Desensitisation of neutrophil responses by systemic interleukin 8 in cystic fibrosis

Yalei Dai, Taraneh P Dean, Martin K Church, John O Warner, Janis K Shute

Abstract

Background - Inflammation associated with neutrophil infiltration is a commonly observed feature of children with cystic fibrosis. Production of the major neutrophil chemotactic cytokine interleukin 8 (IL-8) is potentially of great importance in the pathology of cystic fibrosis. Concentrations of IL-8 in both sputum and bronchoalveolar lavage fluid have been found to be higher in children with cystic fibrosis than in controls. The IL-8 induced chemotactic response and numbers of IL-8 receptors on peripheral neutrophils obtained from children with cystic fibrosis have been compared with a control group of children.

Methods - Cells were isolated from 18 patients with cystic fibrosis (aged 4–20 years) and 13 controls (aged 5–12 years) by dextran centrifugation followed by separation on Lymphoprep. Chemotaxis was assayed using multiwell microchemotaxis chambers and 5 μm polycarbonate filters. Filters were fixed and stained with Haema-Gurr for counting. Results were expressed as numbers of neutrophils per high power field (HPF).

Results - At the optimum concentration (1 x 10^{-8} M) the number of cells migrating were similar for controls (150 (12)/HPF) and for the cystic fibrosis group (140 (14)/HPF). At lower concentrations the numbers of neutrophils migrating were lower for the cystic fibrosis group. Scatchard analysis of [125]I-labelled IL-8 binding revealed lower numbers of receptors on neutrophils from patients with cystic fibrosis (22 000 per cell) than from controls (75 000 per cell).

Conclusions - Reduced responsiveness to IL-8 of neutrophils from patients with cystic fibrosis is associated with receptor desensitisation as a result of exposure to high systemic levels of IL-8.

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Respiratory failure is the ultimate cause of death for most patients with cystic fibrosis and is the result of chronic pulmonary infection and progressive lung injury. The relation between the underlying genetic defect1 and repeated endobronchial infection is not clear, but results in a sustained airway inflammatory response.2,3 Neutrophils are the predominant cells of the alveolar space in patients with cystic fibrosis. It appears that the normally protective and self-limiting inflammatory response is overwhelmed by the continued presence of bacteria and their antigenetically active products in the airways. Activation of neutrophils in endobronchial tissues leads to the release of reactive oxidants and proteases resulting in tissue damage.

Several neutrophil chemoattractants are likely to be involved in the neutrophil infiltration observed in patients with cystic fibrosis. However, the major neutrophil chemotactant in the lung is interleukin 8 (IL-8),4 an 8 kD peptide cytokine. We have recently shown5 that elevated levels of IL-8 in sputum and bronchoalveolar lavage fluid samples from children with cystic fibrosis correlate significantly with disease severity. We also measured significantly higher levels of IL-8 in serum samples from children with cystic fibrosis than normal controls. IL-8 in the blood may derive from stimulated endothelial cells,6 circulating monocytes,7 or T cells.8 Alternatively, IL-8 may diffuse into the circulation from sites of inflammation. The synthesis of IL-8 by various pulmonary cell types including alveolar macrophages,9 bronchial epithelial cells,10 and fibroblasts11 has been demonstrated in vitro. The cellular sources of IL-8 in the airways of patients with cystic fibrosis is, however, unknown. The alveolar macrophage may play a central part in the recruitment of neutrophils to the lung since it produces IL-8 in response to either an exogenous stimulus – that is, bacterial-derived cell wall lipopolysaccharide – or autocrine stimuli such as TNF-α and IL-1β. The latter induces IL-8 production by bronchial epithelial cells and pulmonary fibroblasts, cells which do not respond to lipopolysaccharides. This stimulus specificity indicates a self-amplifying cytokine network for the production of IL-8 in the lung which is dependent on an initial lipopolysaccharide challenge.

The in vitro chemotactic response of peripheral neutrophils from patients with cystic fibrosis has previously been shown to be normal for C5a12 and decreased for LTB413 as the chemoattractant. The hyporesponsiveness to LTB4 in vitro was thought to result from LTB4-induced deactivation of specific surface receptors in vivo. Since IL-8 dynamically regulates its own receptor expression on human neutrophils,14 we investigated the hypothesis that chronic production of IL-8 in cystic fibrosis induces a specific dysfunctional chemotactic response as a result of receptor downregulation.

Methods

MATERIALS
Chloramine-T, sodium metabisulphite, Sephadex G-15, disodium hydrogen phosphate, so-
dium dihydrogen phosphate, and bovine serum albumin (BSA) were purchased from Sigma Chemicals (Poole, UK), and Macrodex from Pharmacia (Milton Keynes, UK). Lymphoprep and Hanks balanced salt solution were obtained from GibCo (Paisley, Scotland). Human recombinant IL-8 for iodination was purchased from Bachem UK Ltd (Saffron Walden, Essex), and human recombinant IL-8 for the determination of non-specific binding was the kind gift of Dr Ivan Lindley (Sandoz Forschungsinstitut, Vienna). 125-labelled iodine in sodium hydroxide solution was purchased from Amersham International (Little Chalfont, UK).

PATIENTS

Chemotaxis experiments

Eighteen patients with cystic fibrosis (10 males, eight females) of mean age 10 (range 4–20) years participated in the study. Thirteen normal controls were recruited from the paediatric day surgery units (12 males and one female) of mean age 7 (range 5–12) years. Control subjects were not enrolled if they had any form of respiratory disorder or current bacterial, viral, or parasitic infection.

IL-8 receptor assays

Six patients with cystic fibrosis (two males, four females) of mean age 10 (range 5–12) years and nine controls (eight males, one female) of mean age 12 (range 6–15) years were recruited for this part of the study.

NEUTROPHIL ISOLATION

Heparinised peripheral venous blood was obtained from all subjects, 5 ml for chemotaxis experiments, 10 ml for receptor assays. Whole blood was mixed with Macrodex (6% dextran in normal saline) in the volume ratio 2:1 and allowed to sediment for 45 minutes at room temperature. The leucocyte rich plasma was collected and layered over an equal volume of Lymphoprep. After centrifugation at 450g for 30 minutes at room temperature the mononuclear cell layer was discarded and the granulocyte pellet was harvested from the tube and washed. Erythrocytes were removed by hypotonic lysis using 0-2% sodium chloride for 15 seconds. Isotonicity was restored by adding an equal volume of 1-6% sodium chloride. The cells were then washed twice more with Hanks’ balanced salt solution (HBSS) without calcium and magnesium and finally resuspended in HBSS with calcium and magnesium for neutrophil chemotaxis and in ice cold PBS/1% BSA without calcium and magnesium for receptor assays. The resulting cell suspension contained more than 96% neutrophils with a viability of more than 95% as assessed by trypan blue exclusion.

NEUTROPHIL CHEMOTAXIS

Chemotaxis assays were performed in duplicate using 48-well microchemotaxis chambers (Neuro Probe, Cabin John, Maryland, USA) with 5 μm pore size polycypryrole-done-free polycarbonate membranes. Twenty five μl of the chemoattractant f-Met-Leu-Phe (fMLP) (10^-6–10^-10 mol/l) or IL-8 (6 × 10^-6–10^-10 mol/l) diluted in HBSS with calcium and magnesium were placed in the bottom chamber and 50 μl of neutrophil suspension (10^6 cells/ml) were added to the top chamber. The chambers were incubated for one hour at 37°C in a humidified incubator with 5% CO2. After incubation, non-migrated cells on top of the filter were scraped off and the filter was fixed in methanol for 10 seconds. The filter was air dried and migrated cells stained with Haema-Gurr stain. Neutrophils which migrated to the lower side of the filter were counted using a 400 × magnification in five random high power fields. Experiments were performed in duplicate for each variable and the mean determined. The results were expressed as number of neutrophils/high power field.

IL-8 RECEPTOR ASSAYS

Human recombinant IL-8 was iodinated by the chloramine-T method as previously described. Adult Iodinated IL-8 was eluted from a 1 ml bed of Sephadex G-15 in PBS/1% BSA, and remaining unbound removed by dialysis overnight against one litre PBS, with one change of buffer, in a small sac of benzoylated dialysis tubing with a molecular weight cut off at 3500 (Sigma Chemicals, UK). Covalent binding of the label was checked by trichloroacetic acid (10%) precipitation and in all preparations there was no free 125I. Iodinated IL-8 has a specific activity of 350–500 Ci/mmol.

Aliquots of neutrophil suspension (10^6 cells) in PBS/1% BSA were pipetted into Eppendorf tubes, and increasing concentrations (0-06–0-8 nmol/l) of 125I-labelled IL-8 in the absence and presence of 1000-fold excess unlabelled IL-8 were added to a final volume of 200 μl. Incubations were for 60 minutes at 4°C to avoid ligand internalisation. After incubation, cells were washed three times with ice cold PBS. Radioactivity was determined in an LKB Wallac multimamma counter (counting efficiency 76%). Non-specific binding was determined in the presence of unlabelled IL-8 and specific binding was defined as total binding minus non-specific binding.

DATA ANALYSIS

All experiments were carried out in duplicate. Statistical analysis was performed on the Minitab computer using the Mann-Whitney test.

Results

CHEMOTAXIS

Chemotactic responses of peripheral neutrophils to IL-8 and fMLP in patients with cystic fibrosis and controls are depicted in
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**Figure 1.** Dose-response curves of human neutrophils (expressed as number of cells/high power field (HPF)) to a range of concentrations of (A) IL-8 and (B) fMLP in the multiwell chemotaxis assay. The results are expressed as mean (SE) from 18 patients with cystic fibrosis (●) and 13 controls (○).

The dose-response curves of both the cystic fibrosis and control group showed maximum migration occurring at $10^{-8}$ mol/l IL-8. The mean (SE) number of cells/high power field migrating at this optimum concentration was not significantly different between the cystic fibrosis and control group (140 (14) and 150 (12), respectively). However, at suboptimal concentrations ($1 \times 10^{-9}$ and $6 \times 10^{-9}$ mol/l) the number of migrating neutrophils from patients with cystic fibrosis was significantly decreased compared with the control group ($p<0.05$).

Figure 1B shows the dose-response curves of both the cystic fibrosis and the control group towards fMLP with maximum migration occurring at $10^{-7}$ mol/l. The number of cells migrated at this optimum concentration was not significantly different between the cystic fibrosis and control group (182 (12) and 182 (16), respectively).

**RECEPTOR ASSAYS**

The dissociation binding constants ($K_d$) and the numbers of IL-8 receptors on the surface of the neutrophils were determined by Scatchard analysis. Figure 2 shows typical Scatchard analyses for neutrophils obtained from one of the patients with cystic fibrosis. It illustrates the binding of $^{125}$I-labelled IL-8 to neutrophils as a function of the radioligand concentration, and it is clear that non-specific binding was less than 5% of the total bound. Curve fitting, assuming the presence of a single class of binding sites, gave mean $K_d$ of 12.7 (4.7) × $10^{-10}$ mol/l and 19.3 (7.2) × $10^{-10}$ mol/l for the cystic fibrosis and control group, respectively. There was no significant difference in the IL-8 receptor affinity between the two groups (fig 3). On the other hand, the patients with cystic fibrosis had a significantly lower number of IL-8 receptors (22 198 (2884) receptors/cell) than the controls (75 698 (15 637) receptors/cell) (fig 4).
Discussion

Chemotactant-induced desensitisation of cellular responses occurs by several mechanisms including decrease in receptor number or downregulation, modification of G protein or effector function, or depletion of messengers or their precursors. Our experiments have shown that the chemotactic responsiveness of neutrophils from patients with cystic fibrosis compared with normal subjects is significantly decreased with respect to IL-8, but not fMLP. We have also shown that the number and Kd of specific IL-8 receptors on neutrophils from normal children is of the same order (75 000/cell with Kd 2 × 10⁻⁸ mol/l) as previously reported – that is, 75 000/cell with Kd 4 × 10⁻⁹ mol/l, and that neutrophils from children with cystic fibrosis express a lower number (22 000/cell) with a single type of high affinity binding which is not significantly different from normal. IL-8 binding to neutrophils rapidly downregulates receptor expression via internalisation of the ligand-receptor complex. Removal of IL-8 from the medium resulted in the rapid reappearance of receptors on the cell surface, and the rapid recycling of IL-8 receptors was therefore proposed to be essential for the chemotactic response of neutrophils.

We have previously shown elevated levels of IL-8 (5 × 10⁻¹⁰) in the serum of patients with cystic fibrosis, indicating that neutrophils in the circulation of patients with cystic fibrosis are chronically exposed to higher concentrations of this cytokine. We speculate, therefore, that the chemotactic hyporesponsiveness of peripheral neutrophils in patients with cystic fibrosis is due to downregulation of surface receptor expression, without a change to a lower affinity state, as a result of prior exposure to elevated levels of free IL-8 in the circulation.

The previously reported decrease in neutrophil chemotactic responsiveness to LTβ₄ in cystic fibrosis was similarly proposed to be consistent with a specific decrease of receptor-mediated response to LTβ₄ rather than a generalised abnormality in neutrophil function, as a result of exposure to local high concentrations of LTβ₄ as cells pass through the pulmonary circulation. The greatest difference between cystic fibrosis and normal neutrophil responses was seen at the optimum concentration of LTβ₄ (10⁻⁹ mol/l). In contrast, we observed a significantly reduced response to IL-8 only at suboptimal concentrations. The accuracy and sensitivity of gradient detection is reduced when receptor density is reduced; therefore, in low attractant concentrations the response of neutrophils from patients with cystic fibrosis is reduced compared with normal. However, at the highest concentration of IL-8 the response of normal and cystic fibrosis neutrophils is not significantly different since detection of steep concentration gradients is independent of the number of receptors, but correlates with the difference in receptor occupancy across the cells as it orientates in the gradient. In view of the steep IL-8 concentration gradient across the lung tissue of patients with cystic fibrosis it is unlikely that receptor downregulation by systemic IL-8 will limit neutrophil recruitment into the lung. As in previous studies we measured no difference in chemotactic responsiveness to fMLP, indicating a specific effect of IL-8. Desensitisation to different chemotactants at moderate concentrations is independent and specific to the chemotactants. Non-specific desensitisation occurs only at high concentrations of ligand – that is, at serum concentrations of IL-8 higher than we measured in cystic fibrosis.

In addition to systemic effects, evidence that in situ generated mediators regulate neutrophil function in inflamed tissue has been reported. Neutrophils infiltrating skin pustules in a patient with relapsing bullosus staphyloderma were dysfunctional compared with circulating neutrophils when chemotaxis, degranulation, respiratory burst, and LTβ₄ production were measured in response to IL-8 and C5a. It was suggested that the impaired responsiveness to proinflammatory stimuli of neutrophils migrating into pustules was the result of specific deactivation for receptor-dependent stimuli. The responses of circulating neutrophils in this patient were normal, indicating that the peripheral (intraepidermal) location of the disease restricted the pathological events to the skin. Intraepidermal accumulation of neutrophils is also characteristic of psoriasis, and several recent studies indicate a role for IL-8 in the inflammatory cell infiltration in this disease. A study of IL-8 receptors in normal and psoriatic neutrophils found a slightly raised receptor density in psoriatic neutrophils which was proposed to contribute to neutrophil accumulation in psoriatic skin. The implication, as for pustulosis, is that IL-8 generated locally influences neutrophil function, and that systemic IL-8 concentrations are not elevated in these conditions.

Free IL-8 is undetectable (10⁻¹² mol/l) in normal serum samples where it is complexed with a specific high affinity autoantibody of the IgG class. IL-8 binding to blood neutrophils would inhibit the neutrophil response to a chemotactic gradient; however, the auto-

![Figure 4](http://www.thorax.bmj.com/content/61/10/554.f4)

**Figure 4** Comparison of the number of IL-8 receptors/cell in six children with cystic fibrosis and six controls (p<0.05). Bars indicate median values.
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antibody prevents the binding of IL-8 to neutrophils after it diffuses into the circulation from the site of production. Following intravenous injection of lipopolysaccharide serum concentrations of free IL-8 have been found to rise to a peak of 500–900 pg/ml (~10–100 mol/l) within three hours. It appears that the rapid response to such a powerful stimulus for IL-8 secretion overwhelms the protective IgG mechanism normally limiting free IL-8 levels. Levels of autoantibody have been shown to adapt to an increase in the circulating load of IL-8 – for example, in rheumatoid arthritis. In cystic fibrosis it appears, however, that either autoantibody levels which are sufficient to limit free IL-8 in normal individuals are ineffective in the presence of sustained release of IL-8 from inflamed lung tissue, or increased levels of autoantibody are induced in patients with cystic fibrosis which are nevertheless unable to neutralize the circulating IL-8 load. We are currently investigating these possibilities.

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