Signalling through neutrophil FcγRIII, FcγRII, and CD59 is not impaired in active rheumatoid arthritis

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Abstract
Objective—To compare neutrophil Fc receptor (FcγR) and CD59 signalling responses in normal healthy subjects and patients with active rheumatoid arthritis (RA).
Methods—Intracellular free calcium concentrations were measured in neutrophils loaded with the fluorescent calcium indicator fura-2, using a spectrofluorimeter.
Results—Basal intracellular calcium ion concentrations were similar in both groups when no primary antibody, CD59, or CD32 (FcγRII) antibody was added. When CD16 (FcγRIII) antibody was added, there was a significantly greater basal calcium concentration in the patient group compared with the control group. Transient cytosolic calcium ion fluxes were observed after binding FcγRII, FcγRIII, or CD59 with specific monoclonal antibodies and cross linking with the F(ab)2 fragment of sheep antihuman IgG. Peak concentrations of intracellular free calcium, [Ca\(^{2+}\)], after cross linking each of the three receptors, were comparable between normal healthy donors and patients with RA. The lag period between addition of cross linking antibodies and the increase in calcium was also similar between normal individuals and patients.
Conclusion—Contrary to previous reports, these results demonstrate that Ca\(^{2+}\) signalling responses of cross linked Fc receptors in blood neutrophils from patients with RA are identical to those in neutrophils of normal subjects. Signalling responses of cross linked CD59 are also unaltered.

Patients and methods
PAtIENTS
Peripheral blood (20 ml) was obtained from eight normal healthy donors and eight patients with RA. The eight patients were classified as having active RA with synovitis, nine or more swollen joints, and an erythrocyte sedimentation rate > 35 mm/1st h. The median and range of age in the control and patient groups were 42 (24–55) years and 60 (45–75) years, respectively.

PREPARATION OF NEUTROPHILS
Blood from donors was collected into heparinised universal containers. Neutrophils were isolated essentially as described previously. Briefly, leucocytes were separated from erythrocytes by dextran sedimentation using 0-6% w/v dextran (Fisons, Loughborough, UK). The leucocyte rich upper layer was then fractionated by layering on Histopaque (Sigma, Poole, Dorset, UK) followed by centrifugation at 220 g for 25 minutes. Residual erythrocytes in the polymorphonuclear rich cell pellet were removed by hypotonic lysis. All these procedures were carried out at ambient temperature. Cell viability, assessed by exclusion of the vital stain trypan blue, showed that the cells were greater than 95% viable. Microscopy showed less than 5% contamination by mononuclear cells. After purification, the neutrophils were suspended in Krebs-HEPES buffer containing (mMol/l): sodium chloride 120; potassium chloride 4-8; monobasic potassium phosphate 1:2; calcium chloride 1-5; magnesium sulphate
Neutrophil signalling responses in RA

1:2; (N-2-hydroxyethylpiperazine-N'-2-ethanesulphonic acid) (HEPES) 25.

ANTIBODIES
FcγRIII antibody (CD16; MEM-154, IgG1 isotype) was a kind gift from Dr Vaclav Horejsi, Academy of Sciences of the Czech Republic, Prague. CD59 antibody (BRIC 229, IgG2b isotype) was obtained from the International Blood Group Reference Laboratory, Elistree, Herts, UK. FcγRII antibody (CD32; AT10, IgG1 isotype) was from Dr Martin Glennie, Tenovus, Southampton, UK. Polyclonal antiamouse IgG was raised in sheep, the IgG fraction purified, and F(ab)2 fragments prepared by standard methods.

ANALYTICAL METHODS
Intracellular calcium ion concentrations ([Ca²⁺]) were measured using the fluorescent calcium binding probe, fura-2 (Molecular Probes, Junction City, Oregon). Population studies were performed rather than single cell studies, for ease of experimentation and in order to allow correlation with the results of Goulding and Guyre. Neutrophils (1 × 10⁷ - 1 × 10⁸/ml) were incubated with fura-2 acetoxyethyl ester (final concentration 1 μmol/l) for 30 minutes at ambient temperature in Krebs-HEPES buffer containing 0-1% bovine serum albumin. After loading, the cells were washed once then resuspended at 5 × 10⁶/ml in the same medium. The cells were incubated with primary antibody (CD16, CD32, or CD59 at a final concentration of 20 μg/ml) for 30 minutes on ice. The cells were washed twice with Krebs-HEPES buffer and kept on ice. Cells (200 μl, 1 × 10⁶) were added to 2 ml Krebs-HEPES buffer prewarmed to 37°C and placed in a stirred quartz fluorimeter cuvette in a dual wavelength fluorimeter (Spex II fluorolog system, Glen Spectra, Stanmore, Middx, UK) with the temperature set at 37°C. Simultaneous excitation at 340 nm and 380 nm was achieved using two lamps and monochromators together with a mirrored chopper system. Fluorescence emission was detected at 505 nm. Basal fluorescence was monitored for one minute before primary antibodies were cross linked by the addition of F(ab)_2 fragment of sheep antiamouse IgG (final concentration 60 μg/ml). The fluorescence intensity of emission was measured for a further four minutes. After completion of the measurements, digitonin (50 μmol/l final concentration) followed by EGTA (20 mmol/l final concentration) were added to the cuvette to provide Ca²⁺ bound (Rmax) and Ca²⁺ free (Rmin) signals, respectively. The ratio (R) of the fluorescent intensities at 340 nm and 380 nm is related to Ca²⁺ concentration by the equation:

\[ \text{Ca}^{2+} = K (R - R_{\text{Rmin}})/(R_{\text{Rmax}} - R) \]

where K = 0.2865 μmol/l⁻¹.

STATISTICAL ANALYSIS
Parameters were compared between normal subjects and patients and the significance of differences expressed using the Mann-Whitney U test, assuming a non-parametric data distribution.

Results
Figure 1 shows representative Ca²⁺ fluxes triggered by cross linking the three different primary antibodies. For a few samples there were insufficient cells to obtain results for all antibodies, but results from a minimum of six samples were analysed per antibody per group.

Figure 2 shows two measurements summarising the data from each recorded trace for each subject. The basal value is the average [Ca²⁺],
from three readings taken during the first 60 seconds of the trace before cross linking antibody was added. The peak [Ca²⁺], is the average of two readings taken at the height of the calcium response. The lag time, defined as the time interval between addition of the cross linking antibody and the start of any increase in calcium concentration, was also measured for each trace (table).

Incubation of fura-2 loaded cells with monoclonal antibodies to CD59 or CD32 had no effect on the basal [Ca²⁺] value in either group, compared with cells that had been incubated without antibody. These basal values did not differ significantly between patients and controls. There was, however, a small but significant increase in basal [Ca²⁺], of cells from patients with RA compared with controls after incubation with CD16 antibody (mean basal value in controls 186 nmol/l and in RA patients 254 nmol/l; p = 0.0007) (fig 2A, B).

Addition of F(ab)₂ fragment of sheep antime IgG to neutrophils that had not been incubated with a primary monoclonal antibody had no effect on [Ca²⁺], (fig 1A, E); we have shown previously that not all intact IgG antibodies to neutrophil surface antigens cause a change in [Ca²⁺]. When cells were preincubated with CD16, CD59, or CD32 antibodies, the addition of the cross linking second antibody resulted in a transient increase in [Ca²⁺], in all donors (figs 1, 2). There was no significant difference in either the lag time or the peak [Ca²⁺], for any of the three antibodies between normal control subjects and patients with RA (table).

**Discussion**

The ability of neutrophils to signal correctly in the presence of external stimuli is an important facet in their armoury, which includes the phagocytosis of immune complexes and opsonised particles, and the production of reactive oxygen species and other effector molecules. Any abnormality in signalling could contribute to the pathogenesis of inflammatory diseases such as RA, and it has been shown previously that subtle but potentially important abnormalities do exist in the calcium signalling of neutrophils from patients with RA: after stimulation, the amount of the cell cytosol occupied by increased amounts of Ca²⁺ was significantly increased in RA blood and synovial fluid neutrophils compared with normal blood neutrophils, while another report described a low basal concentration of [Ca²⁺], and low or no response to cross linking of either CD16 or CD32 on neutrophils from patients with active RA. Our present data strongly suggest that antibodies do not induce signalling by binding of the Fc portion of intact antibody to the Fc receptors, but bind via their F(ab)₂ portions.

Our particular interest in complement regulation and inflammatory disease led us to examine whether cell activation, induced by cross linking the complement regulatory molecule CD59 and other GPI anchored molecules, was abnormal in active RA. In contrast to the report of Goulding and Guyre, our results demonstrate that blood neutrophils from patients with active RA show normal intracellular calcium changes in response to cross linking of the Fc receptors CD16 (GPI anchored) and CD32 (transmembrane), and of the GPI anchored complement regulatory protein, CD59. The reasons for the conflicting results are unclear, but there were several differences in methodology between the two studies. First, the neutrophils in our study were isolated using dextran and Histopaque—a method chosen to cause minimal cell activation—whereas Goulding and Guyre used a

![](http://ard.bmj.com/Downloaded from http://ard.bmj.com/)
Ficoll meglumine M85 gradient method. Second, different monoclonal antibodies to CD16 and CD32 were used: the CD16 and CD32 antibodies used in our study were both IgG1 subclass, while the antibodies used by Goulding and Guyre (IV.3 to FcyRII and 3G8 to FcyRIII) were IgG2b and IgG2a, respectively. The antibodies used in both studies block the ligand binding activity of their respective FcR (confirmed for IV.3 and 3G8 by Medered, and for MEM-154 by Dr V Horejsi, personal communication; reported by Greenman et al for AT10). Third, we labelled cells with monoclonal antibodies at 4°C rather than 37°C to prevent the internalisation or shedding that can occur at 37°C. This may be significant if neutrophils from RA patients and normal control donors differ in their cell surface protein turnover. Fourth, we used fura-2 for measuring [Ca2+]i, because it can be used in a ratiometric way, alleviating the problems caused by background noise that cannot be addressed using indo-1 in the non-ratiometric approach used by Goulding and Guyre. Our study thus does not support previously reported findings of reduced resting [Ca2+]i and markedly reduced intracellular Ca2+ fluxes upon cross linking FcyRII or FcyRIII in RA neutrophils.

Three patients with RA showed a considerably greater increase in [Ca2+]i on cross linking CD16. However, these patients showed no distinctive clinical features, nor did this trend follow on cross linking CD32 or CD59.

Our finding of a small but significant increase in basal [Ca2+]i in RA neutrophils after incubation with CD16 antibody in the absence of cross linking antibody suggests that this antibody may cause some activation in the RA neutrophils. We have previously shown that there is no difference in the surface expression of CD16 on blood neutrophils from normal controls and patients with active RA. It is unlikely, therefore, that the observed increase in basal [Ca2+]i in this study is related to differences in CD16 surface expression between the normal and RA patient group, while it may reflect ‘priming’ of RA neutrophils for response through CD16.

In conclusion, we found no difference in the signalling induced by cross linking CD16, CD32, or CD59 with secondary antibody in peripheral blood neutrophils in patients with active RA compared with neutrophils from control subjects. This suggests that, at least in vitro, peripheral blood neutrophils from patients with RA have a normal capacity to interact with immune complexes, producing an increase in intracellular calcium concentration.

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