Lyme neuroborreliosis in cases of non-specific neurological symptoms

BACKGROUND Analysis of cerebrospinal fluid is required in order to diagnose Lyme neuroborreliosis. We investigated the symptoms of patients in a highly endemic area who were referred for evaluation of possible Lyme neuroborreliosis, and explored whether cerebrospinal fluid analysis confirmed or ruled out the diagnosis.

METHOD We reviewed the medical records of all patients who underwent lumbar puncture at Sørlandet Hospital Arendal in the period 1 January 2013 to 31 December 2013.

RESULTS A total of 140 patients were referred with suspected Lyme neuroborreliosis. Of these, 110 patients had non-specific neurological symptoms (e.g. fatigue, dizziness and headache), only one of whom received a diagnosis of *possible* Lyme neuroborreliosis. Thirty patients had symptoms typical of the condition (such as radiculitis or peripheral facial nerve palsy). Six of these were diagnosed with *definite* Lyme neuroborreliosis, and one with *possible* Lyme neuroborreliosis. None of those diagnosed with Lyme neuroborreliosis had had symptoms lasting more than six months.

INTERPRETATION The probability of Lyme neuroborreliosis is low in the absence of typical symptoms of the condition, even when anti-*Borrelia* antibodies are detected in serum and especially when the symptoms are of long duration.

Lyme disease is a vector-borne zoonosis that is caused by infection with the spirochete *Borrelia burgdorferi* following a tick bite (1). Early localised disease manifests as erythema migrans, and approximately half of those infected have non-specific symptoms such as fever, myalgia, lymphadenopathy and headache (2).

Lyme neuroborreliosis is the most widespread form of disseminated borreliosis in Europe. The condition is reported in 3-12%of those with untreated early localised disease, usually four to eight weeks after a tick bite, but both earlier and later onset occur (3–7). The most common clinical manifestation of Lyme neuroborreliosis is Bannwarth's syndrome, which consists of peripheral facial nerve palsy or oculomotor nerve palsy, radiculitis with symptoms including radiating pain, changes in sensation and/or paralysis, and the presence of lymphocytes in the cerebrospinal fluid (7, 8). In children, typical presentation is facial nerve palsy with low-grade meningitis symptoms without radiculitis (9). In rarer cases of Lyme neuroborreliosis, there may be peripheral neuropathy, cognitive impairment, cerebellar ataxia, encephalitis and myelitis (1, 7).

In Norway, Lyme neuroborreliosis is diagnosed on the basis of European guidelines. These require neurological symptoms consistent with the disease, as well as lumbar puncture showing pleocytosis and a positive antibody index as evidence of intrathecal anti-*Borrelia* antibody production (Box 1) (10). In 10–30% of those with symptom

duration of less than six weeks, *Borrelia* antibodies are not detected in the cerebrospinal fluid (CSF) or in serum, which makes early diagnosis challenging (11).

Between 28 and 50% of those who undergo treatment for Lyme neuroborreliosis develop post-treatment Lyme disease (PTLD), which is marked by fatigue, neuropsychological symptoms and reduced quality of life (12, 13). Diagnostic criteria have been proposed for this condition, but its aetiology, diagnosis and clinical significance remain uncertain and controversial (14–16).

In 2015, a total of 425 cases of disseminated borreliosis, neuroborreliosis, Lyme carditis and Lyme arthritis were recorded in Norway (17). The coastal regions of the Agder counties are considered highly endemic for *Borrelia*, with up to 31.1% of ticks being carriers of *Borrelia* species (18).

At Sørlandet Hospital the number of referrals for possible Lyme neuroborreliosis has increased in recent years. We have investigated the symptoms shown by these patients and the duration of those symptoms, the results of CSF analysis, and the final diagnosis.

Material and method

Patients

The medical records of all outpatients and inpatients who underwent lumbar puncture in the Department of Neurology, Sørlandet Hospital Arendal in the period 1 January 2013 to 31 December 2013 were reviewed. The following information was recorded

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MAIN POINTS

There was a low probability of Lyme neuroborreliosis in cases of prolonged non-specific neurological symptoms, including when anti-Borrelia antibodies were detected in serum

Lyme neuroborreliosis should be suspected in patients with painful radiculitis that worsens at night, especially when accompanied by numbness in a dermatomal distribution and peripheral facial nerve palsy

Patients with symptoms of Lyme neuroborreliosis should be assessed with lumbar puncture

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BOX 1

European guidelines for the diagnosis of Lyme neuroborreliosis (10)

Criteria

Unexplained neurological symptoms consistent with Lyme neuroborreliosis

Cerebrospinal fluid pleocytosis

Positive antibody index indicative of intrathecal anti-Borrelia antibody production

Interpretation

3 of 3 criteria: Definite Lyme neuroborreliosis 2 of 3 criteria: Possible Lyme neuroborreliosis 1 of 3 criteria: Negative for Lyme neuroborreliosis

from referral letters, outpatient notes, laboratory results and discharge summaries: reason for referral, patient's account of the tick bite, symptoms and symptom duration, results of CSF analysis, final diagnosis, treatment and complications.

Patients were classified into two groups. The first group had presented with symptoms typical of Lyme borreliosis, including radiculitis, peripheral facial nerve palsy and other symptoms suggesting involvement of the central nervous system. The second group had presented with non-specific symptoms, such as tiredness/fatigue, dizziness, difficulties with concentration, myalgias, non-specific paraesthesias and tension-type headache. The classification was performed by the lead author.

Dataset

In total, 240 lumbar punctures were performed over the specified period, of which 140 (58%) were in connection with testing for possible Lyme neuroborreliosis. These 140 patients do not include patients whose CSF was tested for *Borrelia* antibodies as part of the workup for other diseases, such as dementia, multiple sclerosis and chronic

fatigue syndrome (CFS/ME). A total of 100 patients were excluded.

Serological testing for *Borrelia* antibodies was performed, and all CSF samples were sent for analysis for *Borrelia* antibodies and the antibody index to check for intrathecal antibody production, and for analysis of cells and proteins in the CSF. All analyses were conducted at the Department of Medical Microbiology at Sørlandet Hospital, which is the reference laboratory for *Borrelia* analyses in Norway.

Serum and CSF testing were conducted using Siemens Enzygnost Lyme link VlsE/IgG and Siemens Enzygnost Borreliosis IgM. The IgG index is calculated in accordance with the Rieber method (11). Borrelia IgG in serum is given as a percentage of the cutoff, where the cutoff is the lowest possible positive value. This value varies and is calculated for each individual setup. With low values, e.g. 200% of cutoff, there is a greater chance that the antibodies detected are not specifically directed against Borrelia than there is with higher values, such as 800% of cutoff (19). Data were analysed using IBM SPSS 22.0.

Ethical approval

The study was presented to the Regional Committee for Medical and Health Research Ethics and was declared exempt from the requirement for formal approval. The study was approved by the Research Director at Sørlandet Hospital.

Results

Of the 140 patients, six received a diagnosis of definite Lyme neuroborreliosis and two of possible Lyme neuroborreliosis. The average age of this group was 48 years, and 53 % were women. There were no significant differences in age or sex with respect to symptoms and final diagnosis.

Of the 30 patients with typical symptoms, six were diagnosed with definite Lyme neuroborreliosis and one with possible Lyme neuroborreliosis. Of the 110 patients with non-specific neurological symptoms, only one was considered to have possible Lyme neuroborreliosis (Table 1).

Of the patients who did not receive a diagnosis of Lyme neuroborreliosis, 52 % had myalgias, 43 % tiredness/fatigue and 42 % non-specific paraesthesias. Eight per cent had been bitten by ticks within the last three months and 5 % had had a rash suspected of being erythema migrans. In this group, 45 % had had symptoms for less than three months and 30% for more than 12 months (Table 2). Of the eight who received a diagnosis of definite or possible Lyme neuroborreliosis, 63 % had peripheral facial nerve palsy, and 50% had radiculitis and paraesthesias in a dermatomal distribution. None reported a history of erythema migrans. Two of the eight described tick bites within the last three months.

In all, 24% of patients who were referred for evaluation of possible Lyme neuroborreliosis on the basis of new-onset peripheral facial nerve palsy received a diagnosis of definite or possible disease. All six patients who were diagnosed with definite Lyme neuroborreliosis tested positive for *Borrelia* IgG antibodies in serum, with values exceeding 650% of cutoff.

Discussion

Our study revealed that 79 % of those referred for lumbar puncture upon suspicion of Lyme neuroborreliosis did not have typical symptoms of the disease – they were referred because of non-specific neurological symptoms such as prolonged tiredness, non-specific paraesthesias, difficulties with concentration, and myalgias. Patients in this group were unlikely to be diagnosed with Lyme neuroborreliosis.

Many of those who were diagnosed with definite or possible Lyme neuroborreliosis had had symptoms for less than three months, and none had had symptoms for more than six months. This may indicate that prolonged non-specific symptoms render a diagnosis of Lyme neuroborreliosis less likely. The symptoms of the disease are often such that patients seek medical assistance relatively quickly. A large percentage of the patients with non-typical symptoms had had them for more than a year, and none received a diagnosis of Lyme neuroborreliosis.

Table 1 Number of persons receiving a diagnosis of Lyme neuroborreliosis among those with(out) typical symptoms who were referred for evaluation with lumbar puncture at the Department of Neurology, Sørlandet Hospital Arendal in the period 1 January 2013–31 December 2013

	Negative for Lyme neuroborreliosis	Possible Lyme neuroborreliosis	Definite Lyme neuroborreliosis	Total
Non-specific neurological symptoms	109	1	0	110
Typical symptoms of Lyme neuroborreliosis	23	1	6	30
Total	132	2	6	140

Table 2 Number of patients with various symptoms/symptom duration with definite/possible Lyme neuroborreliosis/no Lyme neuroborreliosis among those who were referred for evaluation with lumbar puncture at the Department of Neurology, Sørlandet Hospital Arendal in the period 1 January 2013–31 December 2013

	Definite or possible Lyme neuroborreliosis (n = 8)	Negative for Lyme-neuroborreliosis (n = 132)
Symptoms		
Cranial nerve deficit	5	16
Radiculitis	4	11
Paraesthesias	4	55
Myalgias	3	69
Tension-type headache	3	45
Tiredness	1	57
Dizziness	1	39
Meningitis symptoms	0	1
Difficulty concentrating	0	28
Tick bite within last 3 months	2	8
Erythema migrans	0	6
Symptom duration		
< 1 month.	4	36
1-3 months	2	23
3-6 months	1	10
6-12 months	0	15
> 12 months.	0	40
Unknown	1	8

Whether lumbar puncture is necessary to diagnose Lyme neuroborreliosis is subject to international debate. Lumbar puncture is not required in the USA, whereas positive CSF results are required in Europe (7, 10, 20, 21). Certain doctors and patients argue that Lyme neuroborreliosis can be diagnosed on the basis of symptoms alone, including non-specific symptoms such as tiredness, myalgias and headache, and with the use of non-validated tests (15, 22). However, non-specific symptoms are common in the general population and there is little evidence that these are necessarily caused by Lyme neuroborreliosis; such diagnostic criteria are therefore not accepted by the major international academic communities (8, 14-16, 23, 24). An increased focus on the possible under-diagnosis of Lyme neuroborreliosis in persons with medically unexplained symptoms may have led to the increased use of lumbar puncture observed in recent years.

In our dataset, we found that all patients

with definite Lyme neuroborreliosis had relatively high titres of Borrelia IgG antibodies in serum. Similar values were seen in 27 % of those who did not receive the diagnosis. In Sogn og Fjordane county, 10% of healthy blood donors are Borrelia-IgG-seropositive - without this being associated with health problems (24, 25). In Vest-Agder county, 18 % of the population have serum antibodies (26). Exposure to Borrelia can lead to seropositive status for several years, for both IgM and IgG (27). A number of those with nonspecific neurological symptoms may have been referred for evaluation for Lyme neuroborreliosis after testing positive for *Borrelia* antibodies at their GP surgery. The detection of anti-Borrelia antibodies in serum does not necessarily mean that a patient's health problems are caused by Lyme neuroborreliosis, but a positive result may support the diagnosis in those who do have symptoms of the disease (24).

Peripheral facial nerve palsy was present

in 63% of patients with possible or definite Lyme neuroborreliosis, and 24% of those assessed for peripheral facial nerve palsy received a diagnosis of either possible or definite Lyme neuroborreliosis. None of the eight patients with definite or possible Lyme neuroborreliosis had had a rash suspected of being erythema migrans, and only two recalled a tick bite within the last three months. It is known that about half of patients with Lyme neuroborreliosis do not recall either a tick bite or erythema migrans (7, 28).

Lyme neuroborreliosis is thus an important differential diagnosis in cases of newonset peripheral facial nerve palsy, irrespective of whether the patient can recall a tick bite or erythema migrans. Sørlandet Hospital has introduced lumbar puncture for all patients with peripheral facial nerve palsy, and our results support this practice in an area highly endemic for Borrelia. None of those who received a diagnosis of possible or definite Lyme neuroborreliosis had symptoms of meningitis (headache with hypersensitivity to light and sound, and nausea or vomiting). The symptoms of Borrelia meningitis may be pronounced but often resemble those of viral meningitis - headache of varying severity and fluctuating fever, but with no other signs of meningitis

One limitation of this study is that it is based on a review of medical records and not on direct discussions with, and examination of, patients. The dataset is relatively small, with only six patients with definite Lyme neuroborreliosis. Most of the records and referrals provided a good picture of symptoms and medical history, but it was not possible to resolve any ambiguities in the selfreported medical history and clinical results. We assumed that if a symptom was not described in the referral or medical records, then the patient had not had that symptom however, this cannot be ruled out for certain. The individual who recorded symptoms was not formally blinded with respect to the results of the CSF analysis.

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References

- Stanek G, Wormser GP, Gray J et al. Lyme borreliosis. Lancet 2012; 379: 461–73.
- Nadelman RB, Nowakowski J, Forseter G et al. The clinical spectrum of early Lyme borreliosis in patients with culture-confirmed erythema migrans. Am J Med 1996; 100: 502–8.
- Stanek G, Strle F. Lyme disease: European perspective. Infect Dis Clin North Am 2008; 22: 327–39, vii. vii.
- Bacon RM, Kugeler KJ, Mead PS. Surveillance for Lyme disease–United States, 1992–2006. MMWR Surveill Summ 2008; 57: 1–9.
- Halperin JJ. Nervous system Lyme disease. Infect Dis Clin North Am 2015; 29: 241–53.
 Hansen K, Lebech AM. The clinical and epidemiolo-
- Hansen K, Lebech AM. The clinical and epidemiological profile of Lyme neuroborreliosis in Denmark 1985–1990. A prospective study of 187 patients with Borrelia burgdorferi specific intrathecal antibody production. Brain 1992; 115: 399–423.
- Oschmann P, Dorndorf W, Hornig C et al. Stages and syndromes of neuroborreliosis. J Neurol 1998; 245: 262–72.

- Ogrinc K, Lotrič-Furlan S, Maraspin V et al. Suspected early Lyme neuroborreliosis in patients with erythema migrans. Clin Infect Dis 2013; 57: 501–9.
 Øymar K, Tveitnes D. Clinical characteristics of
- Øýmar K, Tvěitnes D. Clinical characteristics of childhood Lyme neuroborreliosis in an endemic area of northern Europe. Scand J Infect Dis 2009; 41:88-94
- Mygland A, Ljøstad U, Fingerle V et al. EFNS guidelines on the diagnosis and management of European Lyme neuroborreliosis. Eur J Neurol 2010; 17: 8–16, e1–4.
- 11. Djukic M, Schmidt-Samoa C, Lange P et al. Cerebrospinal fluid findings in adults with acute Lyme neuroborreliosis. J Neurol 2012; 259: 630–6.
- Eikeland R, Mygland A, Herlofson K et al. European neuroborreliosis: quality of life 30 months after treatment. Acta Neurol Scand 2011; 124: 349 – 54.
- Dersch R, Sommer H, Rauer S et al. Prevalence and spectrum of residual symptoms in Lyme neuroborreliosis after pharmacological treatment: a systematic review. J Neurol 2016; 263: 17–24.
- tematic review. J Neurol 2016; 263: 17–24.

 14. Wormser GP, Weitzner E, McKenna D et al. Longterm assessment of fatigue in patients with culture-confirmed Lyme disease. Am J Med 2015; 128: 181–4.
- Feder HM Jr, Johnson BJ, O'Connell S et al. A critical appraisal of «chronic Lyme disease». N Engl J Med 2007: 357: 1422–30.
- J Med 2007; 357: 1422–30.

 16. Wormser GP, Dattwyler RJ, Shapiro ED et al. The clinical assessment, treatment, and prevention of lyme disease, human granulocytic anaplasmosis, and babesiosis: clinical practice guidelines by the Infectious Diseases Society of America. Clin Infect Dis 2006; 43: 1089–134.
- Ocampo JMF, Jore S, Vold I et al. Årsrapport. Flått og flåttbårne sykdommer i 2015. Oslo: Folkehelseinstituttet, 2016.
- Kjelland V, Stuen S, Skarpaas T et al. Prevalence and genotypes of Borrelia burgdorferi sensu lato infection in Ixodes ricinus ticks in southern Norway. Scand J Infect Dis 2010; 42: 579–85.
- Grude N. Laboratoriediagnostikk ved borreliose. Strategirapport. Oslo: Folkehelseinstituttet, 2011.

- 20. Rupprecht TA, Pfister HW. What are the indications for lumbar puncture in patients with Lyme disease? Curr Probl Dermatol 2009; 37: 200–6.
- Stanek G, Fingerle V, Hunfeld KP et al. Lyme borreliosis: clinical case definitions for diagnosis and management in Europe. Clin Microbiol Infect 2011; 17: 69–79.
- Cameron D, Gaito A, Harris N et al. Evidence-based guidelines for the management of Lyme disease.
 Expert Rev Anti Infect Ther 2004; 2 (suppl): S1-13.
 Ljøstad U, Mygland Å. Chronic Lyme; diagnostic
- Ljøstad U, Mygland Å. Chronic Lyme; diagnostic and therapeutic challenges. Acta Neurol Scand Suppl 2013; 127: 38–47.
- Hjetland R, Reiso H, Ihlebæk C et al. Subjective health complaints are not associated with tick bites or antibodies to Borrelia burgdorferi sensu lato in blood donors in western Norway: a crosssectional study. BMC Public Health 2015; 15: 657.
- Hjetland R, Nilsen RM, Grude N et al. Seroprevalence of antibodies to Borrelia burgdorferi sensu lato in healthy adults from western Norway: risk factors and methodological aspects. APMIS 2014; 122: 1114–24.
- Mygland A, Skarpaas T, Ljøstad U. Chronic polyneuropathy and Lyme disease. Eur J Neurol 2006; 13: 1213–5.
- Kalish RA, McHugh G, Granquist J et al. Persistence of immunoglobulin M or immunoglobulin G antibody responses to Borrelia burgdorferi 10–20 years after active Lyme disease. Clin Infect Dis 2001; 33: 780–5.
- Ljøstad U, Skogvoll E, Eikeland R et al. Oral doxycycline versus intravenous ceftriaxone for European Lyme neuroborreliosis: a multicentre, noninferiority, double-blind, randomised trial. Lancet Neurol 2008: 7: 690-5.
- Stanek G, Strle F. Lyme borreliosis. Lancet 2003; 362: 1639–47.

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