Role of neutrophils in cutaneous leishmaniasis

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Neutrophils are always present in the cytomorphologic process of leishmaniasis but their role is still not fully understood. Microscopic examination was done on smears from 56 cases of clinically diagnosed cutaneous leishmaniasis. Neutrophils were the predominant cells in the smear (>35% of the total cell count including neutrophils, macrophages and lymphocytes) in 7% of cases, a minority (10%–35% of total cells) in 36% of cases and rare (<10% of total cells) in 57% of cases. Microscope images confirmed that neutrophils appeared to have an important role in leishmania elimination through phagocytosis of amastigotes in the later stages of the disease process.

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Introduction

Cutaneous leishmaniasis is endemic in over 70 countries, with an estimated annual incidence of 1.5 million cases [1]. Neutrophils are always present in the cytomorphologic process of leishmaniasis, but their role is still not fully understood. Polymorphonuclear (PMN) neutrophils have been reported to have a crucial role in the destruction of the *Leishmania major* parasites at an early stage of infection [2,3]. A rapid and sustained neutrophilic infiltration localized at the site of the sand-fly bite has been observed using dynamic intravital microscopy and flow cytometry [4]. Studies showing that PMN neutrophils are present in vivo at the sites of the lesion and can kill the parasites in vitro have suggested that these cells might be involved in inhibiting multiplication of the parasite. Neutrophils also appear to play a major role in the development of protective immunity [5,6]. One study with C57BL/6 mice demonstrated that the interaction of *L. major*-infected macrophages with dying neutrophils induced parasite destruction mediated by neutrophil elastase and tumour necrosis factor-α production from neutrophils [5].

Most of these studies focused on the very early stages of infection; little has been mentioned about the role of neutrophils in the later stages of cutaneous leishmaniasis. The pathological features in the later stages of the disease indicate a gradual decrease in the number of amastigotes and macrophages, leaving a granulomatous infiltrate consisting of lymphocytes, epithelioid cells and multinucleated giant cells. At this stage it is difficult or even impossible to detect the amastigotes in haematoxylin and eosin or Giemsa-stained sections [7]. In a dry nodular type of lesion, there is a tendency to form granuloma with fewer lymphocytes and scanty plasma cells [8]. Previous studies have not claimed any role for neutrophils or demonstrated the appearance of neutrophils during the late stages of the disease process, although they mention the disappearance of the amastigotes without giving any rationale for their disappearance.

This cytomorphological study of the role of neutrophils in cutaneous leishmaniasis aimed to investigate the appearance of neutrophils together with the phagocytosis function on the amastigotes in infected lesions from humans with cutaneous leishmaniasis at a later stage of the disease.

Methods

The sample for the study was all 56 cases of cutaneous leishmaniasis (50 males and 6 females) referred for investigation to a medical laboratory in Damascus, Syrian Arab Republic over the period October 2006 to February 2008. All the cases were clinically diagnosed as cutaneous leishmaniasis by an expert dermatology consultant at a dermatology clinic in Damascus. All cases referred over the time period were included and the stage of the disease had not been determined.

Two microscope slides were prepared from each patient, stained with Wright stain and examined under the microscope at ×400 magnification. A total of 50 fields were studied in each slide and the neutrophil, phagocyte and lymphocyte counts per field were noted. For each case the cell counts were calculated as mean percentages.

For each case the neutrophils were defined as: +++ if they constituted >35% of the different cells microscopically present in the smear including macrophages and lymphocytes; ++ if they were between 10% and 35% of total; or + if they were <10% of the total.

The presence of amastigotes in the smear was noted in relation to the neutrophils cell counts.

Table 1 shows the neutrophil counts as a percentage of the total neutrophils, phagocytes and lymphocytes in the light microscopy slides. Neutrophils were present in a high concentration among the 3 cell types (neutrophils, lymphocytes and phagocytes) in only 7% of the cases studied, while in 93% of cases neutrophils were not the predominant cells in the slides and in 57% of cases neutrophils were rarely seen (<10%).

<table>
<thead>
<tr>
<th>Cells</th>
<th>Neutrophils present</th>
<th>No. of patients</th>
<th>% (n = 56)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Neutrophils present*</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>+++ (&gt;35% of cells)</td>
<td>4</td>
<td>7</td>
<td></td>
</tr>
<tr>
<td>++ (10%–35% of cells)</td>
<td>20</td>
<td>36</td>
<td></td>
</tr>
<tr>
<td>+ (&lt;10% of cells)</td>
<td>32</td>
<td>57</td>
<td></td>
</tr>
<tr>
<td>Amastigotes present</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Heavily concentrated in extracellular fluid</td>
<td>+++</td>
<td>4</td>
<td>7</td>
</tr>
<tr>
<td>Present in intracellular fluid, low in extracellular fluid</td>
<td>++ or +</td>
<td>23</td>
<td>41</td>
</tr>
<tr>
<td>Not seen in intra- or extracellular fluid</td>
<td>+</td>
<td>29</td>
<td>52</td>
</tr>
</tbody>
</table>

*In relation to total cells: neutrophils, lymphocytes and phagocytes.
Table 1 also shows the relationship between the appearance of amastigotes in the smear and the neutrophil concentrations in the smear. When the amastigotes were heavily concentrated in the extracellular fluid, the neutrophils were more concentrated, while when the amastigotes were present intracellularly with very low appearance in the extracellular fluid, the neutrophils showed low presence (0%–35%) among the cells and when the amastigotes disappeared from the screen, the neutrophils had very low counts in the microscope slide (< 10%).

It appears from the sample microscope images that once the neutrophils became the predominant cell type in the field, their function in phagocytosis of the amastigotes becomes apparent (Figures 1 and 2). When the neutrophils were in lower concentrations, phagocytosis of amastigotes was not seen (Figure 3). The greater appearance of neutrophils when the amastigotes were present in the extracellular fluid, as shown in the cell counts, is evident in Figure 4.

Discussion

The cases in this study were referred with clinical signs of cutaneous leishmaniasis and no definite disease stage was determined for any of them. We can assume that none of the cases were at the very beginning of the disease process (as presented in other studies [2–4]) as studies indicate that no immediate clinical symptoms appear at the very early stage of the disease soon after the sand-fly transmits the parasite to the host [7,8]. When a patient presents clinically with a skin ulcer measuring 2–5 cm in diameter with a wet exudate and the pathological features show a dermal infiltrate mainly composed of macrophages filled with amastigotes, lymphocytes and plasma cells, then this is an early stage of the disease process. When the lesions clinically appear smaller in size, dry and nodular with dried exudates and the microscopic features show a decrease in the number of, or a disappearance of, the amastigotes and their macrophages, leaving a granulomatous infiltrate consisting of epithelioid cells, multinucleated giant cells, fewer lymphocytes and scanty plasma cells [7,8], then this is a later stage of the disease process.

In the microscope images, neutrophils are shown phagocytosing the released amastigotes in the extracellular fluid of infected lesions. Neutrophils are believed to be specialized in the phagocytosis process, while macrophages are considered less efficient than PMN neutrophils at killing foreign microorganisms, and the mechanism is not as well understood [9]. Accordingly, we can assume that those PMN neutrophils have a strong role to play in phagocytosis and destruction of amastigotes released after membrane rupture of the infected macrophages. Otherwise it is not possible to explain the disappearance of the amastigotes from the infected tissues after their release in huge numbers into the extracellular fluid later in the healing process of cutaneous leishmaniasis [7].

We can hypothesize that once the amastigotes are released from the ruptured phagocytes into the extracellular fluid in the late stage of the disease process, they are recognized by the neutrophils as foreign bodies and an acute-phase reaction is established. More neutrophils are attracted, as seen by an increase in the neutrophil concentration in the damaged area. The neutrophils start the process of phagocytosis of the released amastigotes and consequent limitation of disease.

In conclusion, this study has found that PMN neutrophil cells were at no time absent from the disease process of cutaneous leishmaniasis, in either the early or late stages. While previous
studies have noted the role of PMN neutrophils in the early stages of the disease process, this study concentrated on their role in the later stages. The neutrophils seemed to be highly effective in phagocytosis of amastigotes released from ruptured phagocytes to the extracellular fluid. The PMN neutrophils seemed to establish their role by phagocytosis of the amastigotes as demonstrated by the disappearance of amastigotes when neutrophil counts drop.

References

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