Cerebrospinal fluid findings in adults with acute Lyme neuroborreliosis

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Abstract Presence of BB-specific antibodies in the cerebrospinal fluid (CSF) with evidence of their intrathecal production in conjunction with the white cell count in the CSF and typical clinical symptoms is the traditional diagnostic gold standard of Lyme neuroborreliosis (LNB). Few data are available on the CSF lactate concentration in European adults with the diagnosis of acute LNB. The objective of the study was to investigate the CSF changes during acute LNB. Routine CSF parameters [leukocyte count, protein, lactate and albumin concentrations, CSF/serum quotients of albumin (Q Alb), IgG, IgA and IgM, and oligoclonal IgG bands] and the Borrelia burgdorferi (BB)-specific antibody index were retrospectively studied in relation to the clinical presentation in patients diagnosed with acute LNB. A total of 118 patients with LNB were categorized into the following groups according to their symptoms at presentation: group 1: polyradiculoneuritis (Bannwarth’s syndrome), group 2: isolated facial palsy and group 3: predominantly meningitic course of the disease. In addition to the CSF of patients with acute LNB, CSF of 19 patients with viral meningitis (VM) and 3 with neurolues (NL) were analyzed. There were 97 patients classified with definite LNB, and 21 as probable LNB. Neck stiffness and fever were reported by 15.3% of patients. Most of these patients were younger than 50 years. Polyradiculoneuritis was frequently found in patients older than 50 years. Lympholecytosis was found in all patients. Only 5 patients had a CSF lactate \( \geq 3.5 \text{ mmol/l} \), and the mean CSF lactate level was not elevated \( 2.1 \pm 0.6 \text{ mmol/l} \). The patients with definite LNB had significantly higher lactate levels than patients with probable LNB. Elevated lactate levels were accompanied by fever and headache. In the Reiber nomograms, intrathecal immunoglobulin synthesis was found for IgM in 70.2% followed by IgG in 19.5%. Isoelectric focussing detected an intrathecal IgG synthesis in 83 patients (70.3%). Elevated BB AIs in the CSF were found in 97 patients (82.2%). Patients with VM showed lower CSF protein concentration and CSF/serum quotients of albumin than LNB patients. In acute LNB, all patients had elevated cerebrospinal fluid (CSF) leukocyte counts. In contrast to infections by other bacteria, CSF lactate was lower than 3.5 mmol/l in all but 5 patients. The CSF findings did not differ between polyradiculoneuritis, facial palsy, and meningitis. The CSF in LNB patients strongly differed from CSF in VM patients with respect to protein concentration and the CSF/serum albumin quotient.

Keywords Acute lyme neuroborreliosis · CSF findings · Clinical symptoms
Introduction

Lyme neuroborreliosis (LNB) is the neurological manifestation of systemic infection caused by the spirochete *Borrelia burgdorferi* (*BB*) sensu lato. Clinical features of LNB are diverse and differ in European and American patients—most probably because of variations in the *Borrelia* species [1, 2].

The diagnosis of LNB is based on a combination of history, clinical examination, analysis of cerebrospinal fluid (CSF), and the detection of antibodies against *BB* in serum and CSF.

According to the criteria of the German Society of Neurology (DGN) LNB is probable when typical clinical symptoms, pleocytosis in the CSF, and positive serum tests for *Borrelia*-specific IgG or IgM are present [3]. The detection of intrathecal synthesis of specific anti-*Borrelia* antibodies is considered necessary to diagnose definite LNB in Germany and other European countries [1, 3, 4], whereas American criteria do not require an elevated anti-*Borrelia* antibody index (AI) [5, 6].

Laboratory confirmation of LNB is hampered by the low sensitivity of culture and polymerase chain reaction (PCR) in CSF [7]. Presence of *BB*-specific antibodies in the CSF with evidence of intrathecal production is the traditional diagnostic gold standard but has limitations such as low sensitivity in the very early phase of the disease [1, 2, 4] and can persist for years after eradication of the infection [8, 9]. In recent years, the B lymphocyte chemotactant chemokine CLCX13 has been identified as a potentially important biomarker for the diagnosis of acute LNB [10, 11]. Reported sensitivities and specificities are high (in a similar range as for AIs), suggesting that CSF CLCX13 might be a useful addition to the diagnostic armamentarium, especially in the very early stage of LNB, where AIs might still be negative. However, as patient numbers in the published studies are rather low, further corroboration of these results is needed. Several other, more or less validated laboratory tests have been developed to improve diagnosis. Few data are available on the CSF lactate levels in adults with definite acute LNB. The aim of this study was to evaluate CSF changes in patients with acute LNB.

Methods

Patients

All files of patients who were coded as “probable” or “definite” acute LNB upon admission to the Neurological Department of the University Medical Center Goettingen during the past 14 years were screened. The diagnosis of acute LNB was made if in addition to the typical neurological symptoms and CSF lymphopleocytosis, either IgM or IgG antibodies for *BB* were identified in serum and/or CSF. Only patients ≥16 years of age who had received a lumbar puncture and with a sufficient clinical documentation were eligible for this investigation. Exclusion criteria were other putative reasons for inflammation in the central nervous system (CNS) at presentation. Files were screened for a study on long-term outcome of patients after acute NB.

To compare the CSF findings of patients with LNB to CSF findings of patients with CSF pleocytosis of other origins, files of patients with viral meningitis (VM) (*n* = 19) and neurolues (NL) were also screened. Moreover, all files of patients with NL without concomitant HIV infection from the last 10 years were included.

The study was approved by the local ethics committee.

Clinical examination

For clinical symptoms, we applied the Scripps Neurological Rating Scale (SNRS) [12], which met our needs for thorough documentation and reproducibility. In addition, age, fever, number of days from the onset of symptoms to hospitalization, information on tick bites, and erythema migrans (EM) were registered.

LNB patients were categorized into the following groups according to their symptoms at presentation: group 1—polyradiculoneuritis (Bannwarth’s syndrome), group 2—isolated facial palsy, group 3—predominantly meningitic course of the disease.

Laboratory tests

IgM and IgG antibody response to antigens of *BB* was determined in serum by sandwich enzyme-linked immunosorbent assays (ELISA) according to the instructions of the manufacturer (Genzyme Virotech, Rüsselsheim, Germany). The IgM ELISA was based on antigens of *Borrelia afzelii* (analytical sensitivity 99%, specificity 98.8%, manufacturer’s information). The antigens of the IgG ELISA were *Borrelia afzelii* lysate plus VlsE (analytical sensitivity 99%, specificity 97%, manufacturer’s information). In patients with a positive ELISA response, Western blots (Borrelia line, Genzyme Virotech, Rüsselsheim, Germany) were performed to confirm the specificity of the antibodies [13]. The antigens investigated were OspC, VlsE, BmpA, and DbpA for the detection of specific antibodies (analytical sensitivity 99%, specificity 98%, manufacturer’s information).

The CSF was examined for leukocyte count (pleocytosis defined as >4 leukocytes/μl), levels of protein, albumin, and lactate (pathological ≥2.1 mmol/l). Intrathecal
concentrations of IgG, IgA, and IgM were measured immunonephelometrically, and the CSF/serum ratios of IgG, IgA, IgM, and albumin were calculated. IgG, IgA, and IgM ratios were plotted versus albumin ratios according to the nomogram of Reiber and Felgenhauer [14].

The presence of an intrathecal production of BB-specific antibodies was evaluated by calculation of the BB-specific antibody index (AI). For this purpose, a modified immunonassay was used, based on the commercially available immunonassay plates Enzygnost Borreliosis IgG/IgM of Siemens Healthcare Diagnostics GmbH, Marburg, Germany. To increase the sensitivity, the secondary antibodies supplied with the test kit were replaced by anti-IgG or anti-IgM antibodies from DAKO, Hamburg, Germany.

Arbitrary concentration units were calculated based on a standard curve derived from seven 2-fold serial dilutions of a positive standard serum pool to fit an absorbance range of 0.05–2.0. The highest standard concentration (absorption of approximately 2.0) was defined as 100 arbitrary units. CSF and serum pairs of every patient had to be analyzed on the same plate, and the CSF and serum sample were diluted to yield antibody concentrations that were in similar parts of the standard curve. Therefore, standard predilution ratios were 1:30 for CSF and 1:6000 for serum, in case of an intrathecal immunoglobulin production modified predilution ratios were used. Only those extinctions were used for AI calculation that lay more than 0.1 above the blank of approximately 2.0. The mean age of all patients was 50±17.1 years and the gender ratio was 40/78 (F:M).

In the cohort, 15.3% of patients presented with headache and fever. Sixteen of these patients were younger than 50 years (34.6±11.2 years). Patients with headache and fever displayed more often neck stiffness (n=18) than patients with polyradiculitis (5 out of 53) (p=0.015). Polyradiculitis was frequently found in patients older than 50 years (age 54.1±15.1 years). The frequency of facial palsy was not significantly different in patients younger (n=28) and older than 50 years (n=19), respectively, (p=0.4). One patient presented with stroke, one with paresis of N. abducens, one with an epileptic seizure, and two with vertigo.

Clinical symptoms and CSF parameters of the different groups are summarized in Table 1 and Fig. 1.

Only 5 patients with LNB showed CSF lactate concentrations ≥3.5 mmol/l, and the mean CSF lactate concentration was not elevated (2.1±0.6 mmol/l). The mean CSF lactate levels of patients with definite LNB (2.3±0.7 mmol/l) were slightly but significantly higher (p=0.002) than the mean CSF lactate levels of patients with probable LNB (1.8±0.3 mmol/l). Patients with elevated lactate levels (≥2.1 mmol/l) ([group 1—32.1%; group 2—46.8%; group 3—55.6%], n=49) presented significantly more often with fever and headache (n=43; Fisher’s exact test p=0.02).

Elevated BB AIs in CSF were found in 97 patients (82.2%). BB IgM AIs were elevated in 72 patients and BB IgG AIs in 82 patients. Of the 23 patients pre-treated with antibiotics, 15 patients had elevated BB AIs (65.2%) in CSF. Symptoms developed in 107 patients within 6 weeks. The BB AIs of 21 patients were <1.5 (17.8%). Eleven
patients reported symptoms for 6 weeks or longer prior to lumbar puncture. Of these patients, 10 had elevated BB AIs (90.9%).

Intrathecal immunoglobulin synthesis as estimated by Reiber–Felgenhauer nomogram was found for IgM in 70.2%, followed by IgG in 19.5%. Intrathecal synthesis of IgM, IgG, and IgA was found in 11 patients.

Isoelectric focussing, more sensitive than the Reiber–Felgenhauer plots, detected oligoclonal IgG bands indicating IgG synthesis in the CSF in 83 patients (70.3%). The positive oligoclonal IgG bands were significantly more frequent in patients with polyradiculoneuritis and facial palsy than in patients with a meningitic course of acute LNB. This may be a sign of stronger immune responses in these patients. The differences of BB-specific AIs and oligoclonal bands between the groups are summarized in Table 2.

Neurolues (NL) was diagnosed during the last 10 years in only 6 patients in our department. Three of them had concomitant HIV encephalitis. The CSF samples of the remaining 3 patients with isolated neurolues were analyzed. All 3 patients were male (100%). The mean age of patients with NL was 46.2 ± 13.8 years.

Fig. 1  Symptoms in patients younger versus patients older than 50 years. Polyradiculoneuritis was significantly more frequent in patients older than 50 years. In contrast, a meningitic course of the disease was more common in patients under 50 years

Table 1  Comparison of laboratory parameters between patients with polyradiculoneuritis (group 1), facial palsy (group 2), and patients with a meningitic course of LNB (group 3)

| Parameter                  | Polyradiculoneuritis (n = 53) | Facial palsy (n = 47) | Meningitic course (n = 18) | P      |
|----------------------------|--------------------------------|-----------------------|---------------------------|--------|
| CSF leukocytes [μl⁻¹]      | 171.5 (54.5; 354)              | 210.5 (90.3; 411.8)   | 345.5 (100.8; 543.3)      | 0.12   |
| CSF protein [mg/l]         | 1,339 (779; 2,087)             | 1,372 (887; 2,003)    | 942 (553.0; 1,543)        | 0.16   |
| QAlbumin × 10⁻³            | 19.8 (9.8; 31.7)               | 21.0 (12.0; 28.4)     | 12.6 (8.1; 23.3)          | 0.23   |
| CSF lactate [mmol/l]       | 2.0 (1.7; 2.6)                 | 2.0 (1.7; 2.6)        | 2.4 (1.8; 2.9)            | 0.56   |

Descriptive values are median and interquartile range (Q25 and Q75). After using the D’Agostino–Pearson normality test, the Kruskal–Wallis test was used for statistical comparisons of not normally distributed data (cell count, protein level, and QAlbumin). ANOVA was used for statistical analysis of normally distributed data (lactate concentration). Between the groups, we could not find any significant differences.

Table 2  Numbers of patients with positive oligoclonal IgG bands in CSF and BB-specific antibody index (AI) in 118 patients diagnosed with acute LNB

| Parameter                  | Polyradiculoneuritis (n = 53) | Facial palsy (n = 47) | Meningitic course (n = 18) | P      |
|----------------------------|--------------------------------|-----------------------|---------------------------|--------|
| Positive oligoclonal IgG bands | 40                            | 35                    | 8                         | 0.03*  |
| Positive BB AI IgG         | 45                            | 37                    | 11                        | 0.10   |
| Positive BB AI IgM         | 39                            | 34                    | 9                         | 0.15   |

Group 1 represents patients with polyradiculoneuritis, group 2 patients with facial palsy and group 3 patients with a meningitic course of acute LNB. Positive oligoclonal IgG bands were significantly more frequent in patients with polyradiculoneuritis and facial palsy (Chi square test).
Table 3  CSF finding in patients with acute lyme neuroborreliosis (LNB), viral meningitis (VM) and neurotuberculosis (NL)

| Parameter                  | LNB (n = 118) age (50 ± 17.1) | VM (n = 19) age (38.3 ± 17.3) | NL (n = 3) age (46.2 ± 13.8) |
|----------------------------|-------------------------------|-------------------------------|-------------------------------|
| CSF leukocytes [μl⁻¹]      | 170.5 (57.0; 369)             | 97.0 (21.0; 210.0)            | 65 (7.0; 100.7)              |
| CSF protein [mg/l]         | 1.232 (697; 1.926)            | 628 (493.0; 969)*             | 614 (328.0; 800.0)           |
| QAlbumin × 10³             | 17.2 (9.7; 28.4)              | 9.1 (7.2; 14.7)*              | 7.6 (4.0; 11.7)              |
| CSF lactate [mmol/l]       | 2.0 (1.6; 2.6)                | 2.1 (1.9; 2.3)                | 1.5 (1.3; 1.6)               |

Descriptive values are median and interquartile range (Q25 and Q75)

*p ≤ 0.05 LNB versus VM (Mann–Whitney U test)

patients. Of these 12 patients, 9 had positive BB AI IgM, BB AI IgG, or both in CSF.

In the remaining 23 patients, blood samples were not tested for antibodies against BB in a separate quantitative test, as definite evidence for acute LNB had been established previously by typical clinical signs and positive AI against BB for IgG, IgM, or both in CSF and serum samples measured simultaneously in the same assay.

Serum CRP was measured in 81 patients, and elevated levels were found in 11. WBC was measured in all patients; 14 patients showed leukocytosis (≥11,000/μl). Between the different clinical presentations, we could not find significant differences for these laboratory parameters.

Discussion

European LNB is associated with elevated cell count in the CSF, typically 10–1,000 leukocytes/mm³ (60% between 30 and 300 cells/mm³), mainly B lymphocytes and plasma cells. Together with the intrathecal immunoglobulin production with intrathecal IgM dominance and blood–CSF barrier dysfunction, these typical CSF changes allow the diagnosis of acute LNB with a specificity of 96% and sensitivity of 70% [17]. Published data compare well with our findings of intrathecal IgM synthesis in 70.2% of the patients and lymphocytosis in all patients. The diagnostic sensitivity of BB-specific antibody production in CSF is about 80% [18]. In our study, the sensitivity of the elevated BB-specific antibody index was 82.2%.

A positive AI may persist for years after successful therapy [8, 9]. In some cases, the AIs were ≥1.5 while in serum BB-specific antibodies were below the cut-off of the ELISA [19]. This phenomenon was found in 9 patients in our study.

After effective treatment of LNB, BB-specific antibodies concentrations frequently decrease more rapidly in serum than in the CSF. This can lead to an increased AI during recovery.

BB AI negativity in CSF in patients with early LNB has been documented in series of patients with LNB confirmed by culture, PCR, or BB antigen demonstration [20, 21]. The frequency of 18.7% BB AI negativity in our patients with symptoms lasting <6 weeks is in accordance with frequencies of 10–20% early BB AI negativity reported from other European centers [15, 17, 22]. These findings confirm that BB AIs in the acute phase of LNB have only moderate sensitivity.

Early antibiotic therapy can abort the humoral immune response resulting in undetectable or normal BB AI or may impede the immune switch from IgM to IgG in plasma [23, 24]. On the other hand, one study showed an increased percentage of B cells in CSF and of intrathecally produced BB antibodies immediately after initiation of antibiotic treatment [25]. Our data demonstrate the necessity to determine the AIs both for BB IgM and IgG.

Patients with LNB with normal leukocyte counts have been described exceptionally. These patients were either in the very early stage of the disease or were immunosuppressed individuals or were children with an isolated facial palsy [19]. In our study, all patients had CSF lymphocytosis, and we have not seen such a case in adults during our clinical practise.

Studies which investigated the role of CSF lactate levels for diagnosis of LNB are very rare. For other types of bacterial meningitis, such as pneumococcal or meningococcal meningitis, a CSF lactate level ≥3.5 mmol/l is typical and found in 90% of patients during the acute stage [17]. In our study, a normal mean CSF lactate level was found for all patients. However, patients with definite LNB showed slightly elevated CSF lactate concentrations compared to the patients with probable LNB. All 5 LNB patients with CSF lactate levels ≥3.5 mmol/l had a meningitic course of the diseases. Elevated lactate levels were significantly more often accompanied by fever and headache. With respect to the CSF abnormalities, the most difficult differential diagnosis of LNB is tuberculous meningitis. The typical CSF changes during tuberculous meningitis are lymphocytosis (80% between 30 and 300 cells/mm³), severe blood–CSF barrier dysfunction and predominant intrathecal IgA synthesis. In contrast to LNB, patients with neurotuberculosis in general show CSF lactate levels ≥3.5 mmol/l [17].
The most common neurological manifestation of LNB in Europe in adults is polyradiculoneuritis (Bannwarth’s syndrome) [26]. In our population polyradiculoneuritis was also the predominant syndrome which could be found significantly more frequently in patients older than 50 years. During LNB headache occurs in about 40%, but headache as the prominent symptom without (pseudo) radicular pain or paresis, i.e., a meningitic syndrome, is rare in adults [27]. In our study, isolated headache and neck stiffness was observed in 15.2% of patients younger and in 5 patients older than 50 years. These were the patients in whom fever and elevated lactate concentrations were also measured. In another study, headache and stiff neck was found in 13% of patients but differences in clinical manifestation and age of patients were not considered [28]. To our knowledge, this is the first study investigating the influence of age on clinical manifestation in an adult patient cohort with definite and probable LNB.

In conclusion, acute LNB is a clinical diagnosis supported by characteristic CSF abnormalities, in particular elevated CSF leucocyte counts, blood–CSF barrier dysfunction, predominant intrathecal IgM synthesis and elevated BB-specific AI for IgM or IgG. In contrast to other forms of bacterial meningitis, the mean CSF lactate level in our study was not elevated. Lactate can provide rapid and reliable information, in particular, with respect to the differential diagnosis tuberculosis meningitis. Therefore, we recommend routine measurement of CSF lactate in adults with suspected acute LNB. Lactate is unsuitable for the differentiation of LNB and VM; CSF protein content and the CSF/serum albumin quotient are more appropriate.

Conflict of interest None.

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