Differential Expression of Proinflammatory Cytokines and Their Inhibitors during the Course of Meningococcal Infections

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Circulating concentrations of tumor necrosis factor-α (TNF), interleukin (IL)-1β, IL-6, IL-1 receptor antagonist (IL-1ra), and soluble TNF receptors p55 (sTNFr-55) and p75 (sTNFr-75) and ex vivo production of TNF, IL-1, IL-6, and IL-1ra using a whole blood culture system were measured during the acute and convalescent stages of meningococcal infection. Circulating TNF and IL-1 were below detection level, whereas IL-6 and IL-1ra, sTNFr-55, and sTNFr-75 were increased at admission. The ex vivo production of proinflammatory cytokines TNF, IL-1, and IL-6 was suppressed at admission and restored gradually during recovery. On the contrary, the production of the antiinflammatory IL-1ra was increased at admission. The elevated concentrations of both IL-1ra and sTNFr early in the course of infection suggest a regulatory role for these antiinflammatory compounds. The observed down-regulation of the ex vivo production of TNF, IL-1, and IL-6 and up-regulation of the production of IL-1ra in the acute stage may indicate a protective regulation mechanism.

The principle sources of these cytokines are blood monocytes and tissue macrophages. On incubation with LPS, isolated monocytes or whole blood taken from healthy donors produce ex vivo cytokines in a spectrum similar to that observed after LPS injection in volunteers. Without LPS stimulation, there is no or minimal ex vivo cytokine production [16, 17]. Several investigators have reported that during the acute stage of serious infections, the ex vivo production of TNF and IL-1 is depressed [18–21]. The cause of this impaired production is unknown, but it has been considered as a down-regulated state of the cytokine-producing cells, possibly reflecting a protective mechanism by averting high concentrations of these cytokines [19, 21]. This hypothesis would be supported if the production of the antiinflammatory cytokine IL-1ra were inversely regulated and increased during the acute stage.

We describe the pattern of circulating TNF, IL-1, IL-6, IL-1ra, sTNFr-55, and sTNFr-75 and the ex vivo production of TNF, IL-1, IL-6, and IL-1ra during meningococcal infection.

Patients and Methods

Five patients with bacteriologically proven acute meningococcal infections, admitted to our intensive care unit, were studied. Four patients had meningitis without severe hemodynamic complications, and 1 (patient 2) had mild sepsis without meningitis. Some clinical parameters indicating the severity of disease and prognosis are summarized in table 1. All patients received antibiotics. Dexamethasone was given in different doses over 1–7 days (table 1). Patient 2 was also treated with two exchange transfusions [22]. All patients recovered completely, except patient 3 who developed sensorineural deafness.
Serial plasma and serum samples were collected shortly after admission and daily for 6 days. For endotoxin measurements, 2 mL of blood was drawn into 5-mL pyrogen-free plastic vials (Falcon; Becton Dickinson Labware, Lincoln Park, NJ) containing 50 IU of pyrogen-free heparin and centrifuged at 200 g for 10 min. Plasma for cytokine assays was drawn into 4-mL tubes (Vacutainer System; Becton Dickinson, Rutherford, NJ) containing 48 µL of 15% EDTA(K3) and 250 µL of aprotinin (10,000 kallikrein-inactivating units/mL; Bayer, Leverkusen, Germany). The tubes were centrifuged immediately at 2250 g for 10 min and then at 15,000 g for 5 min to remove the platelets. For IL-6 measurements serum was used. Aliquots were stored at -20°C until assay.

The ex vivo production of cytokines was measured in whole blood using similar 4-mL tubes [23]. One tube was incubated without LPS: in the other, 50 µL of LPS (final concentration, 10 µg/mL; Escherichia coli O55:B5; Sigma, St Louis) was added under sterile conditions. After 24 h of incubation at 37°C, both tubes were centrifuged and handled as described above.

Endotoxin was measured in platelet-rich plasma by a chromogenic limulus amoebocyte lysate assay (KabiVitrum, Stockholm). TNF was determined by RIA as described by Van der Meer et al. [24] (detection level, 100 pg/mL). IL-1 was measured by RIA according to Lisi et al. [25] (detection level, 80 pg/mL). IL-6 was measured by ELISA as delineated by Barrera et al. [26] (detection level, 20 pg/mL). IL-1ra was determined by RIA according to Poutsika et al. [17] (detection level, 300 pg/mL). sTNFr were measured by an enzyme-linked immunobinding assay (Hoffmann-La Roche; detection level, 80 pg/mL for sTNFr-55 and 300 pg/mL for sTNFr-75); normal values measured in 19 healthy volunteers were 1470 ± 190 pg/mL (median ± SD) for sTNFr-55 and 2520 ± 660 pg/mL for sTNFr-75. To minimize analytical errors, all samples from the same patient were analyzed in the same run in duplicate.

### Results

**Circulating cytokines and sTNFr.** The plasma concentrations of TNF and IL-1 were below detection level in all samples. At admission, IL-6 (range, 365–2550 pg/mL; median, 860), IL-1ra (range, 2840–4680 pg/mL; median, 3740), sTNFr-55 (range, 2782–5215 pg/mL; median, 3873), and sTNFr-75 (range, 5700–17282 pg/mL; median, 12910) were increased (figure 1). Within 2 days, these concentrations fell to normal levels.

**Ex vivo cytokine production.** The ex vivo production of TNF without LPS stimulation was below detection level in all samples. With LPS stimulation, TNF production was below detection level during the acute stage of the disease. After 3 days, during convalescence, this production gradually recovered to 810–3430 pg/mL (median, 2260) at day 6 (figure 2).

The course of IL-1 production showed a similar pattern: no measurable production at admission and restored LPS-stimulated production during recovery at day 6 (range, 945–4300 pg/mL; median, 2325). IL-6 was detectable at admission in unstimulated and in LPS-stimulated tubes. However, after correction for the circulating serum concentrations, the ex vivo production of IL-6 at admission was negligible. Similar to the pattern of ex vivo production of TNF and IL-1, the LPS-stimulated production of IL-6 was restored during recovery, reaching 165–9000 pg/mL (median, 2850) at day 6.

The ex vivo production of IL-1ra followed a different pattern. At admission, the production of IL-1ra in both unstimulated and LPS-stimulated cultures appeared to be increased. The concentration at day 0 in the unstimulated tubes ranged from 4450 to 6510 pg/mL (median, 5570); in the LPS-stimulated tubes, this was 7080–12,990 pg/mL (median, 11,140). The LPS-stimulated production decreased during the next 2 days toward a stable median of 6710 pg/mL. For this pattern, the LPS-stimulated production of IL-6 was restored during recovery, reaching 165–9000 pg/mL (median, 2850) at day 6.

**Discussion**

We followed the concentrations of circulating cytokines and their ex vivo production during the course of meningococcal infections at admission and their dexamethasone therapy.

<table>
<thead>
<tr>
<th>Patient no...</th>
<th>Disease period before hospitalization (h)</th>
<th>Blood pressure (mm Hg)</th>
<th>Arterial HCO₃⁻ concentration (mmol/L)</th>
<th>Leukocytes in cerebrospinal fluid (x10⁶/L)</th>
<th>Leukocytes in peripheral blood (x10⁶/L)</th>
<th>Plasma endotoxin concentration (pg/mL)</th>
<th>Dexamethasone therapy</th>
</tr>
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<tbody>
<tr>
<td>1. F/3</td>
<td>18</td>
<td>100/55</td>
<td>20.0</td>
<td>24,320</td>
<td>30.9</td>
<td>&lt;12.5</td>
<td>3.5 (days) 1.000 (mg/kg/day)</td>
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<tr>
<td>2. F/5</td>
<td>16</td>
<td>80/50</td>
<td>16.9</td>
<td>11</td>
<td>7.3</td>
<td>162</td>
<td>4 (days) 0.60 (mg/kg/day)</td>
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<tr>
<td>3. M/6</td>
<td>36</td>
<td>110/85</td>
<td>17.0</td>
<td>13,800</td>
<td>19.2</td>
<td>&lt;12.5</td>
<td>7 (days) 0.80 (mg/kg/day)</td>
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<td>110/80</td>
<td>16.0</td>
<td>10,000</td>
<td>20.0</td>
<td>21</td>
<td>1.5 (days) 0.40 (mg/kg/day)</td>
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<tr>
<td>5. F/20</td>
<td>24</td>
<td>120/78</td>
<td>15.3</td>
<td>13,500</td>
<td>29.8</td>
<td>283</td>
<td>1 (days) 0.13 (mg/kg/day)</td>
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<th>Duration Dose</th>
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<td></td>
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Cytokines in Meningococcal Infections

The onset of acute meningococcal infections is abrupt, with a rather short disease period before hospitalization. As such, meningococcal infections resemble experimental models using endotoxin challenge.

In our patients, circulating TNF and IL-1 concentrations were all below detection level. Several factors may be responsible for this negative finding. It may be possible that we missed the initial cytokinemia. The disease period before admission varied from 16 to 36 h, and it is known that after LPS injection in human volunteers, TNF and IL-1 levels peak within 90–180 min [1, 2]. From clinical studies, we know that high TNF concentrations occur early during severe meningococcal infections [8–10]. The relatively low endotoxin concentrations at admission, the good prognostic score, and the absence of shock in 4 of the 5 patients may also explain the low TNF and IL-1 concentrations [8–10, 27, 28]. Finally, a slight increase in TNF and IL-1 could have been missed because of the relatively high detection level of our assays. Recently, we have been able to detect rapidly declining TNF and IL-1 concentrations early in the course of very severe and lethal meningococcal disease (unpublished data). In the present series, actual activation of the cytokine network was documented by elevated concentrations of IL-6. Furthermore, the increased sTNFr-55 and sTNFr-75 levels are compatible with TNF activity on target cells in tissues, because binding of TNF to its receptors on these cells leads to shedding of the extramembranous part of the receptor [5, 29]. IL-1ra concentrations were also elevated during the acute stage. The high concentrations of both sTNFr and IL-1ra early in the course of the infection suggest a regulating role for these antiinflammatory compounds.

The LPS-stimulated ex vivo production of the proinflammatory cytokines TNF, IL-1, and IL-6 was suppressed during the acute stage. Gradually the production capacity was restored during reconvalescence. In contrast, the capacity to produce ex vivo IL-1ra was regulated inversely (figure 2). The LPS-stimulated IL-1ra production was maximal during the acute stage and reached stability after 2 days. Moreover, in the unstimulated cultures at admission, a spontaneous production of IL-1ra was observed, which gradually decreased during convalescence.

We used a whole blood culture system to measure the ex vivo cytokine production. In vitro studies with isolated peripheral blood mononuclear cells (PBMC) or monocytes...
have yielded a substantial part of our knowledge of the cytokine response to infectious stimuli. However, isolation of PBMC requires large blood volumes and is laborious and difficult to organize. Whole blood culture systems are a suitable alternative in these circumstances [23, 30-32]. In addition, the whole blood system represents the cytokine response of all types of cells present in blood at a certain time point and may therefore be a more realistic assessment than measurements using a fixed number of isolated and cultured cells.

Impaired ex vivo production of the proinflammatory cytokines TNF and IL-1 during acute infection has been reported by others [18-21]. Our study confirms these observations but also shows that the impaired capacity to produce proinflammatory cytokines does not indicate that the cells present in blood are completely refractory in terms of cytokine production. Rather, they seem to have switched from a mainly proinflammatory action (as in healthy individuals) to an antiinflammatory action. During recovery, the cells gradually switch again to the balanced proinflammatory status.

Further research should be done on the mechanism behind this programmed switch. Our findings in meningococcal infection may represent a process similar to endotoxin tolerance, the phenomenon that survival in experimental animals is markedly increased if a lethal dose of endotoxin is preceded by a smaller dose. Inhibition of the production of TNF, IL-1, and IL-6 is regarded as crucial [33]. Similarly, in human volunteers receiving endotoxin intravenously, the ex vivo production by CD14+ cells of TNF, IL-1, and IL-6 is significantly decreased [34]. IL-4 and IL-10 may play a role in this process. IL-10 and IL-4 both down-regulate the LPS-stimulated production by human PBMC and monocytes of TNF, IL-1, and IL-6, whereas preincubation with IL-4 increases the synthesis of IL-1ra [35-37]. Better insight into these regulatory mechanisms may enable us to understand why in some patients the invasion in the bloodstream of Neisseria meningitidis induces high plasma concentrations of TNF, IL-1, and IL-6 with severe shock, whereas in others a localized meningitis develops with low systemic concentrations of these cytokines [8-10]. Further study on these regulatory mechanisms may provide new therapeutic interventions.

In conclusion, antiinflammatory compounds such as sTNFr and IL-1ra are present in the circulation during early meningococcal infection. The down-regulation of the production of TNF, IL-1, and IL-6 and up-regulation of the production of IL-1ra during acute infection could serve as a mechanism of protection.

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References


