**ABSTRACT**

Since 2007, the Netherlands is faced with the largest outbreak of Q fever ever reported. In the last four years, over 4000 cases have been reported. The course of the epidemic and possible factors associated with this sudden surge in cases of Q fever is described and the preventive measures in the veterinary sector and the outbreak management of this unique epidemic are summarised. Finally, the latest data on clinical presentation and diagnostic and therapeutic dilemmas of Q fever in the Netherlands are reviewed.

**KEYWORDS**

Coxiella, Q fever, the Netherlands

**INTRODUCTION**

Since 2007, the Netherlands is faced with the world’s largest outbreak of Q fever with over 4000 notified cases. Q fever is caused by *Coxiella burnetii*, a small, Gram-negative obligate intracellular bacterium, phylogenetically related to Legionellales. Transmission occurs through inhalation of infected aerosols. The reservoir consists mainly of dairy goats and sheep, but excretion of *Coxiella* has also been described in other cattle and rodents. Small outbreaks associated with parturient pets such as cats and dogs have also been reported. The animals shed the bacterium in urine and faeces, and in very high concentrations in birth by-products. *C. burnetii* is extremely infectious as was illustrated by an experiment demonstrating that inhalation of a single bacterium could cause seroconversion in humans. It was classified as a category B bioterrorism agent.
Acute Q fever occurs two to six weeks after exposure depending on the infective dose. The infection remains asymptomatic in up to 60% of patients. Patients with symptomatic disease usually present with fever and mild flu-like symptoms. Because these symptoms are very non-specific, under-reporting is probably quite substantial. A case-control study investigating the first outbreak of Q fever in the Netherlands in 2007 showed that all patients with symptomatic infection experienced fever. Headache and cough were reported by 92 and 85%, respectively. Smoking was found to be an important risk factor, as had been shown by others previously. Male sex has also been identified as a risk factor for symptomatic disease. In 2007 and 2008, the female-to-male ratio of acute Q fever was 1:1.7. The mean age was 51 years. Although hospitalisation rates of 2% have been described in literature, hospital admission was much more frequent in the Netherlands. In 2007, almost 50% of Q fever cases were admitted. This high percentage could have been influenced by active case finding in a retrospective study among hospitalised patients. In the subsequent years, admission rates stabilised to around 20%, still considerably higher than reported in literature.

Chronic Q fever
Chronic Q fever develops in 1 to 2% of patients after acute Q fever. Some patients with chronic Q fever do not recall having had an acute infection. It usually develops insidiously, months or even years after acute infection and patients often present with non-specific symptoms such as low-grade fever, night sweats and weight loss. In a large retrospective study from France, endocarditis was found to be the predominant manifestation of chronic Q fever, constituting 73% of chronic Q fever cases. Other manifestations were vascular infection (8%), chronic infection in pregnancy (6%), and chronic hepatitis (3%).

In the Netherlands however, a substantially higher percentage of chronic Q fever cases consists of patients with infected aneurysms and vascular prostheses. In a recent report, 12 out of 22 chronic Q fever patients in the Netherlands had vascular infection. Four of these patients were diagnosed by screening 52 patients with an aortic aneurysm. The authors advocate screening all patients with symptomatic aortic aneurysms and vascular prostheses. In a nationwide database on chronic Q fever will be established to collect these data and facilitate epidemiological research.

Q fever and pregnancy
Literature on chronic Q fever during pregnancy is limited. A case series of 53 pregnant women diagnosed with Q fever showed obstetric complications in 81% of patients not treated with long-term cotrimoxazole therapy compared with 44% in patients who did receive cotrimoxazole. An important pitfall in this observational study, as indicated by the authors, is the fact that serology for Q fever was
performed only after delivery in 28% of patients, often because of obstetric complications, creating a selection bias. Interpretation of these results is therefore difficult. Two large seroprevalence studies found no significant association between seropositivity for Coxiella and adverse pregnancy outcome. A study among pregnant women in the area of the first outbreak in the Netherlands in 2007 showed evidence of recent infection in three out of 19 women (16%). This was significantly higher than in the surrounding regions. A retrospective study in the highly endemic area in the southern part of the Netherlands showed no significant correlation of seropositivity for Q fever during early pregnancy and adverse pregnancy outcome. To further investigate the effect of Q fever on pregnancy, a randomised controlled trial was started in the spring of 2010 evaluating the effect of a screen and treat policy of pregnant women in this area.

**Post Q fever fatigue syndrome**

Following acute Q fever, patients frequently report a long-lasting and debilitating fatigue. A study performed after an outbreak of acute Q fever in the United Kingdom showed 20% of patients suffered from chronic fatigue syndrome after ten years of follow-up, compared with 4% of controls. A study among abattoir employees in Australia showed that 28% of patients with proven acute Q fever still fulfilled the CDC criteria of chronic fatigue syndrome at five years after infection compared with none of seronegative controls. A case-control study among 54 patients who contracted acute Q fever in the first year of the epidemic in the Netherlands showed that after one year, over 50% still reported severe fatigue compared with 26% of controls. The aetiology of this severe fatigue, referred to as QFS (Q fever Fatigue Syndrome), still remains to be elucidated. Cytokine dysregulation resulting from chronic immune stimulation and modulation by persistence of Coxiella or its antigens has been hypothesised. Genotyping of patients suffering from QFS showed significant differences in HLA-DRB1*11 and interferon-γ intron 1 microsatellite compared with controls. Some reports suggest persistence of Coxiella or antigenic non-viable cell residues in bone marrow. Studies evaluating antibiotic treatment for QFS have shown conflicting results. QFS leads to considerable morbidity and a high socioeconomic burden related to increased use of healthcare facilities and absence from work.

**Diagnosis of Q Fever**

Analysis of specimens from various infected dairy farms has shown that 14 different strains circulate in the Netherlands, but one is predominantly present in the highly endemic area (Roest HJ, unpublished data). Isolation of Coxiella from blood culture specimens of Q fever patients is difficult since it is an obligatory intracellular organism. In addition, chronic infection often resides in tissues (e.g., heart valves or vascular aneurysms) and shedding into peripheral blood occurs in very low concentrations. Culture of Coxiella requires very specific procedures and a biosafety level 3 laboratory, which is not available to most hospitals. Until now, culture of the pathogen has been successful in only one patient in the Netherlands, in whom Coxiella was cultured from a resected heart valve (Roest HJ, personal communication). Diagnosis of acute Q fever is based on serology, the reference method being immunofluorescence assay (IFA). A seroconversion of a fourfold rise in antibody titre is diagnostic for acute Q fever. An important drawback in diagnosis based on serology is that antibody production does not usually occur until a few weeks after onset of clinical symptoms. PCR on serum has been shown to have a high sensitivity (98%) for acute Q fever in seronegative patients and is a useful diagnostic tool for early diagnosis. An algorithm for the diagnosis of acute Q fever in the Netherlands, designed by the Dutch working group on diagnostics of acute Q fever (an initiative of the National Institute for Public Health and the Environment (RIVM) and the Dutch Association for Medical Microbiology), has been published very recently. There has been a substantial reduction of the diagnostic delay in the Netherlands from 82 days in 2007 to 20 days in 2009. Because treatment has to be started before laboratory confirmation, physicians have to rely on clinical signs to guide the decision to start empiric therapy for Q fever. Antibiotic treatment of community acquired pneumonia in a high endemic region should include an agent active against Coxiella burnetii. Diagnosis of chronic Q fever remains difficult. Because the infection persists intracellularly, PCR on peripheral blood is not always positive. Imaging techniques can be useful to localise infection. Infected aneurysms or vascular prostheses can be identified by ¹⁸F-fluorodeoxyglucose (FDG) positron emission tomography (PET) or CT. In case of endocarditis, however, diagnosis is often more complex. The original Duke criteria developed for diagnosing infective endocarditis include vegetations and positive blood cultures as major criteria. However, vegetations are often absent in Q fever endocarditis and as mentioned above, Coxiella does not grow in conventional blood culture media. Therefore a revision of the Duke criteria has been proposed in which a serological profile compatible with chronic Q fever has been added to the major criteria. Serology is therefore an important tool in the evaluation of the development of chronic disease. Coxiella burnetii displays a unique antigenic variance in surface polysaccharides (phase 1 and phase 2 antigens). This can be used to distinguish between acute and chronic infection.
In acute infection, mainly phase 2 antibodies develop and convalescent sera show low titres of phase 1 antibodies, whereas chronic infection is characterised by high titres of phase 1 antibodies. Most literature on determination of cut-off values for establishing the diagnosis of chronic Q fever originates from the French National Reference Centre for Rickettsial Diseases (NRC) and this group has proposed a cut-off value for IgG to phase 1 proteins of 1:800 for the diagnosis of chronic Q fever. This cut-off value was also adopted by the revised Duke criteria. In the Netherlands, however, a substantial percentage of patients showed much higher titres of IgG1 during the first months after acute infection. There seems to be a considerable difference between the serological method used in the Netherlands (Focus diagnostics) and the method used in the NRC in France. This was recently illustrated by a case report of serological follow-up after acute Q fever, which showed high titres to IgG1 comparable with those found in Dutch patients but much lower when tested in the NRC in France. In 2009, a provisional guideline was published in the Netherlands proposing a new cut-off value for IgG1 of 1:4096 (or an IgG1 equal to IgG2) (www.medischcontact.artsennet.nl). However, when using this algorithm, 40% of patients with proven chronic Q fever (signs and or symptoms compatible with chronic Q fever and persistently positive PCR on blood or positive PCR on resected valves or aneurysms) who presented at the Radboud Expertise Centre for Q fever do not fulfill these criteria (table 1). The Netherlands Society for Medical Microbiology (NVMM) is currently developing a new guideline for the diagnosis of chronic Q fever in the Netherlands.

## Treatment of Q Fever

Comparative trials regarding the optimal antibiotic treatment for acute Q fever are sparse, often retrospective and sometimes show conflicting results. Doxycycline is the preferred choice, but the new-generation quinolones such as moxifloxacin are also active against C. burnetii. Clarithromycin has also been shown to be effective and co-trimoxazole is the treatment of choice in children younger than 8 years of age. Although the national guidelines for treating community acquired pneumonia (CAP) issued by the Dutch Working Party on Antibiotic Policy (SWAB) and the Dutch College of General Practitioners (NHG) recommend doxycycline for first-line treatment of CAP, the alternative regimens in these guidelines do not routinely cover C. burnetii. Nevertheless, most general practitioners in the highly endemic region are aware of this problem and treat their patients with either doxycycline or moxifloxacin.

A survey among general practitioners in this region showed that 95% would consider Q fever as a possible pathogen when suspecting a pneumonia and would start empiric treatment with doxycycline. The move away from doxycycline in the proposed update of the NHG guidelines for treatment of pneumonia by general practitioners seems inappropriate for endemic regions and may lead to increased numbers of chronic infections.

The optimal treatment of chronic Q fever consists of doxycycline and hydroxychloroquine. The latter increases the intralysosomal pH and thereby achieves a bactericidal effect in vitro when combined with doxycycline, whereas doxycycline alone is only bacteriostatic. Based on a retrospective study, a minimum duration of 18 months of combination therapy has been recommended with target levels of doxycycline of 5 mg/l. This long-term therapy is associated with significant adverse effects and photosensitisation has been described in 100% of patients on long-term therapy. Other frequently reported side effects include nausea, headache and dizziness. Regular evaluation by an ophthalmologist is recommended because of possible irreversible maculopathy due to hydroxychloroquine. Maintaining optimal adherence to therapy therefore requires intensive counselling and therapeutic drug monitoring.

<table>
<thead>
<tr>
<th>Patient</th>
<th>Localisation of infection</th>
<th>IgG1 titre at diagnosis</th>
<th>IgG2 titre at diagnosis</th>
<th>PCR Q fever Blood</th>
<th>PCR Q fever Tissue</th>
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<td>32268</td>
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</table>
PREVENTION OF Q FEVER IN HUMANS

The preventive measures taken in the veterinary sector have been aimed at reducing the spread of *C. burnetii* in the environment and thereby limiting the transmission to humans. Human vaccination is a different approach in preventing Q fever in individuals with a high risk of exposure to *Coxiella*. An effective whole-cell vaccine is available in Australia and has been extensively used in persons with high occupational risks such as abattoir employees. In this population, it has been proven to be highly effective. Because administering this vaccine to patients with pre-existing immunity increases the risk of local and systemic inflammatory reactions, prior infection needs to be excluded by skin testing and serology. Recently, the Health Council of the Netherlands issued an advice on vaccinating patients with increased risk of chronic Q fever with this whole cell vaccine (http://www.gezondheidsraad.nl/nl/advies/en-van-mensen-tegen-Q-koorts-eerste-advies). The target population has been defined as patients with underlying cardiac conditions (the same category of patients who would also qualify for endocarditis prophylaxis according to current guidelines), as well as patients with a known (aortic) aneurysm or vascular prosthesis.

Since Q fever is highly endemic in the southern part of the Netherlands and infection can be asymptomatic, there is a possible risk of transmission through blood transfusion. Preliminary results indicate that in 2009 *C. burnetii* DNA was present in a small percentage of blood donations in this area (indicated by positive PCR on donated blood). Sanquin, the Dutch blood supply foundation, has instituted screening of donated blood in the high-incidence area as a precautionary measure.

CONCLUSION

Although it appears that the epidemic of Q fever in the Netherlands is now subsiding, physicians are still faced with growing numbers of patients suffering from long-term sequelae of Q fever such as chronic infection and Q fever fatigue syndrome. Optimal management of these conditions is still unclear and further research is needed to improve diagnostic strategies, to evaluate treatment, and to prevent chronic infections.

REFERENCES


