

OXA-198, an Acquired Carbapenem-Hydrolyzing Class D β -Lactamase from *Pseudomonas aeruginosa*[∇]

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A carbapenem-resistant *Pseudomonas aeruginosa* strain (PA41437) susceptible to expanded-spectrum cephalosporins was recovered from several consecutive lower-respiratory-tract specimens of a patient who developed a ventilator-associated pneumonia while hospitalized in an intensive care unit. Cloning experiments identified OXA-198, a new class D β -lactamase which was weakly related (less than 45% amino acid identity) to other class D β -lactamases. Expression in *Escherichia coli* TOP10 and in *P. aeruginosa* PAO1 led to transformants that were resistant to ticarcillin and showed reduced susceptibility to carbapenems and cefepime. The *bla*_{OXA-198} gene was harbored by a class 1 integron carried by a ca. 46-kb nontypeable plasmid. This study describes a novel class D β -lactamase involved in carbapenem resistance in *P. aeruginosa*.

Carbapenem drugs are often used for treating infections due to multidrug-resistant isolates (2). However, resistance to these antibiotics increases, leading to an ever-restricted therapeutic choice. In *Pseudomonas aeruginosa*, resistance to β -lactam agents can be due to the overproduction of chromosome-encoded cephalosporinase (23), to the alteration of the outer membrane protein OprD (36, 37), to the overexpression of the efflux system (36, 37), and to the acquisition of exogenous β -lactamases (28). Among these, class B carbapenemases (i.e., VIM and IMP) and to a lesser extent some class A extended-spectrum β -lactamases (ESBLs) (i.e., GES, KPC) have been involved in the resistance of *P. aeruginosa* to carbapenems and are considered the transferable resistance determinants having the highest clinical impact on antimicrobial therapy in hospitals worldwide (41).

Carbapenem-hydrolyzing class D β -lactamases (OXA-23, OXA-40, OXA-58, OXA-143) (CHDLs) are almost exclusively found in multidrug- and carbapenem-resistant *Acinetobacter baumannii* (6, 14, 16, 18, 34). In *P. aeruginosa*, to the best of our knowledge, only OXA-40 has been detected in two clonally unrelated clinical isolates resistant to imipenem in Spain (38, 39). In these two isolates, the *bla*_{OXA-40} gene was located on a 32-kb plasmid also found in the *A. baumannii* SM28 strain isolated in the same hospital (25, 38, 39).

In this study, we have identified a new CHDL, OXA-198, that belongs to a new group of class D β -lactamases in a clinical isolate of *P. aeruginosa*.

MATERIALS AND METHODS

Bacterial strains and antimicrobial susceptibility. A *P. aeruginosa* clinical isolate (PA41437) was recovered from the culture of a sputum specimen of an elderly patient hospitalized at the UCL Mont-Godinne University Hospital (Yvoir, Belgium) in April 2010. Bacterial species identification was confirmed by

matrix-assisted laser desorption ionization–time of flight mass spectrometry (MALDI-TOF MS) with a Microflex LT mass spectrometer (Bruker Daltonik GmbH, Leipzig, Germany). *P. aeruginosa* reference strain PAO1 (40), *P. aeruginosa* PU21 (19), and *Escherichia coli* TOP10 (Invitrogen, Merelbeke, Belgium) were used for plasmid transfer and/or cloning experiments. Susceptibility to antimicrobials was determined by Etest (bioMérieux, Brussels, Belgium) on Mueller-Hinton (MH) agar plates (Bio-Rad, Nazareth, Belgium) and interpreted according to CLSI breakpoints (10).

PCR sequencing of *bla* and of porin-coding genes. Detection of OXA β -lactamase genes (*bla*_{OXA-1} group, *bla*_{OXA-2} group, *bla*_{OXA-10} group, *bla*_{OXA-9}, *bla*_{OXA-20}, *bla*_{OXA-18}), penicillinase genes (*bla*_{PSE/CARB}), and genes encoding class A ESBL enzymes (GES, VEB, PER, BEL) and metallo- β -lactamases (VIM, IMP) was achieved by PCR as previously described (4, 5). The genetic context of the *bla*_{OXA-198} gene was assessed by PCR sequencing of the variable region of the integron by using primers designed on the basis of the 5' and 3' conserved segments (CS) of class 1 integron (22) and by primer walking sequencing performed directly on the purified p41437 plasmid using an external sequencing service (Macrogen, Seoul, South Korea).

Nucleotide sequences were analyzed with the BLASTN algorithm available from the National Center for Biotechnology Information (<http://www.ncbi.nlm.nih.gov/BLAST/>). PCR sequencing of the gene encoding the outer membrane protein OprD was achieved with primers oprD-FW (5'-CTTCCTTTATAGGC GCGTTG-3'), oprD-RV (5'-AACATAAGACATGCCGTGGA-3'), and oprD1 to oprD4 (15).

Plasmid analysis, mating out, and electroporation experiments. Plasmid DNA was extracted by the Kieser method (20) or by the QIAfilter plasmid midikit (Qiagen, Venlo, The Netherlands). *E. coli* NCTC50192, harboring four plasmids of 154, 66, 38, and 7 kb, was used as a plasmid size marker. Plasmid DNA was analyzed by electrophoresis with a 0.7% agarose gel. Determination of the incompatibility groups of plasmids was done as described by Carattoli et al. (7), as the authors were able to type some *P. aeruginosa* plasmids belonging to IncP and IncA/C. The transfer of the resistance marker from *P. aeruginosa* (PA41437) to *P. aeruginosa* PU21 (rifampin resistant) was attempted by solid and liquid mating out assays at 37°C (24). Selection was performed on brain heart infusion (BHI) agar plates supplemented with ticarcillin at 100 μ g/ml and rifampin at 100 μ g/ml. Moreover, plasmid DNA extracts from *P. aeruginosa* PA41437 were electroporated into *P. aeruginosa* PAO1 and in *E. coli* TOP10 using a Gene Pulser II (Bio-Rad, Marnes-la-Coquette, France) as previously described (3, 8), and transformants were selected on BHI agar plates containing 100 μ g/ml of ticarcillin.

Cloning of OXA-198 β -lactamase. For cloning of OXA-198, the *bla*_{OXA-198} gene was amplified with primers OXA-41437FW (5'-CTCGAATTCATGCATA AACACATGAGTAAG-3' [EcoRI site underlined]) and OXA-41437RV (5'-CTCAAGCTTTTATTCGATGATCCCCCTTT-3' [HindIII site underlined]) and cloned in the shuttle vector pUCP24 (43) in chemically competent *E. coli* TOP10 cells (Invitrogen, Merelbeke, Belgium). The sequence of the cloned *bla*_{OXA-198}

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TABLE 1. Antimicrobial susceptibility determination for *P. aeruginosa* clinical isolate, wild-type *P. aeruginosa* PAO1, PAO1 harboring natural plasmid p41437, PAO1 harboring recombinant plasmid pOXA-198, *E. coli* reference strain TOP10, and TOP10 harboring pOXA-198 or p41437

Antibiotic ^a	MIC ^b (μ g/ml)						
	<i>P. aeruginosa</i>				<i>E. coli</i>		
	PA41437	PAO1	PAO1 p41437	PAO1 pOXA-198	TOP10	TOP10 p41437	TOP10 pOXA-198
TIM	>256	24	>256	>256	2	>256	>256
PIP	24	6	32	24	1.5	4	32
TZP	16	3	24	24	1	4	16
FEP	24	3	12	12	0.064	0.064	0.125
CAZ	1.5	2	2	2	0.125	0.125	0.125
IPM	>32	1.5	3	3	0.38	0.5	0.5
MEM	12	0.38	1.5	1.5	0.032	0.047	0.032
ATM	4	4	4	4	0.032	0.047	0.047
GEN	12	1.5	6	>256 ^c	0.25	0.5	32 ^c
TOB	96	0.5	32	0.75	0.38	6	0.75
AMK	96	3	64	3	0.75	12	0.75
CIP	>32	0.094	0.094	0.094	<0.002	<0.002	<0.002
CST	1.5	1.5	2	2	0.5	0.5	0.5

^a TIM, ticarcillin-clavulanic acid; PIP, piperacillin; TZP, piperacillin-tazobactam; FEP, cefepime; CAZ, ceftazidime; IPM, imipenem; MEM, meropenem; ATM, aztreonam; GEN, gentamicin; TOB, tobramycin; AMK, amikacin; CIP, ciprofloxacin; CST, colistin.

^b MICs were determined by Etest. Values in boldface correspond to significant changes of MICs.

^c High gentamicin MICs are due to the selecting marker (AAC(3)-I) of the pUCP24 shuttle vector.

PCR-generated DNA fragments was confirmed by sequencing on both strands. The obtained plasmid, pOXA-198, was then electroporated in the wild-type reference *P. aeruginosa* PAO1 as previously described (3, 8), and transformants were selected on BHI agar plates containing 100 μ g/ml of ticarcillin and 20 μ g/ml of gentamicin (selecting marker of pUCP24).

Isoelectric focusing analysis. Analytical isoelectric focusing (IEF) was performed on freeze-thaw crude culture extracts with an ampholine polyacrylamide gel on pH 3 to 9 PhastGel medium with Phastsystem (Amersham Biosciences, Uppsala, Sweden) according to the manufacturer's protocol, with the only exception being that the migration step was performed at 500 V instead of 2,000 V.

β -Lactamase extraction and purification. OXA-198 β -lactamase was extracted and purified as previously described (18, 33, 34) with slight modifications. Briefly, *E. coli* TOP10 harboring recombinant plasmid pOXA-198 was grown until A_{600} reached 1 at 37°C in 4 liters of BHI broth (BD, Erembodegem, Belgium) containing 100 μ g/ml of amoxicillin and 15 μ g/ml of gentamicin. The bacterial suspension was pelleted, resuspended in 100 mM phosphate buffer (pH 7.0), and lysed using a cell disrupter Emulsiflex-C3 (Avestin, Mannheim, Germany). The crude extract was then treated with benzonase endonuclease (Merck, Darmstadt, Germany) as recommended by the supplier and centrifuged at 16,000 \times g for 30 min at 4°C. The supernatant was dialyzed against 20 mM Tris-H₂SO₄ buffer (pH 8.0), filtered through a 0.45- μ m filter, and loaded on a Q-Sepharose ion exchange chromatography column equilibrated with the same buffer. The β -lactamase was recovered in the flowthrough. The extract was then dialyzed in 20 mM CHES (*N*-cyclohexyl-2-aminoethanesulfonic acid) buffer (pH 10.0) and loaded again onto the Q-Sepharose column equilibrated with the same buffer. Elution was performed with a K₂SO₄ gradient. The fractions containing the highest β -lactamase activities were pooled and dialyzed against 100 mM phosphate buffer (pH 7.0). Enzyme purity was estimated by sodium dodecyl sulfate-polyacrylamide gel electrophoresis. The protein content was measured by the Bio-Rad DC protein assay (Bio-Rad, Nazareth, Belgium).

Kinetic studies. Purified β -lactamase kinetic parameters (k_{cat} and K_m) were determined by UV spectrophotometry at 30°C in 100 mM sodium phosphate (pH 7.0) as previously described (13, 31). The 50% inhibitory concentration (IC₅₀) was determined as the inhibitor (clavulanate or NaCl) concentration that reduced the hydrolysis rate of 100 μ M nitrocefin by 50% under conditions in which the enzyme was preincubated with various concentrations of inhibitor for 3 min at 30°C before the addition of the substrate (31, 33). Specific activity of the β -lactamase OXA-198 was defined as the amount (unit of enzyme) that hydrolyzed 1 μ mol of benzylpenicillin/minute/milligram of protein.

Microbiological assay of carbapenemase activity. Carbapenem hydrolysis activity of OXA-198 was phenotypically investigated on crude extract of PA41437 with a modified method of Masuda et al., as previously described (6, 26). Briefly, an MH agar plate was inoculated with the ATCC 25922 *E. coli* strain, and a disk of imipenem (10 μ g) or meropenem (10 μ g) was placed at the center of the plate.

Four filter paper disks containing 20, 10, and 5 μ l of crude extract or 20 μ l of sodium phosphate buffer (pH 7.0) were placed 15 mm and 25 mm from the imipenem and meropenem disks, respectively. Plates were incubated overnight at 37°C, and inactivation of imipenem or meropenem was shown by growth of the ATCC 25922 strain within the expected inhibition zone.

Nucleotide sequence accession number. The nucleotide sequence data reported in this paper have been assigned to the EMBL/GenBank nucleotide sequence database under accession number HQ634775.

RESULTS

Origin of the clinical isolate and antibiotic susceptibility.

The *P. aeruginosa* PA41437 strain was isolated from consecutive lower-respiratory-tract specimens of a 79-year-old man with no travel history and who had been admitted at the UCL Mont-Godinne University Hospital in April 2010 for acute exacerbation of chronic bronchitis. Initial sputum specimens yielded the presence of a multidrug-resistant *Enterobacter aerogenes* strain in pure culture, and the patient was treated for 8 days with meropenem (1 g three times a day [TID]). Subsequently, the patient had to be transferred to the intensive care unit (ICU), where he had to be intubated and mechanically ventilated because of severe hypoxemia and major acute respiratory failure. During his stay in the ICU, he developed an aspiration pneumonia, and repeated quantitative cultures of endotracheal aspirates yielded the presence of *P. aeruginosa* (PA41437) in significant amounts (>10⁶ CFU/ml). The *in vitro* susceptibility pattern of the *P. aeruginosa* PA41437 isolate is shown in Table 1. PA41437 was resistant to most β -lactams, including imipenem, but it presented an intermediate susceptibility to meropenem. However, it remained susceptible to ceftazidime and to aztreonam. This strain was also resistant to aminoglycosides (gentamicin, amikacin, and tobramycin) and to ciprofloxacin. Following obtention of the microbiological results, the patient was treated with ceftazidime (2 g TID) for one additional week, but he eventually died due to acute renal failure and cardiac insufficiency.

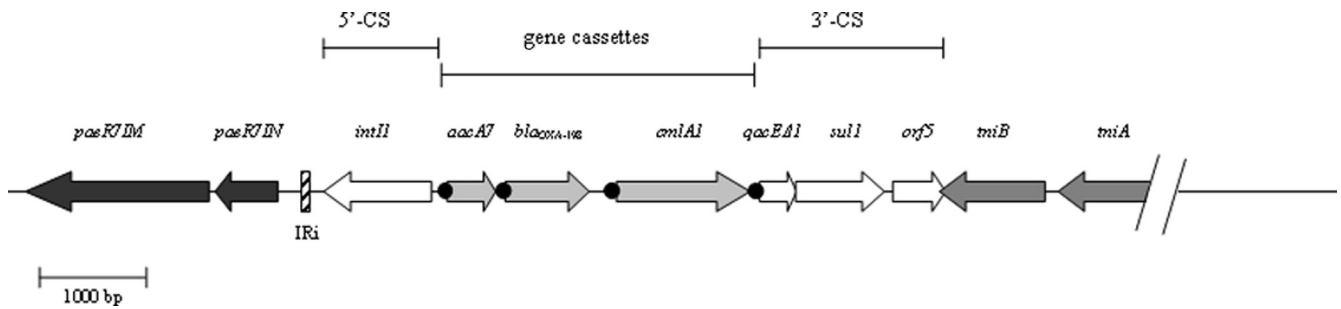


FIG. 1. Schematic representation of the genetic environment of the *bla*_{OXA-198} gene. The 5' conserved segment (5'-CS) contains the integrase gene *intI1*, while the 3'-CS contains the *qacEΔ1*, *sulI*, and *orf5* genes typical of class I integrons. Filled circles indicate *attC* sites. Arrows indicate the direction of transcription of the coding regions. IRI of the Tn6060-like sequence is shown. *paeR7IM* and *paeR7IN* code for an adenine-specific methyltransferase and a DNA invertase, respectively. Genes *tniB* and *tniA*, involved in transposition, are located downstream of *orf5*.

Plasmid content, conjugation experiment, and β -lactamase analysis. Extraction of plasmid content of clinical strain PA41437 by the Kieser method revealed the presence of a single nontypeable plasmid, p41437, of ca. 46 kb (data not shown). Transfer of p41437 in wild-type *P. aeruginosa* PAO1 or in wild-type *E. coli* TOP10 by electroporation led to transformants resistant to ticarcillin, tobramycin, and amikacin and with reduced susceptibility to piperacillin, piperacillin-tazobactam, carbapenems, cefepime, and gentamicin (Table 1). A conjugation experiment with clinical strain PA41437 or PAO1 p41437 as the donor and rifampin-resistant *P. aeruginosa* PU21 as the recipient failed to yield transconjugants. β -Lactamase content analysis of culture extracts of the PA41437 clinical strain or of PAO1 transformants harboring p41437 by isoelectric focusing (IEF) showed a β -lactamase band with a pI of 6.4. All PCR screening assays were negative for metallo- β -lactamases, ESBLs, and class D β -lactamases (4, 5) (data not shown), leading to the hypothesis that the p41437 plasmid harbored an unknown β -lactamase.

Cloning and sequencing of OXA-198. PCR was done on p41437 plasmid DNA with consensus primers targeting 5' CS and 3' CS of class 1 integrons (22). A 3,000 bp PCR fragment was amplified and yielded three open reading frames (ORFs): (i) an *aacA7* gene which encodes the AAC(6')-Ib enzyme; (ii) an undescribed 789-bp *bla*_{OXA} gene (GC content of 47.8%), named *bla*_{OXA-198}, which encodes a β -lactamase with 83% amino acid identity with a β -lactamase of *Chlorobaculum parvum* NCIB 8327; and (iii) a *cmlA1* gene which encodes a chloramphenicol efflux protein. Analysis of the genetic environment of *bla*_{OXA-198} was performed by primer walking on p41437. Figure 1 shows that *bla*_{OXA-198} was part of a class 1 integron very similar to that described in the transposon Tn6060 of a clinical strain of *P. aeruginosa* (9). Upstream of the *aacA7* gene, the *intI1* gene, coding for an integrase, was found. Upstream of the 5' CS end of the integron, we found IRI inverted repeats flanked by *paeR7IM* and *paeR7IN* genes (accession number DQ839391) coding for an adenine-specific methyltransferase and a DNA invertase, respectively. Downstream of the *bla*_{OXA-198} gene cassette and the *cmlA1* gene, we found the 3' CS fragment containing *qacEΔ1*, *sulI*, and *orf5* genes. The *tniB* and *tniA* genes involved in transposition were located downstream of *orf5* (Fig. 1).

*bla*_{OXA-198} encodes a 262-amino-acid protein with a theoretical molecular mass of 30,073 Da, including the signal peptide.

Within the deduced amino acid sequence, the five typical sequence motifs of class D β -lactamases were found, namely, ₇₀STFK₇₃, ₁₁₈SXV₁₂₀, ₁₄₄YGX₁₄₆, _W₁₆₄, and ₂₁₆KTG₂₁₈ (21, 35, 42) (Fig. 2). OXA-198 was weakly related to other carbapenem-hydrolyzing class D β -lactamases sharing 35%, 32%, and 30% amino acid identity with OXA-48, -58, and -143, respectively, and 29% with OXA-51, -23, and -40 (Fig. 3). Cloning of *bla*_{OXA-198} in the shuttle vector pUCP24 and expression in *E. coli* TOP10 or *P. aeruginosa* PAO1 led to transformants with reduced susceptibility to ticarcillin, piperacillin, piperacillin-tazobactam, carbapenems, and cefepime (Table 1). However, the MIC values of carbapenems for the transformants remained lower than that observed for the clinical isolate PA41437, suggesting that other mechanisms could account for the resistance to carbapenems. Sequencing of the gene coding for the porin OprD in the clinical strain compared to that of PAO1 revealed several mutations and a deletion of 88 nucleotides, which led to a truncated protein lacking the first 110 amino acids (data not shown).

Biochemical properties of β -lactamase OXA-198. After purification from extracts of *E. coli* TOP10 harboring recombinant plasmid pOXA-198, the specific activity of OXA-198 against benzylpenicillin was 2.2 U/mg of protein, and its purification factor was 210-fold. The protein purity was estimated to be >95% by sodium dodecyl sulfate-polyacrylamide gel electrophoresis analysis (data not shown). OXA-198 has a narrow-spectrum hydrolysis profile, including mostly penicillins (Table 2). No activity could be detected against ceftaxime, ceftazidime, cefepime, and aztreonam, as observed for other CHDLs. The rates of imipenem and meropenem hydrolysis were low, with k_{cat} values of 0.1 s⁻¹ and 0.01 s⁻¹ for imipenem and meropenem, respectively (Table 2), although the MICs of both carbapenems for OXA-198-expressing PAO1 were 2- to 3-fold increased (Table 1). As previously described for other CHDLs, hydrolysis of imipenem was faster than that of meropenem (35). Determination of K_m values for imipenem and meropenem (0.15 and 0.006 μ M, respectively) showed that OXA-198 has a strong affinity for carbapenems even if hydrolysis is low, resulting in a high k_{cat}/K_m ratio (670 and 1,670 mM⁻¹ s⁻¹, respectively). In comparison to values obtained for benzylpenicillin, the k_{cat}/K_m ratios for meropenem and imipenem were 1.5-fold higher and 2-fold lower, respectively (Table 2). A positive modified Masuda test (6, 26) performed on a crude sonicated extract of *E. coli* TOP10 expressing OXA-

	10		20		30		40
OXA-198	MHKHMSKLF I	AFLAFLLSVP	AAA-----	-----	EDQTLAELFA	QQGIDGTIVI	
OXA-48	-MRVLALSAV	FLVASIIGMP	AVAKE-----	-----WQ	ENKSNWAHFT	EHKSQGVVVL	
OXA-58	-MKLLKILSL	VCLSSISIGAC	AEHMSRAKT	STIPQVNSI	IDQNVQALFN	EISADAVFVT	
OXA-143	-MKKF-ILPI	LSISTLLSVS	ACSSIQTKFE	DTFHTS-NQQ	HEKAIKSYFD	EAQTQGVIII	
OXA-51	-MN---IKTL	LLITSAIFIS	ACSPYIVTAN	PNHSASKSDE	KAEEKIKLNF	EVHTTGLVVI	
OXA-23	-MNKY-FTCY	VVAS--LFLS	GCT-VQHNLI	NETPSQIVQG	HNQVIHQYFD	EKNTSGVLVI	
OXA-40	-MKKF-ILPI	FSISILVSL	ACSSIQTKSE	DNFHIS-SQQ	HEKAIKSYFD	EAQTQGVIII	
	50	60	70	80	90	100	
OXA-198	SSLHNGKTFI	HNDPRAKQRF	STASTFKILN	TLISLEEKAI	SGKDDVLKWD	GHIYDFPDWN	
OXA-48	WNENKQQGFT	NNLKRANQAF	LPASTFKIPN	SLIALDLGVV	KDEHQVFKWD	GQTRDIATWN	
OXA-58	YDGGNIKKYG	THLDRAKTAY	IPASTFKIAN	ALIGLENHKA	TST-EIFKWD	EKPRFFKAWD	
OXA-143	KKGKNIITYG	NNLTRAHTEY	VPASTFKMLN	ALIGLENHKA	TTT-EIFKWD	GKKRSYPMWE	
OXA-51	QQGQTQQSYG	NDLARASTEY	VPASTFKMLN	ALIGLEHKA	TTT-EVFKWD	GQKRLPEWE	
OXA-23	QTDKKINLYG	NALSRANTEY	VPASTFKMLN	ALIGLENQKT	DIN-EIFKWD	GEKRSFTAWE	
OXA-40	KEGKNLSTYG	NALARANKEY	VPASTFKMLN	ALIGLENHKA	TTN-EIFKWD	GKKRTYPMWE	
	110	120	130	140	150	164	170
OXA-198	RDQTLESFAFK	VSCVWCYQAL	ARQVGAEKYR	NYLRSKVYGE	LREPFEETTF	WLDGSLQISA	
OXA-48	RDHNLITAMK	YSVVPVYQEF	ARQIGEARMS	KMLHAFDYGN	EDISGMVDSF	WLDGGIRISA	
OXA-58	KDFTLGEAMQ	ASTVPVYQEL	ARRIGPSLMQ	SELQIRIGYGN	MQIGTEVDQF	WLKGPLTITP	
OXA-143	KDMTLGDAMA	LSAVPVYQEL	ARRTGLDLMQ	KEVKRVGFNG	MNIGTQVDNF	WLVGPKLITP	
OXA-51	KDMTLGDAMK	ASAIPVYQDL	ARRIGLELMS	KEVKRVGYGN	ADIGTQVDNF	WLVGPKLITP	
OXA-23	KDMTLGEAMK	LSAVPVYQEL	ARRIGLDMQ	KEVKRIGFNG	AEIGQVDNF	WLVGPKLITP	
OXA-40	KDMTLGEAMA	LSAVPVYQEL	ARRTGLELMQ	KEVKRVNFGN	TNIGTQVDNF	WLVGPKLITP	
	180	190	200	210	220	230	
OXA-198	IEQVNFLKKV	HLRTLPPFSAS	SYETLRQIML	IEQTPAFTLR	AKTGWATRVK	PQVGWYVGHV	
OXA-48	TEQISFLRKL	YHNKLHVSEK	SQRIVKQAML	TEANGDYIIR	AKTGYSTRIE	PKIGWVVGWV	
OXA-58	IQEVKVVYDL	AQQQLPFKPE	VQQQVKEMLY	VERRGENRLY	AKSGWGMADV	PQVGWYVGFV	
OXA-143	IQEVNFADDF	AMNRLPFKLE	TQEEVKKMLL	IKEFNGSKIY	AKSGWGMADV	PQVGWLTGWV	
OXA-51	QQEAQFAYKL	ANKTLPPSPK	VQDEVQSMFL	IEEKNGNKIY	AKSGWGDVDV	PQVGWLTGWV	
OXA-23	IQEVFVSQL	AHTQLPFSEK	VQANVKNMML	LEESNGYKIF	GKTGWAMDIK	PQVGWLTGWV	
OXA-40	VQEVNFADDF	AHNRLPFKLE	TQEEVKKMLL	IKEVNGSKIY	AKSGWGMGVT	PQVGWLTGWV	
	240	250	260	270			
OXA-198	ETPTD-VWFF	ATNIEVRDEK	DLPLRQKLTR	KALQAKGII	--		
OXA-48	ELDDN-VWFF	AMNMDMPTSD	GLGLRQAITK	EVLKQEKIIP	--		
OXA-58	EKADGQVAV	ALNMQMKAGD	DIALRKLQSL	DVLDKLGWVF	YL		
OXA-143	EKSNGEKVAF	SLNIEMKQGM	PGSIRNEITY	KSLENLGI	--		
OXA-51	VQPQGNIVAF	SLNLEMKGI	PSSVRKEITY	KSLEQLGIL	--		
OXA-23	EQPDGKIVAF	ALNMEMRSEM	PASIRNELLM	KSLKQLNII	--		
OXA-40	EQANGKIPF	SLNLEMKEGM	SGSIRNEITY	KSLENLGI	--		

FIG. 2. Comparison of the OXA-198 amino acid sequence with those of carbapenem-hydrolyzing class D β-lactamases OXA-48, -58, -143, -51, -23, and -24/40. Conserved residues are shaded. β-Lactamases are numbered according to the Ambler class D β-lactamase numbering system (11).

198 and obtained with imipenem or meropenem confirmed imipenem or meropenem hydrolysis by OXA-198 (Fig. 4). In general, the catalytic activities of OXA-198 were in the same range as those of others CHDLs, such as OXA-58 or OXA-40 (34). Studies of activity inhibition, as measured by IC₅₀ determination, showed that OXA-198 was inhibited by NaCl (IC₅₀, 37 mM) like many other class D β-lactamases (30, 35). Interestingly, OXA-198 activity was also inhibited by clavulanic acid (IC₅₀, 7 μM), with concentrations in the same range as that reported for OXA-163, another CHDL with a weak activity against carbapenems (32).

DISCUSSION

In this study, we have characterized a new carbapenem-hydrolyzing class D β-lactamase, OXA-198, in *P. aeruginosa*. This class D β-lactamase was only weakly related to other CHDLs, and it constitutes a novel subclass of CHDLs. OXA-198 β-lactamase hydrolyzed penicillins and to a lesser extent carbapenems but had no significant hydrolytic activity against

expanded-spectrum cephalosporins, as also observed with other CHDLs (6, 17, 18, 34). It is, however, likely that OXA-198 may contribute to decreased susceptibility to carbapenem, as confirmed by the modified phenotypic Masuda test (6, 26). Comparison of the resistance phenotype of *P. aeruginosa* clinical strain PA41437 and of its transformants highlighted that additional mechanisms were responsible for the high level of carbapenem resistance. This was substantiated by the observation of various mutations and of a deletion of 88 nucleotides in the *oprD* gene which led to a truncated protein lacking the first 110 amino acids (data not shown). Our study constitutes the second description of a carbapenem-hydrolyzing class D β-lactamase in *P. aeruginosa*, after that of Sevillano et al., who reported the presence of OXA-40 on a plasmid also carried by *A. baumannii* (38).

Motifs YGN or FGN at Ambler positions 144 to 146 are characteristic of class D β-lactamase and are involved in the formation of the so-called “oxyanion hole,” a region in space responsible for the stability of enzymatic intermediates and

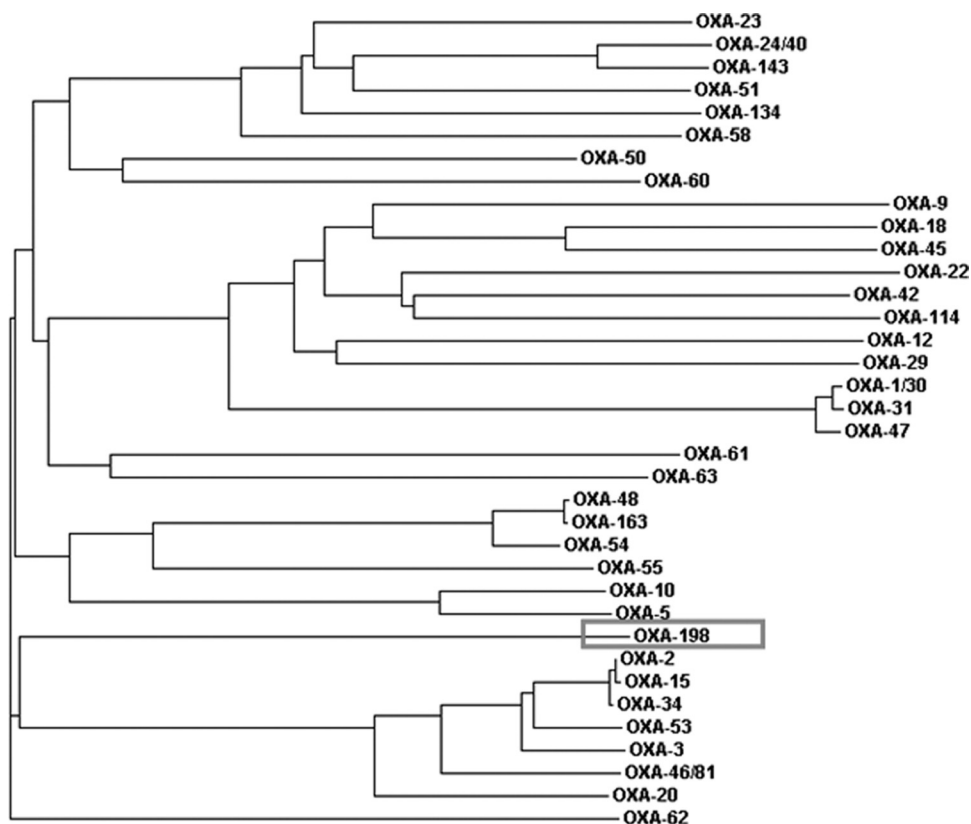


FIG. 3. Dendrogram obtained for representative class D β -lactamases. The alignment used for tree calculation was performed with the ClustalW2 multiple sequence alignment program. Branch lengths are drawn to scale and are proportional to the number of amino acid changes. The distance along the vertical axis has no significance.

also described as responsible for much of the catalytic efficiency of the serine enzyme (12, 27, 29). For the first time, in OXA-198, a YGE motif was found instead of the classical motif YGN or FGN. Site-directed mutagenesis experiments are needed to explore the catalytic consequences of the replacement of the polar asparagine (N) by the glutamic acid (E) at this position.

OXA-198 is relatively distant from other carbapenem- and

non-carbapenem-hydrolyzing class D β -lactamases. Interestingly, it shares 83% sequence amino acid identity with the native β -lactamase of *Chlorobaculum parvum*, a member of the green sulfur bacteria of the phylum *Chlorobi*, which are characteristically found in stratified lakes, microbial mats, and sul-

TABLE 2. Kinetic parameters of purified β -lactamase OXA-198

Substrate	OXA-198 ^a		
	k_{cat} (s ⁻¹)	K_m ^b (μ M)	k_{cat}/K_m (mM ⁻¹ s ⁻¹)
Benzylpenicillin	15	14	1,070
Ampicillin	37	216	170
Piperacillin	2.6	35	74
Oxacillin	25	30	830
Cephalothin	0.19	12	16
Cefotaxime	ND	—	—
Ceftazidime	ND	—	—
Cefepime	ND	—	—
Aztreonam	ND	—	—
Imipenem	0.1	0.15	670
Meropenem	0.01	0.006	1,670

^a Data are means from three independent experiments. Standard deviations were within 10% of the means. ND, no detectable hydrolysis (<0.01 s⁻¹); —, not determinable.

^b Determined as K_i using nitrocefim as the substrate (13).

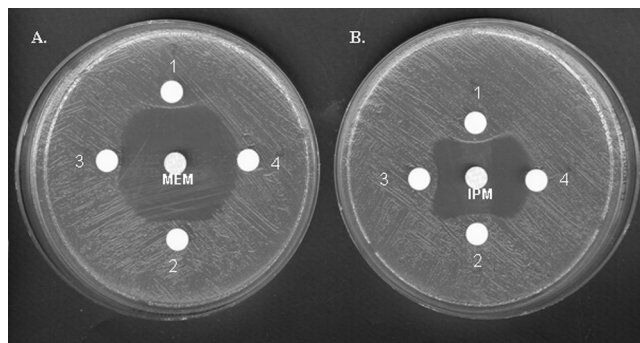


FIG. 4. Microbiological assay plate showing inactivation of meropenem (A) (central disk) or imipenem (B) by the OXA-198 enzyme. MH agar plates were inoculated with the carbapenem-susceptible *E. coli* ATCC25922 strain. A central disk of meropenem (10 μ g) or imipenem (10 μ g) was put on the dish with 4 satellite disks containing 20 μ l of crude sonicated extract of *E. coli* TOP10 expressing OXA-198 (nitrocefim-specific activity, 1.05 μ mol/min/mg) (1), 10 μ l of extract (2), 5 μ l of extract (3), and 20 μ l of phosphate-buffered saline (4). MEM, meropenem; IPM, imipenem.

vide-rich hot springs (1), suggesting the possible environmental acquisition of this new class D β -lactamase by *P. aeruginosa*. Acquisition of this β -lactamase by *P. aeruginosa* on a class 1 integron carried by a plasmid which can easily be transformed in *P. aeruginosa* or *E. coli* could lead to the dissemination of this class D β -lactamase. Moreover, this integron is very similar to that carried by the Tn6060 transposon found on a genomic island in a VIM-1-positive clinical strain of *P. aeruginosa* and easily transposable to a conjugative plasmid (9).

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