

# Nationwide Investigation of Extended-Spectrum $\beta$ -Lactamases, Metallo- $\beta$ -Lactamases, and Extended-Spectrum Oxacillinases Produced by Ceftazidime-Resistant *Pseudomonas aeruginosa* Strains in France<sup>∇</sup>

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**A nationwide study aimed to identify the extended-spectrum  $\beta$ -lactamases (ESBLs), metallo- $\beta$ -lactamases (MBLs), and extended-spectrum oxacillinases (ES-OXAs) in a French collection of 140 clinical *Pseudomonas aeruginosa* isolates highly resistant to ceftazidime. Six ESBLs (PER-1,  $n = 3$ ; SHV-2a,  $n = 2$ ; VEB-1a,  $n = 1$ ), four MBLs (VIM-2,  $n = 3$ ; IMP-18,  $n = 1$ ), and five ES-OXAs (OXA-19,  $n = 4$ ; OXA-28,  $n = 1$ ) were identified in 13 isolates (9.3% of the collection). The prevalence of these enzymes is still low in French clinical *P. aeruginosa* isolates but deserves to be closely monitored.**

*Pseudomonas aeruginosa* could potentially become resistant to any of the antibiotics used to treat Gram-negative nosocomial infections. The development of resistance to  $\beta$ -lactams in this opportunistic pathogen results from mutations leading to stable overexpression of intrinsic cephalosporinase AmpC, overproduction of efflux systems, reduced permeability, acquisition of transferable genes coding for a variety of secondary  $\beta$ -lactamases, or a combination of these mechanisms (21). A growing number of Ambler class A extended-spectrum  $\beta$ -lactamases (ESBLs), class B carbapenemases (metallo- $\beta$ -lactamases [MBLs]), and class D extended-spectrum oxacillinases (ES-OXAs) have been reported in clinical strains of *P. aeruginosa* (14, 18, 19, 34, 40). The present multicenter study gave a snapshot of these acquired enzymes in a French collection of 140 *P. aeruginosa* isolates highly resistant to ceftazidime.

During a 1-month period (June 2007), 85 hospital laboratories participating in the surveillance networks affiliated with ONERBA (Observatoire National de l'Epidémiologie de la Résistance Bactérienne aux Antibiotiques) collected nonredundant strains of *P. aeruginosa* resistant to ceftazidime (Caz<sup>r</sup>) (as defined by the Comité de l'Antibiogramme de la Société Française de Microbiologie [CA-SFM] in 2006 [12]), except those obtained from screening samples and cystic fibrosis patients. The susceptibility tests were performed in each laboratory according to their routine testing methods. All isolates showing an inhibition zone of <15 mm around the ceftazidime-containing disk (30  $\mu$ g) or with a MIC of ceftazidime of >32  $\mu$ g/ml were sent to a central laboratory for further investigation. In addition, the total number of patients with at least one clinical specimen positive for *P. aeruginosa* as well as the number of hospitalization days was recorded in each partici-

pating center during the study period. The central laboratory confirmed bacterial identification by using API32GN strips (bioMérieux, Craponnes, France) and determined the MICs of eight antipseudomonal antibiotics by the conventional 2-fold dilution method in agar (26). The  $\beta$ -lactamase contents of the strains were first analyzed by isoelectric focusing (IEF) (23) and then confirmed by gene sequencing with consensus primers targeting the *bla*<sub>TEM</sub>, *bla*<sub>PSE</sub>, *bla*<sub>SHV</sub>, *bla*<sub>PER</sub>, *bla*<sub>VEB</sub>, *bla*<sub>GES</sub>, *bla*<sub>BEL</sub>, *bla*<sub>CTX-M</sub>, *bla*<sub>VIM</sub>, *bla*<sub>SPM</sub>, *bla*<sub>OXA-I</sub> group, *bla*<sub>OXA-II</sub> group, *bla*<sub>OXA-III</sub> group, and *bla*<sub>OXA-18</sub> genes (1, 3, 5, 6, 24, 25, 28, 30–32, 35, 38). Genes *bla*<sub>IMP</sub>, *bla*<sub>GIM</sub>, and *bla*<sub>OXA-9</sub>, respectively, were also specifically amplified with primers IMP2004-A and IMP2004-B (5'-ACAYGGYTTGG TTGTTCTTG-3' and 5'-GGTTTAAAYAAAACAACCACC-3', respectively), GIM-A and GIM-B (5'-GGAGTATATCTT CATACCTCC-3' and 5'-TTCCAACCTTGCCATGCCCC-3', respectively), and OXA9A and OXA9B (5'-CCGAGAGATC GCACATACAA-3' and 5'-CCCATCAACACGGGTAATT C-3', respectively). Class 1 integrons were amplified in the isolates producing ESBLs, MBLs, and ES-OXAs with consensus primers (20) for content analysis and *bla*<sub>ESBL</sub>, *bla*<sub>MBL</sub>, and *bla*<sub>ES-OXA</sub> localization. Purified amplicons were sequenced on both strands, and their nucleotide sequences were compared and aligned with reference sequences using the NCBI BLAST program (2). Clonality of the Caz<sup>r</sup> isolates was investigated by pulsed-field gel electrophoresis (PFGE) of DraI macrorestricted genomic DNA (36, 37).

**Incidence of *P. aeruginosa* infections.** Eighty-five hospital laboratories from 70 cities in France were enrolled in the study (Fig. 1). With 58,022 beds, the participating hospitals accounted for a total annual activity of 17 million hospital days. The total catchment area population was 8 million people, which corresponds to 13% of the French population. Public (university-affiliated or general) hospitals accounted for 95% of the hospital beds. During the 1-month study, the participating centers isolated 2,326 nonredundant isolates of *P. aeruginosa*.

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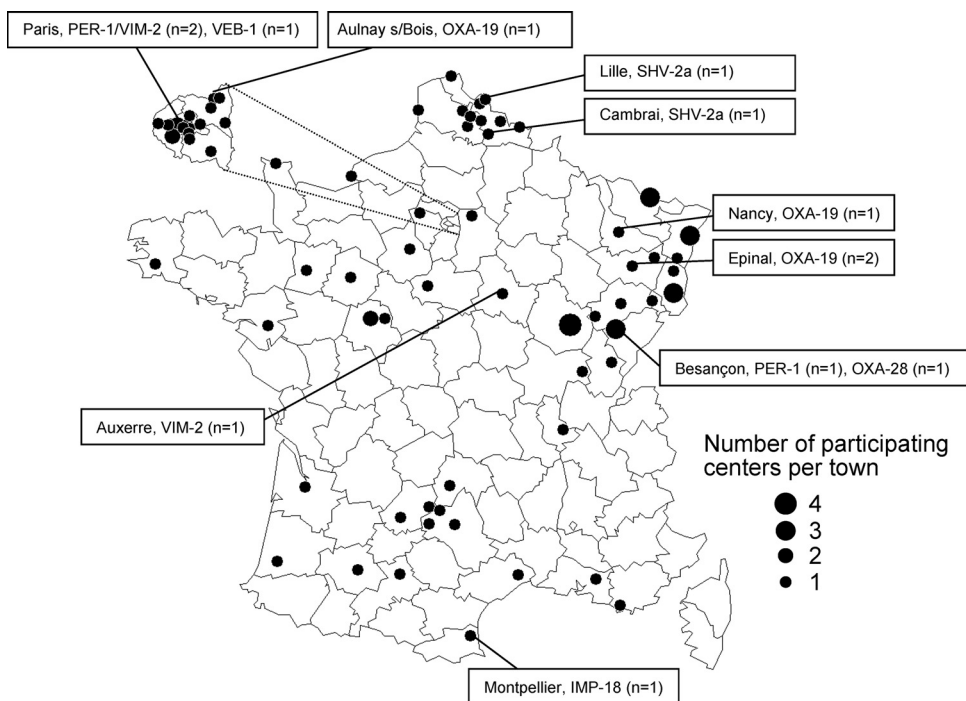


FIG. 1. Map of France, showing the 85 sites included in the study and the localization of isolates of *P. aeruginosa* producing ESBLs, MBLs, and ES-OXAs. An enlarged map of the Ile-de-France region is provided at the upper left. Labels indicate the town of isolation, the nature of the enzymes, and the number of isolates.

*nosa*, giving an attack rate of 0.76 cases per 100 admissions or a global incidence of 1.58 per 1,000 patient days. One hundred forty of these isolates (6.0%) appeared to be resistant to ceftazidime (MIC of >32  $\mu\text{g/ml}$ ). The resistance rates were similar between the university-affiliated (6.4%) and general (5.3%) hospitals, for a global incidence of *Caz<sup>r</sup> P. aeruginosa* isolates of 0.095 per 1,000 patient days.

**Secondary  $\beta$ -lactamases in *Caz<sup>r</sup> P. aeruginosa*.** The  $\beta$ -lactamases detected in the 140 *Caz<sup>r</sup>* isolates are indicated in Table 1. Six ESBLs, four MBLs, and five ES-OXAs were identified in 13 isolates, for an overall prevalence of 9.3% of the 140 *Caz<sup>r</sup>* isolates and 0.6% of the total isolates. Table 2 gives the resistance levels to antipseudomonal compounds and characteristics of the isolates producing these enzymes.

In our series, the overall prevalence of *P. aeruginosa* strains producing ESBLs or MBLs remains relatively low, far below that observed in some Asian countries (4, 10), Latin America (9), or Turkey (27) but in concordance with those observed in previous studies in France (7, 8, 13, 15–17, 22, 29, 39).

However, we showed here an unexpected proportion of *P. aeruginosa* strains producing ES-OXAs (OXA-19 and OXA-28). As expected, all of the *bla<sub>ES-OXA</sub>* genes were borne by class 1 integrons (Table 2) (33). Some new extended-spectrum oxacillinases have recently been described in several European countries (19, 33). Altogether, these data suggest the possible emergence of this class of enzymes in *P. aeruginosa*. *bla<sub>VIM</sub>* and *bla<sub>VEB</sub>* genes are usually carried on class 1 integrons (34, 40). However, *bla<sub>VIM-2</sub>*, in isolates P19 and P22, and *bla<sub>VEB-1</sub>*, in isolate P151, have not been associated with such genetic determinants.

**Genotyping.** One hundred thirty-seven isolates (3 isolates were nontypeable using PFGE) clustered in 113 PFGE patterns as follows: 98 unique patterns, 12 patterns including isolates from two patients, 1 pattern including isolates from 3 patients, 1 pattern including isolates from 4 patients, and 1 pattern including isolates from 8 patients. In most cases, the clonally related isolates were recovered from the same hospital

TABLE 1. Secondary  $\beta$ -lactamases detected in the collected *P. aeruginosa* isolates

Secondary $\beta$ -lactamase(s) <sup>a</sup>	No. of isolates
None .....	110
TEM-2 .....	5
PSE-1 .....	4
<b>OXA-19</b> .....	3
OXA-56 .....	3
OXA-10 .....	2
OXA-9 .....	2
SHV-2a .....	2
PSE-1, OXA-10 .....	1
<b>OXA-19, OXA-2</b> .....	1
<b>OXA-28</b> .....	1
<b>PER-1</b> .....	1
<b>VEB-1a, OXA-10</b> .....	1
<b>VIM-2</b> .....	1
<b>IMP-18</b> .....	1
OXA-30, PSE-1, <b>VIM-2, PER-1</b> .....	1
OXA-10, PSE-1, <b>VIM-2, PER-1</b> .....	1

<sup>a</sup> Secondary  $\beta$ -lactamases with an extended spectrum are shown in boldface type. In a given strain, the  $\beta$ -lactamases are ordered according to decreasing pI.

TABLE 2. Epidemiological data, clonal lineages, and resistance phenotypes of *P. aeruginosa* isolates producing ESBLs, MBLs, and ES-OXAs

Isolate	Origin	$\beta$ -Lactamase(s) (pI) <sup>a</sup>	Isolation site	MIC ( $\mu$ g/ml) <sup>b</sup>								PFGE pattern
				Tic	Tzpc	Caz	Fep	Atm	Ipm	Amk	Cip	
P9	Epinal	<b>OXA-19</b> (7.5)	Wound	512	128	256	64	16	16	<b>16</b>	64	A
P11	Epinal	<b>OXA-19</b> (7.5)	Wound	256	<b>64</b>	256	32	16	8	<b>8</b>	32	A
P122	Nancy	<b>OXA-19</b> (7.5)	Sputum	256	<b>64</b>	256	32	16	8	<b>8</b>	64	A
P66	Aulnay sous Bois	<b>OXA-2</b> (7.7), <b>OXA-19</b> (7.6)	Blood	512	<b>32</b>	64	16	16	<b>1</b>	128	128	C
P174	Besançon	<b>OXA-28</b> (7.8)	Blood	128	128	256	32	32	<b>4</b>	<b>16</b>	128	E
P19	Paris	OXA-30 (7.2), PSE-1 (5.7), VIM-2 (5.6), PER-1 (5.3)	Urine	>512	256	256	128	128	128	64	256	F
P22	Paris	OXA-10 (6.3), PSE-1 (5.7), VIM-2 (5.6), PER-1 (5.3)	Urine	>512	256	256	128	256	128	64	256	F
P170	Besançon	PER-1 (5.1)	Urine	512	128	256	64	128	<b>4</b>	32	64	G
P60	Cambrai	SHV-2a (7.4)	Sputum	>512	128	64	64	32	<b>1</b>	32	32	H
P102	Lille	SHV-2a (7.4)	Urine	>512	128	64	64	32	16	128	32	H
P151	Paris	VEB-1a (7.3), OXA-10 (6.3)	ETA <sup>c</sup>	>512	256	>512	>512	>512	32	128	512	I
P67	Auxerre	<b>VIM-2</b> (5.6)	Urine	>512	<b>32</b>	128	64	32	512	32	128	J
P85	Montpellier	<b>IMP-18</b> <sup>d</sup>	Wound	>512	128	>512	256	16	64	>512	128	K

<sup>a</sup> *bla* genes borne by class 1 integrons are indicated in boldface type. The gene cassettes and order for these isolates are as follows, with GenBank accession numbers given in parentheses: for P9, P11, and P122, *aacA4* and *bla*<sub>OXA-19</sub> (FJ906752); for P66, *bla*<sub>OXA-2</sub> and *bla*<sub>OXA-19</sub>; for P174, *aacA4* and *bla*<sub>OXA-28</sub> (FJ374756); for P67, *bla*<sub>VIM-2</sub>; and for P85, *dfrA22* and *bla*<sub>IMP-18</sub>.

<sup>b</sup> Tic, ticarcillin; Tzp, piperacillin-tazobactam; Caz, ceftazidime; Fep, cefepime; Atm, aztreonam; Ipm, imipenem; Amk, amikacin; Cip, ciprofloxacin. Drug susceptibility according to current NCCLS/CLSI breakpoints (26) is shown in boldface type: for Tic, MICs of  $\leq 64$   $\mu$ g/ml; for Tzp, MICs of  $\leq 64$   $\mu$ g/ml; for Caz, MICs of  $\leq 8$   $\mu$ g/ml; for Fep, MICs of  $\leq 8$   $\mu$ g/ml; for Atm, MICs of  $\leq 8$   $\mu$ g/ml; for Ipm, MICs of  $\leq 4$   $\mu$ g/ml; for Amk, MICs of  $\leq 16$   $\mu$ g/ml; and for Cip, MICs of  $\leq 1$   $\mu$ g/ml. The susceptible reference strain *P. aeruginosa* ATCC 27853 was used as the internal quality control.

<sup>c</sup> MIC of piperacillin with a fixed concentration (4  $\mu$ g/ml) of tazobactam.

<sup>d</sup> The pI value for IMP-18 has not been reported in the literature. IEF experiments for this isolate showed a smear between pI 5.0 and 7.5.

<sup>e</sup> ETA, endotracheal aspiration.

or from hospitals in the same region. Regarding the isolates producing ESBLs, MBLs, or ES-OXAs, genotypic analysis revealed that a clone (PFGE pattern A) producing OXA-19 had spread in two hospitals (Nancy and Epinal, France, 70 km apart). The spread of this clone has been described in a recent publication (11). A second clone (PFGE pattern F), producing both PER-1 and VIM-2, was isolated in different wards of the same university hospital in Paris, France, while a third clone (PFGE pattern H), producing SHV-2a, was detected in two other hospitals in the north of France (Lille and Cambrai, 68 km apart) (Fig. 1).

Since most ES-OXAs are poorly inhibited by clavulanate, used in screening tests (14), *Pseudomonas aeruginosa* strains expressing these enzymes remain difficult to recognize in routine testing and require genotypic methods. ES-OXAs have been described to occur sporadically, but their spread in the clinical setting remains poorly understood and probably underestimated. Our data stress the need for a simple and reliable routine test able to detect ESBLs, MBLs, and ES-OXAs produced by clinical *P. aeruginosa* strains. This test will be helpful to rapidly implement control measures for preventing the spread of multidrug-resistant strains harboring emerging resistance mechanisms.

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## REFERENCES

- al Naiemi, N., B. Duim, and A. Bart. 2006. A CTX-M extended-spectrum  $\beta$ -lactamase in *Pseudomonas aeruginosa* and *Stenotrophomonas maltophilia*. *J. Med. Microbiol.* **55**:1607–1608.
- Altschul, S. F., W. Gish, W. Miller, E. W. Myers, and D. J. Lipman. 1990. Basic local alignment search tool. *J. Mol. Biol.* **215**:403–410.
- Arllet, G., and A. Philippon. 1991. Construction by polymerase chain reaction and use of intragenic DNA probes for three main types of transferable  $\beta$ -lactamases (TEM, SHV, CARB). *FEMS Microbiol. Lett.* **66**:19–25.
- Azim, A., M. Dwivedi, P. B. Rao, A. K. Baronia, R. K. Singh, K. N. Prasad, B. Poddar, A. Mishra, M. Gurjar, and T. N. Dhole. 2010. Epidemiology of bacterial colonization at ICU admission with emphasis on extended-spectrum  $\beta$ -lactamases and metallo- $\beta$ -lactamase producing gram-negative bacteria—an Indian experience. *J. Med. Microbiol.* doi:10.1099/jmm.0.018085-0.
- Babini, G. S., and D. M. Livermore. 2000. Are SHV  $\beta$ -lactamases universal in *Klebsiella pneumoniae*? *Antimicrob. Agents Chemother.* **44**:2230.
- Bert, F., C. Branger, and N. Lambert-Zechovsky. 2002. Identification of PSE and OXA  $\beta$ -lactamase genes in *Pseudomonas aeruginosa* using PCR-restriction fragment length polymorphism. *J. Antimicrob. Chemother.* **50**:11–18.
- Brasme, L., P. Nordmann, F. Fidel, M. F. Lartigue, O. Bajelet, L. Poirel, D. Forte, V. Vernet-Garnier, J. Madoux, J. C. Reveil, C. Alba-Sauvati, I. Baudinat, P. Bineau, C. Bouquigny-Saison, C. Eloy, C. Lafaurie, D. Simeon, J. P. Verquin, F. Noel, C. Strady, and C. De Champs. 2007. Incidence of class A extended-spectrum  $\beta$ -lactamases in Champagne-Ardenne (France): a 1 year prospective study. *J. Antimicrob. Chemother.* **60**:956–964.
- Cavallo, J. D., D. Hocquet, P. Plésiat, R. Fabre, and M. Roussel-Delvallez. 2007. Susceptibility of *Pseudomonas aeruginosa* to antimicrobials: a 2004 French multicentre hospital study. *J. Antimicrob. Chemother.* **59**:1021–1024.
- Celenza, G., C. Pellegrini, M. Caccamo, B. Segatore, G. Amicosante, and M. Perilli. 2006. Spread of *bla*<sub>CTX-M-type</sub> and *bla*<sub>PER-2</sub>  $\beta$ -lactamase genes in clinical isolates from Bolivian hospitals. *J. Antimicrob. Chemother.* **57**:975–978.
- Chayakulkeeree, M., P. Junsriwong, A. Keerasuntonpong, C. Tribuddharat, and V. Thamlikitkul. 2005. Epidemiology of extended-spectrum  $\beta$ -lactamase producing gram-negative bacilli at Siriraj Hospital, Thailand, 2003. *Southeast Asian J. Trop. Med. Public Health* **36**:1503–1509.
- Cholley, P., D. Hocquet, C. Alauzet, A. Cravoisy, D. Talon, A. Nejl, P. Plésiat, and X. Bertrand. 2010. Hospital outbreak of *Pseudomonas aeruginosa* producing extended-spectrum oxacillinase OXA-19. *J. Med. Microbiol.* doi:10.1099/jmm.0.019364-0.
- Comité de l'Antibiogramme de la Société Française de Microbiologie. 13 November 2007. Guidelines 2006. [http://www.sfm.asso.fr/doc/download.php?doc=DiU8C&fic=casfm\\_2006.pdf](http://www.sfm.asso.fr/doc/download.php?doc=DiU8C&fic=casfm_2006.pdf).
- Corvec, S., L. Poirel, J. W. Decousser, P. Y. Allouch, H. Drugeon, and P. Nordmann. 2006. Emergence of carbapenem-hydrolysing metallo- $\beta$ -lactamase VIM-1 in *Pseudomonas aeruginosa* isolates in France. *Clin. Microbiol. Infect.* **12**:941–942.
- Danel, F., M. G. P. Page, and D. M. Livermore. 2007. Class D  $\beta$ -lactamases, p. 163–194. *In* R. A. Bonomo and M. E. Tolmasey (ed.), *Enzyme-mediated resistance to antibiotics*. ASM Press, Washington, DC.
- David, M., J. F. Lemeland, and S. Boyer. 2008. Emergence of extended-spectrum  $\beta$ -lactamases in *Pseudomonas aeruginosa*: about 24 cases at Rouen University Hospital. *Pathol. Biol.* **56**:429–434.
- De Champs, C., C. Chanal, D. Sirot, R. Baraduc, J. P. Romaszko, R. Bonnet, A. Plaidy, M. Boyer, E. Carroy, M. C. Gbadamassi, S. Lалуque, O. Oules, M. C. Poupart, M. Villemain, and J. Sirot. 2004. Frequency and diversity of class A extended-spectrum  $\beta$ -lactamases in hospitals of the Auvergne, France: a 2 year prospective study. *J. Antimicrob. Chemother.* **54**:634–639.
- Dubois, V., C. Arpin, V. Dupart, A. Scavelli, L. Coulangue, C. André, I. Fischer, F. Grobost, J. P. Brochet, I. Lagrange, B. Dutilh, J. Jullin, P. Noury, G. Larribet, and C. Quentin. 2008.  $\beta$ -Lactam and aminoglycoside resistance rates and mechanisms among *Pseudomonas aeruginosa* in French general practice (community and private healthcare centres). *J. Antimicrob. Chemother.* **62**:316–323.
- Fournier, D., D. Hocquet, B. Dehecq, P. Cholley, and P. Plésiat. 2010. Detection of a new extended-spectrum oxacillinase in *Pseudomonas aeruginosa*. *J. Antimicrob. Chemother.* **65**:364–365.
- Juan, C., X. Mulet, L. Zamorano, S. Alberti, J. L. Perez, and A. Oliver. 2009. Detection of the novel extended-spectrum  $\beta$ -lactamase OXA-161 from a plasmid-located integron in *Pseudomonas aeruginosa* clinical isolates from Spain. *Antimicrob. Agents Chemother.* **53**:5288–5290.
- Levesque, C., L. Piche, C. Larose, and P. Roy. 1995. PCR mapping of integrons reveals several novel combinations of resistance genes. *Antimicrob. Agents Chemother.* **39**:185–191.
- Livermore, D. M. 2002. Multiple mechanisms of antimicrobial resistance in *Pseudomonas aeruginosa*: our worst nightmare? *Clin. Infect. Dis.* **34**:634–640.
- Llanes, C., C. Neuwirth, F. El Garch, D. Hocquet, and P. Plésiat. 2006. Genetic analysis of a multiresistant strain of *Pseudomonas aeruginosa* producing PER-1  $\beta$ -lactamase. *Clin. Microbiol. Infect.* **12**:270–278.
- Mathew, A., A. M. Harris, M. J. Marshall, and G. W. Ross. 1975. The use of analytical isoelectric focusing for detection and identification of  $\beta$ -lactamases. *J. Gen. Microbiol.* **88**:169–178.
- Mendes, R. E., K. A. Kiyota, J. Monteiro, M. Castanheira, S. S. Andrade, A. C. Gales, A. C. C. Pignatari, and S. Tufik. 2007. Rapid detection and identification of metallo- $\beta$ -lactamase-encoding genes by multiplex real-time PCR assay and melt curve analysis. *J. Clin. Microbiol.* **45**:544–547.
- Naas, T., P. Bogaerts, C. Bauraing, Y. Degheldre, Y. Glupczynski, and P. Nordmann. 2006. Emergence of PER and VEB extended-spectrum  $\beta$ -lactamases in *Acinetobacter baumannii* in Belgium. *J. Antimicrob. Chemother.* **58**:178–182.
- NCCLS/CLSI. 2009. Method for dilution antimicrobial susceptibility tests for bacteria that grow aerobically. Approved standard, 8th ed. M7-A7. NCCLS/CLSI, Wayne, PA.
- Ozgunus, O. B., R. Caylan, I. Tosun, C. Sandalli, K. Aydin, and I. Koksul. 2007. Molecular epidemiology of clinical *Pseudomonas aeruginosa* isolates carrying IMP-1 metallo- $\beta$ -lactamase gene in a university hospital in Turkey. *Microb. Drug Resist.* **13**:191–198.
- Philippon, L., T. Naas, A. Bouthors, V. Barakett, and P. Nordmann. 1997. OXA-18, a class D clavulanic acid-inhibited extended-spectrum  $\beta$ -lactamase from *Pseudomonas aeruginosa*. *Antimicrob. Agents Chemother.* **41**:2188–2195.
- Poirel, L., L. Brinas, N. Fortineau, and P. Nordmann. 2005. Integron-encoded GES-type extended-spectrum  $\beta$ -lactamase with increased activity toward aztreonam in *Pseudomonas aeruginosa*. *Antimicrob. Agents Chemother.* **49**:3593–3597.
- Poirel, L., L. Brinas, A. Verlinde, L. Ide, and P. Nordmann. 2005. BEL-1, a novel clavulanic acid-inhibited extended-spectrum  $\beta$ -lactamase, and the class 1 integron In120 in *Pseudomonas aeruginosa*. *Antimicrob. Agents Chemother.* **49**:3743–3748.
- Poirel, L., I. Le Thomas, T. Naas, A. Karim, and P. Nordmann. 2000. Biochemical sequence analyses of GES-1, a novel class A extended-spectrum  $\beta$ -lactamase, and the class 1 integron In52 from *Klebsiella pneumoniae*. *Antimicrob. Agents Chemother.* **44**:622–632.
- Poirel, L., M. Magalhaes, M. Lopes, and P. Nordmann. 2004. Molecular analysis of metallo- $\beta$ -lactamase gene *bla*<sub>SPM-1</sub>-surrounding sequences from disseminated *Pseudomonas aeruginosa* isolates in Recife, Brazil. *Antimicrob. Agents Chemother.* **48**:1406–1409.
- Poirel, L., T. Naas, and P. Nordmann. 2010. Diversity, epidemiology, and genetics of class D  $\beta$ -lactamases. *Antimicrob. Agents Chemother.* **54**:24–38.
- Rossolini, G. M., and J. D. Docquier. 2007. Class B  $\beta$ -lactamases, p. 115–144. *In* R. A. Bonomo and M. E. Tolmasey (ed.), *Enzyme-mediated resistance to antibiotics*. ASM Press, Washington, DC.
- Sutcliffe, J. G. 1978. Nucleotide sequence of the ampicillin resistance gene of *Escherichia coli* plasmid pBR322. *Proc. Natl. Acad. Sci. U. S. A.* **75**:3737–3741.
- Talon, D., V. Cailleaux, M. Thouvez, and Y. Michel-Briand. 1996. Discriminatory power and usefulness of pulsed-field gel electrophoresis in epidemiological studies of *Pseudomonas aeruginosa*. *J. Hosp. Infect.* **32**:135–145.
- Tenover, F. C., R. D. Arbeit, R. V. Goering, P. A. Mickelsen, B. E. Murray, D. H. Persing, and B. Swaminathan. 1995. Interpreting chromosomal DNA restriction patterns produced by pulsed-field gel electrophoresis: criteria for bacterial strain typing. *J. Clin. Microbiol.* **33**:2233–2239.
- Vahaboglu, H., R. Ozturk, G. Aygun, F. Coskuncan, A. Yaman, A. Kaygusuz, H. Leblebicioglu, I. Balik, K. Aydin, and M. Otkun. 1997. Widespread detection of PER-1-type extended-spectrum  $\beta$ -lactamases among nosocomial *Acinetobacter* and *Pseudomonas aeruginosa* isolates in Turkey: a nationwide multicenter study. *Antimicrob. Agents Chemother.* **41**:2265–2269.
- Vettoretti, L., N. Floret, D. Hocquet, B. Dehecq, P. Plésiat, D. Talon, and X. Bertrand. 2009. Emergence of extensive-drug-resistant *Pseudomonas aeruginosa* in a French university hospital. *Eur. J. Clin. Microbiol. Infect. Dis.* **28**:1217–1222.
- Weldhagen, G. F., L. Poirel, and P. Nordmann. 2003. Ambler class A extended-spectrum  $\beta$ -lactamases in *Pseudomonas aeruginosa*: novel developments and clinical impact. *Antimicrob. Agents Chemother.* **47**:2385–2392.