Case study

Disseminated *Mycobacterium abscessus* infection in a peritoneal dialysis patient

Vincent H.J.F. Moorena,b,⁎, Michiel W.P. Bleekea,a, Jakko van Ingena, Mirjam H.A. Hermansa, Peter C. Wevere

⁎ Corresponding author at: Department of Internal Medicine, Radboudumc, P.O. Box 9101, Route 463, 6500 HB, Nijmegen, The Netherlands.
E-mail address: vincent.mooren@radboudumc.nl (V.H.J.F. Mooren).

A 73-year-old Caucasian woman with end-stage renal disease who had been treated with continuous ambulatory peritoneal dialysis (CAPD) for 12 months presented on routine evaluation with local tenderness and erythema at the exit-site of the PD catheter. Her medical history was significant for recurrent episodes of pyelonephritis due to nephrolithiasis, ischaemic heart disease, and two caesarean sections.

On physical examination there was no purulent discharge from the exit-site. The peritoneal fluid was clear and the patient had no fever. Patient history and physical examination revealed no further abnormalities. According to protocol she was treated as an outpatient for an exit-site infection (ESI) with oral flucloxacillin. After ten days of treatment there was no clinical improvement. As the exit-site culture seemed to grow *Corynebacterium* species, therapy was switched to oral amoxicillin-clavulanic acid. On day 15, it was reported that the isolate was resistant to all tested antimicrobial agents including vancomycin. Since there was still no sign of clinical improvement the infected catheter was surgically removed on day 24 and in the same session a new catheter was inserted in a different abdominal region. She was switched to haemodialysis (HD) via a tunnelled catheter. On day 29, 16S ribosomal RNA polymerase chain reaction (16S rRNA PCR) identification performed on the presumed *Corynebacterium* isolate revealed 100% homology with *Mycobacterium chelonae-abscessus* complex. Further species identification by the National Institute for Public Health and the Environment (RIVM) revealed *M. abscessus*. As the PD catheter had been replaced, no further therapeutic actions were undertaken.

Ten days after surgical PD catheter-replacement the patient presented with fever and abdominal complaints. This time the new PD exit-site was inflamed. Laboratory studies showed an elevated C-reactive protein level: 130 mg/L (ref. < 6 mg/L) and white blood cell count: 12.7 × 10⁹/L (ref. 4.0–10.0 × 10⁹/L). Ultrasonography showed no evidence for abdominal or tunnel abscesses. Fluclxacinil was commenced intravenously and the patient was admitted to the nephrology ward. Bacterial cultures taken from the exit-site, peripheral blood and jugular venous catheter remained negative. Despite prolonged antimicrobial therapy the patient had recurrent fever and chills, especially during HD sessions. While initial blood cultures obtained on admission and on day 5 remained negative, blood cultures obtained on day 11...
after admission ultimately revealed growth of *M. abscessus*. Immediate removal of both the new PD catheter and the HD catheter was ordered. Flucl oxidacin was discontinued and switched to tigecycline and clari thor mycin. Unfortunately, bilateral thrombosis of the jugular veins occurred, which caused major access problems for adequate dialysis. The patient was, therefore, transferred to a nearby academic hospital. Broth microdilution susceptibility testing of the bloodstream isolate revealed amikacin susceptibility, intermediate imipenem susceptibility (minimum inhibitory concentration [MIC] 8 mg/L), inducible macrolide resistance and a tigecycline MIC of 0.25 mg/L [1]. Based hereupon, clari thor mycin was switched to imipenem. Additional imaging studies showed no source of infection explaining the bacteraemia. Follow up blood cultures remained negative. The patient unexpectedly died because of ventricular fibrillation directly after insertion of a central venous catheter in the left subclavian vein.

### Discussion

Among the rapidly growing nontuberculous mycobacteria (RGM), *Mycobacterium fortuitum, M. chelonae* and *M. abscessus* are considered clinically most relevant. They are ubiquitous in the environment, including water, soil and dust, and survive nutritional deprivation and extreme temperatures. The most common clinical manifestations comprise skin and soft tissue infections, catheter-related infections and pulmonary infections. RGM are infrequent causative agents of ESIs and PD-peritonitis; *M. abscessus* infections are particularly rare in this context with only eight previous reports comprising 23 cases (14 peritonitis; 9 ESIs) (Table 1) [2–9]. The vast majority of cases are described in Asian countries. To the best of our knowledge, this is the second case in literature of a *M. abscessus* bloodstream infection as a complication of PD-related infection [3].

Predisposing factors for development of PD-related infections with RGM include additional immunosuppressive conditions, recurrent episodes of ESIs or peritonitis with multiple courses of broad-spectrum antimicrobials, and underdialysis [5]. In addition, the use of topical gentamicin cream is postulated as a risk factor in two reports [2,10]. None of these applied to our patient.

The microbiological similarities of RGM and Corynebacterium species impose potential for misidentification in the laboratory. This phenomenon, which also happened in our case, has been previously reported [11,12]. In order to diagnose RGM infection, performance of auramine or Ziehl-Neelsen staining and mycobacterial culture are essential. An earlier auramine or Ziehl-Neelsen staining might in our case, have led to earlier adequate antimicrobial therapy and possibly a better clinical outcome.

*M. abscessus* is intrinsically resistant to most antimicrobials [1]. Treatment consists of combination therapy, usually with cefoxitin or imipenem, amikacin and clarithromycin [1]. The minimum duration and optimal regimen for treatment for PD-related RGM infection are not known, but a prolonged course of antimicrobial treatment is recommended, especially when the plan is to salvage the PD catheter [13]. There is no guideline recommending catheter removal following diagnosis of PD-related RGM infection. High rates of catheter loss and mortality are reported in PD-related RGM disease [3]. Extensive comorbidities, as in our case, contribute to fatal outcomes [13].

### Conclusion

We present a case with a disseminated PD-related *M. abscessus* infection. To prevent delay in optimal antimicrobial treatment, RGM infections should be considered when initial therapy does not lead to clinical improvement.

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### Conflicts of interest

None.

### References


