Hindawi Case Reports in Infectious Diseases Volume 2019, Article ID 3715404, 4 pages https://doi.org/10.1155/2019/3715404

Case Report

A Case of Serratia marcescens Endocarditis in a Nonintravenous Drug-Using Male Patient and Review of Literature

Achilleas Nikolakopoulos (1), ¹ Nikolaos Koutsogiannis, ² Panagiota Xaplanteri (1), ³ Charalambos Gogos, ^{1,4} Fevronia Kolonitsiou, ³ and Alexandra Lekkou ^{1,4}

Correspondence should be addressed to Achilleas Nikolakopoulos; achylles@otenet.gr

Received 14 February 2019; Accepted 25 March 2019; Published 20 June 2019

Academic Editor: Gloria Taliani

Copyright © 2019 Achilleas Nikolakopoulos et al. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Introduction. Serratia marcescens is a rare cause of infective endocarditis and has almost exclusively been associated with intravenous drug use and hospital-acquired infections. Here, we present a case of infective endocarditis caused by Serratia marcescens in an otherwise healthy, nonintravenous drug-using male patient. Case Report. A 41-year-old man presented with hypertension and hemoptysis. Blood cultures were obtained that showed bacteremia by Serratia marcescens. An echocardiogram was carried out that revealed severe mitral regurgitation accompanying ruptured mitral chordae tendineae. The patient received the appropriate antibiotic treatment, without further surgical intervention. Discussion. Hospital-acquired infections by Serratia species are a common problem in medical practice and have been attributed to specialized interventional procedures. Taking into consideration the patient's immunocompetence and lack of intravenous drug use, it is possible that bacteremia could be attributed to a medical procedure. Moreover, in contrast to most cases described in the literature, no surgery was performed.

1. Introduction

Serratia marcescens, an aerobic (facultative anaerobe), oxidase-negative, non-lactose-fermenting Gram-negative bacillus, is a rare cause of infective endocarditis and has almost exclusively been associated with intravenous drug use and hospital-acquired infections. Here, we present a case of infective endocarditis caused by Serratia marcescens in an otherwise healthy, nonintravenous drug-using male patient.

2. Case Report

The patient is a 41-year-old man, without any history of disease or medication. He is slightly obese and a current smoker (>50 pack-years) and reports moderate alcohol consumption. He was admitted in a regional hospital with sudden onset of hemoptysis and headache. There, the patient

was diagnosed with hypertension (systolic blood pressure: 240 mmHg, diastolic blood pressure: 110 mmHg), and hemoptysis was confirmed. A brain and chest computed tomography (CT) scan was performed that showed no findings from the brain and alveoral hemorrhage, respectively (Figure 1). In addition, the patient underwent a formal transthoracic echocardiogram (TTE), which revealed mild to moderate mitral valve regurgitation and prolapse. Blood tests were within normal rates. The patient was subsequently transported to the Pulmonary Department of our hospital for further investigation and treatment.

Upon his admission, the patient's temperature was 37.6° C, heart rate was 98 bpm, respiratory rate was 22 breaths per minute, blood pressure was 177/89 mmHg, and an arterial blood gas showed mild hypoxemia in room air (pH = 7.47; pCO₂ = 34 mmHg; pO₂ = 68 mmHg; HCO₃ = 18 mmol/L; O₂ saturation = 93%). Blood tests were normal again, except for

¹Department of Internal Medicine, University Hospital of Patras, 26504 Patras, Greece

²Department of Cardiology, University Hospital of Patras, 26504 Patras, Greece

³Department of Microbiology, University Hospital of Patras, 26504 Patras, Greece

⁴Department of Infectious Diseases, University Hospital of Patras, 26504 Patras, Greece

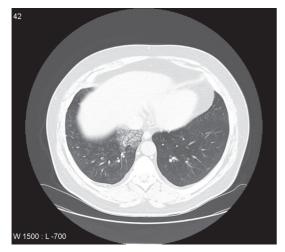


FIGURE 1: Chest CT scan showing alveoral hemorrhage and consolidation in the right middle lobe.

elevated uric acid levels. Blood cultures were collected, and he was initially started on empiric broad-spectrum antibiotic treatment of piperacillin-tazobactam and oxygen supplementation due to hypoxemia. Chest CT was performed once again, which identified findings compatible with alveoral hemorrhage and consolidation in the right middle lobe. An abdominal CT scan resulted in findings such as slightly enlarged liver without focal lesions and bilateral, well-defined adrenal lesions, possibly adenomas. The patient refused to undergo a new brain CT or MRI scan. Pulmonary function tests showed obstructive lung disease, and bronchoscopy revealed blood clots in the right middle lobe. His HIV-screening test was negative, as was Mantoux tuberculin skin test (TST).

Two blood cultures revealed bacteremia by *Serratia marcescens*, and this finding was confirmed with one more blood culture, taken 5 days later. The *Serratia marcescens* isolate was susceptible to cefepime, meropenem, gentamicin, sulfamethoxazole-trimethoprim, and ciprofloxacin. An urgent transesophageal echocardiogram (TEE) was performed which demonstrated severe mitral regurgitation accompanying ruptured mitral chordae tendineae (Figures 2 and 3). The patient was transported to the Infectious Diseases Department, and antibiotic treatment was subsequently changed to meropenem 2 g tid, ciprofloxacin 400 mg bid, and gentamicin 80 mg tid intravenously, according to antibiogram. Of notice, he reported a diagnostic arthrocentesis in his left knee joint a few weeks ago.

Three days later, the patient was afebrile, hypoxemia and hemoptysis were resolved, blood tests remained within normal limits, including inflammation markers such as C-reactive protein (CRP) and WBC, and multiple subsequent blood culture results came back negative. A new transesophageal echocardiogram was performed 20 days after the previous one, without further deterioration.

The patient, after having stayed at our hospital for 4 weeks, was transported back to the regional hospital in order to continue his intravenous antibiotic treatment with meropenem and ciprofloxacin for a total of 6 weeks. He had already completed 2 weeks of gentamicin treatment.



FIGURE 2: Echocardiogram showing ruptured mitral chordae tendineae.



FIGURE 3: Echocardiogram showing severe mitral regurgitation.

Both cardiologist and cardiac surgeons suggested surgery for valve repair or replacement, but the patient refused. Therefore, close surveillance with repeat echocardiograms every 3–6 months was recommended. Moreover, pulmonary surveillance with a repeat bronchoscopy was advised after completion of therapy. After 24 months, he is currently asymptomatic, without limitations in physical activity.

3. Discussion

Community-acquired infective endocarditis due to *Serratia* species is almost exclusively seen in intravenous drug users. Most commonly, infection implicates the aortic and mitral valves, frequently occurring on previously undamaged cardiac valves. Septic embolization is also common. Antibiotic treatment alone is usually insufficient, and surgical replacement of the involved valve is needed [1–7].

Hospital-acquired infections by *Serratia* species are a common problem in medical practice and have been attributed to specialized interventional procedures such as surgery, bronchoscopy, and foreign body placement, as well as routine hospitalization practices, through contaminated vials and other materials [8–11]. Its ability to survive and grow under extreme conditions, including in disinfectants,

antiseptics, and double-distilled water, probably facilitates nosocomial infections. Contamination can occur whenever between medication and material manufacturing by pharmaceutical manufacturers or compounding pharmacies and administration to the patient. Use of single-use medication vials on multiple patients, use of a common syringe for multiple medications/patients, and use of multidose vials have all been implicated in outbreaks of nosocomial infections [12-14]. In most cases, lack of adequate sterile conditions and hand hygiene, in particular, is the common cause. Patients most at risk are those in intensive care units who are subjected to medical devices, especially central venous catheters, and treated with broad-spectrum antimicrobial drugs. Outbreaks in neonatal wards have frequently been reported, as well [15, 16]. Contamination of materials and devices, such as bronchoscopes, could be attributed to inadequate or erroneous cleaning/sterilization procedures, or even manufacturing defects [17, 18].

Serratia marcescens possesses a number of virulence factors, such as fimbriae or fimbria-like adhesins, which mediate surface attachment and biofilm formation and likely increase the opportunities of this organism to infect humans through attachment to abiotic surfaces [19].

Infections caused by Serratia marcescens may be difficult to treat because of resistance to a variety of antibiotics, including ampicillin and first-, second-, and third-generation cephalosporins. Moreover, imipenem-resistant strains exhibiting β -lactamase production have also been reported, that may become prevalent in the near future, further hindering treatment. Resistance to β -lactams has been described to be mediated through two distinct mechanisms: first, high-level production of chromosomal AmpC cephalosporinases combined with substantially decreased outermembrane permeability, and second, synthesis of β -lactamases able to hydrolyse carbapenems. Aminoglycosides have good activity against Serratia marcescens, but resistant strains have also been reported. Alterations in the cell envelope prevent uptake of the drug, and also the drug itself can be modified by inactivating enzymes that adenylate, acetylate, or phosphorylate the hydroxyl or amino groups of aminoglycosides. This type of resistance is commonly mediated by plasmids and is often transferable [20].

Taking into consideration the patient's immunocompetence and lack of intravenous drug use, it is possible that bacteremia could be attributed to a medical procedure. One possible scenario is that the infection occurred during the arthrocentesis procedure he was subjected to, a few weeks before his hospitalization. This presumes that the first echocardiogram performed 2 weeks before his arrival, at the regional hospital, misidentified ruptured mitral chordae tendineae for mitral valve prolapse. Another scenario is that the infection occurred during his current hospitalization, where bacteremia resulted in infective endocarditis and ruptured mitral chordae tendineae of an already faulty mitral valve with prolapse. Although Serratia species have been associated with bronchoscopes and the procedure of bronchoscopy, in this case, bronchoscopy was preceded by the blood culture test, so it is unlikely that this could be the route of entry.

Finally, it is of notice that, in contrast to most cases described in the literature, no surgery was performed. Endocarditis by *Serratia* species has a poor prognosis, and surgical management is often required. In this case, the patient received only antibiotic treatment and survived, while remaining asymptomatic.

Conflicts of Interest

The authors declare that they have no conflicts of interest.

References

- J. Mills and D. Drew, "Serratia marcescens endocarditis: a regional illness associated with intravenous drug abuse," Annals of Internal Medicine, vol. 84, no. 1, pp. 29–35, 1976.
- [2] B. E. Reisberg, "Infective endocarditis in the narcotic addict," Progress in Cardiovascular Diseases, vol. 22, no. 3, pp. 193–204, 1979.
- [3] F. E. Berkowitz, P. Colsen, and K. Raw, "Serratia marcescens endocarditis treated with ceftazidime. A case report," South African Medical Journal, vol. 64, no. 3, pp. 105-106, 1983.
- [4] Y. Hadano, T. Kamiya, and N. Uenishi, "A fatal case of infective endocarditis caused by an unusual suspect," *Internal Medicine*, vol. 51, no. 11, pp. 1425–1428, 2012.
- [5] V. Phadke and J. Jacob, "Marvelous but morbid: infective endocarditis due to *Serratia marcescens*," *Infectious Diseases in Clinical Practice*, vol. 24, no. 3, pp. 143–150, 2016.
- [6] C. G. Meyer, T. P. Vacek, A. Bansal, R. Gurujal, and A. Parikh, "Dynamic course of *Serratia marcescens* pulmonic valve endocarditis resulting in submassive PE and valve replacement," *Journal of Investigative Medicine High Impact Case Reports*, vol. 6, article 2324709618759128, 2018.
- [7] H.-M. Yeung, B. Chavarria, and D. Shahsavari, "A complicated case of *Serratia marcescens* infective endocarditis in the era of the current opioid epidemic," *Case Reports in Infectious Diseases*, vol. 2018, Article ID 5903589, 5 pages, 2018.
- [8] A. Khanna, M. Khanna, and A. Aggarwal, "Serratia marcescens—a rare opportunistic nosocomial pathogen and measures to limit its spread in hospitalized patients," Journal of Clinical and Diagnostic Research, vol. 7, no. 2, pp. 243–246, 2013.
- [9] M. Šiširak and M. Hukić, "An outbreak of multidrug-resistant Serratia marcescens: the importance of continuous monitoring of nosocomial infections," Acta Medica Academica, vol. 42, no. 1, pp. 25–31, 2013.
- [10] I. Wilhelmi, J. C. Bernaldo de Quirós, J. Romero-Vivas, J. Duarte, E. Rojo, and E. Bouza, "Epidemic outbreak of Serratia marcescens infection in a cardiac surgery unit," Journal of Clinical Microbiology, vol. 25, no. 7, pp. 1298–1300, 1987.
- [11] E. R. van der Vorm and C. Woldring-Zwaan, "Source, carriers, and management of a *Serratia marcescens* outbreak on a pulmonary unit," *Journal of Hospital Infection*, vol. 52, no. 4, pp. 263–267, 2002.
- [12] R. F. Chemaly, D. B. Rathod, M. K. Sikka et al., "Serratia marcescens bacteremia because of contaminated prefilled heparin and saline syringes: a multi-state report," American Journal of Infection Control, vol. 39, no. 6, pp. 521–524, 2011.
- [13] P. C. Chiang, T. L. Wu, A. J. Kuo et al., "Outbreak of Serratia marcescens postsurgical bloodstream infection due to contaminated intravenous pain control fluids," International Journal of Infectious Diseases, vol. 17, no. 9, pp. e718–e722, 2013.

- [14] A. Pan, L. Dolcetti, C. Barosi et al., "An outbreak of Serratia marcescens bloodstream infections associated with misuse of drug vials in a surgical ward," Infection Control & Hospital Epidemiology, vol. 27, no. 1, pp. 79–82, 2006.
- [15] J. M. Muñoz and A. E. Macías, "Nosocomial outbreak of *Serratia marcescens* in a neonatal intensive care unit," *Infection Control & Hospital Epidemiology*, vol. 24, no. 5, p. 312, 2003.
- [16] J.-H. Kim, W.-H. Choi, S.-W. Yun, S.-A. Chae, and B.-H. Yoo, "An outbreak of Serratia marcescens sepsis in a pediatric ward," Clinical Pediatrics, vol. 49, no. 10, pp. 1000–1002, 2010.
- [17] S. F. Webb and A. Vall-Spinosa, "Outbreak of Serratia marcescens associated with the flexible fiberbronchoscope," Chest, vol. 68, no. 5, pp. 703–708, 1975.
- [18] D. L. Kirschke, T. F. Jones, A. S. Craig et al., "Pseudomonas aeruginosa and Serratia marcescens contamination associated with a manufacturing defect in bronchoscopes," New England Journal of Medicine, vol. 348, no. 3, pp. 214–220, 2003.
- [19] R. M. Q. Shanks, N. A. Stella, E. J. Kalivoda et al., "A Serratia marcescens OxyR homolog mediates surface attachment and biofilm formation," Journal of Bacteriology, vol. 189, no. 20, pp. 7262–7272, 2007.
- [20] A. Hejazi and F. R. Falkiner, "Serratia marcescens," Journal of Medical Microbiology, vol. 46, no. 11, pp. 903–912, 1997.