Inflammatory cytokines in CSF in bacterial meningitis: association with altered blood flow velocities in basal cerebral arteries

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Abstract

Objective—To investigate the association between release of humoral inflammatory mediators in CSF and blood and alterations of cerebral blood flow in patients with bacterial meningitis.

Methods—Immunomodulatory (interleukin-1β (IL-1β), interleukin-6 (IL-6), and tumour necrosis factor-α (TNFα)) and vasoactive (thromboxane A, prostacyclin, endothelin-1) molecules of probable or confirmed leucocyte origin were determined in CSF and venous blood from 20 patients with bacterial meningitis, and matched control subjects. Their concentrations were related to the presence of increased blood flow velocities in the middle cerebral arteries, as recorded by transcranial Doppler sonography.

Results—Concentrations of proinflammatory cytokines and prostacyclin and leucocyte counts were significantly increased in meningitis, but concentrations of the vasoconstrictors thromboxane and endothelin-1 were not. Patients with high blood flow velocities (>140 cm/s) had significantly increased concentrations of IL-1β and IL-6 and raised cell counts in CSF.

Conclusion—The increases of key mediators of inflammation and immunomodulation and of leucocyte count in the CSF of patients with high cerebral blood flow velocities suggest a role of excessive compartmentalised host defence in pathogenesis of disorders of cerebral blood flow in bacterial meningitis.

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Cerebrovascular disorders have been documented as one of the most common intracranial complications in adults with bacterial meningitis.1 Narrowing of basal cerebral arteries has been shown by angiography at sites where they are bathed within the subarachnoidal exudate in the cisterns and subarachnoidal space, and has been discussed as a pathomechanism of cerebral infarctions seen in this disease.2 4 An increased cerebral blood flow velocity (CBFV) in the basal cerebral arteries has been shown in bacterial meningitis by transcranial Doppler sonography.5-7 An inverse relation between vessel diameter on angiography and CBFV on transcranial Doppler sonography has been found,9 and there is considerable evidence that these alterations reflect changes in calibre of the insonated vessels as a result of transient or persistent narrowing of cerebral vessels.5-7 The non-invasive character of this method facilitates studies of disorders of cerebral blood flow in patients with meningitis.

Most humoral factors considered to play a part in the pathogenesis of vasospasms in subarachnoidal haemorrhage—for example, erythrocyte or platelet derived products—are absent in bacterial meningitis. The hallmark of bacterial meningitis is a massive leucocyte infiltration into the perivascular spaces and the CSF.5 These cells act through synthesis of proinflammatory cytokines (interleukin-1β (IL-1β), interleukin-6 (IL-6), and tumour necrosis factor-α (TNFα)), which orchestrate the local and systemic aspects of host response to infection and tissue damage.10-11 Huge amounts of these cytokines have recently been detected in CSF in bacterial meningitis.12 Arachidonic acid metabolites are also important mediators of inflammation.13 An imbalance between the potent vasoconstrictor thromboxane A, and the vasodilator prostacyclin has been implicated in the pathogenesis of cerebral vasospasms after subarachnoidal haemorrhage.14 Their synthesis also belongs to the repertoire of activated leucocytes.15 Endothelin-1 induces extremely sustained vasoconstriction, and is discussed as a causal factor in vasospasms secondary to subarachnoidal haemorrhage.15-17 Its synthesis by leucocytes after bacterial lipopolysaccharide stimulation,18 the induction of synthesis by polymorphonuclear leucocytes19 and cytokines,20 21 and the presence of a consensus sequence for acute phase elements in its gene22 prompted us to include investigation of endothelin-1 in this study.

By contrast with the many studies on subarachnoidal haemorrhage, humoral factors have, to our knowledge, not been investigated in patients with alterations of cerebral blood flow in meningitis. The aim of this study was to characterise the release of immunocompetent and vasoactive leucocyte products within the CSF in relation to alterations of cerebral blood flow in patients with bacterial meningitis.

Patients and methods

PATIENTS

Twenty patients (seven women and 13 men) aged between 18 and 85 (median 45) with
pyogenic meningitis, admitted within 48 hours after onset of first symptoms, were included in this study. Diagnosis was based on evidence from cultures of a predominant microorganism from a CSF sample and/or pleocytosis of > 1000 cells/mm³, > 60% polymorphonuclear CSF leucocytes, pronounced increase in protein concentration, clinical signs and symptoms of bacterial meningitis (fever, nuchal rigidity, headache, photophobia), and evidence for a systemic inflammatory response (increased white cell count and concentrations of C reactive protein). The causative organisms were Streptococcus pneumoniae (n = 5), Staphylococcus aureus (n = 4), Escherichia coli (n = 2), Neisseria meningitidis (n = 2), Haemophilus influenzae (n = 2), Mycobacterium tuberculosis (n = 1), and Listeria monocytogenes (n = 1). In three patients, no aetiological microorganism was isolated. These patients had received antibiotics before admission to hospital. Moreover, they showed a rapid response to antibiotic treatment. In 12 patients, the infection was community acquired, and in eight nosocomial. Other diseases known to be associated with alterations of blood flow in basal arteries (for example, subarachnoid haemorrhage) were excluded by clinical and laboratory examinations and by cranial CT.

Samples of CSF and serum from 20 subjects matched in age and sex (10 women and 10 men) aged between 26 and 71 (median 43) were used for controls. These were examined for exclusion of possible inflammatory disease, but turned out to have no neurological disease after extensive clinical examination and laboratory tests, including standard blood and CSF analyses.

TRANSCRANIAL DOPPLER SONOGRAPHY
The CBFV was recorded with transcranial Doppler ultrasound in parallel with collection of blood and CSF samples. Transcranial Doppler examinations were performed with a DWL Multidop X TCD device (DWL Sipplingen, Germany). Maximum systolic CBFV values were measured transtemporally in the proximal segment of the middle cerebral arteries. Reference values were derived from our transcranial Doppler readings in normal volunteers (table 1). Patients with systolic blood flow velocities in one or both middle cerebral arteries exceeding 140 cm/s (exceeding normal values by 3 SD) were considered to have raised CBFV. This cut off value had been shown to be useful in earlier studies.

SAMPLE HANDLING AND ROUTINE ANALYSES
Paired samples of blood and CSF from patients with meningitis were obtained at admission (within 48 hours after the onset of symptoms). Routine CSF analysis included cell count, cytology, albumin, and the albumin quotient ([CSF albumin] × 10³/[serum albumin]). For analysis, the collected CSF and blood samples were centrifuged at 3000 rpm for five minutes and the supernatant was stored at −80°C until used.

PROINFLAMMATORY CYTOKINES
Concentrations of IL-1β, IL-6, and TNFα were determined in CSF and serum with quantitative “sandwich” enzyme immunoassays (R and D Systems, Minneapolis, MN, USA), with antibodies specific for these cytokines coated on to microtitre plates and enzyme linked polyclonal specific antibodies, added after washing. The lower limits of detection for IL-1β, IL-6, and TNFα were 0·3 pg/ml, 0·7 pg/ml, and 4·4 pg/ml, respectively.

EXTRACTION AND QUANTIFICATION OF EICOSANOIDS
Eicosanoids were extracted from CSF and plasma by addition of chloric acid and ethyl acetate. The organic phase was evaporated under N2 and reconstituted with radiolmmunoassay buffer. The stable metabolites thromboxane B2 and 6-keto-prostaglandin F1α were used to estimate the amounts of thromboxane A2 and prostacyclin respectively. Thromboxane B2 and 6-keto-prostaglandin F1α were both determined by radiolmmunoassays (PerSeptive Diagnostics, Cambridge, MA). Briefly, these are based on competition of analyte in the samples with [14C] labelled analyte for a limited number of sites on a specific rabbit antianalyte antibody. Antibody bound analyte is separated from unbound analyte with goat antirabbit IgG through centrifugation, and the antibody bound labelled analyte is quantified in a gamma counter. The sensitivity of determination of thromboxane B2 and PGF1α was < 6·0 pg/ml and < 3·3 pg/ml respectively.

EXTRACTION AND ASSAY OF ENDOTHELIN-1
Endothelin-1 was extracted from acidified samples on C18 columns by the addition of acetic acid and evaporated under nitrogen gas. After reconstitution in an assay buffer, the extracted endothelin was measured in a radioimmunoassay (Nichols Institute Diagnostics, San Juan Capistrano, CA). Radio-labelled endothelin-1 competes with endothelin in the test samples for a limited number of rabbit antianalyte sites. Antibody bound endothelin is separated from free endothelin using antirabbit IgG antibody coated cellulose in suspension as the solid phase. The unbound endothelin is decanted after centrifugation and the endothelin is measured in a gamma counter. The sensitivity of the assay was 2 pg/ml.

STATISTICS
Because considerable amounts of prostaglandins and endothelin are present in normal serum, a “protein index”—by analogy with the IgG index—was used to correct for passage in CSF through a disrupted blood brain barrier. Results are expressed as mean (SE). Values were log transformed before statistical analysis, because their frequency distributions were asymmetric. For statistical analysis, the Student’s t test for unpaired data with a Bonferroni correction was employed to assess the significance of the difference between
groups. Thus values of P < 0.05/10 (10 being the number of the different mediators investigated in CSF) were required for differences to be considered significant. For the comparison of the differences between patients with high and low CBFV, values of P < 0.05/5 (five being the number of the different mediators tested) were required to be considered significant.

Results

Cerebral Blood Flow Velocity

Table 1 presents values of CBFV in patients with meningitis, as well as reference values. Eleven of 20 patients with bacterial meningitis had high CBFVs. A considerable asymmetry was noted. Only seven of 11 of the patients in the subpopulation with high CBFVs showed simultaneously increased values in both middle cerebral arteries.

Cell Counts and Albumin Quotient

As expected, cell counts were significantly increased in the CSF of patients with meningitis (table 2). Cell counts were significantly increased in patients with high CBFVs (figure). Albumin quotients were massively increased in bacterial meningitis (mean 30.7-70 (SE 7.27)).

Proinflammatory Cytokines

Concentrations of the cytokines IL-1β, TNFα, and IL-6 were significantly increased in CSF of patients with meningitis (table 2). Concentrations of IL-1β and IL-6 were significantly raised in patients exhibiting increased CBFVs (figure), whereas the concentrations of TNFα tended to be increased in this subpopulation, but without reaching statistical significance.

Thromboxane and Prostacyclin

Concentrations of the stable metabolites of prostacyclin (6-keto prostaglandin F1α) were significantly increased in the CSF of patients with bacterial meningitis. However, concentrations of this prostaglandin did not significantly differ in the study groups with higher or lower CBFV (figure). Although the highly variable CSF concentrations of the stable metabolite of thromboxane A2 (thromboxane B2) tended to be raised in meningitis, a significant difference could not be found. The indices of both eicosanoids studied were not significantly increased.

Endothelin-1

Concentrations of this vasoconstrictor were highly variable in patients with meningitis (table 2). Four patients with bacterial meningitis exhibited high CSF concentrations (13.58, 14.08, 18.77, and 19.52 pg/ml). However, when the study populations with and without meningitis were compared, neither CSF concentrations nor the corresponding index differed significantly (table 2).

Table 2  Mean concentrations (SE) of immunomodulatory and vasoactive mediators in patients with meningitis and controls.

<table>
<thead>
<tr>
<th>Variable</th>
<th>Control subjects</th>
<th>Patients with meningitis</th>
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<tbody>
<tr>
<td></td>
<td>Serum</td>
<td>Index†</td>
</tr>
<tr>
<td>Cells/mm³</td>
<td>0.9 (0.3)</td>
<td>—</td>
</tr>
<tr>
<td>IL-1β (pg/ml)</td>
<td>0.0 (0.0)</td>
<td>0.0 (0.0)</td>
</tr>
<tr>
<td>IL-6 (pg/ml)</td>
<td>6.1 (2.5)</td>
<td>1.3 (0.2)</td>
</tr>
<tr>
<td>TNFα (pg/ml)</td>
<td>0.2 (0.1)</td>
<td>2.1 (0.4)</td>
</tr>
<tr>
<td>TXB (pg/ml)</td>
<td>15.8 (6.4)</td>
<td>120.1 (378.1)</td>
</tr>
<tr>
<td>PGI2 (pg/ml)</td>
<td>2.4 (1.1)</td>
<td>8.2 (2.5)</td>
</tr>
<tr>
<td>ET-1 (pg/ml)</td>
<td>1.8 (1.0)</td>
<td>3.4 (0.4)</td>
</tr>
</tbody>
</table>

*P < 0.05 (after Bonferroni correction).
†Indices for correction of passage of proteins from peripheral blood in CSF according to the formula:

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\text{Protein index} = \frac{\text{CSF protein}}{\text{serum protein}} = \frac{\text{CSF albumin}}{\text{serum albumin}}
\]
Discussion

Many factors discussed as inducers of vasospasms in subarachnoid haemorrhage (for example, erythrocyte or platelet derived substances) are absent in bacterial meningitis. This investigation of humoral factors possibly involved in pathogenesis of cerebrovascular complications in bacterial meningitis focused on leucocytes and inflammatory mediators. The results show significantly higher leucocyte counts and a significantly increased compartmentalisation of IL-1β and IL-6 in the CSF of patients with high CBFVs. Concentrations of keto-prostaglandin F₂α—stable metabolite of prostacyclin—but not of thromboxane B₂—the stable metabolite of thromboxane A₂—were significantly increased in meningitis. A clear shift of the equilibrium between the antagonistic acting eicosanoids towards the vasoconstrictor thromboxane A₂ could not be demonstrated. Concentrations of endothelin-1 in CSF were highly variable, as were concentrations of the eicosanoids.

Possible reasons may be a considerable variation in the degree of blood-brain barrier disruption, or a different extent of systemic inflammation and infection in these patients. Concentrations of endothelin tended to be increased, but not significantly, in patients with meningitis. Interestingly, four patients exhibited high endothelin concentrations in CSF but not in venous blood, suggesting intrathecal synthesis in some patients. The cause of relatively high indices of eicosanoids and endothelin in normal controls is unclear. Concentrations in a similar range have been reported. Possible explanations may be a higher rate of proteolytic degradation in venous blood than in CSF, or an intrathecal synthesis even in the normal condition.

As in subarachnoid haemorrhage, in which the research for one single factor causing vasospasm has been disappointing, a multifactorial origin of vasospasms is most likely in meningitis. The increased compartmentalisation of proinflammatory cytokines and raised leucocyte counts in patients with high CBFVs suggests a role of excessive host defence to bacterial outer cell membrane components in pathogenesis of vasospasms. The interleukins IL-1β and IL-6 play a key part in inflammation and early immunooactivation, orchestrating a cascade of further proinflammatory and contrainflammatory immunomediators. High concentrations of these immunomediators in CSF may exert vasoactive effects—for example, IL-1β (MW 17 000 Da) or IL-6 (MW 26 000 Da) could easily get access to the walls of contiguous basilar arteries from their adventitial side—as even larger molecules (for example, horseradish peroxidase, MW 40 000 Da) pass from the cisterna magna through the vessel wall to the basal membrane within minutes. This is possible because, by contrast with other arteries, the surface of the major cerebral arteries is not confined by collagen or fibroblasts, but is in direct contact with the CSF. At this location, proinflammatory cytokines could alter vessel tone either directly or indirectly—for example, by induction of synthesis of endothelin-1 by endothelial or other cells, as has recently been shown in vitro. It is still controversial whether inflammatory exudate is harmful or helpful in infection. Vasocostrictive properties of leukocytes and their products could also be, within certain limits, of benefit in acute inflammation—for example, in impairment of spreading of infectious agents or pluriopotent immunomediators. Earlier studies showed that adjuncts to antibiotic treatment that reduce inflammation—for example, corticosteroids or treatments aimed to decrease the accumulation of leukocytes in CSF—have the potential to reduce mortality, although further research is necessary. Such strategies could be useful in prophylaxis and treatment of cerebrovascular complications of bacterial meningitis. Because IL-6 and other cytokines have recently been detected in the CSF of patients with subarachnoid haemorrhage, investigations of immunomediators may also be an interesting approach in elucidation of the pathogenesis of vasospasms in subarachnoid haemorrhage.

In conclusion, the increase of key mediators of immunoactivation and of leucocyte counts in the CSF of patients with high CBFVs suggests a role of excessive host defence compartmentalised in the subarachnoidal space in pathogenesis of vasospasms in bacterial meningitis.

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14. Chan RC, Durity FA, Thompson GB, Nugent RA, Kendall M. The role of the prostacyclin-thromboxane system in cerebral vasospasm following induced sub-
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Tinel's sign of formalization

Few signs have caused more controversy than the sign of Tinel.

Julius Tinel (1879–1952) was son of the professor of anatomy at Rouen. He worked with Dejerine, and became chief de clinique in 1911 and chief of the laboratory at Salpêtrière in 1913. He was very active with his son Jacques in the French resistance movement in the second world war, and was imprisoned in Bordeaux at Fort de Ha. His book Nerf Wound records 600 patients with nerve injuries sustained in the first world war. He called his sign the sign of formalization.

"The all important sign is formalization. We find that sudden pressure or percussion of the nerve trunk, below the lesion, calls forth a tingling sensation in the cutaneous region of the nerve... It appears about the fourth or sixth week... Then it gradually becomes more pronounced and it is possible to follow, week by week, in the course of the nerve, the progress of this provoked formalization, pari passu with the advance of the axis cylinders. The formalization sign is thus of supreme importance since it enables us to see whether the nerve is interrupted, or in the course of regeneration; whether a nerve suture has succeeded or failed, or whether regeneration is rapid and satisfactory, or reduced to a few significant fibres."

Tinel's sign fell into disrepute when a positive sign was elicited in a patient in whom it was shown that there was an anatomical gap at the site of nerve injury. The salient feature is "peripheral reference" of the tingling sensation on percussion of the nerve that occurs in normal nerves; but in a pathological nerve it is more easily provoked and may persist longer. It may be used to localise the site of injury and suggests the presence of regeneration. Now widely used in nerve entrapment syndromes, it remains as a "soft sign". Second world war experience proved that a negative Tinel's sign is of no diagnostic value.

The translation of the two papers of Paul Hoffman, published in the same year that Tinel wrote his article (1915) has caused some authorities to refer to the Hoffman-Tinel sign.

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References: