Clinical features and prognostic factors in adults with bacterial meningitis
van de Beek, D.; de Gans, J.; Spanjaard, L.; Weisfelt, M.; Reitsma, J.B.; Vermeulen, M.

Published in:
The New England journal of medicine

DOI:
10.1056/NEJMoa040845

Link to publication

Citation for published version (APA):

General rights
It is not permitted to download or to forward/distribute the text or part of it without the consent of the author(s) and/or copyright holder(s), other than for strictly personal, individual use, unless the work is under an open content license (like Creative Commons).

Disclaimer/Complaints regulations
If you believe that digital publication of certain material infringes any of your rights or (privacy) interests, please let the Library know, stating your reasons. In case of a legitimate complaint, the Library will make the material inaccessible and/or remove it from the website. Please Ask the Library: http://uba.uva.nl/en/contact, or a letter to: Library of the University of Amsterdam, Secretariat, Singel 425, 1012 WP Amsterdam, The Netherlands. You will be contacted as soon as possible.
Clinical Features and Prognostic Factors in Adults with Bacterial Meningitis

Diederik van de Beek, M.D., Ph.D., Jan de Gans, M.D., Ph.D., Lodewijk Spanjaard, M.D., Ph.D., Martijn Weisfelt, M.D., Johannes B. Reitsma, M.D., Ph.D., and Marinus Vermeulen, M.D., Ph.D.

BACKGROUND
We conducted a nationwide study in the Netherlands to determine clinical features and prognostic factors in adults with community-acquired acute bacterial meningitis.

METHODS
From October 1998 to April 2002, all Dutch patients with community-acquired acute bacterial meningitis, confirmed by cerebrospinal fluid cultures, were prospectively evaluated. All patients underwent a neurologic examination on admission and at discharge, and outcomes were classified as unfavorable (defined by a Glasgow Outcome Scale score of 1 to 4 points at discharge) or favorable (a score of 5). Predictors of an unfavorable outcome were identified through logistic-regression analysis.

RESULTS
We evaluated 696 episodes of community-acquired acute bacterial meningitis. The most common pathogens were *Streptococcus pneumoniae* (51 percent of episodes) and *Neisseria meningitidis* (37 percent). The classic triad of fever, neck stiffness, and a change in mental status was present in only 44 percent of episodes; however, 95 percent had at least two of the four symptoms of headache, fever, neck stiffness, and altered mental status. On admission, 14 percent of patients were comatose and 33 percent had focal neurologic abnormalities. The overall mortality rate was 21 percent. The mortality rate was higher among patients with pneumococcal meningitis than among those with meningococcal meningitis (30 percent vs. 7 percent, P<0.001). The outcome was unfavorable in 34 percent of episodes. Risk factors for an unfavorable outcome were advanced age, presence of otitis or sinusitis, absence of rash, a low score on the Glasgow Coma Scale, and tachycardia, a positive blood culture, an elevated erythrocyte sedimentation rate, thrombocytopenia, and a low cerebrospinal fluid white-cell count.

CONCLUSIONS
In adults presenting with community-acquired acute bacterial meningitis, the sensitivity of the classic triad of fever, neck stiffness, and altered mental status is low, but almost all present with at least two of the four symptoms of headache, fever, neck stiffness, and altered mental status. The mortality associated with bacterial meningitis remains high, and the strongest risk factors for an unfavorable outcome are those that are indicative of systemic compromise, a low level of consciousness, and infection with *S. pneumoniae*.
THE EPIDEMIOLOGY OF BACTERIAL MENINGITIS HAS CHANGED. Meningitis due to Haemophilus influenzae type b has been nearly eliminated in the Western world since vaccination against H. influenzae type b was initiated, and the introduction of conjugate vaccines against Streptococcus pneumoniae is expected to reduce the burden of childhood pneumococcal meningitis significantly. Although vaccination with a pneumococcal conjugate vaccine is producing herd immunity among adults, the age distribution of meningitis has now shifted to older age groups. Several studies of clinical features and prognostic factors in adults with bacterial meningitis have been performed; however, all were retrospective and relatively small in size. We performed a nationwide prospective study of clinical features and prognostic factors in adults with community-acquired bacterial meningitis in the Netherlands.

METHODS

We identified adults (defined as patients older than 16 years of age) who had bacterial meningitis and were listed in the database of the Netherlands Reference Laboratory for Bacterial Meningitis from October 1998 to April 2002. This laboratory receives cerebrospinal fluid and blood isolates from approximately 85 percent of all patients with bacterial meningitis in the Netherlands (population, 16.2 million). The laboratory provides daily updates of the names of hospitals where patients with bacterial meningitis have been admitted in the preceding two to six days and the names of physicians, usually neurologists. Physicians were informed about the study by telephone. Subsequently, patients or their legal representatives received written information concerning the study and were asked to give written informed consent for participation; only patients for whom consent was obtained were enrolled. Case-record forms were used to collect data on patients’ history, symptoms and signs on admission, laboratory findings at admission, clinical course, outcome and neurologic findings at discharge, and treatment.

Patients were categorized as having either community-acquired or hospital-acquired meningitis (the latter was defined as meningitis that occurred during hospitalization or within one week after discharge). Patients with an altered immune status owing to the use of immunosuppressive drugs or splenectomy, diabetes mellitus, or alcoholism were considered immunocompromised, as were patients infected with the human immunodeficiency virus. At discharge, all patients underwent a neurologic examination performed by a neurologist, and the outcome was graded according to the Glasgow Outcome Scale. A score of 1 on this scale indicates death; a score of 2, a vegetative state (the patient is unable to interact with the environment); a score of 3, severe disability (the patient is unable to live independently but can follow commands); a score of 4, moderate disability (the patient is capable of living independently but unable to return to work or school); and a score of 5, mild or no disability (the patient is able to return to work or school). A favorable outcome was defined as a score of 5, and an unfavorable outcome was as a score of 1 to 4. The Glasgow Outcome Scale is a well-validated instrument with good interobserver agreement.

The susceptibility of meningococci to penicillin was determined by inoculating strains onto chocolate agar containing 0.1 mg of penicillin per liter. A 1-µg oxacillin disk was used to identify penicillin-resistant strains of pneumococci. Whenever a strain showed antibiotic resistance, the E rosette assay was used to determine the minimal inhibitory concentration of the antibiotic. The inoculation procedure and susceptibility testing were performed as described elsewhere. Parts of this cohort study have been reported previously.

This observational study used patient data that had been rendered anonymous and was carried out in accordance with Dutch privacy legislation. The study was announced in the journal of the Dutch Neurologic Society, followed by periodic reminders. Before the study, all Dutch neurologists received information about the study, including a case-record form.

Parametric and nonparametric tests were used to identify differences between groups in continuous outcomes, and chi-square tests were used to compare categorical outcomes. We used logistic regression to examine the association between potential predictors and the likelihood of an unfavorable outcome. Odds ratios and 95 percent confidence intervals were used to quantify the strength of these associations. On the basis of previous research and pathophysiologic interest, 20 potentially relevant predictors were chosen.

Despite the low median percentage of missing values for individual variables (2 percent), only 320 of the 696 patients had complete data on all predictors — which presented a considerable
limitation for multivariate models. Therefore, we used multiple imputation techniques to reduce this loss. All predictors together were used to impute missing values on the basis of multivariate normal distributions. The coefficients of five rounds of imputation were combined to obtain the final estimates for the multivariate model. The low rate of missing values per variable, the large number of predictors, and the large sample size maximized the benefits of the multiple imputation methods. All statistical tests were two-tailed, and P values of less than 0.05 were considered to indicate statistical significance.

Statistical analyses were performed with use of SAS software, version 8.02 (SAS Institute). The study was designed, conducted, and analyzed independently of the sponsor.

**RESULTS**

A total of 1108 episodes of bacterial meningitis were identified by the reference laboratory. A total of 994 case-record forms were sent out, and the response rate was 76 percent (754 of 994) (Fig. 1). The demographic characteristics of patients with meningitis identified by the laboratory and those included in the analysis were similar for each causative organism. Fifty-eight patients were excluded — 50 patients with hospital-acquired meningitis, 3 patients with a recent history of neurosurgery, and 5 patients with a neurosurgical device — leaving a total of 696 episodes of community-acquired meningitis in 671 patients. The annual incidence of community-acquired bacterial meningitis was 2.6 cases per 100,000 adults.

Characteristics of the study population are shown in Table 1. Seizures occurred before admission in 32 of 666 episodes (5 percent). Predisposing conditions were present in 48 percent of episodes, the most common of which were otitis or sinusitis in 25 percent, pneumonia in 12 percent, and an immunocompromised state in 16 percent. Patients with pneumococcal meningitis were more likely to have distant foci of infection than were patients with meningococcal meningitis (62 percent vs. 9 percent, P<0.001).

Classic symptoms and signs of bacterial meningitis were present in a large proportion of patients (Table 1). Headache occurred in 87 percent of episodes, neck stiffness in 83 percent, fever in 77 percent, and a change in mental status (defined by a Glasgow Coma Scale score below 14) in 69 percent. Only 44 percent of episodes were characterized by the classic triad of fever, neck stiffness, and a change in mental status. At least two of four signs (the classic triad plus headache) were present in...
664 episodes (95 percent), only one in 28 episodes (4 percent), and none in 4 episodes (1 percent). The classic triad was significantly more likely to be present in patients with pneumococcal meningitis than in those with meningococcal meningitis (58 percent vs. 27 percent, \( P<0.001 \)). Rash was present in 176 of 683 episodes (26 percent); the causative species was *Neisseria meningitidis* in 162 episodes, *S. pneumoniae* in 8, *Staphylococcus aureus* in 2, group B streptococcus in 2, *H. influenzae* in 1, and *Listeria monocytogenes* in 1. The rash was petechial in 157 of these 176 episodes (89 percent).

### Table 1. Characteristics of the Study Population.

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Episodes of Meningitis (N=696)</th>
<th>Characteristic</th>
<th>Episodes of Meningitis (N=696)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Symptoms on presentation</strong></td>
<td></td>
<td><strong>Indexes of CSF inflammation</strong></td>
<td></td>
</tr>
<tr>
<td>Headache — no./no. evaluated (%)</td>
<td>544/626 (87)</td>
<td>-opening pressure — mm of water††</td>
<td>370±130</td>
</tr>
<tr>
<td>Nausea — no./no. evaluated (%)</td>
<td>449/610 (74)</td>
<td>Mean — cells/mm&lt;sup&gt;3&lt;/sup&gt;</td>
<td>7753±14,736</td>
</tr>
<tr>
<td>Neck stiffness — no./no. evaluated (%)</td>
<td>569/685 (83)</td>
<td>&lt;100/mm&lt;sup&gt;3&lt;/sup&gt; — no./no. evaluated (%)</td>
<td>47/645 (7)</td>
</tr>
<tr>
<td>Rash — no./no. evaluated (%)</td>
<td>176/683 (26)</td>
<td>100–999/mm&lt;sup&gt;3&lt;/sup&gt; — no./no. evaluated (%)</td>
<td>93/645 (14)</td>
</tr>
<tr>
<td>Systolic blood pressure — mm Hg</td>
<td>144±33</td>
<td>&gt;999/mm&lt;sup&gt;3&lt;/sup&gt; — no./no. evaluated (%)</td>
<td>505/645 (78)</td>
</tr>
<tr>
<td>Diastolic blood pressure — mm Hg</td>
<td>79±20</td>
<td>Protein — g/liter</td>
<td>4.9±4.5</td>
</tr>
<tr>
<td>Body temperature</td>
<td></td>
<td>CSF:blood glucose ratio</td>
<td>0.2±0.2</td>
</tr>
<tr>
<td>Mean — °C</td>
<td>38.8±1.2</td>
<td>Positive blood culture — no./no. evaluated (%)</td>
<td>404/611 (66)</td>
</tr>
<tr>
<td>≥38°C — no./no. evaluated (%)</td>
<td>522/678 (77)</td>
<td>Blood chemical tests</td>
<td></td>
</tr>
<tr>
<td>Score on Glasgow Coma Scale&lt;sup&gt;§&lt;/sup&gt;</td>
<td></td>
<td>ESR — mm/hr††</td>
<td>46±37</td>
</tr>
<tr>
<td>Mean</td>
<td>11±3</td>
<td>C-reactive protein — mg/liter†‡‡</td>
<td>225±132</td>
</tr>
<tr>
<td>&lt;14 (indicating change in mental status) — no. (%)</td>
<td>477 (69)</td>
<td>Thrombocyte count — platelets/mm&lt;sup&gt;3&lt;/sup&gt;§§</td>
<td>198,000±100,000</td>
</tr>
<tr>
<td>&lt;8 (indicating coma) — no. (%)</td>
<td>96 (14)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

* The study included 671 patients who had a total of 696 episodes of community-acquired meningitis. Plus–minus values are means ±SD.
† Immunocompromise was defined by the use of immunosuppressive drugs, a history of splenectomy, or the presence of diabetes mellitus or alcoholism, as well as patients infected with the human immunodeficiency virus (HIV).
‡ The number of patients tested for HIV infection is unknown.
§ Scores on the Glasgow Coma Scale can range from 3 to 15, with 15 indicating a normal level of consciousness. Glasgow Coma Scale scores were evaluated in 694 patients; 1 patient with pneumococcal meningitis and 1 with meningococcal meningitis were not evaluated.
¶ The CSF pressure was measured in 216 patients.
** Blood culture was performed in 611 patients.
†† The erythrocyte sedimentation rate (ESR) was determined in 549 patients.
‡‡ C-reactive protein levels were determined in 394 patients.
§§ The thrombocyte count was determined in 653 patients.
cent). The rash was petechial in 13 of 14 episodes that were characterized by a rash and a nonmenin-
gococcal cause of meningitis.

In 14 percent of episodes, the patients were co-
matose on admission, and in 33 percent of epi-
sodes, focal neurologic deficits were present on ad-
mission. Patients with pneumococcal meningitis
had more severe disease than did patients with me-
ingococcal meningitis, as reflected by a higher fre-
frequency of seizures (P=0.001) and focal neurologic
deficits (P<0.001) and a lower level of conscious-
ness (P<0.001). Results of funduscopic examina-
tion were recorded for 386 episodes; 13 episodes
were characterized by papilledema (3 percent).

Lumbar puncture was performed in all patients.
The time between admission and lumbar puncture
was not recorded. Cerebrospinal fluid pressure was
evaluated in 216 episodes. Opening pressures were
normal (less than 200 mm of water) in 38 episodes
(18 percent) and exceeded 400 mm of water in 85
(39 percent). Patients with opening pressures of
more than 400 mm of water were more likely to be
admitted in a coma than were those with lower
opening pressures (14 of 131 [11 percent] vs. 20 of
85 [24 percent], P=0.01). However, the percentage
of patients with an unfavorable outcome was simi-
lar in the two groups (30 percent and 35 percent,
respectively). The mean opening pressures were
similar among patients with papilledema and those
without papilledema. At least one individual cere-
brosplinal fluid finding predictive of bacterial men-
ingitis (a glucose level of less than 34 mg per deci-
liter [1.9 mmol per liter], a ratio of cerebrospinal
fluid glucose to blood glucose of less than 0.23,
a protein level of more than 220 mg per deciliter,
or a white-cell count of more than 2000 per cubic
millimeter)31 was present in 567 of 645 episodes
(88 percent).

Cranial computed tomography (CT) was per-
formed on admission in 496 episodes (71 percent);
the results were normal in 325. Abnormalities were
recorded in 171 episodes (34 percent): cerebral edema
was identified in 48 of 496 episodes (10 percent),
sinusitis or otitis in 48 (10 percent), evidence of
recent infarction in 30 (6 percent), and hydro-
cephalus in 15 (3 percent). Thirty-eight episodes
involved other abnormalities on cranial CT; old vas-
cular lesions in 12, cerebral atrophy in 6, skull frac-
ture in 5, pneumatocepha!us in 5, arachnoid cyst in
3, vascular aneurysm in 2, subarachnoid hemor-
rhage in 1, brain-parenchyma hemorrhage in 1, sub-
dural empyema in 1, meningioma in 1, and Dandy-
Walker malformation in 1; in 8 episodes, two ab-
normalities were identified.

Cranial CT was performed before lumbar punc-
ture in 337 of 696 episodes (48 percent). Focal neu-
rologic deficits (not including cranial-nerve abnor-
malities), a score of less than 10 on the Glasgow
Coma Scale, or both were present in 313 of 696 epi-
sodes (45 percent), and cranial CT was performed
before lumbar puncture in 197 of these 313 epi-
sodes (63 percent). Therapy was initiated before
CT in 35 percent of episodes in which CT was per-
formed before lumbar puncture.

The most common microorganism was S. pneu-
moniae, accounting for 51 percent of isolates cul-
tured from cerebrospinal fluid (Fig. 1). The distri-
bution of serotypes among 352 pneumococci is
shown in Table 2. N. meningitidis was responsible
for 37 percent of the episodes, with group B identi-
fied in 173 episodes, group C in 79, group Y in 3,
group H in 1, and group W135 in 1. Gram’s stain-
ing of cerebrospinal fluid revealed the microor-
ganism in 524 of 652 episodes (sensitivity, 80 per-
cent; specificity, 97 percent). The yield of Gram’s
staining was similar in patients who had previously
received antimicrobial therapy and those who had
not. Antibiotic susceptibility was tested in 351 pneu-
omococcal and 256 meningococcal strains; 2 pneu-
omococcal and 4 meningococcal strains showed in-
termediate resistance to penicillin; all other strains
were sensitive to penicillin. Initial antibiotic treat-
ment consisted of penicillin or amoxicillin in 43 per-
cent of episodes, third-generation cephalosporins
in 16 percent, and a combination of penicillin or
amoxicillin with a third-generation cephalosporin
in 25 percent; other regimens were used in 16 per-
cent of episodes.

During the clinical course, focal neurologic ab-
normalities were found in half the episodes (Ta-
ble 3); most were present on admission. Seizures
occurred in 15 percent of episodes, and cardiores-
piratory failure in 29 percent. Complications were
significantly more likely to develop among patients
with pneumococcal meningitis than among pa-
tients with meningococcal meningitis (P<0.001).

The mortality rate was 21 percent (Table 3) and
varied depending on the causative organism; it was
30 percent for pneumococcal meningitis, as com-
pared with 7 percent for meningococcal meningi-
tis (P<0.001) (Table 3) and 20 percent for menin-
gitis due to other pathogens (P=0.05). Thirty-four
percent of episodes had an unfavorable outcome.
A neurologic examination was performed at dis-
Corticosteroids were administered in 118 episodes (17 percent), and episodes involving corticosteroid therapy were more likely to have an unfavorable outcome than were episodes that did not involve corticosteroid therapy (51 of 118 [43 percent] vs. 186 of 578 [32 percent], P=0.03). Episodes in which corticosteroids were administered before antibiotics were less likely to have an unfavorable outcome than episodes in which corticosteroids were administered after antibiotics (3 of 24 [12 percent] vs. 48 of 94 [51 percent], P=0.001). Opening pressures of more than 400 mm of water were recorded in 46 percent of patients who received corticosteroids and 38 percent of those not receiving corticosteroids.

In the multivariate model several characteristics were significantly associated with an unfavorable outcome (Table 4): advanced age, the presence of otitis or sinusitis, a heart rate of more than 120 beats per minute, a low score on the Glasgow Coma Scale, a cerebrospinal fluid white-cell count of fewer than 1000 per cubic millimeter, a positive blood culture, an elevated erythrocyte sedimentation rate, and a reduced platelet count. Six other characteristics tended toward statistical significance: the presence of symptoms for less than 24 hours before admission, seizures, pneumonia, an immunocompromised state, a heart rate below 60 beats per minute, and hypotension (defined as a diastolic blood pressure of less than 60 mm Hg).

The causative organism had an independent effect when it was added to the multivariate model. The odds of an unfavorable outcome was six times as high (95 percent confidence interval, 2.61 to 13.91; P<0.001) among patients infected with S. pneumoniae as among patients infected with N. meningitidis, even after adjustment for other clinical predictors.

During episodes of meningococcal meningitis, hypotension was more common among patients with a cerebrospinal fluid white-cell count of fewer than 100 cells per cubic millimeter than among patients with a higher cerebrospinal fluid white-cell count (8 of 18 [44 percent] vs. 24 of 211 [11 percent], P<0.001). This association was absent during episodes of pneumococcal meningitis.

The prevalence of the classic triad of fever, neck stiffness, and an altered mental status is low among adults with community-acquired bacterial meningitis. However, almost all patients (95 percent) presented with at least two of the four symptoms of headache, fever, neck stiffness, and an altered mental status. In addition, a high percentage of patients (33 percent) were admitted with focal neurologic deficits. These percentages are similar to those found in retrospective studies; however, we found an exceptionally high prevalence of aphasia.4-6,9-23

The majority of patients who presented with moderate or severe impairment of consciousness, neurologic deficits (not including cranial-nerve abnormalities), or both, which are contraindications to performing lumbar puncture, underwent CT before lumbar puncture (63 percent). However, a large number of patients without these “red flags” also underwent CT first, and many patients with such red flags did not undergo CT before lumbar puncture.

Table 2. Capsular Pneumococcal Serotypes from 352 Adults with Meningitis.

<table>
<thead>
<tr>
<th>Serotype</th>
<th>No. of Patients (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>3</td>
<td>36 (10)</td>
</tr>
<tr>
<td>14</td>
<td>34 (10)</td>
</tr>
<tr>
<td>19F*</td>
<td>29 (8)</td>
</tr>
<tr>
<td>7F</td>
<td>28 (8)</td>
</tr>
<tr>
<td>9V</td>
<td>27 (8)</td>
</tr>
<tr>
<td>6B</td>
<td>21 (6)</td>
</tr>
<tr>
<td>10A</td>
<td>21 (6)</td>
</tr>
<tr>
<td>8</td>
<td>16 (5)</td>
</tr>
<tr>
<td>4*</td>
<td>16 (5)</td>
</tr>
<tr>
<td>23F*</td>
<td>14 (4)</td>
</tr>
<tr>
<td>6A</td>
<td>10 (3)</td>
</tr>
<tr>
<td>19A</td>
<td>10 (3)</td>
</tr>
<tr>
<td>12F</td>
<td>10 (3)</td>
</tr>
<tr>
<td>Other†</td>
<td>80 (23)</td>
</tr>
</tbody>
</table>

* This serotype is included in the seven-valent conjugated vaccine (coverage, 149 of 352 isolates [42 percent]).† Other types were as follows: 22F in eight patients, 18C (included in the seven-valent conjugated vaccine) in eight, 35F in six, 1 in five, 9N in five, 17F in five, 38 in four, 15B in four, 16F in four, 18B in four, 33F in four, 23B in four, 24F in three, 34 in three, 5 in two, 15A in two, 12C in two, 20 in two, 22A in two, 9A in one, 18F in one, and 23A in one.
In most patients who underwent CT before lumbar puncture, therapy was not initiated before CT was performed. Cranial CT has been recommended as a precaution before lumbar puncture to predict the likelihood and avoid the possibility of brain herniation. The withdrawal of cerebrospinal fluid removes counterpressure from below, thus increasing the effect of compression from above and exacerbating the brain shift already present.

In patients with red flags indicating space-occupying lesions and in those with moderate or severe impairment of consciousness (as indicated by a score of less than 10 on the Glasgow Coma Scale), lumbar puncture should be preceded by cranial CT. However, a delay in the initiation of antimicrobial therapy may lead to a poor outcome, especially in patients with clinical deterioration. Therefore, in patients with suspected meningitis in whom cranial CT is performed before lumbar puncture, CT should be preceded by blood cultures and the initiation of antibiotic therapy and corticosteroids.

Community-acquired bacterial meningitis has a high rate of an unfavorable outcome in adults (34 percent). Our multivariate model identified several unfavorable prognostic factors, most of which pointed to systemic compromise. A low level of consciousness on admission was predictive of an unfavorable outcome, as was a low cerebrospinal fluid white-cell count. Finally, factors predictive of pneumococcal infection were associated with an unfavorable outcome (advanced age; presence of otitis or sinusitis, pneumonia, or immunocompromised status; and absence of rash).

In our cohort, one of every six patients received corticosteroids. The European Dexamethasone Study showed adjunctive treatment with dexamethasone to be beneficial in adults with bacterial meningitis.

Table 3. Clinical Course, Outcome, and Neurologic Findings at Discharge.

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>All Episodes of Meningitis (N=696)</th>
<th>Episodes of Pneumococcal Meningitis (N=352)</th>
<th>Episodes of Meningococcal Meningitis (N=257)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>number of episodes (percent)</td>
<td>number of episodes (percent)</td>
<td>number of episodes (percent)</td>
</tr>
<tr>
<td>Clinical course</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Focal neurologic deficits</td>
<td>351 (50)</td>
<td>227 (65)</td>
<td>85 (33)</td>
</tr>
<tr>
<td>Seizures</td>
<td>107 (15)</td>
<td>85 (24)</td>
<td>12 (5)</td>
</tr>
<tr>
<td>Cardiorespiratory failure</td>
<td>201 (29)</td>
<td>134 (38)</td>
<td>45 (18)</td>
</tr>
<tr>
<td>Mechanical ventilation</td>
<td>160 (23)</td>
<td>109 (31)</td>
<td>36 (14)</td>
</tr>
<tr>
<td>Score on Glasgow Outcome Scale</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1 (death)</td>
<td>143 (21)</td>
<td>107 (30)</td>
<td>19 (7)</td>
</tr>
<tr>
<td>2 (vegetative state)</td>
<td>3 (&lt;1)</td>
<td>3 (1)</td>
<td>0</td>
</tr>
<tr>
<td>3 (severe disability)</td>
<td>24 (3)</td>
<td>17 (5)</td>
<td>4 (2)</td>
</tr>
<tr>
<td>4 (moderate disability)</td>
<td>67 (10)</td>
<td>50 (14)</td>
<td>7 (3)</td>
</tr>
<tr>
<td>5 (mild or no disability)</td>
<td>459 (66)</td>
<td>175 (50)</td>
<td>227 (88)</td>
</tr>
<tr>
<td>Neurologic findings at discharge</td>
<td>number/number evaluated (percent)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cranial-nerve palsy</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>3rd nerve</td>
<td>7/550 (1)</td>
<td>6/243 (2)</td>
<td>0/237</td>
</tr>
<tr>
<td>6th nerve</td>
<td>14/550 (3)</td>
<td>6/243 (2)</td>
<td>1/237 (&lt;1)</td>
</tr>
<tr>
<td>7th nerve</td>
<td>8/550 (1)</td>
<td>4/243 (2)</td>
<td>2/237 (1)</td>
</tr>
<tr>
<td>8th nerve</td>
<td>78/550 (14)</td>
<td>53/243 (22)</td>
<td>18/237 (8)</td>
</tr>
<tr>
<td>Aphasia</td>
<td>11/550 (2)</td>
<td>8/243 (3)</td>
<td>1/237 (&lt;1)</td>
</tr>
<tr>
<td>Hemiparesis</td>
<td>24/550 (4)</td>
<td>18/243 (7)</td>
<td>2/237 (1)</td>
</tr>
<tr>
<td>Quadriparesis</td>
<td>6/550 (1)</td>
<td>4/243 (2)</td>
<td>2/237 (1)</td>
</tr>
</tbody>
</table>

* The study included 671 patients with a total of 696 episodes of community-acquired meningitis.
† Neurologic examination was performed at discharge in 550 of 553 surviving patients: 243 of 245 patients after pneumococcal meningitis, 237 of 238 patients after meningococcal meningitis, and 70 of 70 patients after other types of meningitis.
Dexamethasone reduced the rate of an unfavorable outcome from 25 percent to 15 percent (relative risk, 0.59; 95 percent confidence interval, 0.37 to 0.94; P=0.03). A post hoc analysis showed that the beneficial effect of dexamethasone on pneumococcal meningitis was attributable to a reduction in systemic rather than neurologic complications.

A systemic review showed that adjunctive corticosteroid therapy also reduced the frequency of neurologic sequelae among survivors.

The increased likelihood of an unfavorable outcome among those receiving corticosteroids in our cohort was probably due to the fact that corticosteroids were used more frequently in patients with clinical deterioration or those with factors associated with a poorer prognosis.

A low cerebrospinal fluid white-cell count was associated with an adverse outcome. This association has been described before. In patients with meningococcal meningitis, a low cerebrospinal fluid white-cell count may be found in those with sepsis and early meningitis. Studies of animals with pneumococcal meningitis showed a relation among a large bacterial cerebrospinal fluid load, lack of response of cerebrospinal fluid leukocytes, and intracranial complications. It probably indicates excessive bacterial growth and lack of a leukocyte response in the cerebrospinal fluid.

In our study, patients with pneumococcal meningitis were at risk for an unfavorable outcome, even after correction for other clinical predictors. Thus, for clinicians, knowledge of the causative organism of meningitis is important in predicting the risk of an unfavorable outcome. Gram’s staining of cerebrospinal fluid permits rapid and accurate identification of the causative bacteria in patients with bacterial meningitis and should therefore be routine.

Table 4. Multivariate Analysis of Factors Associated with an Unfavorable Outcome.*

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Favorable Outcome (N=459)</th>
<th>Unfavorable Outcome (N=237)</th>
<th>Odds Ratio (95% CI)‡</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age — yr</td>
<td>45±20</td>
<td>60±18</td>
<td>1.19 (1.06–1.35)</td>
<td>0.005‡</td>
</tr>
<tr>
<td>Duration of symptoms &lt;24 hr — no./no. evaluated (%)</td>
<td>222/440 (50)</td>
<td>95/221 (43)</td>
<td>0.70 (0.46–1.07)</td>
<td>0.10</td>
</tr>
<tr>
<td>Seizures — no./no. evaluated (%)</td>
<td>19/453 (4)</td>
<td>13/213 (6)</td>
<td>0.44 (0.17–1.12)</td>
<td>0.09</td>
</tr>
<tr>
<td>Pretreated with antibiotics — no./no. evaluated (%)</td>
<td>41/455 (9)</td>
<td>23/237 (10)</td>
<td>0.61 (0.31–1.21)</td>
<td>0.16</td>
</tr>
<tr>
<td>Coexisting conditions — no. (%)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Otitis or sinusitis</td>
<td>100 (22)</td>
<td>76 (32)</td>
<td>1.80 (1.13–2.84)</td>
<td>0.01</td>
</tr>
<tr>
<td>Pneumonia</td>
<td>37 (8)</td>
<td>46 (19)</td>
<td>1.76 (0.96–3.21)</td>
<td>0.07‡</td>
</tr>
<tr>
<td>Immunocompromise§</td>
<td>51 (11)</td>
<td>63 (27)</td>
<td>1.63 (0.95–2.79)</td>
<td>0.08</td>
</tr>
<tr>
<td>Symptoms at presentation</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Headache — no./no. evaluated (%)</td>
<td>396/434 (91)</td>
<td>148/192 (77)</td>
<td>1.34 (0.71–2.54)</td>
<td>0.37‡</td>
</tr>
<tr>
<td>Nausea — no./no. evaluated (%)</td>
<td>333/429 (78)</td>
<td>116/181 (64)</td>
<td>1.03 (0.61–1.75)</td>
<td>0.91</td>
</tr>
<tr>
<td>Neck stiffness — no./no. evaluated (%)</td>
<td>397/455 (87)</td>
<td>172/230 (75)</td>
<td>1.37 (0.70–2.70)</td>
<td>0.36‡</td>
</tr>
<tr>
<td>Rash — no./no. evaluated (%)</td>
<td>146/451 (32)</td>
<td>30/232 (13)</td>
<td>0.46 (0.25–0.86)</td>
<td>0.01‡</td>
</tr>
<tr>
<td>Heart rate — no./no. evaluated (%)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;60 beats/min</td>
<td>6/430 (1)</td>
<td>9/222 (4)</td>
<td>4.05 (0.95–17.24)</td>
<td>0.06‡</td>
</tr>
<tr>
<td>60–90 beats/min</td>
<td>166/430 (39)</td>
<td>49/222 (22)</td>
<td>1.00 —‡</td>
<td>—‡</td>
</tr>
<tr>
<td>&gt;90–120 beats/min</td>
<td>190/430 (44)</td>
<td>91/222 (41)</td>
<td>1.42 (0.85–2.36)</td>
<td>0.18</td>
</tr>
<tr>
<td>&gt;120 beats/min</td>
<td>65/430 (15)</td>
<td>76/222 (34)</td>
<td>2.67 (1.46–4.89)</td>
<td>0.002‡</td>
</tr>
<tr>
<td>Diastolic blood pressure &lt;60 mm Hg — no./no. evaluated (%)</td>
<td>39/443 (9)</td>
<td>22/227 (10)</td>
<td>1.99 (0.95–4.20)</td>
<td>0.07</td>
</tr>
<tr>
<td>Body temperature ≥38°C — no./no. evaluated (%)</td>
<td>333/448 (74)</td>
<td>189/230 (82)</td>
<td>1.26 (0.67–2.37)</td>
<td>0.48‡</td>
</tr>
<tr>
<td>Score on Glasgow Coma Scale¿</td>
<td>10±3</td>
<td>12±3</td>
<td>0.83 (0.76–0.90)</td>
<td>&lt;0.001‡</td>
</tr>
<tr>
<td>Triad of fever, neck stiffness, and change in mental status — no. (%)</td>
<td>186 (41)</td>
<td>119 (50)</td>
<td>0.58 (0.30–1.15)</td>
<td>0.12‡</td>
</tr>
<tr>
<td>Cerebral abnormality — no. (%)**</td>
<td>79 (17)</td>
<td>78 (33)</td>
<td>1.07 (0.56–2.04)</td>
<td>0.84‡</td>
</tr>
<tr>
<td>Cranial-nerve palsy — no. (%)</td>
<td>47 (10)</td>
<td>42 (18)</td>
<td>1.50 (0.81–2.80)</td>
<td>0.20‡</td>
</tr>
</tbody>
</table>
Several studies of prognostic factors in bacterial meningitis have been performed; however, all were retrospective and relatively small in size.\textsuperscript{4-6,10-22} Our study was nationwide and, therefore, we were able to study a representative sample of adults with acute bacterial meningitis. Furthermore, our prospective approach allowed us to collect comprehensive data on signs and symptoms, clinical course, and outcome. In addition, our large sample gave us the statistical power to perform multivariate analysis.

Our study has one important limitation: only patients who underwent lumbar puncture and who had a positive cerebrospinal fluid culture were included. Negative cerebrospinal fluid cultures occur in 11 to 30 percent of patients with bacterial meningitis.\textsuperscript{4-23} No significant differences in clinical presentation have been reported between patients with culture-positive bacterial meningitis and those with culture-negative bacterial meningitis.\textsuperscript{4,5,9} Therefore, it is unlikely that this factor confounded our results. Patients with space-occupying lesions on cranial CT do not undergo lumbar puncture,\textsuperscript{32,33} and patients with meningitis and a florid rash or septic shock also may not undergo lumbar puncture initially. Thus, an unknown number of such patients were excluded from the cohort, which may have resulted in an underestimation of the mortality rate, especially among patients with meningococcal meningitis.

Rates of antibiotic resistance among meningococcal and pneumococcal isolates were very low. Similar rates have been found in other studies in the Netherlands.\textsuperscript{26,38} In the United States, France, Spain, and other countries, antibiotic-resistant \textit{S. pneumoniae} strains are highly prevalent and have emerged as a major problem in the treatment of patients with bacterial meningitis.\textsuperscript{39}

\begin{table}
\centering
\caption{Clinical features and prognostic factors of bacterial meningitis}  
\begin{tabular}{|l|l|l|l|l|}
\hline
\textbf{Characteristic} & \textbf{Favorable Outcome (N=459)} & \textbf{Unfavorable Outcome (N=237)} & \textbf{Odds Ratio (95\% CI)} & \textbf{P Value} \\
\hline
\textbf{Indexes of inflammation in the CSF} & \textbf{White-cell count — no./no. evaluated (\%)} & & & \\
<100/mm\textsuperscript{3} & 23/428 (5) & 39/217 (18) & 3.43 (1.64–7.20) & 0.001‡ \\
100–999/mm\textsuperscript{3} & 56/428 (13) & 66/217 (30) & 2.82 (1.59–4.78) & <0.001‡ \\
1000–10,000/mm\textsuperscript{3} & 238/428 (56) & 87/217 (40) & 1.00 — & — \\
>10,000/mm\textsuperscript{3} & 111/428 (26) & 25/217 (12) & 0.55 (0.30–1.01) & 0.05‡ \\
\textbf{Protein — g/liter} & 4.8±4.7 & 5.4±3.9 & 1.03 (0.99–1.07) & 0.17‡ \\
\textbf{CSF:blood glucose ratio — mg/dl} & 0.18±0.2 & 0.15±0.2 & 0.91 (0.70–1.17) & 0.44‡ \\
\textbf{Positive blood culture — no./no. evaluated (\%)} & 238/403 (59) & 166/208 (80) & 2.24 (1.24–4.03) & 0.009‡ \\
\textbf{Blood chemical tests} & & & & \\
\textbf{ESR — mm/hr} & 42±37 & 56±37 & 1.20 (1.03–1.40) & 0.02‡ \\
\textbf{Thrombocyte count — platelets/mm\textsuperscript{3}} & 208,000±100,000 & 180,000±97,000 & 0.92 (0.88–0.97) & 0.003 \\
\hline
\end{tabular}
\end{table}
to this epidemiologic trend, treatment recommenda-
dations for suspected and confirmed cases of bac-
terial meningitis continue to evolve.40 In areas with
high rates of pneumococcal that are resistant to
broad-spectrum cephalosporins, a combination of
a third-generation cephalosporin and vancomycin
should be the initial choice for patients with acute
bacterial meningitis.40 Treatment failures in pa-
tients with meningitis due to multidrug-resistant
bacterial isolates have been described.9,41,42 The
outcome was not significantly influenced by the
presence of antibiotic-resistant pneumococcal iso-
lates in several studies20,23,41,42; however, the re-
lationship of antibiotic resistance to the outcome
of meningitis remains to be elucidated.

Despite the fact that the case fatality rate is de-
creased by the use of adjunctive dexamethasone,
there is still room for improvement. Prompt use
of dexamethasone and appropriate antibiotics,
together with optimal supportive care, can further
reduce the mortality and morbidity associated with
bacterial meningitis. Most risk factors for an un-
favorable outcome that we identified indicated the
presence of systemic compromise, a low level of
consciousness, and infection with S. pneumoniae.
Therefore, aggressive supportive care of patients
with bacterial meningitis and systemic complica-
tions, preferably in specialized care units, is needed.

Supported in part by a grant from Roche Pharmaceuticals.

We are indebted to Professor P. McIntyre for his comments on the
manuscript and to many physicians in the Netherlands for their co-
operation.

REFERENCES
after the introduction of protein-polysac-
348:1737-46.
3. Netherlands Reference Laboratory for
Bacterial Meningitis. Bacterial meningitis in
The Netherlands: annual report 2002. Am-
sterdam: University of Amsterdam, 2003.
328:21-8.
5. Sigurdardottiri B, Bjornsson OM, Jones-
dottiri KE, Eraldsdottiri H, Gudmundsdott
S. Acute bacterial meningitis in adults:
a 20-year overview. Arch Intern Med 1997;
6. Hussein AS, Shafsan SD. Acute bacterial
meningitis in adults: a 12-year review. Medi-
cine (Baltimore) 2000;79:360-8.
8. Aronin SI, Peduzzi P, Quagliarello VJ. Community-acquired bacterial meningitis:
risk stratification for adverse clinical out-
come and effect of antibiotic timing. Ann
9. Tunel AR. Bacterial meningitis. Phila-
10. Olsson RA, Kirby JC, Romansky ML.
Pneumococcal meningitis in the adult: clin-
ical, therapeutic, and prognostic aspects in
11. Dodge PR, Swartz MN. Bacterial menin-
gitis — a review of selected aspects. II. Spe-
cial neurologic problems, postmeningitis
complications, and clinicopathological cor-
12. Weiss W, Figueroa W, Shapiro WH, Flipp-
pin HE. Prognostic factors in pneumococcal
meningitis. Arch Intern Med 1967;120:517-
24.
13. Magnusson CR. Meningitis in adults:
ten-year retrospective analysis at communi-
prognostic factors in 164 cases based on
mortality and on a study of lasting sequelae.
15. Brunn GW, Kremer HPH, de Marie S,
Padberg GW, Hermans J, van Furth R. Clin-
ical evaluation of pneumococcal meningitis
in adults over a twelve-year period. Eur J Clin
Microbiol Infect Dis 1989;8:695-700.
Canton P. Mortality in pneumococcal men-
ingitis: a multivariate analysis of prognostic
17. Kirkpatrick B, Reeves DS, MacGowan
AR. A review of the clinical presentation, lab-
oratory features, antimicrobial therapy and
outcome of 77 episodes of pneumococcal
meningitis occurring in children and adults.
18. Antal J, Hatala R, Cook DJ, Wong KG. The
rational clinical examination: does this adult
patient have acute meningitis? JAMA 1999;
282:175-81.
19. Kragsbjerg P, Kallman J, Olesen P. Pneu-
ococcal meningitis in adults. Scand J Infect
Pneumococcal meningitis in the intensive
care unit: prognostic factors of clinical out-
come in a series of 80 cases. Am J Respir
Community-acquired bacterial meningitis
in adults: the epidemiology, timing of appro-
priate antimicrobial therapy, and prognostic
factors. Clin Neurol Neurosurg 2002;104:
352-8.
22. Kastenbauer S, Pfister HW. Pneumococ-
cal meningitis in adults: spectrum of com-
lications and prognostic factors in a series
of 87 cases. Brain 2003;126:1015-25.
23. Flores-Cordero JM, Amaya-Villar R,
Rincon-Ferrari MD, et al. Acute communi-
ty-acquired bacterial meningitis in adults
admitted to the intensive care unit: clinical
manifestations, management and prognostic
24. Statistics Netherlands. StatLine, Voor-
burg/Heerlen, 2001. (Accessed October 4,
2004, at http://www.cbs.nl.)
25. Jennett B, Teasdale G. Management of
head injuries. 2nd ed. Philadelphia: F.A.
Davis, 1981.
26. van de Beek D, de Gans J, Spanjaard L,
Vermeulen M, Dankert J. Antibiotic guide-
lines and antibiotic use in adult bacterial
meningitis in The Netherlands. J Antimicrob
27. de Gans J, van de Beek D. Dexametha-
sone in adults with bacterial meningitis.
28. Idem. Dexamethasone in adults with
348:955-6.
29. Little RJ, Rubin D. Statistical analysis
30. Arnold AM, Kronmal RA. Multiple im-
putation of baseline data in the Cardiovasc-
ular Health Study. Am J Epidemiol 2003;
157:74-84.
31. Spanos A, Harrell FE Jr, Durack DT.
Differential diagnosis of acute meningitis: an
analysis of the predictive value of initial ob-
Clinical Features and Prognostic Factors of Bacterial Meningitis


Copyright © 2004 Massachusetts Medical Society.

JOURNAL EDITORIAL FELLOW

The Journal’s editorial office invites applications for a one-year research fellowship beginning in July 2005 from individuals at any stage of training. The editorial fellow will work on Journal projects and will participate in the day-to-day editorial activities of the Journal but is expected in addition to have his or her own independent projects. Please send curriculum vitae and research interests to the Editor-in-Chief, 10 Shattuck St., Boston, MA 02115 (fax, 617-739-9864), by January 15, 2005.

Copyright © 2004 Massachusetts Medical Society.
CORRECTION

Clinical Features and Prognostic Factors in Adults with Bacterial Meningitis

Clinical Features and Prognostic Factors in Adults with Bacterial Meningitis. On page 1853, in the right-hand column, lines 5 through 8 of the second full paragraph should have read, `N. meningitidis was responsible for 37 percent of the episodes, with group B identified in 174 episodes, group C in 79, group Y in 3, and group W135 in 1,` rather than, `with group B identified in 173 episodes, group C in 79, group Y in 3, group H in 1, and group W135 in 1,` as printed.