First case of Campylobacter rectus and Solobacterium moorei mixed bacteraemia successfully identified by MALDI TOF-MS

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Abstract

Campylobacter rectus and Solobacterium moorei are anaerobic Gram-negative and Gram-positive rods, respectively, that are occasionally members of the human oral flora. Bacteraemia has rarely been reported. We present the first case of mixed *C. rectus*–S. moorei bacteraemia in an individual with diabetes and human immunodeficiency virus infection. Both bacteria were successfully identified by MALDI-TOF MS.

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Case report

A 70-year-old man was admitted to our institution for a progressive decay of his general conditions: important weight loss and fatigue, and dysphagia associated with fever, cough and dyspnoea. The patient had human immunodeficiency virus type I infection and was being treated: viral load was <50 copies/mL and CD4⁺ T-cell count was 890/µL. After the collection of two sets of blood cultures, empirical treatment with amoxicillin/ clavulanic acid was started. After 72 hours of incubation in a BD

Bactec FX blood culture system, one out of the two anaerobic blood culture bottles became positive (Becton Dickinson, Franklin Lakes, NJ, USA). Gram-staining achieved from the positive blood culture showed the presence of both Gramnegative and Gram-positive rods. Rapid matrix-assisted laser time-of-flight mass desorption/ionization spectrometry (MALDI-TOF MS) identification was performed on the positive blood culture, but provided no reliable identification. The blood was inoculated onto both aerobic and anaerobic media (respectively Columbia agar and Schaedler agar; Becton Dickinson). After 2 days, only anaerobic cultures became positive. Identifications were performed by MALDI-TOF MS, according to the manufacturer's instructions, using the direct smear procedure without protein extraction (Biotype IVD 4.2.80; Bruker Daltonics, Bremen, Germany). Campylobacter rectus and Solobacterium moorei were identified with best-match score values of 1.96 and 2.3, respectively. Considering that a score value of minimum 2 is required for acceptance, the C. rectus

Authors	Case: age; sex	Concomitant infection and possible source	Co-pathogen isolated	Co-morbidities/risk factors	Treatment	Identification
Detry et al., 2006 [5]	l) 67 y; M	Sepsis Dental abscess	_	Multiple myeloma	Cefepime 15 days	PCR 16s rRNA
Lau et al., 2006 [10]	l) 43 y; F	Acute proctitis	_	Carcinoma of the cervix stage III-B	Piperacillin/tazobactam 15 days	PCR 16s rRNA
Martin et al., 2007 [11]	I) 37 y; M	Septic pulmonary embolism Femoral vein thrombophlebitis and abscess	Fusobacterium nucleatum; Bacteroides ureolyticus	Intravenous drug abuse	Benzylpenicillin ? days metronidazole ? days	PCR 16s rRNA
Pedersen et al., 2011 [3]	I) 43 y; M	Tooth abscess	_	Lymphoma, kidney transplantation	Benzylpenicillin 14 days metronidazole 14 days	PCR 16s rRNA
	2) 66 y; F	Pulmonary abscess	Eikenella corrodens			PCR 16s rRNA
	3) 64 y; M	Sepsis	—	Colon cancer	Cefuroxime 28 days metronidazole 28 days	PCR 16s rRNA
	4) 33 y; F	Femoral vein thrombosis and abscess	Actinomyces meyeri	Intravenous drug abuse Chronic HBV infection	Cefuroxime → benzylpenicillin + metronidazole 35 days	PCR 16s rRNA
	5) 77 y; M	Pneumonia	Porphyromonas uenonis	Prostate cancer History of heart disease	Benzylpenicillin → phenoxymethyl-penicillin 10 days	PCR 16s rRNA
Genderini et al., 2019 [current]	I) 70 y; M	Pneumonia HSV-1 oesophagitis	Campylobacter rectus	HIV infection Diabetes	Amoxicillin/clavulanic acid 14 days	MALDI-TOF MS

TABLE I. Summary o	f the cases of Solobacte	rium moorei bacteraemi	a reported in the literature
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identification was confirmed by the partial sequencing (1249 bp) of the 16S rRNA gene using universal primers (27F: 5'-AGAGTTTGATCMTGGCTCAG-3' and 1492R: 5'-TACGGY-TACCTTGTTACGACTT-3'). The yielded sequence (GenBank Accession no.: MK063870.1) had 99.60% homology with С. rectus strain RM3267 (GenBank Accession no.: ACFU01000050) by using EZBIOCLOUD (www.ezbiocloud.net/) and 99% homology with C. rectus strain JCM 6301 (GenBank Accession no.: NR_113247.1) by using the NCBI database (www.ncbi.nlm.nih.gov/). The antimicrobial susceptibility testing performed using E-test gradient strips (bioMérieux, Marcy l'Etoile, France) and following EUCAST v8.0 clinical breakpoints showed full susceptibility to amoxicillin/clavulanic acid (C. rectus MIC 0.047 µg/mL; S. moorei MIC 0.016 µg/mL). An oesophagogastroduodenoscopy showed an erosive oesophagitis and pathological examination of the biopsies indicated the presence of viral inclusions in epithelial cells. The immunostaining was positive for herpes simplex virus type I (HSV-I Polyclonal antibody, Biocare Medical, Pacheco, CA, USA). Dental examination did not reveal any source of infection, only focal signs of periodontitis. ¹⁸F-Fluorodeoxyglucose positron emission tomography-CT was performed and demonstrated abnormal diffuse oesophageal hypermetabolism, suggesting an inflammatory origin but without evidence of malignancy.

This is the first case of a concomitant bacteraemia with *C. rectus* and *S. moorei. Campylobacter rectus*, an anaerobic Gramnegative rod, and *S. moorei*, a Gram-positive anaerobic rod, are both occasional members of the human oral flora and were found in individuals with refractory periondotitis [1]. Invasive infections such as empyema and brain abscess have been mainly reported in individuals with poor oral hygiene [2–4].

Solobacterium moorei bacteraemia is reported especially in individuals with malignancies [3,5]. In Table I a review of cases of S. moorei bacteraemia is presented. In all infections, identification was performed using 16s rRNA sequencing. We identified only two cases of bacteraemia caused by C. rectus; in one identification was performed using I6s rRNA sequencing [4,6]. The present case is the first in which both S. moorei and C. rectus were identified using MALDI-TOF MS. This case illustrates the added value of MALDI-TOF MS, not only in clinical microbiology [7] but also as a fast and reliable alternative to expensive molecular techniques such as 16s rRNA sequencing [8]. The probable source of the mixed bacteraemia was the translocation caused by the HSV-I oesophagitis. Disruptions of the mucosal barrier along with host immunodeficiency are associated with higher risk of bacterial translocation from the gastrointestinal tract [9].

Conflict of interest

None to declare.

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