CASE REPORT

Legionella longbeachae; don’t miss it!

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ABSTRACT

We here report on two immunocompetent patients admitted to our hospital within 3 weeks’ time, both suffering from pneumonia caused by Legionella longbeachae (L. longbeachae). The pathogen was identified in broncho-alveolar lavage (BAL) liquid by Polymerase Chain Reaction (PCR), whereas sputum cultures remained negative. This organism is worldwide still relatively unknown and consequently underdiagnosed. However, with an increasing number of confirmed infections in Europe and more specifically in the Netherlands, early awareness and diagnostic measurements are indicated. As routine laboratory techniques like the urine antigen test do not detect L. longbeachae, we advocate early use of specific tests for non-pneumophila Legionella species such as PCR. Furthermore, we advocate the start of empirical antibiotic therapy (i.e. ciprofloxacin) and continuation in suspected cases.

KEYWORDS

ICU, Intensive Care Unit; legionella; legionella longbeachae; pneumoniae

INTRODUCTION

As one of over 50 species, L. longbeachae belongs to the extensive family of Legionellae. These gram negative, opportunistic bacteria can cause Pontiac fever and Legionnaires’ disease, a multisystem disease including pneumonia with a severe course. It has no particular clinical features that distinguish it from other types of pneumonia. In contrast to L. pneumophila which is linked mainly to contaminated water reservoirs, the general source of contamination for L. longbeachae is considered to be compost and other growing media. Since L. pneumophila is responsible for 80-90% of the Legionella infections in Europe, our diagnostic system is mainly targeting this species. The restricted sensitivity of the urine antigen test and the low sensitivity of culture testing cause the risk of false rejection of the diagnosis Legionella’s pneumonia.

L. longbeachae is increasingly being reported in Europe. Over the last decade the incidence of infections by this pathogen has also increased in Europe, including the Netherlands. L. longbeachae is not detected by the commonly used urine antigen test, and it often does not grow in diagnostic cultures. Awareness of the risk of non-pneumophila Legionella species is often still limited to immune-deficient patients and severe cases of pneumonia without clinical recovery on empirical antibiotic therapy.

What was known on this topic?
L. longbeachae is known to be a major cause of Legionellosis in Australia and South East Asia. Over the last decade the incidence of infections by this pathogen has also increased in Europe, including the Netherlands. L. longbeachae is not detected by the commonly used urine antigen test, and it often does not grow in diagnostic cultures. Awareness of the risk of non-pneumophila Legionella species is often still limited to immune-deficient patients and severe cases of pneumonia without clinical recovery on empirical antibiotic therapy.

What does this add?
After the first L. longbeachae case report in an immunocompetent patient by Diederen et al. in 2005 and recently in a brief photo quiz by Hannivoort et al, there were, to the best of our knowledge, no other cases published in our country. Since 2014 the national incidence seems on the rise. This case series confirms this observation, and emphasizes the need for early awareness and diagnostic interventions for non-pneumophila legionella species.
CASE REPORTS

Patient A is a 79-year-old man admitted in June 2017 with malaise, fever and heavy diaphoresis. Medical history included type 2 diabetes mellitus and hypertension. He had a history of heavy nicotine abuse in the past with 100 pack years until 8 years ago, implying the possibility of undiagnosed chronic obstructive lung disease. Physical examination showed a heart rate of 80/minute, blood pressure of 200/105 mmHg, SpO₂ of 94% without supplemental oxygen, respiration rate of 16/minute and a tympanic temperature of 40°Celsius. C-reactive protein (CRP) was 197 mg/l and there was minimal leucocytosis. Chest X-ray showed consolidation of the left upper lung. Urine antigen tests (Streptococcus pneumoniae and Legionella pneumophila) were negative. We suspected pneumonia and started intravenous treatment with cefuroxim. Two days later there was no clinical improvement and CRP rose to 330 mg/l. Sputum cultures and HIV-testing remained negative. On suspicion of atypical bacteria, antibiotics were switched to azithromycin. On day four of admission a BAL was performed, and subsequent PCR on Legionella species revealed L. longbeachae upon which antibiotics were switched to ciprofloxacin. Despite the positive PCR results, sputum cultures remained negative for Legionella species. On day six a chest CT-scan showed extensive bilateral ground glass opacity. Chest CT-scan showed extensive right-sided ground glass opacity. Urine antigen tests (Streptococcus pneumoniae and Legionella pneumophila) and sputum cultures remained negative. Because of the recent travel and hence potential atypical pneumonia, treatment with ciprofloxacin and amoxicillin was started. The next day he was transferred to the Intensive Care Unit due to respiratory insufficiency. Physical examination showed tachypnoea of 30/minute, SatO₂ of 94% with 10 litres of oxygen per minute on a non-rebreather-mask, tachycardia of 120/minute, blood pressure of 127/71 mmHg and tympanic temperature of 38°Celsius. Laboratory results showed a rise in CRP rise to 318 mg/l and leucocytosis of 25.7 x10⁹/l. Blood gas analysis showed hypoxia (pH 7.41, HCO₃⁻ 20.3 mmol/l, Base Excess -3.9 mmol/l, pCO₂ 4.2 kPa, pO₂ 9.2, SatO₂ 94%). He was intubated and mechanically ventilated. The following days his pulmonary condition worsened and on day four the heart was hypertrophic, with signs of subendocardial infarction of the left ventricular cardiac wall, in addition with signs of swelling of liver and spleen implying acute congestive cardiac failure as the probable cause of death.

Patient B, also a 79-year-old man, was admitted three weeks later with dyspnoea, altered consciousness and confusion. Medical history revealed obstructive sleep apnoea syndrome and – uncomplicated – knee surgery 10 days before this current admission. He was a non-smoker. Notable detail was a recent 2-week vacation in Turkey from which he had returned five weeks before this admission. Physical examination on admission showed a heart rate of 104/minute, blood pressure of 130/70 mmHg, SpO₂ of 93% with 6 litres of oxygen trough an oral-nasal mask, respiration rate of 30/minute and a tympanic temperature of 37.8°Celsius. Chest CT-scan showed extensive right-sided ground glass opacity. Urine antigen tests (Streptococcus pneumoniae and Legionella pneumophila) and sputum cultures remained negative. Because of the recent travel and hence potential atypical pneumonia, treatment with ciprofloxacin and amoxicillin was started. The next day he was transferred to the Intensive Care Unit due to respiratory insufficiency. Physical examination showed tachypnoea of 30/minute, SatO₂ of 94% with 10 litres of oxygen per minute on a non-rebreather-mask, tachycardia of 120/minute, blood pressure of 127/71 mmHg and tympanic temperature of 38°Celsius. Laboratory results showed a rise in CRP rise to 318 mg/l and leucocytosis of 25.7 x10⁹/l. Blood gas analysis showed hypoxia (pH 7.41, HCO₃⁻ 20.3 mmol/l, Base Excess -3.9 mmol/l, pCO₂ 4.2 kPa, pO₂ 9.2, SatO₂ 94%). He was intubated and mechanically ventilated. The following days his pulmonary condition worsened and on day four...
of admission antibiotics were switched to ceftriaxone. CT-scan showed expansion of infiltration of the left lung (figure 1, panel B) upon which a BAL was performed. On day seven ciprofloxacin was restarted, because clinical suspicion of Legionellosis still remained, and his clinical situation under treatment with ceftriaxone worsened. PCR on atypical pathogens in the BAL liquid tested positive for \textit{L. longbeachae} whereas cultures remained negative. Treatment with ciprofloxacin was continued for three weeks. The patient recovered slowly, and after 18 days he was extubated and eventually discharged from the hospital. Retrospectively, patient A frequently worked with potting soils in his garden, potentially being the source of contamination. Patient B didn’t work with compost soils, but he had recently been gardening near his garden pond. Unfortunately the Area Health Authorities performed no source detection investigation, because \textit{L. longbeachae} is not yet recognized as a public health issue.

\textbf{DISCUSSION}

Despite an increase in the last decade, Legionellosis caused by non-pneumophila species is still rare on our continent. From 2011 to 2015, 3645 culture-confirmed cases of Legionella’s infection in Europe were documented, 83\% of which were caused by \textit{L. Pneumophila}.\textsuperscript{9-10} Among the non-pneumophila group, \textit{L. longbeachae} is the dominant species with 35 culture-confirmed cases (1\%) within these five years.\textsuperscript{9-11} In Australia and Eastern Asia \textit{L. longbeachae} is responsible for approximately half of the Legionellosis.\textsuperscript{4,12} In the Netherlands we confirmed a maximum of 1 patient per year in the decade preceding 2014, but from then on the incidence appears to rise (table 1). How can this be explained? As we know, many factors contribute to incidence outcomes, such as geographical distribution, infectious dose, infection type, host susceptibility and the sensitivity of detection techniques. A contributory theory could be that there has been a change in the composition of European compost soils. One of the explanations why \textit{L. longbeachae} occurs in large numbers in Australia and South East Asia derives from the theory that the presence of bark containing sawdust and hammer milled bark makes these soils a natural habitat for the bacterium.\textsuperscript{12} Nowadays Australian growing materials and soils even come with warning labels attached (figure 2). In Europe, peat rather than composted bark is the major component used in growing media, but the composition is always subject to change.\textsuperscript{13}

Another contributing factor could be the wider application of PCR with regard to Legionella species in hospital laboratories nationwide, causing higher detection rates. Formerly incidence rates depended on the availability of appropriate laboratory techniques and skilled employees for analysis. In Scotland, where several outbreaks of \textit{L. longbeachae} were recorded, researchers also described underdetection of these infections.\textsuperscript{5,14} Current diagnostics like the urine antigen test are useful to detect \textit{L. pneumophila} serogroup 1, but are not suitable

\textbf{Table 1. Data from the National Institute for Public Health and the Environment (RIVM) showing an increase of incidence in The Netherlands over the past decade}\textsuperscript{1}

<table>
<thead>
<tr>
<th>Year</th>
<th>Culture confirmed cases</th>
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<tbody>
<tr>
<td>2008</td>
<td>1</td>
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<tr>
<td>2009</td>
<td>0</td>
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<tr>
<td>2010</td>
<td>1</td>
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<td>2011-2013</td>
<td>0</td>
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<td>2014</td>
<td>3</td>
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<td>2015</td>
<td>2</td>
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<tr>
<td>2016</td>
<td>6</td>
</tr>
<tr>
<td>2017</td>
<td>5*</td>
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for non-pneumophila Legionella species.\textsuperscript{15} Whereas bacterial culture remains the golden standard for confirmation of Legionella’s disease is,\textsuperscript{14,15} this technique requires specialized laboratory media as overgrowth with competing flora is a common problem.\textsuperscript{15} As illustrated by both of our cases the cultures remained ‘false’ negative, confirming that PCR is an essential test for the diagnosis. In conclusion, \textit{L. longbeachae} is a sporadically emerging pathogen with a severe course of disease. Over the past decade the number of reports on \textit{L. longbeachae} has increased markedly across Europe. Timing of diagnosis and targeted therapy influences the severity and course of the disease. Recommended antibiotic treatment for \textit{L. longbeachae} (like pneumonia with (suspected) atypical bacteria in general and Legionella specifically) are fluorquinolones like ciprofloxacin. Physicians and guideline committees, like the Dutch SWAB-guidelines committee,\textsuperscript{16} have adopted the concept of using the ‘severity of illness’ at the time of clinical presentation (based on validated scoring systems like CURB-65 and Pneumonia Severity Index (PSI), and the need for ICU admission) in order to choose the type of empirical antimicrobial in community acquired pneumonia (CAP). In category IV (severe CAP with ICU admission) it is always recommended to empirically cover \textit{S. pneumoniae}, Legionella species and Gram-negative bacteria.\textsuperscript{16}

It is difficult to predict atypical bacteria being the cause of CAP based on presentation, signs and symptoms of CAP, medical history, physical examination and laboratory results.\textsuperscript{17} We recommend extending the medical interview on admission routinely from inquiring after travel abroad and visits to swimming pools and sauna’s to questions on recent gardening activity and exposure to gardening soil. We also recommend the early usage of rapid and sensitive PCR techniques on non-pneumophila Legionella species in patients with severe and treatment-resistant pneumonia, especially in the presence of risk factors like recent exposure to gardening soils. This way, the diagnostic process may be instrumental in improving things; more awareness among diagnosing clinicians and laboratory staff of non-pneumophila Legionella species is necessary to reduce underdiagnoses and associated risks.

**DISCLOSURES**

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**REFERENCES**