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## **Case Series Study of Melioidosis, Colombia**

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We report 7 cases of melioidosis in Colombia and comparision of 4 commercial systems for identifying *Burkholderia pseudomallei*. Phoenix systems were not a definitive method for identifying *B. pseudomallei*. For accurate identification, we recommend including this bacterium in the library databases of matrix-assisted laser desorption/ionization mass spectrometry systems in Latin America.

Melioidosis is an infectious disease caused by *Burkholderia pseudomallei*, a saprophytic soil bacterium (1). Recently, an increase in cases outside the Asia–Pacific region, including the Americas, has been reported. It is not clear whether this increased number of cases reflects an increase in incidence of this disease or improvements in its identification by microbiological laboratories and research facilities (2).

#### The Study

We describe 7 cases of melioidosis in the Caribbean coast region of Colombia among patients who sought emergency services or were referred to 3 hospitals in the city of Valledupar (Figure). None of the patients reported travel abroad. Four patients were admitted during 2015 and 2016; we analyzed the corresponding isolates using 4 commercial methods and a molecular identification method. For the other 3 cases, which were diagnosed in 2014, we reviewed patient medical records and microbiological results. These

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isolates were not available. The study was approved by the ethics committees of Clinica Laura Daniela, Clínica Médicos, and Instituto Cardiovascular del Cesar.

Of the 7 patients, 6 (86%) were men; 2 patients were admitted to the intensive care unit (Table 1). The most common clinical presentation was bacteremic pneumonia (in 4 patients); 5 patients had a history of diabetes.

The 4 strains isolated in 2015 and 2016 were identified as *B. pseudomallei* by Vitek Compact 2 (bioMérieux, https://www.biomerieux.com) and Microscan (Walkaway Beckman Coulter, https://www.beckmancíoulter.com) but as *B. cepacia* by Phoenix (Becton Dickinson, http://www. bd.com) and as *B. thailandesis* by MALDI Biotyper v3.1 matrix-assisted laser desorption/ionization time-of-flight (MALDI-TOF) mass spectrometry (Bruker Daltonics, https://www.bruker.com). Internal transcribed spacer (ITS) DNA sequencing identified all the isolates as *B. pseudomallei*. Sequences of these 4 isolates were submitted to Gen-Bank (accession nos. KX898558, KY659330, KY996759, and KY659331). According to patient medical records, the 3 isolates from 2014 were identified as *B. pseudomallei* using the Vitek or Microscan system or both (Table 2).



Figure. Geographic locations of 7 reported cases of melioidosis in Colombia.

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	Patient					Appropriate			
Patient	age, y/	Medical history and	Clinical		Empirical	treatment			
no.	sex	risk factors	manifestation	Type of sample	treatment	(duration)	ICU	Outcome	Relapse
1	72/M	Diabetes, arterial	Bacteremic	Blood culture,	TZP +	IP: MER +	Yes	Died	NA
		hypertension	pneumonia	urine culture,	CLR	TMP/SMX (14			
				endotracheal		d); EP: NA			
				aspirate					
2	60/F	Diabetes, arterial	Bacteremic	Wound	CFZ +	IP: MER (10 d);	No	Alive	No
		hypertension,	SSTI	secretion, blood	CLI	EP: TMP/SMX			
		minor head injury		culture		(3 mo)			
		with open scalp							
3	49/M	wound	Bacteremic	Blood culture,	TZP +		No	Alive	No
3	49/10	Diabetes, leprosy, erythema nodosum	pneumonia	urine culture.	CLR	IP: MER (10 d); EP: TMP/SMX	No	Allve	No
		leprosum, chronic	prieumonia	sputum culture	ULK	(6 mo)			
		use of steroids		sputum culture		(01110)			
4	71/M	Arterial	Bacteremic	Blood culture	TZP +	IP: MER (14 d);	No	Alive	No
		hypertension,	pneumonia		DOX	EP: TMP/SMX			
		multiple myeloma,	•			(3 mo)			
		bone marrow				( )			
		transplant, chronic							
		use of steroids and							
		thalidomide							
5	66/M	Diabetes,	Bacteremic	Blood culture,	TZP	NA	Yes	Died	NA
		arterial hypertension	pneumonia	endotracheal					
_			· · · <u>- · · · · · · · · · · · · · · · ·</u>	aspirate					
6	56/M	Diabetes, chronic	UTI	Urine culture	TZP	IP: MER (10 d);	No	Alive	At 6 mo:
		renal failure				EP: TMP/SMX			bacteremia
7	E 4 / N 4	This and Charles and	0071	14/1	057	(3 mo)	N.L.	A 15	UTI, SSTI
7	54/M	Tibia and fibula open	SSTI	Wound	CFZ +	IP: MER (10 d);	No	Alive	No
		fracture		secretion, blood	CLI	EP: TMP/SMX			
		clindamycin; CLR, clarith		culture		(6 mo)			

 Table 1. Epidemiologic and demographic characteristics, underlying conditions, clinical manifestations, and outcomes of patients with

 melioidosis in the Caribbean coast region of Colombia\*

MIC testing was performed using the system available in the institution where the patient was being treated: Microscan Walkaway (isolates 1, 2, 3, 5) and Phoenix systems (isolates 4, 6, 7). According to published cutoff points for *B. pseudomallei* (Clinical and Laboratory Standards Institute, https://clsi.org/ standards/products/microbiology/documents/m45), all isolates were susceptible to trimethoprim/sulfamethoxazole and meropenem, and 2 isolates (isolates 2 and 7) were resistant to ceftazidime (MIC >16  $\mu$ g/mL); however, isolate 7 was tested 2 more times using Microscan, resulting in a value of <2  $\mu$ g/ mL. Isolate 2 was not available for further analysis.

Within the genus *Burkholderia*, *B. pseudomallei*, *B. mallei*, and the *B. cepacia* complex are the species usually associated with human infection (3, 4). Historically, melioidosis is recognized as a major cause of fatal pneumonia and sepsis in Southeast Asia, South Asia, and northern Australia. It is believed that movements of both humans and cargo have contributed to the dissemination. A phylogenetic reconstruction study determined the African origin of the American isolates and the overlapping of the introduction of *B. pseudomallei* into the New World with the height of the slave trade from Africa to the Americas (5).

In Colombia, melioidosis is not a disease of public health interest and thus could be underreported. However, Colombia is second to Brazil with the highest number of cases reported in South America (1). The actual number of cases is unknown, probably because of the lack of awareness and suspicion of the disease by health professionals, together with the absence of technology for proper diagnosis and the underreporting of diagnosed cases. At least 10 reported cases of melioidosis have been acquired in the Andean region of Colombia. Severiche published a report in 1998 about a patient with pneumonia (6). Montúfar et al. described a series of 7 patients during 1998-2012, of whom 5 had bacteremic pneumonia (7). Since then, other cases of bacteremic pneumonia have been reported (8,9), as well as 1 case of chronic lung melioidosis in a patient on Colombia's west Caribbean coast (10). Guzmán-Gómez et al. described a case of osteoarticular melioidosis acquired in the Colombian rainforest in 2015 (11), the only case confirmed by sequencing (ITS-16S rRNA gene). The areas in Colombia where melioidosis cases have been reported, including these new cases from the Caribbean coast, are consistent with the previously proposed model (2) (Figure 1).

The isolation of *B. pseudomallei* from clinical specimens is the standard for a diagnosis of melioidosis. However, the microorganism is not often recovered and may not be correctly identified even when isolated. *B. pseudomallei* 

 Table 2. Isolate identification results by commercial systems, 16S rRNA sequencing analysis, and antimicrobial drug susceptibility of isolates of Burkholderia pseudomallei obtained from clinical specimens in the Caribbean coast region of Colombia\*

	Com	nmercial bacterial i									
		Walkaway/		GenBank	MIC, μg/mL						
Isolate	Vitek (%	Microscan (%	Phoenix	MALDI-TOF	ITS sequencing	accession			TMP/		
no.	probability)	probability)	(% probability)	MS (score)	segments	no.	CAZ	MER	SMX		
1	ND	B. pseudomallei (93%)	ND	ND	ND	ND	2	<u>&lt;</u> 1	<u>&lt;</u> 2/38		
2	B. pseudomallei (99%)	B. pseudomallei (99.5%)	ND	ND	ND	ND	>16	<u>&lt;</u> 1	<u>&lt;</u> 2/38		
3	B. pseudomallei (94%)	B. pseudomallei (99.8%)	ND	ND	ND	ND	2	<u>&lt;</u> 1	<u>&lt;</u> 1/19		
4	B. pseudomallei (94%)	B. pseudomallei (99.9%)	B. cepacia (99%)	<i>B. thailandensis</i> (1,899)	B. pseudomallei	KX898558	4	1	2/38		
5	B. pseudomallei (94%)	B. pseudomallei (99.5%)	B. cepacia (99%)	<i>B. thailandensis</i> (1,898)	B. pseudomallei	KY659330	4	2	1/19		
6	B. pseudomallei (95%)	B. pseudomallei (99.9%)	B. cepacia (99%)	B. thailandensis (1,898)	B. pseudomallei	KY996759	4	1	1/19		
7	B. pseudomallei (94%)	B. pseudomallei (93%)	B. cepacia (99%)	B. thailandensis (1,898)	B. pseudomallei	KY659331	<2	2	1/19		
*CAZ, ceftazidime; ITS, internal transcribed spacer; MALDI-TOF MS, matrix-assisted laser desorption/ionization time-of-flight mass spectrometry; MER, meropenem; ND, no data; TMP/SMX, trimethoprim/sulfamethoxazole.											

can be identified using commercial identification systems. However, these tests may fail to distinguish B. pseudomallei from B. thailandensis and other members of the B. cepacia complex (12,13). MALDI-TOF mass spectrometry is an accurate and rapid procedure for the identification of B. pseudomallei if the appropriate database is used (14). However, genotyping methods based on rRNA sequencing should be used for a more accurate diagnosis. We compared different phenotyping methods (Vitek, MicroScan Walkaway, Phoenix, and MALDI-TOF mass spectrometry) using 4 isolates from this report. Sequencing of the ITS region confirmed the identification of B. pseudomallei and matched the identifications obtained by the Vitek and Walkaway systems. However, the Phoenix system erroneously identified isolates as B. cepacia, and MALDI-TOF mass spectrometry incorrectly identified isolates as B. thailandensis. Although B. pseudo*mallei* was included in the Phoenix database, we strongly recommend not using Phoenix as a single or final method to identify possible isolates of B. pseudomallei (for example, gram-negative, oxidase-positive, and positive-arginine bacilli). MALDI-TOF mass spectrometry is currently being introduced in Colombia, and we recommend the inclusion of B. pseudomallei in the database.

The treatment of melioidosis is prolonged and includes 2 phases: intensive treatment with intravenous antimicrobial therapy for 10–14 days using ceftazidime, imipenem, or meropenem; and an eradication phase with oral antimicrobial therapy for 3–6 months with trimethoprim/sulfamethoxazole alone or in combination with doxycycline (15). In the cases we report, treatment with meropenem was started, because it is easier to prescribe this antimicrobial drug in Colombia, given the high rate of extended-spectrum  $\beta$ -lactamase–producing *Enterobacteriaceae*; the use of ceftazidime is restricted for the same reason. Because of the severity of illness and the high rate of death from this

disease, along with the required prolonged antimicrobial drug therapy and the small number of drugs available for its treatment, it is necessary not only to strengthen the public health surveillance and clinical suspicion of melioidosis but also to acquire tools that permit an adequate diagnosis, especially in potentially endemic areas, which, in the case of Colombia, could be an extensive geographic area.

#### Conclusions

With the presence of *B. pseudomallei* on Colombia's Caribbean coast confirmed, clinicians should suspect melioidosis in patients with risk factors, suggestive clinical symptoms, and microbiological isolates from clinical specimens of *B. pseudomallei* or other members of the *Burkholderia* genus. In this case, >1 microbiological identification method should be used, especially if the outdated Phoenix or MALDI-TOF mass spectrometry databases are used.

Finally, it is necessary to include melioidosis in a passive surveillance system, especially in those regions of Latin America where the environmental conditions create high probabilities for the presence of the disease. Currently, Colombia has a public health surveillance and control system under the supervision of the National Institute of Health, to which all health institutions must notify diseases of public health interest. To determine the true magnitude of melioidosis in Colombia, it must be included as a notifiable disease and measures established to enable its early diagnosis and treatment.

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