

Bacteremia and Endocarditis Caused by a *Gordonia* Species in a Patient with a Central Venous Catheter

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We report the first case of endocarditis caused by a *Gordonia* species genetically related to *G. sputi* but exhibiting some atypical biochemical features in a 31-year-old woman with a central venous catheter. This unusual pathogen may be a new cause of opportunistic infections in patients with severe underlying diseases.

Gordonia spp. are a gram-positive coryneform bacteria, recently identified in three patients as a cause of systemic infections (1,2), including two associated with indwelling implantable subcutaneous central venous catheters (2). We report *Gordonia* spp.'s capacity to infect a patient's undamaged cardiac valve through an implantable subcutaneous central catheter.

Case Report

The patient was a 31-year-old woman, who had undergone splenectomy at age 9 years, for severe double heterozygous hemoglobinopathy (beta-thalassemia and hemoglobin E disease). Multiple blood transfusions at that time were complicated by hepatitis C with cirrhosis and secondary hemochromatosis, treated at home with twice weekly deferoxamine by subcutaneous central venous catheter. Hemochromatosis was complicated by diabetes, adrenal insufficiency, and peripheral neuropathy. In September 1997, the patient became ill with *Staphylococcus aureus* bacteremia associated with localized renal and cutaneous abscesses; transesophageal echocardiography showed neither valvular vegetation nor a valvular defect suggestive of

endocarditis. The bacteremia was successfully treated with intravenous fosfomycin, cefotaxime and netilmicin, followed by ciprofloxacin and oxacillin for 8 weeks. The subcutaneous central venous port was also changed after 6 weeks of treatment.

The patient remained afebrile until December 1998, when fever and chills developed. Physical examination 1 week after onset of symptoms revealed a temperature of 39°C and a new mitral systolic murmur. The site of the subcutaneous central venous port showed no signs of infection. Results of clinical laboratory tests showed a leukocyte count of $31.4 \times 10^9/L$, an erythrocyte sedimentation rate of 67 mm/h, and a C-reactive protein plasma level <5 mg/L. Transthoracic echocardiography revealed a mitral valvular vegetation (10x5 mm). Two blood cultures (one obtained from the central venous catheter) were performed at admission. One more peripheral blood sample was drawn for culture 12 hours later. All three cultures were positive for a gram-positive coryneform bacterium showing no extensive branching. This organism produced dry, raised, salmon-to-orange colonies without aerial hyphae. As the organism was weakly acid fast, according to the Kinyoun acid-fast stain modified for aerobic actinomycetes, we presumptively identified this organism as *Rhodococcus* sp. sensu lato. According to disk diffusion, the organism was susceptible to

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penicillin G, amoxicillin, cefotaxime, ceftriaxone, imipenem, gentamicin, netilmicin, ciprofloxacin, vancomycin, and erythromycin but was resistant to fosfomycin, ceftazidime, trimethoprim-sulfamethoxazole, and streptogramin. By E-test, the MICs of penicillin G, amoxicillin, and ceftriaxone were 0.047, 0.064 and 0.25 µg/mL, respectively. The patient was successfully treated with intravenous amoxicillin (3 g, 4/day) and intravenous netilmicin (150 mg, 2/day) for 1 week, then with intravenous amoxicillin alone for 3 weeks. This treatment was followed by home treatment with 2 g perfusion of ceftriaxone for 2 weeks. The central venous catheter was left in situ. At 1 year follow-up, the patient was not infected with *Gordonia* sp., and echocardiographic findings were consistent with mitral valve insufficiency without oscillating intracardiac mass on valve.

To accurately identify the organism, we analyzed the p-bromophenacyl esters of mycolic acids of the isolate by using high-performance liquid chromatography, obtaining a pattern consistent with that of *Gordonia* sp. The number of peaks and retention times were similar to those exhibited by the *G. sputi* type strain ATCC 29627. Biochemical test results were positive for the hydrolysis of urease and esculin but negative for the hydrolysis of xanthine, adenine, tyrosine, and hypoxanthine. When inoculated with aerobic, low-peptone carbohydrate slants, the strain produced acid from trehalose but not from L-rhamnose and D-mannitol. These biochemical characteristics fit those of *Gordonia* species, especially *G. bronchialis* (3). To determine partial 16S rRNA gene sequence, the two eubacterial universal primers P8-27 (5'-AGA

GTT TGA TCC TGG CTC AG-3') and P1392-1372 (5'-AAG GCC CGG GAA CGT ATT CAC-3') were used for the amplification, then a direct sequencing method with an internal primer P535-514 (5'-GTA TTA CCG CGG CTG CTG GGC AC-3') 5' labeled with fluorescein isothiocyanate was performed (4). The sequence obtained coincides with the 450 5' base pairs of the 16S rRNA gene and matches totally that of *G. sputi* present in the GenBank-EMBL database. This sequence did not match other bacterial sequences, including those of other *Gordonia* species. The sequence of the isolate differed from the sequence of the *G. aichiensis* type strain by two nucleotides. DNA-DNA similarity experiments, according to the stringent nuclease S1 method, showed 55% DNA relatedness with *G. sputi* type strain and less than 28% with the type strain of other species including *G. aichiensis*. (The genetic definition of a species is more than 70% DNA similarity.) Thus, this isolate does not fit in any of the recognized *Gordonia* species, although it is taxonomically close to *G. sputi*.

Conclusions

The recent differentiation of *Gordonia* sp. as a distinct genus is the outcome of a taxonomic history complicated by several reclassifications (Table 1). Twelve organisms now belong to the genus *Gordonia*, including three species discovered in 1999 (5-9). To review the spectrum of clinical diseases in humans caused by *Gordonia* spp., we performed a Medline search for 1966 to 1999, using all the designations included in Table 1. Only *G. bronchialis* (10), *G. rubropertincta* (11),

Table 1. Classification of the genus *Gordonia*

Present designated species (date)	Former designated species (date)
<i>Gordonia aichiensis</i> ^a (1997)	<i>Gordonia aichiensis</i> (1994) ← <i>Rhodococcus aichiensis</i> (1983)
<i>Gordonia alkanivorans</i> (1999)	--
<i>Gordonia amarae</i> ^a (1997)	<i>Gordonia amarae</i> (1994) ← <i>Nocardia amarae</i> (1980)
<i>Gordonia bronchialis</i> ^a (1997)	<i>Gordonia bronchialis</i> (1989) ← <i>Rhodococcus bronchialis</i> (1980)
<i>Gordonia desulfuricans</i> (1999)	--
<i>Gordonia hirsuta</i> ^a (1997)	<i>Gordonia hirsuta</i> (1996)
<i>Gordonia hydrophobica</i> ^a (1997)	<i>Gordonia hydrophobica</i> (1995)
<i>Gordonia polyisoprenivorans</i> (1999)	--
<i>Gordonia rhizosphaera</i> (1998)	--
<i>Gordonia rubropertincta</i> ^a (1997)	<i>Gordonia rubropertincta</i> (1989) ← <i>Rhodococcus rubropertinctus</i> ^b (1980)
<i>Gordonia sputi</i> ^a (1997)	<i>Gordonia sputi</i> (1989) ← <i>Rhodococcus sputi</i> ^c (1975)
<i>Gordonia terrae</i> ^a (1997)	<i>Gordonia terrae</i> (1989) ← <i>Rhodococcus terrae</i> (1980)

^aThe original spelling, *Gordona*, was changed to *Gordonia* in 1997 (5).

^bOther former designations: *Bacillus rubropertinctus*, *Serratia rubropertincta*, *Mycobacterium rubropertinctum*, *Proactinomyces rubropertinctus*, *Nocardia rubropertincta*.

^cSynonym: *Rhodococcus chubuensis*.

G. sputi (1,12), and *G. terrae* (2,13,14) have been shown to be pathogenic in humans (Table 2). They are derived from soil and may also be isolated in the sputa from patients with chest disorders (15). In an outbreak of sternal-wound infections (10), *G. bronchialis* was isolated from the hand, scalp, and vagina of a nurse as well as from her dog.

Systemic *Gordonia* spp. infections have been described in three patients. Buchman et al. (2) reported two cases of *Gordonia* spp. bloodstream infection associated with a Hickman catheter in two immunocompetent patients receiving long-term parenteral nutrition. Both strains were susceptible to vancomycin and gentamicin. In one case, the *Gordonia* sp. isolated from blood cultures and from a broth culture of the catheter tip was not clearly identified but was close to *G. rubropertincta*. The patient received intravenous vancomycin for 5 days and intravenous gentamicin for 19 days; the catheter was removed after 2 days. In the second case, the microorganism isolated from blood cultures was identified as *G. terrae*, and the patient was treated with intravenous vancomycin for 6 weeks with the catheter left in situ.

The only case of bacteremia caused by *G. sputi* was described in a 34-year-old immunocompromised patient (1) with metastatic melanoma treated with intravenous interleukin-2. The bacterium was thought to have reached the bloodstream through extensive desquamative skin rashes caused by the interleukin-2 treatment or by contamination of the catheter, although cultures from these specimens were

negative. Because *G. sputi* was susceptible to β -lactams, vancomycin, aminoglycosides, doxycycline, and rifampicin, the patient was first treated with amoxicillin-clavulanate (1,000 and 125 mg, respectively, every 3 hours). After 1 week of treatment, a second set of blood cultures yielded the same organism, and the treatment was changed to a combination of amikacin and piperacillin, which was successful (1). As with other *Gordonia* species, *G. sputi* may be isolated in the sputum of patients with pulmonary disease (15). Mediastinitis caused by *G. sputi* after coronary artery bypass grafting was recently described in an immunocompetent patient (12). The patient was treated with cefmetazole sodium (2 g per day for 3 weeks) and piperacillin sodium (2 g per day for 2 weeks) after surgical soft tissue debridement.

In our case, *Gordonia* sp. systemic infection associated with an implantable subcutaneous central venous catheter was complicated by endocarditis. The diagnosis was assessed as definitive on the basis of Duke criteria (one major and three minor, including echographic evidence of an oscillating intracardiac mass with a new regurgitant murmur, two persistently positive blood cultures yielding *Gordonia* sp., fever $>38^{\circ}\text{C}$, and intravenous deferoxamin use). Our patient had neither neutropenia nor immunosuppressive medications, but underlying diseases may have impaired the immune system and facilitated infection. The bacteremia may be caused by manipulations of the implantable subcutaneous central venous catheter during routine home use. However, even if no other

Table 2. Types of infection caused by *Gordonia* species and patients' underlying conditions

Type of infection	Cases (No.)	Age (yrs)	<i>Gordonia</i> species	Underlying conditions	Authors
Sternal wound	7	51-68	<i>G. bronchialis</i>	Surgery	Richet et al. (10)
Mediastinitis	1	54	<i>G. sputi</i>	Surgery	Kuwabara et al. (12)
Brain abscess	1	40	<i>G. terrae</i>	None	Drancourt et al. (14)
	1	3	<i>G. terrae</i>	Cerebral tumor	Drancourt et al. (13)
Lung infection	1	29	<i>G. rubropertincta</i>	Tuberculosis	Hart et al. (11)
Bacteremia due to central venous catheter	2	43 65	<i>Gordonia</i> sp. ^a <i>G. terrae</i>	Breast and ovarian cancer Chronic intestinal pseudo-obstruction syndrome	Buchman et al. (2)
Bacteremia due to cutaneous lesions	1	34	<i>G. sputi</i>	Metastatic melanoma IL2 treatment ^b	Riegel et al. (1)
Skin infection	1	7	<i>G. terrae</i>	None	Martin et al. (16)

^aNot identified.

^bIL2 = interleukin-2.

source for the bacteremia had been evident, no inflammation was observed at the port site, and semiquantitative culture was not available. An environmental investigation was not performed.

We conclude that *Gordonia* spp. may cause opportunistic infections, in particular bacteremia and endocarditis, in patients with severe underlying diseases and indwelling central catheters.

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