Receptor expression in synovial fluid neutrophils from patients with rheumatoid arthritis

Fiona Watson, John J Robinson, Mark Phelan, Roger C Bucknall, Steven W Edwards

Abstract

Objectives—The aim of this study was to determine if neutrophils isolated from the blood and synovial fluid of patients with rheumatoid arthritis had patterns of receptor expression resembling those of blood neutrophils from controls which had been activated and primed in vitro.

Methods—Fluorescence activated cell sorting was used to measure receptor expression in paired blood and synovial fluid neutrophils from patients and in control neutrophils exposed to phorbol myristate acetate and granulocyte-macrophage colony stimulating factor.

Results—There was no significant difference in the patterns of receptor expression in blood neutrophils from patients and healthy controls, but neutrophils in the synovial fluid had been primed and activated within the joint. About 50% of rheumatoid synovial fluid neutrophil samples expressed Fcγ RI, a high affinity receptor for monomeric IgG, which is only expressed in neutrophils exposed to cytokines.

Conclusions—Synovial fluid neutrophils are activated and primed within the inflamed joint and hence their ability to respond to activating factors such as immune complexes will be modulated. As the expression of Fcγ RI requires active biosynthesis, this work indicates that selective gene activation occurs when neutrophils are recruited into rheumatoid joints.


During the active phase of rheumatoid arthritis (RA), neutrophils constitute between 60 and 80% of the total cell population of synovial fluid. Neutrophils are cells of considerable cytotoxic potential and their inappropriate activation has been implicated in the pathogenesis of RA, perhaps through the generation of reactive oxygen metabolites such as hydrogen peroxide. Tissue damage may be promoted indirectly by the extracellular leakage of toxic products from accumulated neutrophils. Indeed, oxidant damaged products are present in synovial fluid and the oxidase function of neutrophils isolated from synovial fluid has been downregulated, indicating that reactive oxidant generation has been activated in vivo. Furthermore, synovial fluid from patients with RA contains myeloperoxidase in a molecular form indicating that it has been co-secreted from neutrophils concomitant with reactive oxidants. The mechanisms by which neutrophils are activated within rheumatoid joints and the synovial fluid factors which result in such activation are incompletely defined.

The function of neutrophils in vivo is regulated by the levels of expression of specific plasma membrane receptors including those for complement components (CR1 and CR3) and IgG molecules (Fc RI, II, and III). On cellular activation the complement receptors are upregulated whereas Fc RIII is shed from the surface of the cell. Fc RII expression remains constant after cell activation whereas Fc RI is only expressed after in vitro culture of the neutrophils with interferon γ. Neutrophil function is also regulated by the activities of cytokines such as granulocyte-macrophage colony stimulating factor (GM-CSF), tumour necrosis factor, and interferon γ which prime the cells into a state of enhanced responsiveness. On priming, reactive oxidant production is increased together with the levels of expression of several surface receptors.

As such cytokines have been found in the synovial fluid of patients with RA, it is likely that these or other similarly acting agents will upregulate neutrophil function in situ within inflamed joints.

In this work we studied the expression of neutrophil plasma membrane receptors using monoclonal antibodies and fluorescence activated cell sorting (FACS) analysis in paired bloodstream and synovial fluid neutrophils from patients with RA. We propose that the changes in membrane expression observed in synovial fluid neutrophils may be caused by the presence of activating factors, perhaps functioning in combination with cytokines within the synovial fluid, to modulate neutrophil responsiveness.

Subjects and methods

PATIENTS

Twelve patients (two men, 10 women) with seropositive RA were studied. Six patients were receiving non-steroidal anti-inflammatory drugs (NSAIDs), one in combination with sulphasalazine. Two patients were receiving prednisolone and three were not receiving any treatment at the time of study. In addition, one patient with seronegative arthritis and another with psoriatic arthritis (both receiving NSAIDs) were studied.
CELL PREPARATION

Isolation from blood

Neutrophils were isolated from the venous blood of healthy volunteers and rheumatoid patients using dextran sedimentation followed by density gradient centrifugation on Ficoll-Paque as described previously.14 Contaminating erythrocytes were eliminated by hypotonic lysis for 25 seconds in glass distilled water, tonicity being restored by the addition of a 9% w/v sodium chloride solution to give a final concentration of 0.9%.

Isolation from synovial fluid

Synovial fluid was filtered through gauze and spun at 2000 rev/min for 10 minutes. Neutrophils were obtained from the resulting cell pellet using the Ficoll-Paque separation procedure. In all instances neutrophils were greater than 95% pure and viability was greater than 95% when assessed by trypan blue exclusion.

CELL STIMULATION

Neutrophils in RPMI (1×10^7/ml) were stimulated by incubation for 15 minutes at 37°C with the appropriate stimuli (0.1 µg/ml phorbol myristate acetate (PMA), 10% v/v cell free synovial fluid) or primed by incubation for 60 minutes at 37°C with 50 U/ml GM-CSF.

RECEPTOR ANALYSES

Monoclonal antibodies

The following monoclonal antibodies were used: Leu11a (CD16), Leu15 (CD11b), and CD35 from Becton and Dickinson; IV.3 (CD32) and CD64 (322) antibodies were obtained from Medarex. Antibodies to CD11b recognise epitopes on the α chain of the CR3 receptor. CD35 is an epitope of the CR1 receptor, whereas CD16 and CD32 recognise the low and intermediate affinity receptors for the Fc portion of IgG respectively. Antibodies to CD64 recognise Fc RI, the high affinity receptor which is usually only expressed on neutrophils after culture in the presence of interferon γ. 31D8 positive cells avidly bind formyl peptides and show membrane depolarisation after cell activation,15 whereas 31D8 negative cells do not respond functionally to fMLP.16

Staining

Neutrophils were suspended in phosphate buffered saline (PBS)/1% bovine serum albumin (BSA) (globulin free)/0.1% sodium azide, pH 7.2 and receptor expression was measured by a standard indirect immunofluorescence technique using FITC labelled goat antismouse immunoglobulin (Becton and Dickinson) as a second layer: both first and second layer antibodies were added at saturating concentrations. In all experiments non-immune mouse immunoglobulin of the appropriate isotype was included as class specific first layer controls. Stained cells were fixed in 1% paraformaldehyde in PBS and analysed using a Becton and Dickinson FACS Analyser I and a Consort 30 computer and software. The modal fluorescence was proportional to the number of antigenic sites on each cell for each individual antibody. Fluorescence distributions represented a total of 5000 gated events.

STATISTICS

Differences between the expression of receptors on neutrophils in the peripheral blood of patients and in control subjects was studied using the Mann-Whitney U test for independent samples. The Wilcoxon matched pairs signed rank test was used to compare the expression of the receptors on the synovial fluid cells with their paired peripheral blood cells from the same patients. Results are expressed as median (interquartile range) responses.

REAGENTS

Recombinant human GM-CSF (97% purity) was a non-glycosylated peptide from Glaxo and had an activity of >1.5 MU/mg protein in the AML-193 proliferation assay. Sodium azide and globulin free BSA were obtained from Sigma. The Ficoll-Paque was from Pharmacia and RPMI was from Flow Laboratories.

Results

EXPRESSION AND MODULATION OF MEMBRANE ANTIGENS IN CONTROL PERIPHERAL BLOOD NEUTROPHILS

When neutrophils isolated from the peripheral blood of healthy controls were primed by incubation in the presence of the cytokine GM-CSF (50 U/ml) an increased expression of the receptor CD11b (the α chain of the complement receptor CR3) was observed as indicated by an increase in fluorescence intensity for each cell (fig 1A). Expression of CD16 and 31D8 was not significantly altered when the cells were exposed to this cytokine in vitro (fig 1B–D).

Activation of neutrophils with PMA resulted in greater increases in CD11b expression than in primed (fig 1A and 1B) cells whereas CD16 (fig 1C and 1D) and 31D8 (fig 1E and 1F) expression decreased. In vitro priming and activation of the cells had no significant effect on the expression of CD64 and CD32 (Fcy RI and II respectively) under the experimental conditions used (maximum incubation time of 4 hours, data not shown). When blood neutrophils were incubated for 18 hours with cell free synovial fluid five of seven synovial fluid samples showed a slight increase in CD64 expression (data not shown).

MEMBRANE ANTIGEN EXPRESSION IN PERIPHERAL BLOOD AND SYNOVIAL FLUID NEUTROPHILS

No significant changes were observed in the percentage of neutrophils expressing the CD32 (Fcy RI) and CD35 (CR1) receptors in peripheral blood neutrophils of the rheumatoid
patients compared with those of healthy controls (table 1). Table 2 indicates that although the median values for CD11b expression are slightly higher and CD16 and 31D8 slightly lower in the peripheral blood neutrophils from rheumatoid patients these differences again did not reach significance.

Neutrophils isolated from the synovial fluid of rheumatoid patients showed a significant decrease in the percentage of cells recognised by antibodies against CD32 (FcγRII; p<0-005) and CD35 (CR1; p<0-005) compared with their peripheral blood counterparts (table 1). The percentage of synovial neutrophils expressing CD11b, CD16, and 31D8 did not differ significantly from the paired peripheral blood neutrophils (data not shown) but there was an increase in the number of CD11b (CR3) receptors expressed on the synovial neutrophils relative to paired bloodstream cells (table 2; p<0-005). In addition, there was a significant decrease in the expression of CD16 (FcγRIII) and 31D8 on the synovial neutrophils compared with paired bloodstream cells (table 2, p<0-005).

The neutrophils from the synovial fluid of a patient with seronegative arthritis showed increased expression of CD11b and a decrease in the expression of CD16 and 31D8 compared with their peripheral blood neutrophils. There was also a decrease in the percentage of cells expressing CD35 and CD32. These changes mirrored those observed for the seropositive patients. The patient with psoriatic arthritis showed a slight decrease in CD16 and 31D8 expression, but no significant changes in the other antigens studied.

CHANGES IN RECEPTOR EXPRESSION OF PERIPHERAL BLOOD NEUTROPHILS IN RESPONSE TO SYNOVIAL FLUID
When control (unprimed) bloodstream neutrophils were incubated for 15 minutes at 37°C with cell free synovial fluid from a patient with RA, expression of CD11b was up-regulated (fig 2A). In contrast, fluorescence distributions after staining for CD16 (Fcγ
R11I), CD32 (Fcγ RII), and 31D8 were all decreased on exposure of the cells to synovial fluid. These decreases may represent decreased expression of receptors as they become internalised or shed (especially the phosphatidylinositol linked CD16 and 31D8) or else because binding of monoclonal antibodies is blocked by ligands present in the synovial fluid (e.g. immunoglobulins interfering with the binding of Fcγ R).

When control blood neutrophils were primed with GM-CSF (fig 2B) for one hour before the addition of synovial fluid, some differences were noted. Firstly, although priming induced an increased expression of CD11b, the addition of synovial fluid resulted in a further increase in expression. Secondly, whereas synovial fluid downregulated CD35 in unprimed cells, it upregulated expression in primed cells. These data indicate that whereas synovial fluid induces some effects on neutrophil receptor expression that resemble those which occur during priming in vitro, there are additional effects which may be attributable only to factors which cause activation through other mechanisms. Similar results were obtained in four other experiments using synovial fluid from different patients and control neutrophils from different donors.

**FCγ R1I expression**

Bloodstream neutrophils do not express CD64 (Fcγ RII), the receptor with a high affinity for monomer IgG. This receptor is only expressed after incubation of cells with interferon γ and the process requires activated de novo biosynthesis: incubation periods in excess of 16 hours are required before the receptor is detectable on the cell surface. In all bloodstream neutrophil preparations from controls or patients with RA, no expression of CD64 expression (using monoclonal antibody 322) could be detected. Furthermore, in five of 11 patients with RA, neutrophils in the synovial fluid expression of this receptor could not be detected (fig 3A). In six of 11 patients expression was detected, however (fig 3B). This observation indicates that these latter neutrophils have been exposed to factor(s) within the synovial fluid which have selectively activated gene expression.

**Discussion**

Neutrophil responses can be evoked by a variety of particulate and soluble stimuli which effect cell activation by first binding to specific cell surface receptors. Neutrophil responses in vivo are regulated by the levels of expression of these surface receptors and how they are coupled to their respective signal transduction systems. Some factors which have specific receptors on the neutrophil surface include C5a, leukotriene B4, platelet activating factor, IgG, and interleukin 8, and these have been implicated in the pathology of inflammatory joint disease. In addition, neutrophil function may be regulated by a variety of cytokines including tumour necrosis factor

![Figure 2](https://example.com/figure2.png)

**Figure 2** Effect of synovial fluid on receptor expression in bloodstream neutrophils. (A) Neutrophils were incubated in the absence (---) or presence (-----) of 10% cell free synovial fluid at 37°C for 15 minutes. (B) Cells were incubated at 37°C for 60 minutes with 50 U/ml granulocyte-macrophage colony stimulating factor and then incubated for a further 15 minutes in the absence (---) or presence (-----) of 10% cell free synovial fluid before staining for fluorescence activated cell sorting analysis. Similar results were obtained in four other experiments using synovial fluid samples from different patients and neutrophils from different controls.

![Figure 3](https://example.com/figure3.png)

**Figure 3** Fcγ R1 expression in neutrophils. Neutrophils were isolated from the synovial fluid of patients with rheumatoid arthritis and stained for the expression of Fcγ R1 using monoclonal antibody 322 (---) or the non-immune isotype control monoclonal antibody (-----). The fluorescence distribution pattern in (A) was obtained in five of 11 patients, whereas the pattern shown in B was observed in six of 11 patients.
and GM-CSF. The synovium itself is the source of a variety of inflammatory mediators such as arachidonic acid metabolites, vasoactive amines, platelet activating factor, and complement cleavage products, and there is also local production of cytokines within the joint. Furthermore, immune complexes present within rheumatoid joints are probably important in the activation of infiltrating neutrophils via Fcγ receptors. Thus the levels of expression of neutrophil receptors within synovial fluid give an indication of the functional status of these cells and will also give an indication of their past history.

Several lines of evidence now point to the fact that neutrophils within the synovial fluid of patients with RA have been exposed to priming and activating factors in vivo. In this study we have used FACs analysis to measure the expression of plasma membrane receptors and compared the results of paired bloodstream and synovial fluid cells with those obtained after bloodstream cells had been primed, activated, or aged in vitro. It has been shown elsewhere that the isolation procedures used to purify neutrophils can sometimes affect the expression of some plasma membrane receptors. Therefore, in this study we used the dextran/Ficoll isolation procedure to isolate blood and synovial fluid neutrophils because other methods (e.g. centrifugation on Mono-Poly-Resolving Medium) are not always successful for the isolation of synovial fluid neutrophils. Comparison of receptor expression in bloodstream neutrophils from healthy controls did not reveal any statistically significant differences with those from patients with RA. Synovial fluid neutrophils, however, showed gross changes in receptor expression compared with paired bloodstream cells from the same patient. The observations for CD11b (CR3) confirm earlier reports, but additionally we have focused on the expression of receptors for Fcγ as these are likely to be the receptors responsible for activation of reactive oxidant generation by synovial fluid neutrophils. Synovial fluid neutrophils showed a decreased expression of CD32 (Fcγ RI) and CD16 (Fcγ RII) compared with paired bloodstream cells. As the expression of CD32 decreases only slightly during priming, activation, or aging, however, decreased staining of synovial fluid neutrophils probably indicates that monoclonal antibody binding is 'blocked' by ligands (e.g. immune complexes) present within the synovial fluid. Such ligand binding may also account, in part, for the decreased staining of CD16, but this receptor (and 31DB8) is also shed from the plasma membrane during activation (fig 1) and this process may contribute to the decreased expression. Upregulation of CD11b in synovial fluid neutrophils is indicative of priming or activation, or both. The observation that a short incubation of primed or unprimed bloodstream neutrophils with cell free synovial fluid induces these patterns of receptor expression indicates that these changes in receptor expression can occur rapidly as newly recruited bloodstream neutrophils infiltrate inflamed joints.

Of considerable interest and importance is the finding that over half of the synovial fluid neutrophil samples express to varying degrees the high affinity IgG receptor, CD64 (Fcγ RI). This receptor is not expressed by bloodstream neutrophils but is induced when protein biosynthesis is selectively upregulated during exposure to agents such as interferon-γ. To determine if factors within synovial fluid could activate the expression of CD64, we incubated blood neutrophils for 18 hours with cell free synovial fluid. Five of seven synovial fluid samples tested induced a slight increase in CD64 expression. Thus it appears that synovial fluid may selectively activate gene expression in newly recruited neutrophils and further work is clearly needed to confirm this observation. Expression of blood neutrophil FcγRIIa to agents such as interferon-γ, GM-CSF, or G-CSF induces a rapid and selective increase in neutrophil protein biosynthesis which results in an enhanced ability to generate reactive oxidants, continued expression of key cellular components such as receptors, extended viability, and the generation of secondary cytokines. We therefore propose that synovial fluid neutrophils have been exposed to agents within synovial fluid which have resulted in such a selective increase in protein biosynthesis. If this phenomenon has occurred within inflamed joints then our current understanding of neutrophil function in disease pathology is grossly underestimated. Further work is necessary to identify the factors within synovial fluid which may be responsible for the regulation of receptor expression and activation of CD64 expression and also to assess the full biosynthetic activity of synovial fluid neutrophils so that their full role in disease pathology can be appreciated.

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Receptor expression in neutrophils from patients with RA

Membrane surface antigen expression on neutrophils: a reappraisal of the use of surface markers for neutrophil activation.