Infective Endocarditis with Negative Cultures: A Defiant Diagnosis

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Abstract

The diagnosis and management of blood culture-negative endocarditis constitute a real clinical challenge and a systemic approach is necessary for a successful outcome. The authors report a case of a female patient aged 26, with previous clinical history of valve disease and heart failure NYHA class II, with decompensation of co-morbidities associated with fever, productive cough, nonselective anorexia and unquantified weight loss with one month of evolution with negative blood cultures. Transesophageal echocardiography revealed the presence of severe pulmonary hypertension and mitral valve vegetations in the context of positive serology for Q fever. Herein, the authors report a case of blood culture-negative endocarditis and present a brief review on the management of this medical condition. We highlighted the diagnostic difficulties of blood culture-negative endocarditis and subacute clinical presentation, which sometimes present with fever of unknown origin and complaints of deterioration of cardiac function, thus creating a challenging differential diagnosis.

Keywords

Coxiella burnetii, Endocarditis, Blood Culture-Negative Endocarditis

1. Introduction

Blood culture-negative endocarditis (BCNE) defines certain or likely endocarditis in which three or more aerobic and anaerobic blood cultures collected over 48 hours persist negative regardless of extended incubation period (superior to one week) [1] [2]. Definitely, it establishes a significant clinical challenge and is frequently connected with higher morbidity and mortality, when compared with blood culture positive endocarditis [3].

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BCNE comprises 2.5% to 31% of all cases of endocarditis [4].

Infective endocarditis (IE), in particular, is a serious disease whose mortality and morbidity have not considerably decreased over the past 30 years [5]-[7], and is potentially fatal if treatment is not appropriate [8].

The epidemiological profile of IE has changed in recent years, especially in industrialized countries. Once considered a disease affecting mostly young adults diagnosed with valvular disease (mainly rheumatic heart disease), in recent years the incidence has increased in the elderly. The incidence is variable, with a record of 3 - 10 episodes per 100,000 individuals per year [5] [7] [8].

The Duke criteria are based upon clinical, microbiological and echocardiographic findings providing high specificity and sensitivity for the diagnosis of IE. Despite the fast development of additional means of diagnosis and the introduction of the Duke criteria, the diagnosis of IE is still a challenge for many clinicians [9] [10] [12], especially when cultures are negative, causing a delay in diagnosis and appropriate treatment [5]. This variation may be explained by several factors such as a difference in diagnostic criteria used; specific epidemiological factors (related to the tedious zoonotic agents); previous use of antibiotics; inadequate techniques of blood cultures or involvement of unknown agents [11] [12].

The proper treatment depends upon the identification of the causative agent, whereby in the presence of BCNE becomes necessary to diversify laboratory techniques [8] [11]. Even with all incremental knowledge of the pathogenesis of BCNE, treatment and outcome of BCNE have not improved substantially over the past 30 years making it important to notify and discuss clinical cases of BCNE, particularly those with special context such as the case presented here.

2. Case Report

A 28-year-old melanodermic female, single, unemployed, born in Cape Verde, living in Portugal for one month previously to hospital admission, with a clinical history of heart failure (HF) NYHA class II and valvular heart disease (i.e., severe mitral stenosis is, moderate aortic stenosis is with aortic regurgitation and tricuspid stenosis). She was admitted to our hospital with fever, profuse nocturnal sweating, cough with pinkish mucous sputum, non-selective anorexia and unquantified weight loss with one month of evolution. Physical examination revealed fever (tympanic temperature 39.2˚C), normal blood pressure and sinus tachycardia (heart rate 140 bpm), rapid breathing rate and dehydrated pale skin and mucous membranes. Systolic murmur grade IV/VI, audible throughout the precordium, hepatosplenomegaly and mild edema of the lower limbs. The investigations yielded the following results: white blood count, 11,050/mm³; neutrophil, 95%; hemoglobin, 8.3 g/dL (VGM: 27.7 fl; HGM: 79.5 pg), platelets: 150,000/mm³. Blood C-reactive protein 0.7 mg/L. Liver and kidney function tests were in the normal range. Urinalysis and thyroid function was normal. Chest roentogram revealed diffuse and bibasilar interstitial infiltrate.

Assuming a provisional diagnosis of pneumonia, empirical antibiotic therapy with ceftriaxone was started with little improvement. Blood cultures were performed in aerobic and anaerobic conditions; *Mycobacteria spp* in blood and sputum were negative. Suspicious mitral vegetations on transthoracic echocardiography (TTE) were revealed. A transesophageal echocardiography (TEE) subsequently performed showed severe pulmonary hypertension and vegetations of the mitral (Figure 1) and aortic valves. In the light of this new data, the previous antibiotic was suspended and replaced by ampicillin and gentamicin, with some clinical improvement. Given the negative blood cultures and by reference to housing in formerly rural areas, serology for *Coxiella burnetii* (*C. burnetii*) was performed and was positive (IgG titer to phase I and phase II > 1/8190, IgM phase I = 1/2048 and IgM phase II = 1/512). The patient started doxycycline (100 mg twice daily) associated with hydroxychloroquine (600 mg once daily) for 21 days with significant clinical and laboratory improvement.

The patient performed a control transthoracic echocardiography at 28 days, which showed clear improvement of pulmonary hypertension, although keeping vegetations of small volume in aortic and mitral valves. She was discharged after one month of hospitalization, continuing with antibiotic therapy with doxycycline and hydroxychloroquine for 18 months and was directed for cardiothoracic surgery consultation.

3. Discussion

The diversity of presentation of BCNE constitutes a challenging diagnosis. The clinical history may be considerably variable according to the etiologic agent, absence or presence of previous heart disease and temporal evolution (acute, sub-acute or chronic). The physical examination, despite plainly useful in establishing the diagnostic
hypothesis of endocarditis, it is improbable to aid in defining the etiology [3]. The risk factors for BCNE include previous antibiotic administration, exposure to slow-growing and fastidious non-bacterial microorganisms, underlying valvular heart disease and endocarditis in patients with vascular or intracardiac devices.

Blood cultures are a primordial element in the diagnosis of infective endocarditis [12] and when these come up negative without recognition of any microorganism, several hypotheses should be considered such as the following scenarios enlightened here. Previous administration of antibiotics three days or longer prior to the blood cultures may be responsible for BCNE caused by common and easily growing streptococci and staphylococci, as verified by positive polymerase chain reaction (PCR) identification of DNA from these microorganisms. This situation includes 45% to 60% of all cases of culture-negative infective endocarditis (CNIE) [3]. Presence of slow-growing or fastidious organisms requiring specific or enriched media or cell culture systems for intracellular bacteria such as subspecies of streptococci (Abiothrophia spp. which require piridoxal), Gram-negative HACEK group (Haemophilus parainfluenzae, H. aphrophilus, H. paraphrophilus, H. influenzae, Actinobacillus, Actinomyces, Cardiobacterium hominis, Eikenellacorrodens, Kingellakingae, and K. denitrificans), Brucella spp., Legionella spp. and fungi (Candida, Histoplasma bacilli or Aspergillus). However, recent studies have shown that prolonging incubation up to five days do not increase the isolation of specific microorganisms due to the high likelihood of contamination. To emphasize that although IE caused by anaerobes uncommon, cultures should be incubated anaerobic and anaerobic environment, to detect any microorganisms such as Bacteroides species or Clostridium spp. Infections by these organisms account for about 40% to 55% of cases of CNIE [3]. Presence of intracellular organisms, such as C. burnetii, Bartonella spp., Chlamydiae, and Tropheryma whipplei (T. whipplei), that requires serologic testing, histologic valvular evaluation or molecular biology to be identified, account for the remaining 5% of the cases of CNIE. Recently, T. whipplei was the most common isolated organism associated to BCNE.

The Table 1 summarizes some characteristics that increase the degree of suspicion of the referred microorganisms and their additional diagnostic tests to be elaborated.

When the investigation is inconclusive, it should also be suspicious of non-infectious causes of endocarditis classified as non-bacterial thrombotic endocarditis. Given a history of chronic disease or cancer, especially adenocarcinoma associated with disseminated intravascular coagulation, one must consider the hypothesis of marantic endocarditis or aseptic vegetations, complications of systemic lupus erythematosus or anti-phospholipid antibodies syndrome [13]. However, undiagnosed infective endocarditis is probably the most common cause of endocarditis in cancer patients with negative blood cultures [3].

Compared to TTE, TEE shows higher sensitivity in the detection of vegetative lesions, peri-valvular infective foci and mural abscesses with 80% to 90% of sensitivity [4].

Regarding the serological methods, they are available for multiple fastidious microorganisms, being included in the Duke criteria for the diagnosis of infective endocarditis, albeit with specific predictive value. Relatively to C.
Table 1. Agents of EICN, suspicious characteristics and complementary means of diagnosis.

<table>
<thead>
<tr>
<th>Microorganism</th>
<th>Suspicious characteristics</th>
<th>Complementary means of diagnosis</th>
</tr>
</thead>
<tbody>
<tr>
<td>Brucella spp.</td>
<td>Ingestion of animal products derived from unpasteurized milk, Contact with animals</td>
<td>Blood culture; serology; cultural examination, immunohistology and surgical sample PCR</td>
</tr>
<tr>
<td>C. burnetii</td>
<td>Contact with animals</td>
<td>Serology; cultural examination, immunohistology and surgical sample PCR</td>
</tr>
<tr>
<td>Bartonella spp.</td>
<td>Nutrition and poor sanitary conditions (B. quintana), Contact with cats (B. henselae)</td>
<td>Blood culture; serology; cultural examination, immunohistology and surgical sample PCR</td>
</tr>
<tr>
<td>T. whipplei</td>
<td>Clinical picture with diarrhea</td>
<td>Histology and surgical sample PCR</td>
</tr>
<tr>
<td>Mycoplasma spp.</td>
<td></td>
<td>Serology; cultural examination, immunohistology and surgical sample PCR</td>
</tr>
<tr>
<td>Legionella spp.</td>
<td>Nosocomial infection in patients with valvular prosthesis</td>
<td>Blood culture; serology; cultural examination, immunohistology and surgical sample PCR</td>
</tr>
<tr>
<td>C. pneumoniae</td>
<td>Association with atherosclerose and valvular disease; Cross-reaction with Bartonella spp.</td>
<td>Microimmunofluorescencetest; Serology; cultural examination, immunohistology and surgical sample PCR</td>
</tr>
</tbody>
</table>

*burnetii*, a single IgG antibody titer 1:800 is considered major criteria (as seen in our case study). Endocarditis Q fever is characterized by high titers of antigen phase I and II of *C. burnetii*, and the anti-phase I IgG antibody with a titer of 1:600 is considered to have a high positive predictive value of 98%. Indirect immune of luorescence tests and complementary attachment tests are the most reliable methods for the detection of the etiological agent [3]. The sample indicated for molecular detection for identification of the etiologic agent is excised valvular tissue that is only available in patients undergoing previous resection for valvular replacement [3]. Histopathological examination of the resected valve remains the gold standard test for the diagnosis and is a major criterion in the classification of Duke, with the Gimenez staining used to detection of *C. burnetii*.

Regarding the treatment of *C. burnetii* BCNE, the combination of doxycycline and hydroxychloroquine (as was established in the patient in question) remains the cornerstone of treatment, confirming a decrease in the duration of antibiotic treatment compared to monotherapy with doxycycline as well as decreased rate of treatment failure or relapse [3]. The surgical valve replacement remains an important option for patients with BCNE, so our patient was referred to cardiothoracic surgery consultation.

4. Conclusion

With the case previously described, we highlighted the diagnostic difficulties of BCNE and subacute clinical presentation, which sometimes present with fever of unknown origin and complaints of deterioration of cardiac function, thus creating a differential diagnosis challenge. The fact that the serological diagnosis of fastidious microorganisms may be delayed, it’s essential to establish an epidemiological history, as in our case, for a better etiological and therapeutic orientation.

References


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