A Case of Infective Endocarditis caused by *Abiotrophia defectiva* in Korea

Seohyun Park¹, Hea Won Ann¹², Jin Young Ahn¹², Nam Su Ku¹², Sang Hoon Han¹², Geu Ru Hong¹, Jun Young Choi¹², Young Goo Song¹², and June Myung Kim¹²

¹Department of Internal Medicine, and ²AIDS Research Institute, Yonsei University College of Medicine, Seoul, Korea

*Abiotrophia defectiva*, a nutritionally variant streptococci can cause bacteremia, brain abscess, septic arthritis and in rare cases, infective endocarditis, which accounts for 5-6% of all cases. *A. defectiva* is characteristically difficult to diagnose and the mortality, morbidity and complication rates are high. Here, we discuss a case of infective endocarditis caused by *A. defectiva*. A 62-year-old female had previously undergone prosthetic valve replacement 6 years prior to admission. She developed infective endocarditis after tooth extraction. Her endocarditis was successfully treated with antimicrobial therapy and mitral valve replacement surgery. This is the first case of infective endocarditis caused by *A. defectiva* reported in Korea. This case shows that *A. defectiva* could be considered as a causative organism of infective endocarditis in Korea.

Key Words: *Abiotrophia defectiva*, infective endocarditis

Introduction

*Abiotrophia defectiva* is part of the normal human microbiota, colonizing the oral, genitourinary, and intestinal tracts [1]. It is a rare, yet important, cause of infective endocarditis, and is estimated to cause approximately 5-6% of all cases of infective endocarditis, including being a major cause of blood culture-negative infective endocarditis [2]. It affects diseased valves in 90% of cases and it is implicated in embolic complications and valvular destruction, despite being sensitive to antibiotics [3]. Previous studies have shown mortality and relapse rates as high as 17% despite antibiotic treatment, and this makes accurate and quick identification important [4-6]. Herein we report the first case of infective endocarditis caused by *A. defectiva* in Korea.

Case Report

A 62-year-old female was admitted to the emergency department (ED) after two consecutive episodes of syncope. The patient was diagnosed with severe rheumatic mitral stenosis, and underwent a mitral valve replacement (MVR). Two months previously, the patient had undergone a simple ex-
traction of her #16 tooth due to secondary dental caries and had taken prophylactic antibiotics (amoxicillin 2,000 mg).

Upon admission to the ED the patient was alert and had a body temperature of 38.6°C, a pulse rate of 66 beats/min, and blood pressure of 149/51 mmHg. Upon physical examination, no cardiac murmur was auscultated and no other evidence, such as clubbed fingers, Janeway lesions or petechiae were found. Laboratory studies showed a white blood cell count of 14,260/mm$^3$ (neutrophil 79.6%), erythrocyte sediment rate of 92 mm/h and C-reactive protein of 111.1 mg/L. Chest X-ray revealed mild cardiomegaly and electrocardiography showed a newly developed complete atrioventricular block. Chest CT and abdominal-pelvic CT was performed to evaluate fever focus, which showed no significant finding.

A transthoracic echocardiogram (TTE) showed that the mechanical prosthetic mitral valve functioned well, and there was no visible vegetation, however, infective endocarditis could not be completely ruled out because of the patient’s previous MVR. Thus, empirical antibiotics (vancomycin 1 g intravenous q12hr, gentamicin 60 mg intravenous q8hr and rifampin 600 mg per oral q24hr) were administered, and a transesophageal echocardiogram (TEE) performed the day after admission showed mitral valve vegetation and mild transvalvular mitral regurgitation (Fig. 1), leading to a conclusive diagnosis of infective endocarditis according to Duke criteria (one major and three minor criteria).

Three sets of blood cultures taken on admission showed tiny, non-hemolytic colonies on a blood agar plate (ASAN Pharmaceutical, Hwaseong, Korea), and pleomorphic Gram-positive cocci from smear preparation of the blood agar plate (Fig. 2). On the third day after admission, A. defectiva was identified by MALDI-TOF-MS (Matrix-assisted laser desorption/ionization time-of-flight mass spectrometry, Bruker Daltonics Inc., MA, USA), not identified in VITEK 2 (bioMérieux, Marcy l’Etoile, France) systems. However, because the amount of bacteria was insufficient, cultures were sub-cultured on a medium containing vitamin B6 to assess antibiotic susceptibility.

Susceptibility to cefotaxime, penicillin G and vancomycin was tested by the E-test method; the isolate tested cefotaxime-sensitive (MIC 0.75 μg/mL), penicillin G-intermediate (MIC 1.0 μg/mL) and vancomycin-sensitive (MIC 0.38 μg/mL). An adjusted antibiotic regimen of vancomycin 1 g q12hr and gentamicin 60 mg q8hr was initiated, after which the patient’s fever subsided. Vancomycin Therapeutic drug monitoring was performed, and vancomycin dosage was adjusted to 400 mg q12hr.

On day 12 of admission, the patient became febrile once more and in the subsequent TEE the mitral valve vegetation on the medial side seemed to have resolved, but there was an increase in the size of the vegetation on the lateral annulus ($0.5 \times 0.3 \text{ cm} - 0.8 \times 0.3 \text{ cm}$) which indicated a perivalvular infection (Fig. 1). Based on these findings, a second MVR was performed on day 19 of admission.

Operation finding showed small vegetation in prosthetic MV and annulus, which was removed, and MVR was conducted using 23 mm ATS medical open pivot heart valve (ATS Medical, Inc., Minneapolis, MN, USA). Pathology finding showed myxoid degeneration, acute and chronic inflammation, fibrosis and calcification, and tissue culture showed no growth of organism (It was on the 19th day on antibiotics, and blood cul-
ture was negatively converted at this time).

On day 23 of admission, a follow-up TTE showed a well-functioning prosthetic mitral valve. The patient was treated with antibiotics for 5 weeks and was discharged on day 36. The patient was followed-up at an out-patient clinic for 3 months after discharge with no significant complications.

Discussion

Organisms of the genus *Abiotrophia* were first classified as nutritionally variant streptococci (NVS) in 1961 [6]. Bouvet et al. identified two species of NVS; *Streptococcus defectivus* and *Streptococcus adjacens* [7]. In 1995, Kawamura et al. identified *Abiotrophia* as a separate genus, based on 16S rRNA analyses [8]. NVS are part of the normal flora of the mouth and urogenital and intestinal tracts [1]. Recently, an increasing number of cases have been described of *A. defectiva* isolates recovered from invasive and non-invasive infections following dental procedures [9].

*Abiotrophia* spp., has fastidious culturing and the unspecific colony morphology that they present on primary detection, such strains have caused major diagnostic difficulties. Thus, it has been supposed that many culture-negative endocarditis could have been caused by these species, which could have lead to an underestimation as pathogens of infective endocarditis [10]. In this case report, we identified *A. defectiva* by MALDI-TOF-MS. MALDI-TOF-MS is a new technology for routine identification of bacteria in clinical microbiology laboratories. Much of the work using MALDI-TOF-MS for microbial identification has focused on demonstrating that reproducible mass spectra can be obtained using intact cells and developing algorithms for interpretation and comparison of these spectra. By testing colonies, it takes only a few minutes to have a correct identification which makes not only possible to identify the microorganisms at the species levels but sometimes at the sub-species and strains levels, allowing the detection of epidemic lineages [11].

*A. defectiva* can cause serious infections such as bacteremia, osteomyelitis, brain abscess, pancreatic abscess, septic arthritis, crystalline keratopathy and in rare cases, infective endocarditis [3, 12]. Endocarditis caused by NVS is implicated in 5–6% of all streptococcal endocarditis cases, and <1% of all endocarditis cases are caused by *A. defectiva* [3]. However, *A. defectiva* has a higher affinity for the endocardium because of its ability secrete exopolysaccharide, enabling it to adhere to fibronectin in the extracellular matrix [2]. Of all the NVS-induced endocarditis patients, 90% suffered from heart disease, and in most cases, bacteremia associated with the pre-existing valvular heart disease led to the development of endocarditis with subacute prognoses [13].

Infective endocarditis caused by *A. defectiva* and other NVS had higher mortality, morbidity and complication rates than those caused by other viridans streptococci [14]. Most deaths were due to refractory congestive cardiac failure or major systemic emboli [15]. A mortality rate of up to 17% has been re-
ported, which is higher compared to endocarditis caused by viridians streptococci (0-12%) [4]. The relapse rate can also be up to 17% in some cases. Previous studies have shown a treatment failure rate as high as 41%, despite the use of appropriate antibiotics [5].

The antibiotic regimen to combat *A. defectiva* endocarditis includes penicillin or ampicillin, plus an aminoglycoside or vancomycin for cases of antimicrobial resistance, taken for 4-6 weeks [16]. Infective endocarditis due to *A. defectiva* progresses slowly, but despite its sensitivity to antimicrobials -50% of cases need surgical management. Surgical treatment combined with concurrent antimicrobial therapy result in better prognoses, and the main indications for surgery are persistent sepsis and vegetation, severe congestive heart failure and recurrent embolism [15, 17, 18].

In Korea, 2 cases of infective endocarditis caused by *S. adjacens* were first reported in 1996. Since then, a case of infective endocarditis by *Grunulicatella adjacens* was reported in 2010. All cases were successfully treated with antibiotics and surgical treatment [19, 20]. After classified as *A. defectiva*, this is the first report as the pathogen of infective endocarditis in Korea.

The case presented here is an example of the successful treatment of infective endocarditis caused by *A. defectiva* following a tooth extraction in a post-MVR patient, and represents the first reported case of infective endocarditis caused by *A. defectiva* in Korea. This case shows that *A. defectiva* could be considered as a causative organism of infective endocarditis in Korea.

**Conflicts of Interest**

No conflicts of interest.

**ORCID**

Seohyun Park  
http://orcid.org/0000-0002-6169-4064

Nam Su Ku  
http://orcid.org/0000-0002-9717-4327

**References**

1. Bouvet A. Human endocarditis due to nutritionally variant streptococci: *Streptococcus adjacens* and *Streptococcus defectivus*. Eur Heart J 1995;16 (Suppl B):24-7.

2. Ruoff KL. Nutritionally variant streptococci. Clin Microbiol Rev 1991;4:184-90.

3. Roberts RB, Krieger AG, Schiller NL, Gross KC. Viridans streptococcal endocarditis: the role of various species, including pyridoxal-dependent streptococci. Rev Infect Dis 1979;1:955-66.

4. Tuazon CU, Gill V, Gill F. Streptococcal endocarditis: single vs. combination antibiotic therapy and role of various species. Rev Infect Dis 1986;8:54-60.

5. Stein DS, Nelson KE. Endocarditis due to nutritionally deficient streptococci: therapeutic dilemma. Rev Infect Dis 1987;9:908-16.

6. Frenkel A, Hirsch W. Spontaneous development of L forms of streptococci requiring secretions of other bacteria or sulphydryl compounds for normal growth. Nature 1961;191:728-30.

7. Bouvet A, Grimont F, Grimont PA. Intraspecies variations in nutritionally variant streptococci: rRNA gene restriction patterns of *Streptococcus defectivus* and *Streptococcus adjacens*. Int J Syst Bacteriol 1991;41:483-6.

8. Kavamura Y, Hou XG, Sultana F, Liu S, Yamamoto H, Ezaki T. Transfer of *Streptococcus adjacens* and *Streptococcus defectivus* to Abiotrophia gen. nov. as *Abiotrophia adjacens* comb. nov. and *Abiotrophia defectiva* comb. nov., respectively. Int J Syst Bacteriol 1995;45:798-803.

9. Akkoyunlu Y, Iraz M, Kocaman G, Ceylan B, Aydin C, Aslan T. *Abiotrophia defectiva* endocarditis presenting with hemiplegia. Jundishapur J Microbiol 2013;6:e8907.

10. Brouqui P, Raoult D. Endocarditis due to rare and fastidious bacteria. Clin Microbiol Rev 2001;14:177-207.

11. Biswas S, Rolain JM. Use of MALDI-TOF mass spectrometry for identification of bacteria that are difficult to culture. J Microbiol Methods 2013;92:14-24.

12. Heath CH, Bowen SF, McCarthy JS, Dwyer B. Vertebral osteomyelitis and discitis associated with *Abiotrophia adjacens* (nutritionally variant streptococcus) infection. Aust N Z J Med 1998;28:563.

13. Okada Y, Kitada K, Takagaki M, Ito HO, Inoue M. Endocardiac infectivity and binding to extracellular matrix proteins of oral *Abiotrophia* species. FEMS Immunol Med Microbiol 2000;27:257-61.

14. Yang YS, Shang ST, Lin JC, Chiu CH, Chang FY. A ruptured cerebral mycotic aneurysm caused by *Abiotrophia defectiva* endocarditis. Am J Med Sci 2010;339:190-1.

15. Takayama R, Motoyasu M, Seko T, Kuroda K, Yamanaka T, Obe T, Yada T, Konishi T, Fujinaga K, Konodoh C, Mizutani T. A case of isolated tricuspid valve infective endocarditis caused by *Abiotrophia defectiva*. Int J Cardiol 2007;118:e3-5.

16. Baddour LM, Wilson WR, Bayer AS, Fowler VG Jr, Bolger AF, Levison ME, Ferrieri P, Gerber MA, Tani LY, Gewitz MH, Tong DC, Steckelberg JM, Baltimore RS, Shulman...
ST, Burns JC, Falace DA, Newburger JW, Pallasch TJ, Takahashi M, Taubert KA; Committee on Rheumatic Fever, Endocarditis, and Kawasaki Disease; Council on Cardiovascular Disease in the Young; Councils on Clinical Cardiology, Stroke, and Cardiovascular Surgery and Anesthesia; American Heart Association; Infectious Diseases Society of America. Infective endocarditis: diagnosis, antimicrobial therapy, and management of complications: a statement for healthcare professionals from the Committee on Rheumatic Fever, Endocarditis, and Kawasaki Disease, Council on Cardiovascular Disease in the Young, and the Councils on Clinical Cardiology, Stroke, and Cardiovascular Surgery and Anesthesia, American Heart Association: endorsed by the Infectious Diseases Society of America. Circulation 2005;111:e394-434.

17. Henry NK, Wilson WR, Roberts RB, Acar JE, Geraci JE. Antimicrobial therapy of experimental endocarditis caused by nutritionally variant viridans group streptococci. Antimicrob Agents Chemother 1986;30:465-7.

18. Lin CH, Hsu RB. Infective endocarditis caused by nutritionally variant streptococci. Am J Med Sci 2007;334:235-9.

19. Chung DR, Kim K, Chae JG, Uhlm WS, Kim YS, Song JH, Song JK, Kim JI, Park SW, Ryu JS, Lee SY, Pai CH. Two cases of infective endocarditis with nutritionally variant streptococci. Korean J Med 1996;50:244-9.

20. Seo MR, Park YS, Kim EJ, Lee HN, Oh KY, Seo YH, and Choi CH. A case of endocarditis due to Granulicatella adiacens. Infect Chemother 2010;42:311-4.