Transdifferentiation of polymorphonuclear neutrophils to dendritic-like cells at the site of inflammation in rheumatoid arthritis: evidence for activation by T cells

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Objectives: To investigate infiltrated cells in the synovial fluid (SF) of inflamed joints of patients with rheumatoid arthritis (RA), with special reference to polymorphonuclear neutrophils (PMN) and their interaction with T cells.

Methods: Expression on PMN of activation associated receptors CD14, CD64, CD83, and major histocompatibility complex (MHC) class II was examined in the SF of 15 patients with RA, as were the infiltrated T cells. SF cytokines were determined by enzyme linked immunosorbent assay (ELISA). To mimic the in vivo situation, co-culture experiments were carried out using PMN and T cells of healthy donors.

Results: The SF contained activated T lymphocytes and abundant PMN. SF PMN expression of CD14 and CD64 was enhanced compared with peripheral blood. Of special interest was the observation that only the SF PMN expressed MHC class II antigens and CD83. Exposure to SF, which contained considerable amounts of cytokines (for example, interferon γ (IFNγ), tumour necrosis factor α, and interleukin 2), induced a similar receptor pattern on blood derived PMN of healthy donors. Furthermore, PMN acquired MHC class II and CD83 within 24 to 48 hours, when co-cultured with autologous T cells or T cell lines. This effect was also achieved by T cell supernatants, was dependent on protein synthesis, and could be inhibited by antibodies against IFNγ.

Conclusions: SF PMN from patients with RA undergo major alterations, including transdifferentiation to cells with dendritic-like characteristics, probably induced by T cell derived cytokines. Because MHC class II positive PMN are known to activate T cells, the mutual activation of PMN and T cells might contribute to the perpetuation of the local inflammatory process, and eventually to the destructive process in RA.
Activation of PMN by T cells in RA

Cytofluorometry

Cells from SF or whole blood were double labelled using a fluorescein isothiocyanate (FITC) labelled antibody to CD66b (Immunotech, Marseilles, France, clone 80H3) as PMN marker and phycoerythrin (PE) labelled antibodies to CD14 (clone UCHM1), CD64 (10.1), major histocompatibility complex (MHC) class II (HLA DR-DP-DQ, clone WR18, all Serotec, Oxford, GB), or CD83 (Immunotech, HB15A). For lymphocytes, antibodies to CD3 (SK7), CD4 (SK3), CD8 (SK1), CD69 (L78; all Becton Dickinson), CD19 (Immunotech, J4.119), and CD25 (Caltag, Burlingame, USA, CD25-3G10) were used. The final concentration of antibodies varied between 1 and 10 μg/test. Isotypic controls IgG1 FITC, IgG2a, and IgG2b PE were used in the same concentration. To reduce non-specific binding Fc receptors were saturated with 10 μg/ml human IgG (Venimmun, Centeno, Marburg, Germany) and with 1% bovine serum albumin before and during incubation. After incubation of SF cells (1×10⁶) or of whole blood (10⁶ μl) with the respective antibodies for 15 minutes at 25°C in the dark, cells were washed twice in FACS buffer, fixed in PBS/1% paraformaldehyde and stored at 4°C until analysed by FACSort and CellQuest software (Becton Dickinson). Whole blood was lysed by FACS lysing solution (BD; 10 minutes in 2 ml; 1:10 in aqua dest.). Results are expressed as the percentage of positive cells in the respective gate or quadrant, or as the mean fluorescence intensity, respectively.

For comparison, receptor expression on peripheral PMN of 20 normal healthy donors (NHD) was measured. Expression of the newly acquired receptors (for example, CD83 and MHC class II) was considered to be positive when the mean plus 2 SD of all NHD was exceeded.

Isolation of PMN and in vitro activation

PMN were separated from 50 ml whole blood by centrifugation on PolymorphPrep (Nycomed, Oslo, Norway), which yielded a highly homogeneous preparation with a purity >98% (judged by cytofluorometry with an antibody to CD66b). PMN were cultivated in AIM (Gibco, Eggenstein, Germany) with 2.5% autologous heat inactivated serum. PMN were either stimulated with the indicated concentration of supernatants of phytohaemagglutinin activated T cells or 100 U/ml IFNγ for 24–48 hours. For co-culture 4×10⁶ PMN and 4×10⁶ T cells were put together for up to 48 hours in 4 ml AIM. In whole blood, PMN (100 μl) were stimulated with 100 μl SF of patients with RA for 30 minutes or 24 hours and used as described above. The T cell clone A37/2 was obtained from Dr Kabelitz/Heidelberg. To prolong the life of the cells they were restimulated with irradiated monocytes and B cells (as feeder cells) every 14–18 days with 1 ng/ml phytohaemagglutinin. Supernatants of T cells were obtained after 48 hours by centrifugation of cells and stored at –20°C until used.

Multiplex enzyme linked immunosorbent assay (ELISA)

Cytokine levels in SF were determined using the Luminex multiplex analyte technology (BioPlex system by BioRad, Hercules, CA, USA) in combination with multiplex cytokine kits. In total, eight different cytokines (cytokines 8-plex-kit BioRad) were measured in 50 μl SF. The detection sensitivity of all cytokines was <10 pg/ml. Data from previous studies (unpublished observation; not shown) suggested that PBS/1% bovine serum albumin/0.1% sodium azide (FACS buffer) and subjected to cytofluorometry. Viability was assessed by trypan blue exclusion and microscopic evaluation. Peripheral blood was taken at the same time from all patients.
detected fluorescence intensity reaches a fixed value. The known input concentration of a plasmid against the calculated from a virtual standard curve, obtained by plotting a and cyclophilin B. The transcript concentration was calculated as the percentage of positive PMN or as mean fluorescence intensity as detected by cytofluorometry by box and whiskers plots. The box contains 50% of the values, the square indicates the mean values, the horizontal bar the median. Differences between the groups were summarised as the percentage of positive PMN or as mean fluorescence intensity per peripheral blood of all 15 patients and of 20 NHD. The data are pretreatment of the SF with hyaluronidase, and filtration of the samples to remove rheumatoid factors, showed no differences compared with untreated samples.

Quantitative reverse transcriptase-polymerase chain reaction RNA was isolated from PMN or T cells by “High Pure” RNA Isolation Kit (Roche Diagnostics, Mannheim, Germany). To prepare the reverse transcriptase the 1st strand cDNA Synthesis Kit (Roche Diagnostics, Mannheim, Germany) was used. The cDNA obtained was diluted 1:25 with water; 10 μl was used in each capillary for amplification. Target sequences were amplified using LightCycler Primer Sets (Sarch-LC, Heidelberg). RNA input was normalised by the average expression of the two housekeeping genes β-actin and cyclophilin B. The transcript concentration was calculated from a virtual standard curve, obtained by plotting a known input concentration of a plasmid against the polymerase chain reaction cycle number at which the detected fluorescence intensity reaches a fixed value. The actual copy number per microlitre of cDNA was calculated as follows: \[ X = e^{(-0.6535 \times CP + 20.62)} \]. The data of two independent analyses for each sample and for each parameter were averaged and presented as adjusted transcripts/μl cDNA.

RESULTS

Analysis of leucocytes in the SF

SF samples of patients with RA contained 0.5–3×10^7 viable leucocytes. In 14/15 patients most cells were identified as PMN (range 34–89.5%; mean (SD) 68.6 (4.8%); n = 14); in the other patient who had longstanding RA (duration 13 years) considerably more lymphocytes than PMN were detected (67% ± 14%). In the other patients, lymphocytes amounted to 2.1–34% (12.4 (9.4)%). Most lymphocytes were identified as T cells (71 (24)%); further analysis showed that CD4 cells were predominant. Expression of CD25 and CD69 on CD4 and on CD8 cells, indicated that T cells were activated. Only a minor portion of B lymphocytes was seen (CD19+ 1.3% (1.59)). Between 4.9% and 32.0% cells were identified as monocytes (CD14+/CD66b– 14.9 (8.3)%).

Transdifferentiation of locally activated PMN to dendritic-like cells

To analyse the SF PMN further, expression of activation associated surface receptors was studied and compared with that in peripheral blood PMN and in PMN of 20 NHD. CD14 was drastically up regulated on the SF infiltrated PMN, but not on peripheral blood PMN of the same patients or of NHD. Similarly, CD64 was up regulated only on the PMN in the SF. To study the transdifferentiation of SF PMN, we looked for the surface expression of MHC class II antigens and of CD83. Neither receptor is constitutively expressed by PMN and the expression after stimulation depends on de novo protein synthesis. On SF PMN, but not on blood PMN, CD83 expression was found in 11/15 patients (mean (SD) 37.9 (37.8)%). Moreover, MHC class II expression was seen on the SF derived PMN in 11/15 patients (mean (SD) 50.1 (44.6)%), whereas the peripheral blood PMN were negative for MHC class II. Of note, MHC class II and CD83 were not necessarily coexpressed: only the PMN of nine patients were double positive for MHC class II and CD83. The amount of both CD83 and MHC class II varied between the patients. Figure 1 shows an example and fig 2 summarises the data for all the patients.

There was, however, no correlation with the number of infiltrated total leucocytes, T cells, PMN, or monocytes, or with the systemic C reactive protein levels of the MHC class II and CD83 expression. Moreover, there was no difference between the patients with early or longstanding RA.

As not all patients were positive for either CD83 or MHC class II, we studied the kinetics of receptor expression of PMN derived from SF ex vivo. Cells with no CD83, but expressing MHC class II were cultivated further for 16 hours; then the viability, CD83 and MHC class II expression were measured. The PMN were still viable, the PMN acquired CD83 and MHC class II varied between the patients. Figure 1 shows an example and fig 2 summarises the data for all the patients.

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Detection of cytokines in SF

By ELISA, T cell derived cytokines, including interferon γ (IFNγ), tumour necrosis factor α (TNFα), granulocyte monocyte-colony stimulating factor (GM-CSF), interleukin (IL) 2, IL4, and IL10, were found in the SF of the majority of patients, as were cytokines which among others are produced by activated PMN (for example, IL8, IL6) or monocytes/macrophages (for example, GM-CSF, TNFα) Figure 4
summarises the data for all patients. There was no correlation between the amount of any cytokine in the SF and the expression of MHC class II antigens or CD83 on PMN. In line with the presence of cytokines in the SF, the short term culture (30 minutes) of PMN derived from NHD with patients’ SF resulted in activation of PMN, seen as an upregulation of CD14 and CD64. Culture for 48 hours induced CD83 expression and, to a lesser extent, MHC class II. Figure 5 shows an example.

**Figure 3** Induction of CD83 and MHC class II expression on SF derived PMN derived cultured ex vivo. (A) Expression of CD83 and MHC class II was measured either directly after recovery of the PMN from the SF (line), or after culture for 16 hours at 37°C (filled peaks). (B) In parallel, activation associated antigens (CD69 and CD25) were determined on T cells after 16 hours (one of three experiments with cells of three patients is shown).

Activation of PMN by T cells in vitro: expression of MHC class II and CD83 on PMN

Because the ex vivo analyses described above suggested a local activation of PMN by the infiltrated T cells or their cytokines, respectively, a series of in vitro experiments were carried out to test the interaction of highly purified T cells with PMN of healthy donors. Firstly, the expression of MHC class II on PMN was measured after incubation with supernatants (SN) of stimulated T cells. SN of the T cell clone A37/2 induced MHC class II on PMN in a concentration dependent manner (fig 6). By an ELISA, IFNγ was detected in the SN; correspondingly, the expression of MHC class II could be inhibited by a neutralising antibody to IFNγ (fig 6). Also, after co-culture with T cells (clone A37/2), PMN expressed MHC class II after 24 hours. Again, the percentage of MHC class II positive PMN increased with the number of T cells. The expression of MHC class II on PMN was significantly decreased when either PMN or T cells were pretreated for 2 hours with a protein synthesis inhibitor (cycloheximide 0.1 µg/ml; the dose did not affect the viability of the PMN or T cells; fig 7), indicative of protein de novo synthesis of MHC class II by PMN, and of a mediator produced by T cells.

By real time polymerase chain reaction, IFNγ mRNA was detected in T cells that had been co-cultivated with autologous PMN for 2 hours; expression was further increased after 6 and 24 hours (table 1).

**DISCUSSION**

Leucocyte infiltration into infected tissues is essential for an efficient host defence; it is, however, also seen at sites of sterile or chronic inflammation—for example, in inflamed joints of patients with RA. In our study the majority of infiltrated leucocytes in the SF of patients with RA were PMN. Although activation of PMN at the inflamed site has been well described by others, the unexpected finding

Figure 4 Cytokines in the SF of all 15 patients: the cytokines were quantified by ELISA and expressed as pg/ml. Data of all patients are summarised as box and whiskers plots. The box contains 50% of the values, the square indicates the mean values, the horizontal bar the median (note, there is a logarithmic y axis). The dots at the bottom symbolise patients who were negative for the respective cytokines. Data from previous studies (unpublished observation; not shown) suggested that pretreatment of the SF with hyaluronidase and filtration of the samples to remove rheumatoid factors is not necessary.
was that a portion of the PMN had acquired CD14, CD83, and MHC class II antigens. Neither CD83 nor MHC class II are stored in resting PMN and normally are only expressed by professional antigen presenting cells—for example, monocytes or dendritic cells. Their expression, however, can be induced by stimulating PMN—for example with IFN-γ, TNF-α, or GM-CSF, and requires de novo protein synthesis. This implies, that the lifespan of the PMN can be extended by appropriate conditions, which in turn leads to a further differentiation, especially obvious when the PMN were further cultured ex vivo. Although this presumption is in contrast with the traditional image of PMN as terminally differentiated cells, there is growing evidence that not only in vitro but also under certain pathological conditions—for example, chronic inflammatory diseases and severe trauma, PMN can escape from apoptosis and differentiate further, thereby acquiring new surface receptors such as MHC class II, CD86, CD80, or CD83.

Figure 5  Activation of PMN in whole blood of NHD by SF of patients with RA. The left panel shows that culture for 48 hours activated PMN as seen by the expression of CD83 (line: cells cultured with PBS alone; filled peak: cultured with 50% SF; PMN were identified by anti-CD66b and gated accordingly). The right panel summarises the results of three independent experiments carried out with the blood of three healthy donors (NHD 1–3) and the SF of two patients with RA (SF-P1 and SF-P2); the CD83 expression after 48 hours stimulation with SF compared with PBS alone is shown.

Figure 6  (A) Induction of MHC class II (HLA DP-DQ-DR) expression on PMN of a healthy donor after stimulation with 20% vol/vol supernatant (SN) of the T cell clone A37/2 for 48 hours (line: culture medium alone, filled peak: 20% SN added). (B) The acquisition of MHC class II was dependent on the SN concentration (10–50% vol/vol) (note: the experiment was performed with cells of a different donor). (C) The effect of the SN could be inhibited by antibodies to IFN-γ and to a lesser extent by anti-GM-CSF (concentration 2 μg/ml each; conditions: 30% SN of the clone A37/2 for 24 hours; mouse IgG as isotype control did not inhibit the induction of MHC class II).
MHC class II antigen expression on PMN in SF and on PMN in the peripheral blood of patients with RA has also been described by Cross et al. 3,24 In contrast with our data, their results showed that PMN were negative for MHC class II after isolation, but expression could be induced when PMN were further cultivated. This discrepancy cannot be fully explained. An explanation may be that in our study a more sensitive detection system (for example, different antibody, different fine specificity) was used, which was supported by the observation that under our culture conditions the surface expression of MHC class II and CD83 was also increased further. Moreover, MHC class II expression was not seen in all of our patients; one possible explanation might be the pronounced differences between patients in the ability of PMN to express the proteins, as reported previously. 25,36 Moreover, it cannot be confirmed as yet whether all PMN can transdifferentiate or whether this is only possible in a subpopulation. Some FACS images imply a subpopulation that can transdifferentiate or whether this is only possible in a subpopulation. Some FACS images imply a subpopulation that

**Figure 7** (A) Induction of MHC class II (HLA DP-DQ-DR) expression on PMN of a healthy donor after co-culture with T cells (clone A37/2; 1 x 10^6 cells each/ml for 24 hours). Line: IgG PE isotype control; filled peak: MHC class II PE (HLA DP-DQ-DR). (B) The cocultivation of T cells with cycloheximide (0.1 μg/ml for 2 hours) led to a decrease in MHC class II positive PMN (−39.6%) (filled peak: MHC class II expression on PMN after culture with untreated T cells; line: MHC class II expression after culture with cycloheximide treated T cells). (C) Number of MHC class II positive PMN with increasing numbers of T cells (clone A37/2) after 24 hours co-culture (1 x 10^6 PMN with 6.25 x 10^6–1 x 10^6 T cells/ml).

Table 1  mRNA expression of IFNγ after coculture of PMN and autologous T cells after different time points (2, 6, and 24 hours)

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<tr>
<th>Time (hours)</th>
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Each 2 x 10^6 cells. IFNγ was not detected in PMN. The data of two independent analyses for each sample were averaged and presented as adjusted copies/μl cDNA.

These arguments suggest that the prolonged lifespan of activated PMN in patients with RA may contribute to the progression of the inflammatory process to chronicity; moreover, the PMN may inflict local tissue damage, and participate in the destruction of cartilage and bone.
REFERENCES


