differ between the groups (table I).

Comparison of eosinophil counts and apoptotic eosinophils

There was no difference in baseline and post treatment eosinophil percentages, apoptotic eosinophil percentages and apoptotic ratio between two groups (p > 0.05). In within group comparisons eosinophil percentage and apoptotic eosinophil percentage did not reach significant levels between baseline and posttreatment evaluations in group 1 (p > 0.05). However apoptotic ratio increased at the end of fluticasone treatment when compared with baseline in the same group (p = 0.05). The ratio of sputum eosinophils decreased significantly (p = 0.02) while there was a significant increase in apoptotic ratio (p < 0.005) in the posttreatment period when compared to baseline values in group 2 (table II).

Comparison of serum soluble Fas Ligand assay

Pretreatment sFas ligand levels were 79.6 ± 64.6 pg/ml, 45.0 ± 55.2 pg/ml and 49.8 ± 60.1 pg/ml in Group 1, Group 2 and control group, respectively. Posttreatment sFas ligand levels were 45.7 ± 62.5 pg/ml and 47.7 ± 59.5 pg/ml in Group 1 and Group 2, respectively. Serum sFasL levels were not different between the two groups both in the pre (p = 0.36) and post (p = 0.36) treatment periods. Furthermore there was no difference in serum sFasL levels between pre and post treatment periods in both groups (Group 1: p = 0.59 and Group 2: p = 0.68). There was also no difference in serum sFasL levels between the control group and study groups at baseline (p > 0.05).

DISCUSSION

In this study we have demonstrated that four weeks treatment with montelukast increased eosinophil apoptosis in induced sputum of asthmatic patients with mild persistant asthma. To our knowledge this is the first report demonstrating the effect of montelukast on eosinophil apoptosis in vivo in human mild persistant asthma. Our results suggest eosinophil apoptosis as a mechanism for the reduction of eosinophilic airway inflammation by montelukast in asthma.

Eosinophils infiltrate the bronchial mucosa of asthmatics and are detectable in bronchial tissue specimens and in sputum even during clinical remission. The overall rate of eosinophils in induced sputum was 3.79 ± 3.48 % in our study group (n = 22) with mild persistant asthma. There are few examples of eosinophil ratios from induced sputum samples of asthmatic patients. Eosinophil ratio in induced sputum of patients with mild asthma has been found to vary between 8.5-10.3 % in different studies. How-

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<tr>
<th>Table I</th>
<th>Demographic features of patients</th>
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<tr>
<td></td>
<td>Group 1 (FP) (n = 10)</td>
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<tr>
<td>Age (year mean ± SD)</td>
<td>36.7 ± 12.3</td>
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<tr>
<td>Sex (Female/Male) n (%)</td>
<td>7/3</td>
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<tr>
<td>Atopy (%)</td>
<td>7/10</td>
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<td>Basal FEV1 (% Pred)</td>
<td>86.9 ± 13.8</td>
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<td>Posttreatment FEV1 (% Pred)</td>
<td>93.7 ± 14.8</td>
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<tr>
<td>Asthma duration (%)</td>
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<tr>
<td>&lt;5 yrs</td>
<td>68.7</td>
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<tr>
<td>5-10 yrs</td>
<td>22.2</td>
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<td>&gt;10 yrs</td>
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<tr>
<th>Table II</th>
<th>Differences of eosinophils and apoptotic eosinophils percentages and apoptotic ratios between Group 1 and 2</th>
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<tr>
<td></td>
<td>Group 1 (FP) (n = 10)</td>
</tr>
<tr>
<td>Pretreatment</td>
<td>Posttreatment</td>
</tr>
<tr>
<td>E (%)</td>
<td>4.7 ± 4.3</td>
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<tr>
<td>AE (%)</td>
<td>3.4 ± 5.0</td>
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<tr>
<td>AR</td>
<td>0.32 ± 0.36</td>
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* p < 0.05: statistically significant
E: Eosinophil; AE: Apoptotic eosinophil; AR: Apoptotic ratio.
ever there was an overlap among asthmatics grouped according to disease severity in these studies, thus there were moderate asthmatics in some study groups. Therefore we believe that our data on induced sputum eosinophil ratios in mild persistent asthma is unique, as firstly the study group was a homogenous one in respect to disease severity and secondly they were patients who were not taking any antiinflammatory drug in the preceding 12 months of the study.

Apoptosis is a physiologic form of cell death and is believed to play a key role in the resolution of inflammation. Apoptosis contributes to the resolution of eosinophilic inflammation in asthma by 1) reducing the activation of airway eosinophils and 2) facilitating their removal. In vitro studies have demonstrated that activation of inflammatory cells is reduced when they undergo apoptosis. Furthermore since the membranes of apoptotic cells remain intact, inadvertent leakage of inflammatory mediators from these cells is prevented. This is in marked contrast to cell death by necrosis, in which the cell membrane ruptures and intracellular mediators are released. This feature of apoptosis is particularly significant for eosinophils which contain cytoxic proteins such as ECP that are capable of damaging airway tissue. Although there are reports on the presence of apoptotic eosinophils in bronchial biopsy tissue specimens from asthmatics with different forms of severity, data on apoptotic eosinophils in induced sputum from asthmatics are very few in the literature. Recently it has been reported that apoptotic eosinophils were also detectable in cytospin slides of induced sputum from patients with seasonal allergic rhinitis. In this study we were able to detect apoptotic eosinophils in induced sputum of patients with mild persistent asthma who were not taking antiinflammatory therapy.

Apoptotic eosinophils have been recently detected in the sputum of asthmatic patients during clinical exacerbations. Further Foresi et al. found that the number of apoptotic eosinophils was greater in patients with symptomatic asthma compared with mild, asymptomatic disease probably reflecting the greater tissue eosinophilia in the former group. However Vignola and coworkers have reported that the apoptotic ratio for eosinophils in bronchial biopsy tissue specimens from asthmatics tends to decrease with the increase in the asthma severity. The ratio of apoptotic eosinophils in our untreated mild asthmatic patients (n = 22) at baseline was similar to that of symptomatic patients in Foresi and coworkers study suggesting that apoptosis operates as a controlling mechanism for eosinophilic inflammation in the airways of patients with symptomatic asthma in need of antiinflammatory treatment.

Corticosteroids have been reported to induce eosinophil apoptosis in vitro with potencies consistent with the potency of their other antiinflammatory action in normal and asthmatic subjects. However in vivo data on the effects of inhaled steroids on eosinophil apoptosis is controversial. A decrease in the number of sputum eosinophils associated with an increased apoptotic index after inhaled beclamethasone has been reported in patients with mild asthma. Furthermore, Kankaanranta and coworkers reported that use of inhaled glucocorticoids reversed the delayed eosinophil apoptosis in peripheral blood in asthma. Conversely Sano et al. demonstrated that in vitro treatment of isolated human eosinophils with fluticasone propionate caused decreased eosinophil survival and LTC4 synthesis by a mechanism not related to cellular apoptosis. The use of 250 µg/day of inhaled fluticasone propionate apoptotic ratio in sputum eosinophils increased approximately two fold in our patients. However the dose of fluticasone propionate was relatively low in our study compared to doses used in studies showing increased apoptosis with inhaled steroids. We believe that higher doses of fluticasone propionate could have a more significant effect on both eosinophil ratios and apoptotic index in the sputum.

Administration of leukotriene receptor antagonists results in improvement of pulmonary function and better control of asthmatic symptoms. It has also been demonstrated that leukotriene antagonists reduce eosinophils in blood by about 50 % and decrease airway eosinophils 50-60 % after 2-4 weeks treatment in addition to improving clinical parameters, but the exact mechanism underlying these effects are not fully understood. We demonstrated that treatment with montelukast for 4 weeks resulted in an increase in apoptotic index of sputum eosinophils in patients with mild asthma. Our findings have relevance to the benefit derived from leukotriene receptor antagonists in the treatment of asthma. The clinical efficacy of these compounds has been attributed primarily to inhibition of leukotriene-mediated chemotaxis and bronchoconstriction. The findings in the present study suggest that inhibition of eosinophil survival by inducing apoptosis should also be considered as a potential additional mode of action of these drugs in asthma.

Fas (APO-1/CD95) is a membrane glycoprotein belonging to the tumor necrosis factor receptor family. Fas antigen is expressed on the surface of a variety of cell types, including eosinophils. The Fas ligand is also a member of the tumor necrosis factor receptor family and induces apoptosis in Fas bearing cells. The membrane bound human Fas ligand was found to be converted to a soluble form (sFas L) by the action of
a matrix metalloproteinase like enzyme. Fas ligand/Fas receptor molecular interactions have been implicated as having an important function for the regulation of eosinophil apoptosis. It was shown that the number of T lymphocytes and other Fas and Fas ligand bearing cells in the bronchial mucosa of steroid untreated asthmatic patients were not different from that of control subjects. However steroid treated asthmatics exhibited increased expression of Fas and Fas ligand in their bronchial epithelium when compared to control subjects and untreated patients. Authors suggested that the decrease in the inflammatory cell infiltrate in the bronchial submucosa of patients treated with steroids might in part be due to the increased expression of Fas on bronchial epithelial cells by the action steroids. However we observed no change in serum sFasL levels following treatment with both montelukast and fluticasone in our study.

In conclusion our findings demonstrate that four weeks treatment with a Cyst LT receptor antagonist – montelukast – resulted in an increase in eosinophil apoptosis comparable to that of fluticasone propionate in mild persistent asthmatic patients, suggesting that induction of apoptosis may be a potential mechanism for mode of action of these drugs in asthma.

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REFERENCES