

resistentes mediante tinción de Zhiel-Nielssen. EL cultivo en el medio LwJ no pudo ser interpretado por encontrarse contaminado. La identificación por MALDI-TOF MS (Bruker Daltonics Inc., Alemania) con un score de 1,909 fue de *M. heckeshornense*. La cepa fue enviada al Centro Nacional de Micobacterias (Instituto Carlos III, Majadahonda, Madrid) para confirmar su identificación por métodos moleculares y la realización de estudios de sensibilidad antibiótica por el método de las proporciones.

La identificación se realizó mediante PRA del gen *hsp65* y digestión con las enzimas BstEII y HaeIII. El patrón que se obtuvo fue de 3 bandas 235/120/100 tras la digestión con la primera enzima y otras 3 bandas tras la digestión con HaeIII 160/105/60. Esta identificación presuntiva se confirmó mediante secuenciación del gen 16S ARNr. La cepa con la que presentó 100% de homología en el genbank fue con la cepa *M. heckeshornense* S369 (NR_028759), comparando 1360pb. El estudio de sensibilidad antibiótica se llevó a cabo mediante CMI en medio sólido; la cepa fue sensible a ciclo-serina, etionamida, rifampicina, capreomicina, estreptomina y kanamicina, y resistente a etambutol, isoniacida, PAS, pirazinamida, TCH y tiosemicarbazona.

No es habitual el aislamiento de micobacterias no tuberculosas en líquido sinovial, presentándose preferentemente en pacientes inmunodeprimidos. En nuestro caso, al igual que en el trabajo de Yokohama et al.⁵, se realizó la identificación por MALDI-TOF, por lo que se puede plantear su utilidad como una herramienta rápida, coste-efectiva y precisa para la identificación de *M. heckeshornense*. Presentamos un quinto caso de afectación osteoarticular añadido a un caso de tenosinovitis² y 3 de espondilodiscitis^{2,3}. Predomina el sexo masculino (4 casos), la existencia de algún factor de riesgo (2 infección por VIH y 2 tratamiento inmunosupresor) y en todos se consiguió la curación. La asociación de fármacos biológicos utilizados en la artritis reumatoide y el mayor riesgo de padecer infecciones graves presentan evidencias controvertidas. Según un metaanálisis publicado en *The Lancet*⁶, los medicamentos biológicos mostraron un incremento significativo del riesgo de sufrir infecciones graves a dosis habituales en comparación con los fármacos antirreumáticos modificadores de la enfermedad (FAME), presentando la combinación de fármacos biológicos el riesgo más alto.

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Parvimonas micra infective endocarditis



Endocarditis infecciosa por Parvimonas micra

Parvimonas micra, formerly *Peptostreptococcus micros* and *Micromonas micros*, are anaerobic, gram-positive cocci that belong to the abdominal, oropharyngeal and genitourinary flora, commonly related with periodontal diseases.¹ Its pathogenicity has been also documented in disseminated diseases such as septic arthritis, spondylodiscitis or abscesses in different organs.^{2–4} In 2015 the first case of endocarditis was described, in a 71-years-old male patient with valve abscess.¹ A literature review showed two cases of infectious endocarditis due to *P. micra*^{1,5} and three cases related to their ancestor *P. micros*.^{6–8}

We present a case of infectious endocarditis in a woman with a pacemaker and prosthetic mitral valve.

A 63-year-old female presented at the emergency department with a history of one month of persistent fever at dawn and noon, with chills, diaphoresis, asthenia and weight loss (5 kg). She denied dental procedures, recent interventions or invasive diagnostic techniques. Oral evaluation was made with no signs of periodontal disease other than edentulism. There were no respiratory, gastrointestinal, musculoskeletal or urinary symptoms and no cutaneous manifestation were found.

She had frequent follow-ups with neurosurgery and neurology for a right cerebral subependymoma and a mid-cerebral artery cardioembolic ictus performed eleven months prior to this episode. She had had a mechanical prosthetic mitral valve replacement in 1990. She had frequent checkups with urology for kidney angiomyolipomas.

On examination, temperature was normal, blood pressure was 130/62 mmHg, pulse of 70 beats per minute, respiratory rate of 18 breaths per minute, and oxygen saturation 100%. Auscultation was clear with no remarkable finding except for a click in the mitral valve. No lymphadenopathies were noted, as well as no edemas, with low muscle mass. The remainder examination was normal.

The patient was hospitalized to study a fever of unknown origin, and an echocardiography and positron emission tomography/computed tomography (PET/CT) were done. The PET/CT showed metabolic signs of infection in mitral prosthetic and periprosthetic tissue with moderate increased tracer uptake at the right carotid artery, aortic arch, thoracic aorta and pulmonary arteries compatible with active vasculitis on the large vessels. No pacemaker or extracardiac deposits were found.

A Transesophageal echocardiography (TEE) showed a vegetation on the medial side of the prosthetic ring of about 8 mm (Fig. 1).

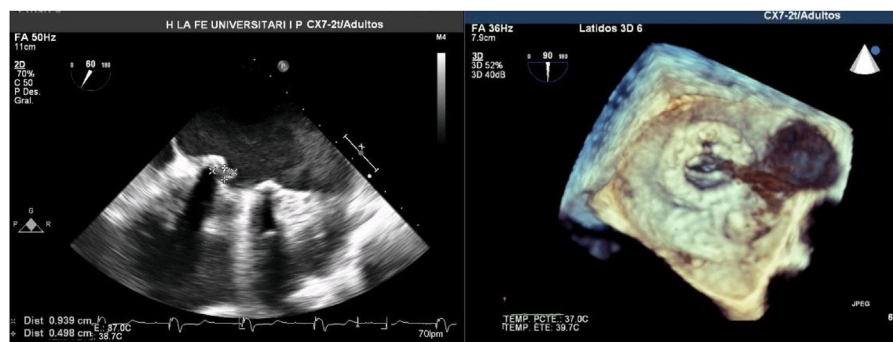


Fig. 1. Transesophageal echocardiography, shows vegetation on the medial side of the prosthetic ring of about 8 mm.

Cardiac and aortic CT discarded the possibility of valvular abscess or aortitis.

During the first 5 days of hospitalization 3 blood cultures were taken and incubated in BACT/ALERT® VIRTUO® (bioMérieux). After 43.94 h of average incubation (44.88 h – 42.82 h – 44.13 h) in the anaerobic media growth of gram-positive cocci identified by MALDI-TOF® as *P. micra*. The antibiotic susceptibility was done using Epsilon test in Columbia blood agar supplemented with 5% lamb defibrinated blood, in anaerobic atmosphere, obtaining the following MIC (mg/L) profile: penicillin 0.016, ampicillin 0.032, amoxicillin-clavulanic 0.047, ceftriaxone 0.125, imipenem 0.012, clindamycin 0.094, vancomycin 0.38, levofloxacin 0.19, rifampicin 0.002.

Empiric treatment was started with ceftriaxone and gentamycin and later modified to penicillin and clindamycin after culture isolation. The first negative blood culture was 4 days after initial treatment. TEE showed vegetation stability without complications, so it was decided to manage the case conservatively, with outpatient orally treatment with ceftriaxone and rifampicin for 6 more weeks. The evolution was favorable with negative control of blood cultures up to one year after the episode. After a year follow up by the Rheumatology Department the conclusion was that, findings on the PET/CT scan were due to the inflammatory response to the infection, which diminished on subsequent scans.

Endocarditis is mainly caused by species of the genus *Streptococcus* and *Staphylococcus*, comprising around two-thirds of the total of infectious endocarditis, while anaerobic organisms are only isolated in 2–16% of the cases.⁹ Predominant anaerobic species are *Cutibacterium acnes*, *Bacteroides fragilis* and *Clostridium* spp.¹⁰ Endocarditis due to *Peptostreptococcus* spp. are infrequent¹⁰ with *P. magnus* being the most common species.⁹ If we focus on *P. micra*, until now five cases have been described in the literature,^{1,5–8} only two of them encompassed within this terminology.^{1,5}

Conflict of interest

None.

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