

Detection of *Achromobacter xylosoxidans* in Hospital, Domestic, and Outdoor Environmental Samples and Comparison with Human Clinical Isolates

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Achromobacter xylosoxidans is an aerobic nonfermentative Gram-negative rod considered an important emerging pathogen among cystic fibrosis (CF) patients worldwide and among immunocompromised patients. This increased prevalence remains unexplained, and to date no environmental reservoir has been identified. The aim of this study was to identify potential reservoirs of *A. xylosoxidans* in hospital, domestic, and outdoor environments and to compare the isolates with clinical ones. From 2011 to 2012, 339 samples were collected in Dijon's university hospital, in healthy volunteers' homes in the Dijon area, and in the outdoor environment in Burgundy (soil, water, mud, and plants). We designed a protocol to detect *A. xylosoxidans* in environmental samples based on a selective medium: MCXVAA (MacConkey agar supplemented with xylose, vancomycin, aztreonam, and amphotericin B). Susceptibility testing, genotypic analysis by pulsed-field gel electrophoresis, and *bla*_{OXA-114} sequencing were performed on the isolates. A total of 50 strains of *A. xylosoxidans* were detected in hospital (33 isolates), domestic (9 isolates), and outdoor (8 isolates) samples, mainly in hand washing sinks, showers, and water. Most of them were resistant to ciprofloxacin (49 strains). Genotypic analysis and *bla*_{OXA-114} sequencing revealed a wide diversity among the isolates, with 35 pulsotypes and 18 variants of oxacillinases. Interestingly, 10 isolates from hospital environment were clonally related to clinical isolates previously recovered from hospitalized patients, and one domestic isolate was identical to one recovered from a CF patient. These results indicate that *A. xylosoxidans* is commonly distributed in various environments and therefore that CF patients or immunocompromised patients are surrounded by these reservoirs.

Cystic fibrosis (CF) is a genetic disorder which affects different organs, including the respiratory system. The presence of abnormal mucus predisposes patients to chronic airway infections. Over the last 2 decades, new emerging bacteria have been found to invade the airways of CF patients, including nontuberculous mycobacteria, *Burkholderia cepacia* complex, *Stenotrophomonas maltophilia*, and *Achromobacter xylosoxidans*. Most of them are multidrug-resistant organisms thought to have been selected by intense antibiotic use over many years (1–4).

A. xylosoxidans is a Gram-negative, aerobic, and oxidase-positive bacillus, often misidentified as *Pseudomonas aeruginosa* (5, 6). It is an opportunistic pathogen that can cause a wide variety of infections in immunocompromised patients (7–10) but is mainly recovered from CF patients' airways. As previously reported, this species is innately resistant to cephalothin, cefoxitin, cefotaxime, aztreonam, and aminoglycosides (11, 12). Acquired resistance to carbapenems, ceftazidime, and ciprofloxacin is frequent, dramatically limiting therapeutic choices (13, 14).

The prevalence of infection or colonization is variable among CF centers, and cases of cross-contamination have been reported (1, 2, 14–16). In France, analysis of the global data from all the centers visited by CF patients revealed an increase in the isolation of *A. xylosoxidans* in recent years, with a rate rising from 2.7% in 2001 to 5.3% in 2011 (http://www.vaincrelamuco.org/e_upload/div/registre_francais_mucoviscidose_2011_15.03.13.pdf). In a recent study, we reported the first epidemiological data about *A. xylosoxidans* in a French CF center (Dijon, Burgundy), with a prevalence of 13.9% among the 120 patients (13). This percentage is, surprisingly, much higher than the one reported in the French global data and does not result from cross-contamination be-

tween the patients. Moreover, we observed an increase in the rate of isolation of *A. xylosoxidans* in hospitalized patients, mainly in intensive care units and hematology wards. This unusually high frequency of *A. xylosoxidans* isolation prompted us to look for specific environmental sources of contamination in the area.

To date, in hospitals, *A. xylosoxidans* has been involved in many procedure-related infections, being associated for instance with contaminated disinfectants (17), dialysis fluids (7), and ultrasound gel (18). There are a few reports of *A. xylosoxidans* being found in outdoor environments, including in some plants (19, 20), in polluted soils (21, 22), and in an indoor swimming pool (23). Nevertheless, the natural habitat of this organism (24, 25) as well as the possible sources of patient contamination remain unknown. Only one report documented the contamination of a patient by an isolate originating from well water (26). The identification of potential reservoirs in the environment might constitute an aid to the prevention of infection in CF patients and in the immunocompromised population. In our laboratory, we have established a collection of all the clinical strains of *A. xylosoxidans* recovered in our hospital, including 808 isolates from CF patients (since 1995) and 32 isolates from non-CF patients (since 2010). For all these isolates, antimicrobial susceptibility tests have been

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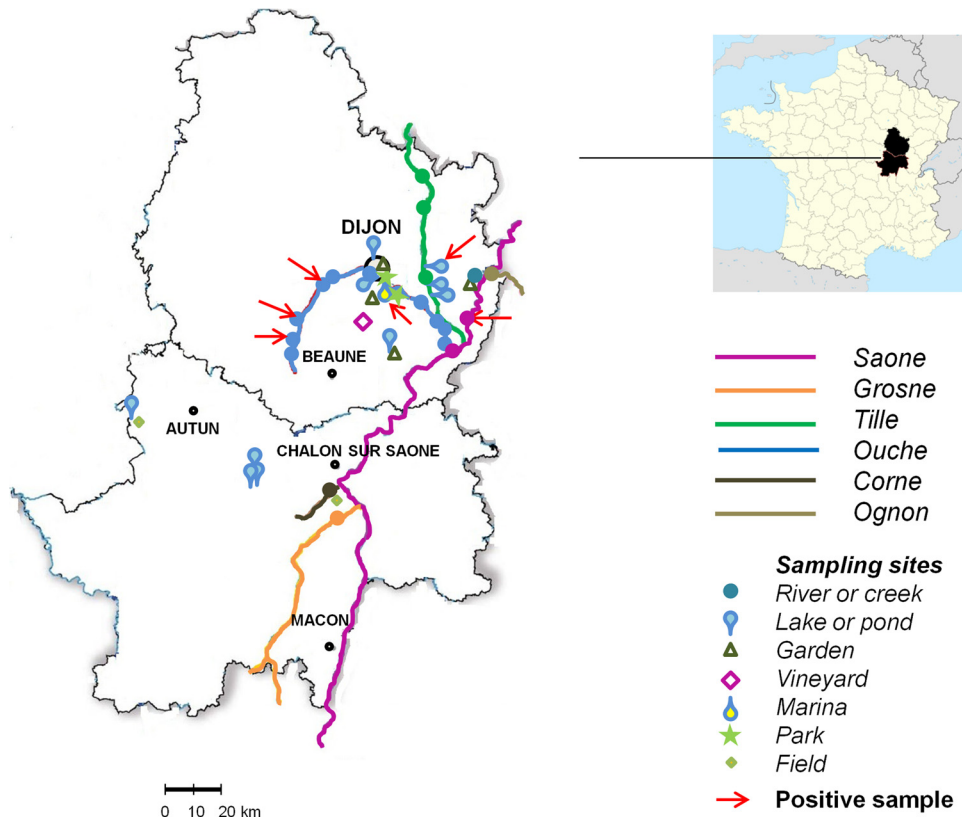


FIG 1 Outdoor sampling sites in Cote d'Or and Saone-et-Loire, France. Red arrows show positive sampling sites. (Map used with permission from CartesFrance.fr.)

performed by the disk diffusion method and genotyping analysis by pulsed-field gel electrophoresis (PFGE).

The aim of the present study was (i) to search different environments (natural, hospital, and domestic) for the presence of *A. xylosoxidans* strains potentially involved in patient contamination, (ii) to study the antimicrobial susceptibility of the isolates, and (iii) to compare the environmental isolates to clinical isolates from our collection by a genotyping method.

MATERIALS AND METHODS

Sampling sites. We chose various sampling sites to get an overview of the presence of *A. xylosoxidans* in diverse environments: hospital, domestic, and outdoor. For outdoor environment sampling sites, we focused our investigation on Cote d'Or and Saone et Loire. Indeed, more than 60% of CF colonized patients visiting our CF center live in these two areas of Burgundy, France. From September 2011 to December 2012, a total of 339 samples were collected. In the University Hospital of Dijon (1,772 beds), 188 distinct samples were obtained in 10 different departments located in 4 different buildings: 4 intensive care units (ICU), a pneumology and nephrology department (first building), a hematology department (second building), a pediatrics department and the CRCM (Centre de Ressources et de Compétences de la Mucoviscidose), where cystic fibrosis patients receive care (third building), and a dentistry department (fourth building, 5 km away). In each department, environmental wet surfaces in patients' rooms (drains of sinks used for hand washing, shower drains, and toilet bowls) and medication preparation rooms (sink drains) and nearby offices (ward sluice sinks) were swabbed. The number of samples per room depended on the department (e.g., ICU rooms are not equipped with showers or toilets). In the dentistry department, we collected water from dental unit waterlines (14 chairs). A total of 58 domestic samples

were collected from 16 residences of healthy volunteers (bathroom sinks, kitchen sinks, shower drains, and toilet bowls). These apartments or houses were located in Dijon ($n = 14$) and in two villages, Domois ($n = 1$) and St Nicolas les Citeaux ($n = 1$) (10 km south and 35 km south of Dijon, respectively). Finally, 93 samples were obtained from outdoor environments: soils ($n = 8$), waters ($n = 36$), mud from rivers or lakes ($n = 29$), and plants ($n = 20$). The sampling sites included 4 lakes with recreational activities (from 7 to 37 ha), 6 rivers (9 to 480 km long), 1 creek, 1 marina, 6 ponds, 2 fields, 1 vineyard, 2 public parks, and 5 domestic vegetable gardens. Plant collection included the aerial parts (leaf, stem, branch, or fruit) of herbs, moss, vines, mushrooms, strawberries, and oaks and the roots of alfalfa, tomato plants, green bean plants, clover, lupine, carrots, convulvulus, and vines. The outdoor sampling sites are indicated on the map in Fig. 1.

Sample collection. Domestic and hospital samples were obtained using sterile cotton swabs. Sink and shower drains were sampled by rotating a cotton swab inserted approximately 5 cm into the sink drain. Toilets were sampled with sterile cotton swabs inserted under the rim of the toilet bowl. Sluice sinks were sampled by inserting swabs in the wastewater and collection bowl and grid surfaces. Outdoor samples were collected in sterile containers. Approximately 1 g of mud or soil and 5 cm² of parts of plants (roots or aerial parts) were disrupted with sterile devices and then vigorously shaken with brain heart infusion enrichment broth (BHI). Water samples were concentrated by filtration of 500 ml through sterile 0.45- μ m membrane filters (Milliflex Plus; Millipore, Billerica, MA). Samples were stored at room temperature for less than 24 h before processing. The protocol for isolation and identification of *A. xylosoxidans* is presented in Fig. 2.

Culture of the samples. In our experience, from previous studies conducted on hospital environment samples, samples such as those collected in this study often contain a wide diversity of Gram-negative bacilli, including nonfermenting Gram-negative rods (NF-GNB) and *Enterobacte-*

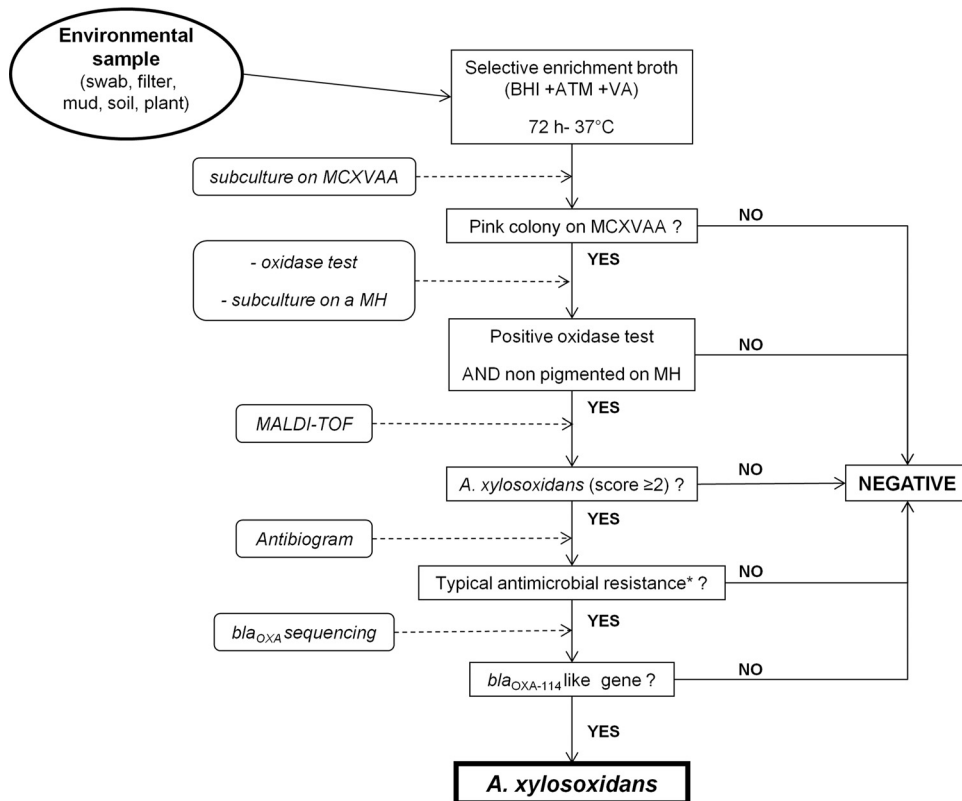


FIG 2 Isolation and identification protocol of *A. xylosoxidans* strains from environmental samples. A typical antibiogram (*) is defined by resistance to all aminoglycosides tested, aztreonam, cefoxitin, cefotaxime, and cefepime. VA, vancomycin; ATM, aztreonam.

riaceae. All these organisms grow on Drigalski agar; therefore, the detection of *A. xylosoxidans* among the environmental flora on this medium is impossible. Given that *A. xylosoxidans* is innately resistant to aztreonam, we tested a Drigalski medium supplemented with this antibiotic. On this medium, we still observed the growth of many different oxidase-positive NF-GNB despite the antibiotic selection. Therefore, we developed a selective enrichment procedure to enhance our ability to detect *A. xylosoxidans* in the samples. Swabs, filters, mud, soil samples, and plants were first enriched with 10 ml of BHI supplemented with aztreonam (32 mg/liter) and vancomycin (32 mg/liter) for 72 h at 37°C. One drop of each of the enrichment cultures was plated on MCXVAA medium and incubated 48 h at 37°C. MCXVAA agar is a medium that we designed for the study: MacConkey agar (selective for Gram-negative rods) supplemented with xylose (5 g/liter), vancomycin (20 mg/liter) (to prevent growth of Gram-positive bacteria), aztreonam (20 mg/liter), and amphotericin B (5 mg/liter) (to prevent fungal growth). In preliminary experiments (data not shown), we checked that *A. xylosoxidans* reference strains (CIP 7132^T, CIP 102236, CIP 101902, and CIP 110540) as well as 15 clinical strains from our previously described collection (13) grew on MCXVAA agar. *A. xylosoxidans* forms pink colonies, resulting from acidification of this medium due to xylose oxidation. Moreover, MCXVAA agar is unable to support the proliferation of Gram-positive bacteria or yeasts/fungi. Nevertheless, it is important to note that other aztreonam-resistant Gram-negative rods are able to grow on this medium. For example *S. maltophilia* forms yellow colonies, and some *Pseudomonas* spp., other *Achromobacter* spp., and *Bordetella* spp. also appear as pink colonies. Therefore, pink colonies must undergo further identification.

Identification. The identification scheme is summarized in Fig. 2. Given that *A. xylosoxidans* is an oxidase-positive bacillus forming pink colonies on MCXVAA agar and nonpigmented ones on Mueller-Hinton agar, we retained for further identification only the strains fulfilling these

3 criteria. These strains were identified by matrix-assisted laser-desorption/ionization time-of-flight mass spectrometry (MALDI-TOF; Bruker). This technique proved to be efficient for identifying *A. xylosoxidans* (27–29). According to the manufacturer's instructions, a score of ≥ 2 indicates a "secure genus identification, and probable species identification." Moreover, such a score was obtained for the reference strains CIP 7132^T, CIP 102236, CIP 101902, and CIP 110540. Therefore, the isolates identified as *A. xylosoxidans* with a score of ≥ 2 were subjected to susceptibility tests to select the strains harboring the typical resistance profile (cefotaxime, cefepime, aztreonam, kanamycin, tobramycin, netilmicin, amikacin, and gentamicin). The last step of identification consisted of detecting and sequencing the naturally occurring *bla*_{OXA} genes of *A. xylosoxidans* (*bla*_{OXA-114} and variants [13, 30, 31] and *bla*_{OXA-243} and variants [13]). We used the primer pair OXA-114A and OXA-114B (30). In cases where PCR was negative, we used the primer pair AXXA-F and AXXA-R (13). Reference strains of *A. xylosoxidans* described above were used as positive controls for PCRs. Positive PCR products were purified with a Millipore centrifugal filter unit (Amicon Microcon PCR kit; Millipore). Double-strand sequencing was then performed using BigDye v1.1 Terminator chemistry and a 3130XL genetic analyzer (Applied Biosystems).

Susceptibility tests. Susceptibility tests of the isolates were performed by the disk diffusion method and interpreted according to recommendations of EUCAST (http://www.eucast.org/fileadmin/src/media/PDFs/EUCAST_files/Breakpoint_tables/Breakpoint_table_v_3.1.pdf). The following antibiotics were tested: β -lactams (ticarcillin, ticarcillin-clavulanic acid, piperacillin, piperacillin-tazobactam, cefoxitin, cefotaxime, ceftazidime, cefepime, imipenem, meropenem, doripenem, and aztreonam), ciprofloxacin, and aminoglycosides (amikacin, gentamicin, kanamycin, netilmicin, and tobramycin).

Genotyping by pulsed-field gel electrophoresis (PFGE). All the strains identified as *A. xylosoxidans* were subjected to genotypic analysis

TABLE 1 Hospital and domestic samples

Source	No. of samples (no. positive)					Total no. of strains	No. of pulsotypes	Pulsotype(s) (OXA variant[s]) ^a
	Handwashing-sink drains	Toilet pans	Shower drains	Sluice sinks	Total			
Hospital								
ICU-1	25 (1)			2	27 (1)	1	1	A (OXA-114h)
ICU-2	11 (1)				11 (1)	1	1	Un. (OXA-114i)
ICU-3	5 (1)			2 (1)	7 (2)	2	1	B (OXA-114p)
ICU-4	7			1	8	0		
Pediatrics	8 (3)	1			9 (3)	3	2	C (OXA-114f), D ^b (OXA-114h)
CF Center	4	1		1 (1)	6 (1)	1	1	E (OXA-114f)
Nephrology	5	1	2 (1)	4 (2)	12 (3)	3	1	F (OXA-114 m)
Pneumology	13 (2)	6	6 (4)	2	27 (6)	7	6	G (OXA-114h), H (OXA-114a), Un. (OXA-114e), Un. (OXA-114g), Un. (OXA-114s), Un. (OXA-114s)
Hematology	43 (7)	6	17 (3)	1	67 (10)	11	6	C (OXA-114f), I (OXA-114c), Un. (OXA-114g), Un. (OXA-114g), Un. (OXA-114l), Un. (OXA-243e)
Dentistry					14 ^c (4)	4	1	J (OXA-114r)
Total	121 (15)	15 (0)	25 (8)	13 (4)	188 (31)	33	19	
Domestic								
	31 (3)	13 (0)	14 (4)		58 (7)	9	9	Un. (OXA-114a), Un. (OXA-114c), Un. (OXA-114c), Un. (OXA-114f), K (OXA-114h), Un. (OXA-114i), Un. (OXA-114j), Un. (OXA-114t), Un. (OXA-114v)

^a Letters designate pulsotypes that were recovered more than once. Un., unique (i.e., the pulsotype was recovered only once). Pulsotypes in bold are identical to clinical strain pulsotypes. Pulsotype C includes strains from 2 departments and clinical strains.

^b Pulsotype also recovered in one sample from the Saone River.

^c Dental chair water samples.

by PFGE as previously described (32). Total DNA was analyzed after digestion with the restriction enzyme XbaI. Electrophoresis was carried out for 20 h at 5.4 V/cm, with pulse times ranging from 5 s to 35 s, using the CHEF-DR II system (Bio-Rad). Restriction patterns were interpreted according to the criteria of Tenover et al. (33), and the isolates were classified into pulsotypes (A, B, C, etc.). These pulsotypes were compared to those of the clinical isolates of our collection gathered since 2006. We have recently described isolates recovered from 21 CF patients (P1 to P21) (13). Clonally related isolates belong to the same pulsotype. Pulsotypes recovered only once and different from all those already identified to date in our hospital were designated “unique.”

Nucleotide sequence accession numbers. OXA-114p, -114q, -114r, -114s, -114t, -114u, and -114v gene sequences have been submitted to

GenBank under accession numbers [KF573363](#), [KF573364](#), [KF573365](#), [KF573366](#), [KF573367](#), [KF573368](#), and [KF573369](#). OXA-243e's gene sequence has been submitted to GenBank under accession number [KF582664](#).

RESULTS

The results are summarized in [Table 1](#) (hospital and domestic), in [Table 2](#) (outdoor) and in [Table 3](#) (susceptibility tests).

Detection of strains. The protocol used in this study allowed us to identify a total of 50 isolates of *A. xylosoxidans* among the 339 samples, mainly in the hospital environment ($n = 33$) but also in domestic ($n = 9$) and outdoor ($n = 8$) environments.

TABLE 2 Characteristics of the strains found at outdoor positives sampling sites^a

Outdoor source and sample type	No. of samples (no. positive)	No. of pulsotypes	Pulsotype(s) (OXA variant) ^b	Description of source
Ouche River (95.4 km)				
Water	11 (3)	3	Un. (OXA-114b) Un. (OXA-114g) Un. (OXA-114g)	Water treatment, port
Mud	13 (2)	2	Un. (OXA-114h) Un. (OXA-114q)	Water treatment
Saone River water (480 km)	2 (1)	1	D ^c (OXA-114h)	Boating, fishing
Lake Arc-sur-Tille water (36 ha)	2 (1)	1	Un. (OXA-114u)	Swimming, water sports
Dijon Marina water (3 ha)	1 (1)	1	Un. (OXA-114a)	Port
Total	29 (8)	8		

^a Only one strain was isolated from each positive sample.

^b Letters designate pulsotypes that were recovered more than once. Un., unique (i.e., the pulsotype was recovered only once).

^c This pulsotype was also recovered in the pediatrics department of the hospital.

TABLE 3 Number of isolates categorized as intermediate or resistant among hospital, domestic, and outdoor strains

Antimicrobial agent ^a	No. of intermediate or resistant isolates			
	Hospital (n = 33)	Domestic (n = 9)	Outdoor (n = 8)	Total (n = 50)
TIC	0	0	0	0
TCC	0	0	0	0
PIP	0	0	0	0
TZP	0	0	0	0
CAZ	1	0	0	1
IPM	10	1	4	15
MEM	0	0	0	0
DOR	2	2	1	5
CIP	32	9	8	49

^a TIC, ticarcillin; TCC, ticarcillin plus clavulanic acid; PIP, piperacillin; TZP, piperacillin plus tazobactam; CAZ, ceftazidime; IPM, imipenem; MEM, meropenem; DOR, doripenem; CIP, ciprofloxacin.

Hospital samples. A total of 33 strains were isolated in 31 of the 188 samples. All studied wards were concerned with the exception of ICU-4. *A. xylosoxidans* has been recovered in hand-washing sink drains, shower drains, and sluice sinks but not in toilet bowls. Water from 4 of the 14 dental chairs was contaminated with *A. xylosoxidans*.

Domestic samples. A total of 9 strains were isolated in 7 out of the 58 samples. These findings concerned 6 of the 16 residences investigated. *A. xylosoxidans* was detected in 4 of 14 shower drains (6 strains), in 1 of 16 bathroom sink drains (1 strain), and in 2 of 15 kitchen sink drains (2 strains). All 13 toilet bowls sampled were negative.

Outdoor samples. A total of 8 strains were isolated from the 93 samples, all in water or mud samples. None of these strains were isolated from plants or soils. The positive samples are indicated with red arrows in Fig. 1 and described in Table 2. Five strains were isolated from the Ouche River (at 3 different sites), one was from Lake Arc-sur-Tille, one was from the Port du Canal marina in Dijon, and the last one was from the Saone River in Auxonne.

Genotyping by PFGE. Among the 50 isolates obtained, we distinguished 35 pulsotypes with PFGE: 19 pulsotypes in hospital samples, 9 in domestic samples, and 8 in outdoor samples, with one pulsotype being identical in hospital and outdoor isolates. Indeed, the isolate detected in water from the Saone River (pulsotype D) was clonally related to isolates recovered in the pediatrics department (2 different hand-washing sink drains). In samples from the hospital environment, we detected isolates belonging to the same pulsotype (pulsotype C) in two departments, pediatrics (one hand-washing sink drain) and hematology (3 samples in a patient's room located on level 0 of the unit and the hand-washing sink drain of a medication preparation room on level 1). We performed repeated samplings at two locations. The first one was the CF center, where we recovered the isolate harboring pulsotype E twice at an 8-month interval (October 2011 and July 2012). The second location was ICU-1, from which the isolate (pulsotype A) was isolated 3 times (September 2011, October 2011, and February 2012). In the dentistry department, the isolates recovered from the 4 dental chairs water belonged to the same pulsotype (pulsotype J). Comparison of environmental with clinical isolates pulsotypes revealed interesting findings. Among the 35 pulsotypes from environmental isolates, 6 were identical to that of patients. In

ICU-1, the strain recovered from a wound of a patient had the same pulsotype as the one isolated in the hand-washing sink of his room during his hospitalization (pulsotype A). Pulsotype C, including 6 environmental isolates, was identical to that of strains isolated in tracheal aspirates of 2 non-CF patients hospitalized in ICU-1 in 2010. The strain recovered from a shower in the pneumology department was clonally related (pulsotype G) to that recovered from the sputum of a CF patient who had been chronically colonized since 1995 (patient P19) but had not previously been hospitalized in that ward. Pulsotype I, including 2 strains recovered in the hematology ward in December 2012 (1 shower and 1 hand-washing sink in two rooms), also included isolates from 6 patients. Among these patients, only one had been hospitalized in the hematology department (blood culture sampled in November 2012). For the other patients, the strains were recovered from sputum samples. Two of them had been hospitalized: one in the pneumology department (March 2012) and one in ICU-2 (January 2013). One outpatient suffering from bronchiectasis regularly attended the pneumology department. (isolate recovered in April 2012). The last two were CF patients (isolates recovered in 2007–2008) P7 and P32. Finally, one domestic isolate recovered in a hand-washing sink drain was identical (pulsotype K) to that of one CF patient who had been chronically colonized since 1997 (patient P8).

Antimicrobial susceptibilities. The majority of the isolates were resistant to ciprofloxacin (49/50). Fifteen isolates were categorized as intermediate or resistant to imipenem (10 hospital, 4 outdoor, and 1 domestic isolate), with one of them (a hospital isolate) also being resistant to ceftazidime and 5 (2 hospital, 2 outdoor, and 1 domestic isolate) being also resistant to doripenem. All isolates remained susceptible to meropenem, ticarcillin, ticarcillin-clavulanic acid, piperacillin, and piperacillin-tazobactam. The results are reported in Table 3.

bla_{OXA} sequencing. On the basis of antimicrobial susceptibility tests and MALDI-TOF analysis, a total of 51 strains were subjected to PCRs. A PCR product was obtained for 50 isolates and sequenced. We could not detect any *bla_{OXA}* gene in the remaining strain; therefore, this strain was not identified as *A. xylosoxidans* and was excluded from the study. Many variants of OXA-114 have been detected in 49 strains. Among them, the variants OXA-114a, -114b, -114c, -114f, -114g, -114h, -114i, -114j, -114l, and -114m had already been identified in clinical isolates. We also detected 7 new variants: OXA-114p, -114q, -114r, -114s, -114t, -114u, and -114v. Finally, only one strain harbored a variant of OXA-243, the new variant OXA-243e.

DISCUSSION

A. xylosoxidans is an important emerging pathogen among CF patients worldwide, commonly described as an environmental pathogen widely distributed in soils and waters (4, 14, 34, 35). Nevertheless, this statement is not supported by any study in natural environments. The scarce reports concern a very limited number of isolates which were recovered mainly from polluted sites. Because of the high prevalence of *A. xylosoxidans* in our CF center and its emergence in immunocompromised patients hospitalized in our university hospital, an investigation of reservoirs was required. To the best of our knowledge, the present study is the first to look for the presence of *A. xylosoxidans* in various environments and to compare the environmental isolates with the clinical ones. For this comparison, we chose the PFGE method,

commonly used to analyze the clinical isolates of this species (14, 16, 31, 32, 36). Very recently (2012) and after the beginning of the present study, several schemes of multilocus sequence typing were proposed for *A. xylosoxidans* (37, 38). This method seems to be interesting for global epidemiological studies and allows inter-laboratory comparisons. Nevertheless, all the isolates from our collection have been subjected to PFGE analysis since 1995, and we took advantage of this large amount of available data.

In the hospital, environmental samplings targeted potential sources of contamination already reported for other pathogens, such as *P. aeruginosa* (39). For outdoor environmental samples, we focused on two areas of Burgundy (Cote d'Or and Saone et Loire) because most of our infected CF patients live there. We searched for *A. xylosoxidans* in different sites related to various common activities: walking (parks and fields), gardening (gardens), and fishing, swimming, or boating (rivers, lakes, and the marina). Because of bacterial species richness in these environments, we designed for this study an identification procedure and a selective medium (MCXVAA) to detect *A. xylosoxidans*. This allowed us to isolate 50 strains of *A. xylosoxidans* from wet environments, including water, mud, and sink and shower drains. Nevertheless, the use of a selective medium may have led to underestimation of the presence of *A. xylosoxidans*, as already described for other media (40). We observed a high diversity among the 50 environmental isolates: they were classified into 35 pulsotypes. It is noteworthy that the 17 domestic and outdoor strains all belonged to different pulsotypes. Moreover, we detected many oxacillinase variants ($n = 19$). These findings strongly suggested that *A. xylosoxidans* is well adapted to the wet environments of hospitals, outdoor sites, and domiciles that act as reservoirs. In the hospital, we found various *A. xylosoxidans* isolates in 9 out of 10 wards studied. Some hospital isolates had the same pulsotype (33 isolates for 19 pulsotypes). For instance, in the nephrology department, the three strains isolated from different sites (in one patient's room and two sluice sinks) were identical. Similarly, a single strain contaminated the water of four dental chairs. In these cases, hand contamination of staff members might explain the spread of the isolate through the ward. In another case, we suspected contamination of the water distribution pipe. Indeed, six isolates belonging to pulsotype C were recovered in the hematology and pediatrics departments, two wards located in adjacent buildings supplied by the same water distribution line. We did not detect *A. xylosoxidans* in the water (data not shown): perhaps the volume of the water was too small, or more likely the contamination occurred months or years ago. Our bacterial findings provided information about the presence of *A. xylosoxidans* in the sink drain at the moment when we obtained the samples but did not allow us to draw conclusions about the temporal and spatial evolution of the colonization. Nevertheless, the results from repeated samplings at two points indicate that isolates can persist in some sites for long periods (at least 8 months), probably thanks to biofilm formation. Surprisingly, we identified 10 hospital environmental isolates that were indistinguishable from patients' isolates. In most cases, it was not possible to determine if the patients acquired *A. xylosoxidans* from the hospital environment or if the environment had been contaminated by an infected patient, because the patients' strains had been isolated mainly before this study. In all cases but one, the patients had not been hospitalized in the wards where the clonally related environmental isolates were detected. For example, the strains harboring pulsotype I

from the hematology department have also been isolated from 6 patients in samples collected over 6 years (in the pneumology department, the hematology department, ICU-2, and the CF center). This reflects a spread of the isolates in all the wastewater system of the hospital, constituting a reservoir of *A. xylosoxidans*. Therefore, hospitalized patients might be contaminated when splash-back occurs, for instance during showering, hand washing, or tooth brushing, as has been described for other pathogens (41, 42). The role of dental chair waters as a source of contamination has to be taken into account for outpatients receiving dental care.

Our study also pointed out that *A. xylosoxidans* is widespread in outdoor aquatic and in domestic environments. Indeed, the organism was detected in 6 samples from bodies of water used for recreational activities and from 6 of 16 residences. For outdoor environments, 4 isolates were recovered from the Ouche River (water and mud) near the points where the effluents from wastewater treatment stations of local villages flow into the river. Moreover, one isolate recovered from the Saone River, 35 km away from Dijon, was identical to one isolate in the pediatrics department of the hospital (pulsotype D). These findings may reflect bacterial pollution of the rivers by the effluents. In domestic environments, *A. xylosoxidans* was detected in hand-washing sinks and shower drains but not in toilet bowls. The colonization of sinks and shower drains by *P. aeruginosa* has already been described in home environment studies (43, 44). These materials are in contact with soap as well as with chemical products used for cleaning, many of them containing alkyldimethylbenzylammonium chloride, a quaternary ammonium compound. The natural resistance of *A. xylosoxidans* to these compounds might explain its presence in the domestic samples (45). We did not detect *A. xylosoxidans* in the toilet bowls, probably because the site sampled (under the rim of the toilet bowl) is easily accessible for regular cleaning and free of stagnant water. Interestingly, one isolate recovered from a residence located in Dijon was identical to that harbored by a CF patient since 1997 who has always lived 37 km away from Dijon. This finding remains unexplained.

These results indicate that in everyday life, CF patients can be exposed to reservoirs of *A. xylosoxidans*. For several authors, the diversity among CF patients isolates suggested the acquisition from various environmental sources (13, 31, 46). This hypothesis of contamination from these different niches out of hospital is corroborated by several of our observations. First, the environmental isolates described here are varied: they belonged to many distinct pulsotypes and harbored many oxacillinase variants. Second, environmental and clinical isolates share common variants of OXA-114. Finally, with regard to antimicrobial resistance, it is interesting that all environmental strains were resistant to ciprofloxacin and remained susceptible to ticarcillin, piperacillin, ticarcillin-clavulanic acid, and piperacillin-tazobactam like most strains (75%) isolated from CF patients' sputa at first positive culture in our CF center. For the first time, we have identified various environmental reservoirs of *A. xylosoxidans* and demonstrated that some of these isolates are indistinguishable from clinical ones. In hospital and domestic environments, *A. xylosoxidans* is widely encountered in sink drains, which are therefore potential sources of contamination for immunocompromised or CF patients. Particular cleaning procedures might be a help in preventing the acquisition of this emerging pathogen. Larger studies in other geographical regions need to be conducted to confirm our findings. A prospective study analyzing the domestic environ-

ments of CF patients would be of particular interest, and our detection procedure may be suitable for this purpose.

In conclusion, there is a need for a global approach, examining medical but also environmental and probably animal aspects of contamination, to better understand the worldwide increased prevalence of *A. xylosoxidans* infections and its possible relationship to anthropogenic activity.

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