Purulent pericarditis-induced intracardiac perforation and infective endocarditis due to *Parvimonas micra*: a case report

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**Background**

Purulent pericarditis, a rare disease with a high associated mortality rate in patients without adequate treatment, can cause serious complications, such as perforation of the surrounding tissue and organs. *Parvimonas micra* is a very rare cause of purulent pericarditis.

**Case summary**

A 70-year-old male patient presented to our emergency room with chest pain of 10 days’ duration. An electrocardiogram showed ST-segment elevation and PR-segment depression on multiple leads. A transthoracic echocardiogram showed normal left ventricular function and a large amount of pericardial effusion. Acute pericarditis was diagnosed, and anti-inflammatory drug therapy was initiated. Due to the lack of improvement in the symptoms, pericardiocentesis was performed on Day 8 and revealed about 800 cc of the bloody fluid. *Parvimonas micra* was detected in a culture of the pericardial effusion and blood. Although intravenous antibiotic therapy was initiated for purulent pericarditis, his fever persisted. Computed tomography of the chest performed on Day 14 showed an abscess cavity in the pericardial space around the right atrium (RA). Furthermore, transoesophageal echocardiography revealed vegetation in the RA. Emergency surgery confirmed the presence of vegetation and minor perforation of the RA with communication to the abscess cavity. After surgical therapy, the patient clinically improved and was discharged on Day 51.

**Discussion**

In cases of acute pericarditis, purulent pericarditis should be considered if clinical improvement is not observed after initial treatment with anti-inflammatory drugs. Once the diagnosis of purulent pericarditis is made, aggressive source control is necessary for improved clinical outcomes.

**Keywords**

Purulent pericarditis • Infective endocarditis • Intracardiac perforation • *Parvimonas micra* • Case report

**Learning points**

- Purulent pericarditis is rare and difficult to diagnose.
- Purulent pericarditis can cause serious complications, such as intracardiac perforation and infective endocarditis.
- Aggressive source control, such as pericardiocentesis and surgical drainage, should be performed immediately to improve clinical outcomes.
Introduction

Purulent pericarditis reportedly accounts for only 0.7% of acute pericarditis cases.1 If appropriate treatment is not performed, the disease can progress very rapidly, and inflammation is likely to spread to surrounding tissues and organs. *Staphylococcus aureus* is the most commonly detected microorganism in purulent pericarditis while *Parvimonas micra* is very rarely implicated. The present report described a case of purulent pericarditis-induced intracardiac perforation and infective endocarditis due to *P. micra*.

Timeline

| Day 0 | The patient presented with chest pain, and acute pericarditis was diagnosed. |
| Day 7 | The patient had a fever, and a blood culture was performed. |
| Day 8 | Pericardiocentesis was performed and revealed bloody fluid. |
| Day 11 | *Parvimonas micra* was detected in the blood and pericardial effusion cultures. |
| Day 14 | Chest computed tomography (CT) revealed an abscess cavity in the pericardial space around the RA and a trabecular shadow suggesting vegetation in the right atrium (RA). |
| Day 15 | Transoesophageal echocardiography revealed vegetation in the RA, and emergency surgery was performed. |
| Day 51 | The patient was discharged after complete recovery. |
| 4 months post-operatively | Chest CT and transthoracic echocardiogram denied any recurrence of the abscess cavity and vegetation. |
| 6 months post-operatively | The patient had no recurrence of infection or signs of constrictive pericarditis. |

Case presentation

A 70-year-old male patient with no remarkable medical history presented to our emergency department with chest pain of 10 days’ duration. His initial vital signs showed sinus tachycardia (100 to 110 b.p.m.), a blood pressure of 134/85 mmHg, and a body temperature of 36.7°C. No heart murmur, no crackle and no rub detected by physical examination during first day of admission. An electrocardiogram (ECG) showed a sinus rhythm with ST-segment elevation and PR-segment depression on multiple leads (Figure 1). A transthoracic echocardiogram (TTE) showed normal left ventricular function and pericardial effusion without vegetations. The amount of pericardial effusion was large (>500 mL). Nevertheless, no compression of the right side of the heart due to pericardial effusion was observed (Figure 2). Laboratory data showed elevated inflammation markers [white blood cell count 13,600/μL; neutrophils 87% (upper limit of normal <9000/μL); C-reactive protein, 16.1 mg/dL (normal range ≤0.3 mg/dL)], and normal cardiac enzyme levels [creatinine kinase (upper limit of normal <150 U/L); high-sensitivity troponin T (normal range ≤0.014 ng/mL)]. Chest X-ray showed mild cardiomegaly. The patient’s symptoms and the TTE and ECG findings led to a diagnosis of viral acute pericarditis, and colchicine and loxoprofen were initiated. His fever worsened on Day 7, and a blood culture was performed. Pericardiocentesis was performed on Day 8, draining 800 cc of bloody fluid, which on analysis showed 15,000 white blood cells (62% polymorphonuclear cells, 38% mononuclear cells), total protein 6.1 g/dL, lactate dehydrogenase 998 U/L, and glucose 20 mg/dL. Cytology, histology (cell block), and mycobacterial culture showed no evidence of malignant pericardial effusion and tuberculous pericarditis. On Day 11, the blood and pericardial fluid cultures were positive for *P. micra*, and purulent pericarditis was diagnosed. A dental examination revealed decayed teeth, and intravenous antibiotic therapy (ampicillin-sulbactam 3 g every 6 h) and dental treatment were initiated. Although pericardial drainage was performed, the spiking fever persisted. Chest computed tomography (CT) performed on Day 14 showed an abscess cavity in the pericardial space around the right atrium (RA) and a trabecular shadow suggesting vegetation in the RA (Figure 3). Transoesophageal echocardiography (TOE) on the following day revealed vegetation in the RA, and emergency surgery was performed via a median sternotomy. The surgical findings...
demonstrated frank pus in the abscess cavity and on the RA surface. After a pericardiotomy was performed and the abscess cavity communicating with the RA was removed, minor bleeding from the RA was observed. The surface of the RA was dented, very fragile, and easily penetrated. A cardiopulmonary bypass was therefore established, and large vegetation was observed in the RA (Figure 5). The vegetation, infected RA wall, and trabeculae carneae were removed, and the tricuspid valve and right ventricle were inspected carefully but showed no vegetations. The culture of the pus in the abscess cavity detected *P. micra*. Post-operatively, the patient improved clinically and was given intravenous ampicillin-sulbactam 3 g every 6 h for 4 weeks. Coronary CT angiography performed due to transient changes in ST-segment on the ECG monitor during the surgery showed no stenotic lesions. During follow-up, chest CT and TTE showed no recurrence of the abscess cavity or vegetations at 4 months post-operatively (Video 3), and the patient experienced no recurrence of infection or signs of constrictive pericarditis at 6 months post-operatively.

**Discussion**

Purulent pericarditis is a rare, life-threatening infection of the pericardial space. Although fever is the most frequent presenting sign, our patient did not have fever at admission. As a result, no blood culture was performed on admission, and diagnosis and appropriate antimicrobial therapy were therefore delayed. Although TOE is strongly recommended in the guidelines for diagnosing infective endocarditis (IE), in the present case the procedure was delayed due to the absence of fever and murmur on admission. Because a pericardial fluid culture is important for early diagnosis and exclusion of purulent pericarditis, early pericardiocentesis should be performed to avoid delaying the diagnosis regardless of whether a fever is present.

The patient’s tooth decay was found to be the cause of the bacteremia and purulent pericarditis via haematogenous dissemination as...
there was no other source of infection. Purulent pericarditis development is most frequently associated with an extension of an infection within the thorax and mediastinum or direct inoculation via trauma. A few cases of purulent pericarditis due to haematogeneous dissemination of pathogens without another primary source of infection have been reported.

Purulent pericarditis is rarely caused by *P. micra*, an anaerobic Gram-positive coccus indigenous to the oral cavity and gastrointestinal tract and often associated with periodontal disease, sinusitis, pulmonary suppuration, and deep tissue infections, such as spinal inflammation. In general, an infection due to anaerobic Gram-positive cocci progresses slowly. Although standard antimicrobial therapy for *P. micra* infection has yet to be established, the organism is generally sensitive to beta-lactam antimicrobials. *Staphylococcus aureus* is the most commonly detected microorganism in purulent pericarditis. Patients with a *S. aureus* infection can experience a rapid development of a broad array of complications. It is likely that the disease developed slowly in our patient because the pathogenic bacterium was *P. micra*. The slow development made diagnosis difficult until the disease worsened sufficiently to cause an intracardiac perforation.

In the present patient, the purulent pericarditis led to an intracardiac perforation and secondary infective endocarditis. At admission, he already had acute pericarditis with signs of intense inflammation, but TTE showed no vegetation. A TTE performed by a sonographer on Day 1 likewise revealed no vegetation. Isolated, right-sided IE accounts for approximately only 5–10% of all IE cases, and there are few case reports of IE with vegetation only at the RA. The major risk factors specific to right-sided IE are intravenous drug use, the presence of an implantable electronic cardiac device or other intravascular device, a prosthetic valve, and congenital heart disease in the right side of the heart, none of which were present in our patient. Furthermore, the vegetation was attached to the abscess cavity, and it was thought that the inflammation from the purulent pericarditis and the abscess cavity spread to the RA, causing perforation and IE. To the best of our knowledge, the present study is the first to report purulent pericarditis due to *P. micra* causing intracardiac perforation.

Figure 5 Intraoperative photos. (A) An abscess cavity was observed around the right side of the heart (white arrows). (B) Pus was removed from the abscess cavity. (C) After abscess cavity removal, subsidence in the right atrium and minor bleeding were observed (yellow arrow). (D) The vegetation (22 mm × 12 mm × 2 mm) removed from the right atrium. Ao, ascending aorta; RA, right atrium; RV, right ventricle.
The standard treatment for purulent pericarditis consists of antimicrobial therapy and source control. The overall survival rate is about 30% with antimicrobial therapy alone and 50% when antimicrobial therapy is combined with early surgical drainage. In the present case, due to the presence of the abscess cavity and vegetation, source control only by pericardiocentesis was inadequate for complete improvement.

**Conclusion**

Because purulent pericarditis can cause serious complications, optimal antimicrobial therapy following detection of the pathogenic organism and immediate source control are indispensable for a favourable clinical outcome.

**Lead author biography**

Dr Hiroaki Morinaga was born in Tokyo, Japan in 1984. He received his MD from Juntendo University School of Medicine (Tokyo, Japan) in 2009 and finished his residency (2009–11) and fellowship in Internal medicine and Cardiology in 2016 in Japan. His research interests are mainly focused on echocardiography and heart failure.

**Supplementary material**

Supplementary material is available at European Heart Journal - Case Reports online.

**Slide sets:** A fully edited slide set detailing this case and suitable for local presentation is available online as Supplementary data.

**Consent:** The authors confirm that written consent for the submission and publication of this case report, including the images and associated text, was obtained from the patient in line with the COPE guidelines.

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