First time identification of *Pandoraea sputorum* from a patient with cystic fibrosis in Argentina: a case report

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**Abstract**

**Background:** *Pandoraea* species are considered emerging pathogens in the context of cystic fibrosis (CF) and are difficult to identify by conventional biochemical methods. These multidrug resistant bacteria remain poorly understood particularly in terms of natural resistance, mechanisms of acquired resistance and impact on the prognosis of the disease and the lung function. Among them, *Pandoraea sputorum* has been previously described in few cases of CF patients from Spain, Australia, France and United States, underlining the need of more clinical data for a better knowledge of its pathogenicity. This is the first report relating to *P. sputorum* in a CF patient in Argentina.

**Case presentation:** *Pandoraea sputorum* was identified in a nine-year-old cystic fibrosis patient from Argentina, after treatment failure during an exacerbation. The isolates were successfully identified by combining molecular techniques based on 16S rRNA sequencing and mass spectrometry (MS) methods, after reassessing previous misidentified isolates by conventional methods. After first isolation of *P. sputorum*, patient’s clinical condition worsened but later improved after a change in the treatment. Although isolates showed susceptibility to trimethoprim–sulfamethoxazole and imipenem, in our case, the antibiotic treatment failed in the eradication of *P. sputorum*.

**Conclusions:** All combined data showed a chronic colonization with *P. sputorum* associated to a deterioration of lung function. We noted that the presence of *P. sputorum* can be underestimated in CF patients and MALDI-TOF MS appears to be a promising means of accurate identification of *Pandoraea* species.

**Keywords:** *Pandoraea sputorum*, Cystic fibrosis, First report, MALDI-TOF MS, Argentina

**Background**

In cystic fibrosis (CF) patients, the main cause of morbidity and mortality is the decline in the pulmonary function subsequent to pathogenic colonization with non-fermenting Gram negative bacteria (NFGNB) [1]. Nevertheless, diverse bacterial species exhibit distinct degrees of pathogenicity and sensitivity patterns, requiring different clinical managements [2]. Regularly, a series of opportunistic pathogens colonizes the CF lung throughout life, often culminating in chronic infection with *Pseudomonas aeruginosa* and, less frequently, with *Burkholderia cepacia* complex (Bcc) organisms. Other opportunistic Gram-negative bacteria are also encountered. They include various Enterobacteriaceae, *Stenotrophomonas maltophilia*, *Achromobacter xylosoxidans* [1, 3] and species that are only occasionally associated with human infections apart from CF, such as *Cupriavidus*, *Ralstonia*, and *Pandoraea* species. Some of these species have been associated with poor outcomes in CF, and the role that others may play in disease progression remains uncertain [2]. Bacteria from the *Pandoraea* genus (*P. pulmonicula*, *P. pnomena*, *P. apista*, *P. norimbergenisis* and *P. sputorum*) are NFGNB that are being considered as emerging...
multi-drug resistant pathogens in the context of CF [4, 5]. As they are still poorly known, particularly in terms of mechanisms of natural or acquired resistance and of their impact on lung function and the prognosis of the disease [5], more information is needed to evaluate their pathogenic role. Conventional microbiology methods remain problematic for the accurate identification of Pandorea species and misidentification with bacteria of the genus Burkholderia or Ralstonia has been reported [6–8]. Moreover, isolates from CF patients with persistent pathogenic colonization often lose their characteristic phenotypes or growth conditions [9]. Therefore, descriptions of infections due to Pandorea species in CF patients could be underestimated. The availability of MALDI-TOF MS in clinical microbiology diagnosis allows a quick and better identification of bacterial species, especially Pandorea species [5, 8, 10–12]. In this study we report for the first time in Argentina, a lung colonization with Pandorea sputorum, in a young CF patient who did not respond to antibacterial empirical treatment. Identification was possible after reassessing historical identification results obtained by conventional methods, using MALDI-TOF MS and sequencing of 16S rRNA gene.

Case presentation
A 9-year-old boy who was diagnosed with CF at 29 months (heterozygous genotype F508del; sweat test positive: Na 74 mEq/l and Cl de 63 mEq/l), was transferred to our CF unit with a 5 years history of chronic bronchopulmonary colonization with Pseudomonas aeruginosa and Staphylococcus aureus. When the first microbiological cultures (always sputum samples) were performed in our laboratory in July 2014, P. sputorum was detected together with S. aureus, but was initially reported as S. maltophilia (Fig. 1). Previously, his pulmonary function test revealed a forced expiratory volume in 1 s (FEV1) of 83%, and the preceding respiratory exacerbations were treated empirically with ceftazidime, amikacin and with aerosolized tobramycin. From the time of the first isolation of P. sputorum and during the next 2 months, his clinical condition worsened and exacerbations with increased cough and secretions were observed. The patient was treated with trimethoprim-sulphamethoxazole (4/20 mg/kg/day) plus ciprofloxacin (15 mg/kg/day) and with aerosolized colistin (2 million units twice daily), however, his lung function declined (FEV1 45%). In June 2015, the patient showed exacerbations with a dramatic deterioration of his pulmonary function (FEV1 27%). A combined therapy for three-weeks of imipenem (80 mg/kg/day), amikacin (40 mg/kg/day), and inhaled colistin was initiated. Consequently, his clinical condition improved with a significant reduction in sputum volume and cough (FEV1 40%). Five months later, his clinical condition was stable. In April 2016, a new respiratory sample was positive for P. sputorum, confirmed by MALDI-TOF MS, together with with S. aureus and P. aeruginosa.

Our laboratory records indicated that sputum cultures during patient’s exacerbations and/or routine follow-up visits until July 2015, showed colonies of NFGNB without any clear phenotypic differences and S. aureus. Conventional identification of purple colonies recovered on B. cepacia complex selective agar plates (BCSA; Britania S.A, Argentina) was performed to some NFGNB
colonies. We used a combination of phenotypic tests like oxidase activity, DNase, carbohydrate oxidation-fermentation, lysine and arginine decarboxylase, and motility tests. After these tests, isolates were reported as NFGNB (n = 2), *S. maltophilia* (n = 2), *B. cepacia* complex (n = 1) and *S. aureus* (n = 7). Later, one of the Gram-negative isolates was identified as *Bordetella hinzii* with a high identification confidence level (99% probability) using VITEK2 automatic system (bio-Mérieux, France). Due to the inaccuracy of biochemical tests, we decided to reassess the identification of all NFGNB isolates (n = 5) by Matrix Assisted Laser Desorption Ionisation-Time of flight (MALDI-TOF) mass spectrometry (MS), using an Ultraflex MALDI TOF/TOF instrument and the MALDI Biotyper 3.1 software (Bruker Daltonics, Bremen, Germany). The five isolates were identified as *Pandorea sputorum* at species-level (score >2). Hierarchical cluster analysis based on MALDI-TOF MS spectra comparison clustered together, clinical isolates and the reference strain *P. sputorum* DSM 21091T, while *P. apista*, *P. pnomenusa*, *P. pulmomica* and *P. norimberegensis* were in a separate branch (Fig. 2). In addition, genetic relatedness was determined by fingerprinting analysis like BOX-PCR (Interspersed repetitive element) and ERIC-PCR (enterobacterial repetitive intergenic consensus) [14], revealing indistinguishable patterns among all our clinical isolates (data not shown). Finally, a subsequent full characterization through PCR and sequencing of 16S rRNA gene was performed using the corresponding universal primers (27 F and 1492R), previously published [15]. Sequence analysis of a 1400-bp fragment of the 16S rRNA gene of one of patient’s isolate (*P. sputorum* HP020) was performed. The corresponding sequence (GenBank accession KX258224) showed 99% of similarity with *P. sputorum* DSM21091T (GenBank accession CP010431), which confirmed the results obtained by MALDI-TOF MS analysis. The phylogenetic relationship among them was investigated by using MEGA 6.0 software (http://www.megasoftware.net) (Fig. 3).

Antibiotic susceptibility testing was performed using a disk diffusion method according to the recommendations of the committee of CLSI [16]. Susceptibility patterns were also identical for all isolates. In the absence of interpretive susceptibility criteria for strains of *Pandorea*, results were interpreted using CLSI criteria for “other non-Enterobacteriaceae” isolates. The microorganism was resistant to amikacin, ceftazidime, cefepime, meropenem, gentamicin, aztreonam and colistin and susceptible to imipenem and trimethoprim-sulfamethoxazole. Minimum inhibitory concentrations (MIC, mg/L) were estimated by Vitek2 Compac® for amikacin (≥64), ceftazidime (≥64), cefepime (≥64), ciprofloxacin (≥4), colistin (≥16), meropenem (≥16), gentamicin (≥64), piperacillin-tazobactam (≥128), imipenem (≥0.25) and trimethoprim-sulfamethoxazole (TMS, ≤1/20), which confirms the susceptibility profile.

**Discussion and Conclusions**

In the last decade, many emerging multidrug resistant pathogens have been described in CF patients including new species of the *Burkholderia cepacia* complex,
Brevundimonas diminuta, Inquilinus limosus, Acetobacter indonesiensis, Achromobacter xylosoxidans, Ochrobactrum anthropi, and bacteria of the genus Pandoraea [17, 18]. The pathogenicity of Pandoraea species remains controversial. These bacteria have been occasionally recovered from the lung transplant and/or respiratory tract of CF patients, as well as from blood cultures in non-CF patients and environmental samples [2, 4, 8, 19]. In our review of the bibliography, we found few cases of CF patients chronically colonized with P. sputorum: one case from Australia [8], two cases from Spain [10, 11], and one case from France [5]. Interestingly, in our CF patient, S. aureus and P. aeruginosa were eventually isolated from respiratory tract cultures together with P. sputorum. This situation makes more difficult to know the real contribution of P. sputorum to the clinical condition of our patient. In this regard, intermittent or persistent colonization with Pandoraea species, that sometimes coincides with a deterioration in lung function, has been reported in addition to clinical evidence of invasive potential [8, 10, 11, 18]. Furthermore, besides the inability of Pandoraea species to produce a biofilm in the respiratory tract [20], it remains still unclear the role of different virulence factors, mechanisms of pathogenicity, the possibility of transmission between patients, and their role in lung damage in CF patients. This situation emphasizes the importance of having more data about clinical cases. The Pandoraea genus is multidrug resistant and treatment may be problematic. In our case, P. sputorum was only susceptible to imipenem and TMS, with a low MIC for imipenem (≤0,25 mg/L), and although the treatment failed to eradicate the bacteria, the clinical condition improved. This pattern of being resistant to carbapenem and meropenem but sensitive to imipenem, appears to be unique to most Pandoraea genus [5, 21], and it can be a practical guideline in the earliest identification steps.

It is documented, when applying only conventional identification phenotypic methods, the microbiology laboratory commonly misidentifies this pathogen as Ralstonia, Stenotrophomonas or Burkholderia species [6–8]. It has been also noted the limitations of the sequences of 16S rRNA and gyrB genes for differentiating the Pandoraea species [8, 22]. The accuracy and usefulness of MALDI-TOF as a routine technique for rapid identification of bacterium was demonstrated. Nevertheless, one factor limiting the use of MALDI-TOF MS is the scant reference data sets for microorganisms that are infrequently isolated from clinical specimens [10]. Due to these limitations, a polyphasic approach for an accurate identification is recommended. In our CF patient, all isolates were identified as P. sputorum, combining 16S rRNA PCR and sequencing, and a MALDI-TOF MS proteomic platform. Lastly, to determine persistence over time of the same P. sputorum isolate, as generally occurs with P. aeruginosa and Bcc in CF patients, a molecular epidemiological analysis was performed [2, 23]. BOX-PCR and ERIC-PCR showed indistinguishable patterns among all isolates of P. sputorum (including last isolate collected in April 2016), indicating a possible chronic colonization, which is defined here as three positive cultures of the same strain isolated within a 6-month period.

In conclusion, we noted that the presence of P. sputorum can be underestimated in CF patients when using conventional biochemical identification methods. We also showed that P. sputorum may have a potential role in pulmonary function deterioration when chronically...
colonized CF patients, emphasizing the need to implement control policies to protect CF community. This is particularly relevant, as in our case, the treatment failed despite sensitivity to imipenem and trimethoprim–sulfa-methoxazole. Finally, MALDI-TOF MS appears to be an excellent tool for an accurate identification of organisms of this genus.

Abbreviations

CF: Cystic fibrosis; CLSI: Clinical Laboratory Standards Institute; FEV1: Forced expiratory volume in one second; MALDI-TOF MS: Matrix-assisted laser desorption ionization–time of flight mass spectrometry; MIC: Minimum inhibitory concentration; NFGNB: Non-fermenting Gram-negative bacteria

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Availability of data and materials

The datasets and materials during and/or analyzed during the current study available from the corresponding author on reasonable request.

Authors’ contributions

PFM, JF, and MVS recognized the novelty of the case and drafted the manuscript. PFM, JF, and MVS analyzed and interpreted the results. JGF and Botteg EC. 16S rRNA gene sequencing versus the API 20 NE system and the VITEK 2 ID-GNB card for identification of nonfermenting gram-negative bacteria in the clinical laboratory. J Clin Microbiol. 2006;44:1359–66.

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PFM and JF are researchers at the National University of Misiones and the National Scientific and Technical Research Council (CONICET).

Competing interests

The authors declare that they have no competing interests.

Consent for publication

Written informed consent was obtained from the parents of the patient for publication in this case report. A copy of the written consent is available for review by the Editor of this journal.

Ethics approval and consent to participate

Ethical approval to report this case was not required.

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