Alveolar macrophage-induced suppression of peripheral blood mononuclear cell responsiveness is reversed by in vitro allergen exposure in bronchial asthma


ABSTRACT: Little information is available on the specific role of alveolar macrophages (AMs) in modulating local cellular reactions to inhaled allergens in atopic asthma.

We investigated the influence of alveolar macrophages obtained by bronchoalveolar lavage (BAL) on the proliferative responses of lavage and peripheral lymphocytes from 12 patients with atopic asthma, 6 nonasthmatic symptomatic atopic subjects, and 6 nonatopic normal volunteers, in the context of in vitro exposure to relevant and nonrelevant allergens.

Fresh nonadherent bronchoalveolar lavage cells from atopic asthmatic patients, depleted of alveolar macrophages, proliferated spontaneously more than nonadherent bronchoalveolar lavage cells from normal subjects. Addition of autologous asthmatic alveolar macrophages reduced this endogenous "activation". Asthmatic and normal alveolar macrophages also inhibited phytohaemagglutinin-stimulated proliferation of both autologous and allogeneic nonadherent peripheral blood mononuclear cells (PBMC). In contrast, autologous asthmatic alveolar macrophages induced strong proliferation of peripheral blood mononuclear cells when stimulated with allergen to which the patient was skin test and radio allergosorbent test (RAST) reactive; however, no response was seen with allergens to which the patient was insensitive. No such allergen-specific proliferation was seen with alveolar macrophages from nonasthmatic atopic subjects.

These data support the presence of functionally-active alveolar macrophages within the airways of atopic asthmatic patients, that under normal stable conditions suppress the induction of peripheral blood mononuclear cell responses, and which on contact with specific allergen appear to switch to inducer alveolar macrophages, with consequent peripheral blood mononuclear cell hyperactivation.


Atopic asthma results from inappropriate cellular immune reactions to nonpathogenic airborne allergens [1]. A multistage inflammatory response develops within the airway, together with the release of biologically active mediators [2–4], and secondary pathological changes in the mucosa [5]. Whilst recognizing the importance of the T-cell system in specific allergen recognition, the critical regulation of these events in allergic asthma may depend on the alveolar macrophage (AM).

The AM has been observed to be involved in a dichotomy of roles in its defence of the local microenvironment. This versatile character allows the macrophage to process inhaled allergen, which it then presents in a "modified and recognizable" form to primed T-lymphocytes [6, 7]. Consequently, the macrophage itself becomes the target of a positive feedback loop through the production of macrophage-activation lymphokines, which heighten its capacity to terminate the offending stimulus [8]. In addition, the AM may also act as a store of allergen through its ability to endocytose, and then return allergen to its surface. The AM therefore possesses the requisite properties for the initiation and prolongation of an allergic and inflammatory reaction, such as bronchial asthma [9, 10].

The question arises as to whether AMs can determine the fate of allergen-driven T-cell activation in asthma. Mounting evidence supports the notion that AMs under steady-state conditions can exert a protective effect on the local milieu by preventing an immunological overreaction to the large amounts of inhaled antigens [11]. Recently, it was shown that the functional capacity of the AM is such that it can regulate the induction and
strength of acquired T-cell responses in the human lung. Thus, AMs can not only induce, but also actively suppress T-cell activation and proliferation in health [12], and during inflammatory states [13].

Overall, little information is available on the precise role of AMs in promoting and regulating the inflammatory response to inhaled allergen that ultimately leads to the asthmatic attack. In the present study, we investigate the influence of AMs obtained from atopic asthmatics and nonasthmatic atopic healthy subjects on T-cell proliferative responses, in the context of in vitro exposure to relevant and nonrelevant allergens.

Material and methods

Subjects

Twelve patients with atopic asthma were recruited, all were nonsmokers, 8 females and 4 males, mean±SEM age 27±3 yrs. Bronchial asthma was diagnosed by clinical history and confirmed by measurement of airways obstruction (reversible by an inhaled beta2-agonist, terbutaline, and bronchial responsiveness to methacholine challenge (mean log provocative concentration of methacholine producing a 20% fall in forced expiratory volume in one second (PC20) 0.27±0.02 mg·ml⁻¹). These patients were free of symptoms at the time of the study, had a resting forced expiratory volume in one second (FEV1) of 74±8.6% predicted, and had not suffered any acute attacks in the preceding 3 months. Seven of the 12 patients were taking inhaled beta2-agonists only; these patients were asked to stop their medication at least one day before bronchoscopy. The remaining five patients were "newly diagnosed" asthmatics, who had not previously received any form of medication. None of the 12 asthmatics was receiving any immunosuppressive or other therapy (inhaled or oral).

Atopy was defined by clinical symptoms, a cutaneous wheal response of >5 mm diameter to at least two of the following aeroallergens (house dust, house dust mite, grass and tree pollens, cat fur and dog hairs, and Aspergillus fumigatus), high total immunoglobulin E (IgE) levels (281±7.5 IU·ml⁻¹) (paper disk radioimmunoassay technique, Pharmacia Laboratories; normal <100 IU·ml⁻¹), and positive specific serum IgE antibodies (radioallergosorbent test (RAST), Pharmacia Laboratories).

The control populations consisted of: 1) six nonasthmatic atopic subjects (4 males and 2 females; 28±2 yrs), who had seasonal symptoms of rhinitis, itchy eyes and/or eczema (mean IgE 165±9.0 IU·ml⁻¹), but were not on any current medication (including immunosuppressants); and 2) six nonatopic normal volunteers, who had no history of asthma or atopy (3 males and 3 females; 29±3 yrs; negative skin prick tests; mean IgE 65.0±2.3 IU·ml⁻¹). All control subjects were nonsmokers with normal pulmonary function tests, and did not reach a PC₂₀ on methacholine challenge; none had suffered from any viral illness in the two weeks preceding the study.

Formal written consent was obtained from all subjects recruited; the study had received prior approval by the local Ethics Committee.

Bronchoalveolar lavage (BAL)

BAL was performed using a 6 mm fiberoptic flexible bronchoscope (Olympus BT-IT20D; Olympus Corp., UK). All subjects were premedicated with intravenous 0.6 mg of atropine sulphate and 2–5 mg midazolam (Hypnovel), as required, 15 min prior to the procedure. After local anaesthesia with 2% lignocaine, the right middle lobe was lavaged with successive 20 ml aliquots of sterile buffered 0.9% isotonic saline to a total volume of 180 ml. The lavage fluid was gently aspirated after each aliquot, and collected into a sterile siliconized glass bottle maintained at 4°C.

Processing of BAL samples

The lavage fluid was filtered through a single layer of coarse gauze and centrifuged at 480×g at 4°C for 5 min. The cell pellet was then washed twice in RPMI 1640 medium (Flow Laboratories, Paisley, Scotland), after which the cells were counted in a modified Neu-bauer haemocytometer and viability assessed by cellular exclusion of trypan blue. The final cell concentration in each sample was adjusted to 1×10⁶ cells·ml⁻¹ using supplemented RPMI 1640, containing 1.25% 200 mM L-glutamine, 10% autologous human serum, 100 µg·ml⁻¹ streptomycin and 100 IU·ml⁻¹ penicillin.

Separation of BAL cells

The above cell suspension was plated onto sterile plastic 85 mm diameter tissue culture grade Petri dishes (Nunc, Denmark), with a total of 4–6×10⁶ cells on each and a medium suspension depth of 3 mm. These were incubated for 2 h at 37°C in an atmosphere of 5% humidified CO₂. The supernatant containing the nonadherent cell population was then collected, and the plate washed three times with medium to remove any further non-adherent cells. The adherent cells were gently scraped off the plates using a sterile "rubber policeman." The adherent and nonadherent cell populations were centrifuged at 480×g, 4°C for 5 min, resuspended in medium at 1×10⁶ cells·ml⁻¹, and kept on ice until required.

Cytospin preparation

Cytospins were prepared on a Shandon Cytospin 2 using 2×10⁶ cells in 100 µl aliquots of each of the above cell suspensions. One cytospin from each sample was stained for morphology, whilst the remainder were air-dried for one hour at room temperature, fixed in a 1:1 mixture of chloroform-acetone for 10 min, wrapped in plastic film and stored at -20°C until use. Cell morphology was determined using a Diff-Quik (Dade Diagnostics, UK) differential white cell stain.
**Immunocytological analysis**

Macroaggregation phenotype in both adherent and non-adherent lavage cell samples was determined by two monoclonal antibodies EBM11 (which identifies all cells of the monocyte-macrophage lineage; Dakopatts, Denmark), and UCHM1 (which identifies antigen present on the majority of blood monocytes; P. Beverley, London, UK). The proportion of AMs expressing human leucocyte antigen-DR (HLA-DR) was investigated in both cell fractions, using a mouse immunoglobulin G (IgG) anti-HLA-DR monoclonal antibody (Becton-Dickinson, UK). A standardized immunoperoxidase technique was used [14].

**Peripheral blood mononuclear cells (PBMC)**

All subjects had 20 ml of peripheral blood taken by venepuncture at the same time as the BAL. PBMC were separated on a Ficoll-Hypaque gradient, washed twice in Hank's balanced salt solution, and then resuspended in supplemented RPMI 1640. The PBMC suspension was counted and viability assessed. Plastic plate adherence was carried out, as with the lavage cells above; the separated PBMC cell fractions obtained were adjusted to a final cell concentration of 1×10⁴ cells·ml⁻¹.

**Cell cultures**

All cultures (a–d) described below were set up in triplicate in flat-bottomed microtitre wells, and incubated at 37°C in an atmosphere of 5% humidified CO₂ for 6 days. Equal volumes of supernatant (50 µl) were gently aspirated from each culture without disturbing the cell pellet, and replaced with an equivalent amount of fresh supplemented RPMI prior to pulsing with 1 µCi ³H-thymidine (³H-Tdr) (Amersham, UK; 5 Ci·mmol⁻¹). The cells were then incubated for a further 6 h, and harvested using a semi-automatic cell-harvester (Titertek-Mark III, Laboratory Inc., Mclean, VA, USA). The amount of incorporated radioactivity was measured in a liquid scintillation counter, and expressed as average counts per minute (cpm) of triplicate cultures. Blank wells consistently gave recordings of less than 40 cpm. The culture supernatants collected were stored at -70°C until further use.

The culture experiments were set up as follows:

a) **BAL cell cultures.** Isolated plastic adherent and non-adherent BAL cells from six asthmatic atopic and six normal nonatopic recruits were cultured on their own and readmixed using a fixed 1:10 adherent:non-adherent cell ratio (1×10⁴ adherent and 1×10⁴ nonadherent cells in each well).

b) **Response to mitogen.** In other cultures, nonadherent PBMC (1×10⁵) were stimulated with the T-cell mitogen phytohaemagglutinin (PHA) at the recommended concentration of 2.5 µg·well⁻¹ (PHA-P, Difco Laboratories); autologous adherent BAL cells (1×10⁴) were added to some of these cultures.

c) **Response to allergen.** In other experiments, control and allergen-treated cultures of adherent and nonadherent BAL cells and PBMC from six other asthmatic atopic and six nonasthmatic symptomatic atopic recruits were set up as above. Cultures of adherent BAL cells (1×10⁴) with autologous nonadherent (1×10⁴) and adherent (1×10⁴) PBMC with or without allergen were also set up. Allergen sensitivity of each subject was determined at the time of recruitment (q.v. above). Dose-response studies for each allergen suspension had been determined in previous experiments (data not shown); the allergen concentration giving the maximum proliferative response was chosen for stimulating the cell cultures in this study (grass pollen 1×10³ U·ml⁻¹; house dust mite 1×10³ U·ml⁻¹; cat fur 1×10³ U·ml⁻¹; *Aspergillus fumigatus* 3 µg·ml⁻¹).

d) **Effect of culture supernatants.** In a separate set of experiments, nonadherent (1×10⁴ cells·well⁻¹) and adherent (1×10⁴ cells·well⁻¹) PBMC obtained from a nonsmoking, nonasthmatic, atopic subject were cultured alone, and in the presence of 100 µl of supernatant collected from the above unstimulated and allergen-stimulated cell cultures of four asthmatic and four nonasthmatic atopic recruits.

**Measurement of prostaglandins**

Prostaglandin E₂ (PGE₂) and the stable product of prostaglandin I₂ (PGI₂) (6-oxo-PGF₁α) were measured in supernatants obtained from asthmatic nonadherent and adherent BAL cells, and co-cultures of adherent BAL cells and autologous nonadherent PBMC, using previously validated and well-established radio-immunooassay techniques [15]. Due to the high specificity and lack of interference of culture medium, no prior extraction and purification was required. The lower limit of prostaglandin detection in these assays was 5 pg. Antibodies against PGE₂ and prostaglandin F₂ (PGF₂α) of high specificity were purchased from Capell Laboratories (West Chester, PA, USA). ³H-PGE₂ (120 Ci·mmol⁻¹) and PGF₂α were purchased from New England Nuclear (Dreieich, West Germany), and unlabelled ligand from Cayman Chemical Co. (Palo Alto, CA, USA).

**Statistical analysis**

The results are expressed as the mean±SEM. Significance between results was determined using Wilcoxon's test or the Mann-Whitney U-test. A probability value of ≤0.05 was taken to indicate statistical significance.

**Results**

**Bronchoalveolar lavage cell recovery**

The percentage return of the total lavage fluid instilled was 69±7.4% in the asthmatic group, 72±4.3% in the
nonasthmatic atopics, and 73±9.6% in normal subjects. The total BAL cell yield in the normal and nonasthmatic atopic volunteers was 9.7±1.1×10^6 cells (absolute number of AMs 9.2±1.1×10^6 cells) and 9.1±0.8×10^6 cells (absolute number of AMs 8.3±0.7×10^6 cells), respectively. The total BAL cell yield in asthmatic patients was 8.2±1.2×10^6, of which the absolute number of AMs was 7.0±0.9×10^6. The mean proportion of lymphocytes in asthmatic BAL was 10.2±5.1%, compared to 5.6±3.2% and 4±3.5%, respectively, in normal and symptomatic atopic lavage. The eosinophil count was higher (p<0.01) in the asthmatic patients (4.6±1.1%) than in the normal (<1%) and nonasthmatic symptomatic atopic (1.3±0.2%) groups.

The viability of all lavage cells throughout all experiments was persistently >90% by trypan blue exclusion. In all groups, plastic adherent cells from both BAL and PBMC were >95% monocytoid in morphology. Adherent lavage cell samples were >97% EBM11 positive but <1% UCHM1 positive; nonadherent lavage cells were <3% EBM11 positive and <1% UCHM1 positive. Over 90% of alveolar macrophages were HLA-DR positive, with no significant differences amongst the three subject populations.

**Cell cultures**

a) **BAL cell cultures.** The endogenous reactivity of nonadherent and adherent lavage cells from asthmatics was significantly higher than in normals (p<0.0001) (fig. 1). In the asthmatic group, nonadherent BAL proliferation was suppressed from 2,100±98 cpm 3H-Tdr uptake to 1,118±53 cpm (p<0.0001) on admixture with autologous adherent lavage cells; this reaction was not apparent in the normal group (fig. 1).

b) **Response to mitogen.** Unstimulated nonadherent PBMC proliferation was higher in the asthmatic group (p<0.0001) (fig. 2). In both test groups, addition of adherent lavage cells to PHA-stimulated autologous nonadherent PBMC produced a reduction in thymidine incorporation (fig. 2). This suppression was more pronounced with asthmatic adherent lavage cells (109%) compared to normal (76%) (calculated as a percentage of the initial PHA-induced proliferation of PBMC) (p<0.0001) (fig. 2).

c) **Response to allergen.** In separate experiments, we compared the effects of adherent BAL cells from six atopic asthmatics on the proliferative responses of autologous readmixed nonadherent and adherent PBMC cultured: 1) with allergen to which the patient was reactive; 2) with allergen to which the patient had no skin test or RAST reactivity; and 3) with any allergen. Similar control cultures were set up with cells from six nonasthmatic symptomatic atopic recruits.

In both groups, addition of adherent BAL cells suppressed the spontaneous proliferation of unstimulated PBMC (fig. 3a and b). In both groups, stimulation with relevant allergen increased autologous PBMC proliferation. However, in the asthmatic but not in the nonasthmatic atopic subjects, the presence of adherent BAL cells in PBMC cultures stimulated with the allergen to which the patients were reactive resulted in enhanced PBMC proliferation by a mean of eight-fold
Fig. 3. – The effect of allergen to which the patient was skin and radioallergosorbent test (RAST) reactive (in this case grass pollen), and irrelevant allergen (negative skin and RAST reactivity, in this case *Aspergillus fumigatus*) on the proliferation of readmixed nonadherent and adherent peripheral blood mononuclear cells (PBMC) is tested in the presence and absence of adherent bronchoalveolar lavage (BAL) cells (AdhBAL). The final concentration of grass pollen used was $1 \times 10^4$ U·ml$^{-1}$, and that of *Aspergillus fumigatus* $3$ µg·ml$^{-1}$. All cultures were incubated for 6 days, with $^{3}$H-Tdr incorporation measured over the last 6 h. Each bar indicates the results of mean±SEM incorporated $^{3}$H-Tdr in counts·min$^{-1}$ for: a) six nonasthmatic atopic patients; and b) six asthmatic atopic patients. Please note the different scales in each figure. 

- PBMC
- PBMC + AdhBAL
- PBMC + *A. fumigatus*
- PBMC + *A. fumigatus* + AdhBAL
- PBMC + grass
- PBMC + grass + AdhBAL

Fig. 4. – The effect of allergen to which the patient was skin and RAST reactive, and irrelevant allergen (negative skin and RAST reactivity) on the proliferation of readmixed nonadherent and adherent peripheral blood mononuclear cells (PBMC) in the presence and absence of adherent BAL cells (AdhBAL) is shown for each of the six atopic asthmatic subjects. Each bar indicates the results of incorporated $^{3}$H-Tdr in counts·min$^{-1}$ for each culture for a single patient (subject number above bar). All cultures were incubated for 6 days, with $^{3}$H-Tdr incorporation measured over the last 6 h. The final concentration of grass pollen used was $1 \times 10^4$ U·ml$^{-1}$, house dust mite (HDM) $1 \times 10^4$ U·ml$^{-1}$, cat fur $1 \times 10^3$·ml$^{-1}$, and *Aspergillus fumigatus* (*A. fum*) $3$ µg·ml$^{-1}$. For abbreviations see legend to figure 3. This figure should be analysed in association with table 1.
This proliferation was more than double that observed in relevant allergen-treated PBMC cultures without adherent BAL cells (p<0.0001) (fig. 3b). In the nonasthmatic group the presence of adherent BAL cells made no difference to relevant allergen-driven PBMC proliferation (p<0.12) (fig. 3a). The dramatic enhancing effect on contact with allergen to which the asthmatic was skin test and RAST reactive was seen consistently in all six asthmatic patients, but not with irrelevant allergen (table 1, and fig. 4 a–d). Interestingly, this reaction to relevant allergen was not observed in stimulated asthmatic non-adherent and adherent BAL cultures in the absence of PBMC (data not shown).

**Prostaglandin analysis**

No PGE$_2$ or PGF$_1$ was detected in any of the supernatants obtained from asthmatic nonadherent and adherent BAL cells, and co-cultures of adherent BAL and autologous nonadherent PBMC. Positive controls set up with "prostaglandin-spiked" preparations all detected the presence of PGE$_2$ or PGF$_1$, as appropriate.

**Table 1.** – Allergen reactivity of asthmatic subjects as studied by cutaneous wheal response and RAST analysis

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<th>Subject</th>
<th>Allergen tested</th>
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<td>No.</td>
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RAST: radioallergosorbent test; +: relevant; -: irrelevant.

**Fig. 5.** – The effect of supernatants (Sup) from allergen (Ag)-stimulated and -unstimulated cultures of asthmatic adherent bronchial lavage (AdhBAL) cells on their own, and co-cultures of asthmatic adherent lavage and autologous peripheral blood mononuclear cells (AdhBAL+PBMC) on the proliferation of allogeneic peripheral blood mononuclear cells (AlloPBMC) from a nonsmoking normal subject. All cultures were incubated for 6 days, with $^3$H-Tdr incorporation measured over the last 6 h. Each bar indicates the results of mean±SEM incorporated $^3$H-Tdr in counts·min$^{-1}$ for four atopic asthmatic patients. As: asthmatic; Sup: supernatant. : AlloPBMC; : AlloPBMC + As (AdhBAL) sup; : AlloPBMC + As (AdhBAL + Ag) sup; : AlloPBMC + As (AdhBAL + PBMC) sup; : AlloPBMC + As (AdhBAL + PBMC + Ag) sup.

**Prostaglandin analysis**

No PGE$_2$ or PGF$_1$ was detected in any of the supernatants obtained from asthmatic nonadherent and adherent BAL cells, and co-cultures of adherent BAL and autologous nonadherent PBMC. Positive controls set up with "prostaglandin-spiked" preparations all detected the presence of PGE$_2$ or PGF$_1$, as appropriate.

**Discussion**

In this study, we have demonstrated that nonadherent lavage cells from stable atopic asthmatic patients proliferate spontaneously more than the corresponding cell fraction from normal donors; and that this can be suppressed by readmixture with the adherent fraction, the majority of the cells being alveolar macrophages (AMs) as determined by morphological and phenotypic examination. This suppressive effect was also observed in asthmatic and control donors, when the AM fraction was added to autologous nonadherent PBMC stimulated in vitro by the mitogen PHA. However, when PBMC from asthmatic patients were cultured with the allergen to which the patient was clinically, as well as skin and RAST test reactive, addition of autologous AMs produced a large increase in cell proliferation in all subjects tested. Such responses did not occur in any of the nonasthmatic control donors, even with allergen to which they were reactive.

Several previous studies have shown that AMs from normal subjects have suppressive activity [16–18]. There is also evidence that under persistent antigen stimulation, such as in smokers [19], and in infective lung disease [20], AMs appear to be more potent suppressors than AMs from healthy subjects. These data suggest that chronic antigenic stimulation favours the expansion of a functionally suppressive AM subpopulation, although AUBAS et al. [21] reported a dose-dependent modulating effect of AMs with decreased suppressor AM activity in asthmatics at certain AM to PBMC cell culture ratios. Supportive evidence for our hypothesis is obtained from recent studies on sarcoid patients, that clearly show the emergence of functional suppressor macrophages within the AM population in proportion to the disease activity [13, 22]. Functional reconstitution experiments performed in vitro show that, whatever the
cell numbers used, these isolated AMs actively down-regulate the induction of T-cell responses set up by other stimulator macrophages [12, 13]. These observations are also supported in animal models [23].

Our study demonstrates that the endogenous reactivity of isolated adherent BAL cells from asthmatics is higher than equivalent normal cells (fig. 1). This observation could reflect a higher "state of intrinsic activation" within the macrophage population in the asthmatic lung [24, 25]. The increased spontaneous proliferation of AM-depleted BAL cells from asthmatics (fig. 1) would be consistent with a chronic inflammatory response occurring in the lungs of asthmatic patients. Interestingly, readmixture with autologous AMs (despite their increased intrinsic activation) reduces both this proliferation and that induced by mitogen. It is, thus, not inconceivable to suggest the existence of an in situ control for local T-cell responses within the asthmatic lung, in which the heterogeneity of AM accessory cell function may be important.

It has been suggested that suppression by AMs is mediated by soluble factors, such as prostaglandins [26]. In this present study, although AM supernatants suppressed PBMC proliferation (fig. 5) we were unable to detect any PGE2 and PGF2α, raising the possibility that other suppressor factors could be involved. In separate studies, we have found that AMs from stable asthmatics spontaneously release sizeable amounts of tumour necrosis factor-α, a mediator known to inhibit T-cell proliferation [27]. This would not preclude a contribution by other inhibitory mediators within the inflammatory cocktail in the asthmatic lung.

Peripheral blood mononuclear cells from asthmatic patients showed small proliferative responses to those allergens to which they were skin test and RAST-reactive, as has been reported elsewhere. However, addition of autologous AMs, rather than suppressing these reactions, resulted in a marked increase in PBMC proliferation. This effect was not observed with AMs from nonasthmatic atopic subjects. Alveolar macrophages are known to be able to present antigens, although there are differences in antigen presenting capacity between AMs and circulating monocytes, which may reflect differences in glycosylation of major histocompatibility complex (MHC) Class II molecules [28]. The apparent selectivity of the enhancing effect seen in atopic asthmatics, but not in nonasthmatic atopics, for the clinically relevant allergens suggests that asthmatic AMs are able to process those same allergens to a highly immunogenic form.

Nonasthmatic control atopics do not show similar stimulation to relevant allergen. This is clearly not due to underlying lack of reactivity to allergen by their PBMC (fig. 3). The results in the nonasthmatic atopic group could reflect the negative clinical expression of atopy in their lungs (symptoms were restricted to the skin, eyes or nose); in addition, they had no evidence of airway hyperreactivity as judged by methacholine challenge. It is interesting that the enhancing effect of asthmatic AMs was only seen with circulating PBMC but not with lavage lymphocytes. The lack of response with lavage lymphocytes may reflect the fact that all our asthmatic patients suffered from seasonal or "contact" asthma, but were studied at times when natural airborne allergen concentrations were low. Moreover, none had experienced an acute asthmatic episode within the preceding 3 months. In other studies, it has been noted that influenza-reactive cells disappear transiently from the blood following influenza immunization, an effect interpreted as being due to the selective localization of antigen-reactive cells at the site of antigen [29]. It may well be, therefore, that in asthma allergen-reactive cells migrate from the airway to the circulation at times when inhaled allergen concentrations are low. In support, sensitized T-cells are readily identified in the peripheral blood of atopic asthmatics, which following bronchial allergen challenge are actively recruited and selectively retained in the lung [30–32].

Evidence shows that functional heterogeneity within AMs may be reflected in cell phenotype and density [13, 33, 34], which may alter with the onset and activity of disease. Chanez et al. [35] recently described a population of hypodense AMs in the lavage of stable asthmatic patients, distinct from the higher density AM fractions found in normal subjects. We have shown phenotypic differences between AMs from patients with active sarcoidosis and normal individuals [36]. Moreover, the phenotypic profile of sarcoidosis varies with disease activity [22], and is modified by corticosteroid therapy [37].

Whilst supernatants of allergen-pulsed AMs also enhanced PBMC proliferation, it is unclear whether this is due to carry-over of highly immunogenic processed allergen or to the production by pulsed AMs of immunoenhancing mediators. Studies on the cytokine profile produced by allergen-pulsed AMs are in progress.

Our observations argue against the suggestion of Aubus et al. [21] of "a general decreased functional activity" within the AM population in asthma. On the contrary, our study suggests that the asthmatic AM population may consist of a dynamic system of functionally suppressor and enhancer AM subpopulations, as demonstrated in other studies [12, 13], capable of serving as "accessory cells" to lung T-cells and modulating the nature of the lymphocyte response to aeroallergens. Future studies will focus on the identification of the specific features of these AM subpopulations in asthmatic patients; this could lead to specific targeting of therapy to the enhancer AM population, with abortion of induction of the cellular hypersensitivity cascade which leads to the clinical asthmatic reaction.

References

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