

Phenotypic and Molecular Detection of CTX-M- β -Lactamases Produced by *Escherichia coli* and *Klebsiella* spp.

Johann D. D. Pitout,¹ Ashfaque Hossain,² and Nancy D. Hanson^{2*}

Division of Microbiology, Calgary Laboratory Services, and Department of Pathology & Laboratory Medicine, University of Calgary, Calgary, Alberta, Canada,¹ and Department of Medical Microbiology and Immunology, Creighton University School of Medicine, Omaha, Nebraska²

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Organisms producing CTX-M- β -lactamases are emerging around the world as a source of resistance to oxyiminocephalosporins such as cefotaxime (CTX). However, the laboratory detection of these strains is not well defined. In this study, a molecular detection assay for the identification of CTX-M- β -lactamase genes was developed and used to investigate the prevalence of these enzymes among clinical isolates of *Escherichia coli* and *Klebsiella* species in the Calgary Health Region during 2000 to 2002. In addition, National Committee for Clinical Laboratory Standards (NCCLS) recommendations were evaluated for the ability to detect isolates with CTX-M extended-spectrum β -lactamases (ESBLs). The PCR assay consisted of four primer sets and demonstrated 100% specificity and sensitivity for detecting different groups of CTX-M- β -lactamases in control strains producing well-characterized ESBLs. Using these primer sets, 175 clinical strains producing ESBLs were examined for the presence of CTX-M enzymes; 24 (14%) were positive for *bla*_{CTX-M-1-like} genes, 95 (54%) were positive for *bla*_{CTX-M-14-like} genes, and the remaining 56 (32%) were negative for *bla*_{CTX-M} genes. Following the NCCLS recommendations for ESBL testing, all of the control and clinical strains were detected when screened with cefpodoxime and when both cefotaxime and ceftazidime with clavulanate were used as confirmation tests.

Resistance to the expanded-spectrum cephalosporins can occur in *Escherichia coli* and *Klebsiella* species via the production of extended-spectrum β -lactamases (ESBLs) that are capable of hydrolyzing the oxyiminocephalosporins and monobactams (11). Recently, a family of ESBLs which preferentially hydrolyze cefotaxime (CTX), the CTX-M- β -lactamases, have been recognized and reported in the literature with increasing frequency (3). This resistance mechanism is widespread throughout the world, with reports of clinical isolates producing these β -lactamases from Europe, Africa, Asia, South America, and most recently North America (3, 27).

CTX-M- β -lactamases are not closely related to TEM or SHV ESBLs (7) but share high amino acid identity with chromosomal β -lactamases from *Kluyvera georgiana* (34), *Kluyvera cryocrescens* (17) and *Kluyvera ascorbata* (24). In fact, the CTX-M-5 enzyme is identical to the chromosomal gene of *K. ascorbata* (3). According to a recent review and new data within GenBank, CTX-M- β -lactamases can be divided into five groups based on their amino acid sequence identities (3). Group I includes CTX-M-1, -3, -10 to -12, -15 (UOE-1), -22, -23, -28, -29, and -30. Group II includes CTX-M-2, -4 to -7, and -20 and Toho-1. Group III includes CTX-M-8. Group IV includes CTX-M-9, -13, -14, -16 to -19, -21, and -27 and Toho-2. Finally group V includes CTX-M-25 and -26. The members of these groups exhibit >94% amino acid identity within the group and \leq 90% amino acid identity between groups (3).

The laboratory detection of organisms producing CTX-M- β -lactamases is not well defined. The guidelines published by

the National Committee for Clinical Laboratory Standards (NCCLS) for the detection of ESBL-producing *E. coli* and *Klebsiella* spp. were first published in 1999 before organisms producing CTX-M- β -lactamases became recognized as an important cause of resistance to the newer cephalosporins (3, 29). It is unclear if these guidelines are capable of detecting organisms producing CTX-M- β -lactamases.

A study was designed to develop and evaluate a molecular detection assay for the identification of strains producing known CTX-M- β -lactamases and to investigate if genes encoding these enzymes were present among ESBL-producing strains of *E. coli* and *Klebsiella* species isolated from clinical specimens. Only *E. coli* and *K. pneumoniae* were examined because NCCLS guidelines for screening and confirmation of ESBL-production are established for these organisms. The ability of these guidelines to detect strains producing CTX-M- β -lactamases was also evaluated.

MATERIALS AND METHODS

Bacterial strains. Strains with well-described β -lactamases were used as positive and negative controls (Table 1). Consecutive nonduplicate isolates of *E. coli* and *Klebsiella* spp. collected at Calgary Laboratory Services during January 2000 to December 2002 were included in this study. These organisms were screened for the presence of ESBLs and then investigated for the presence of CTX-M- β -lactamases. Strains were identified to the species level with Vitek (Vitek AMS; bioMérieux Vitek Systems Inc., Hazelwood, Mo.).

Antimicrobial susceptibility testing. MICs of the following drugs were determined by Vitek (Vitek AMS): piperacillin (PIP), piperacillin-tazobactam (TZP), cefpodoxime (CPD), cefotaxime (CTX), and ceftazidime (CAZ). The quality control strains used for this study were *E. coli* ATCC 25922, *E. coli* ATCC 35218, *Pseudomonas aeruginosa* ATCC 27853, *Staphylococcus aureus* ATCC 29213, and *K. pneumoniae* ATCC 700603. Throughout this study, results were interpreted using NCCLS criteria for broth dilution (30).

Screening for and confirmation of ESBLs. The presence of ESBLs was evaluated in both the control strains and the recent clinical isolates. Screening was

* Corresponding author. Mailing address: Department of Medical Microbiology and Immunology, Creighton University School of Medicine, Omaha, NE 68178. Phone: (402) 280-5837. Fax: (402) 280-1875. E-mail: ndhanson@creighton.edu.

TABLE 1. Control strains producing well-characterized β -lactamases

Strain	Organism	β -Lactamase	Group ^a	Source or reference
CF2	<i>Enterobacter cloacae</i>	CTX-M-1	I	18
Rio-4	<i>Proteus mirabilis</i>	CTX-M-2	II	5
VER-1	<i>E. cloacae</i>	CTX-M-3	I	18
Cfr2525/96	<i>Citrobacter freundii</i>	CTX-M-3	I	21
Eco3553/98	<i>E. coli</i>	CTX-M-15	I	1
34	<i>Salmonella enterica</i> serovar Typhimurium	CTX-M-5	II	9
Rio-3	<i>Enterobacter aerogenes</i>	CTX-M-8	III	5
785D	<i>E. coli</i>	CTX-M-9	IV	36
EC97/38582	<i>E. coli</i>	CTX-M-10	I	31
EC984167	<i>E. coli</i>	CTX-M-14	IV	26
CF1	<i>E. coli</i>	CTX-M-14	IV	18
Rio-6	<i>E. coli</i>	CTX-M-16	IV	4
BM4493	<i>Klebsiella pneumoniae</i>	CTX-M-17	IV	13
ILT-2	<i>K. pneumoniae</i>	CTX-M-18	IV	35
ILT-3	<i>K. pneumoniae</i>	CTX-M-19	IV	35
CT1 ^b	<i>E. coli</i>	Toho-1	II	25
S1 ^b	<i>E. coli</i>	SHV-2		20
S2 ^b	<i>E. coli</i>	SHV-7		8
T1 ^b	<i>E. coli</i>	TEM-3		20
T4 ^b	<i>E. coli</i>	TEM-10		20
T5 ^b	<i>E. coli</i>	TEM-50		37

^a Group I includes CTX-M-1, -3, -10 to -12, -15 (UOE-1), -22, -23, -28, -29, and -30. Group II includes CTX-M-2, -4 to -7, and -20 and Toho-1. Group III includes CTX-M-8. Group IV includes CTX-M-9, -13, -14, -16, to -19, -21, and -27 and Toho-2. Finally, group V includes CTX-M-25 and -26. We were unable to obtain a strain producing any group V enzymes.

^b These strains are part of an unpublished isogenic panel. The original strains producing these β -lactamases were provided to the Center for Research in Anti-Infectives and Biotechnology by the referenced authors listed in the table.

performed with Vitek (Vitek AMS) using 1- μ g/ml CPD, CAZ, and CTX. Screening and disk confirmation tests using CTX (30 μ g) and CAZ (30 μ g) disks in combination with 10 μ g of clavulanate (CLA) were performed and interpreted by NCCLS criteria for ESBL screening and disk confirmation tests (30). Disks for ESBL confirmation tests were obtained from Oxoid, Inc. (Nepean, Ontario, Canada). *K. pneumoniae* ATCC 700603 and *E. coli* ATCC 25922 were used as positive and negative controls, respectively.

β -Lactamase gene identification. DNA template preparation and PCR amplification for CTX-M- β -lactamase genes were carried out on a Thermal Cycler 9600 instrument (Applied Bio-systems, Norwalk, Conn.) as previously described (23). The primers, sizes of the expected amplification product, and annealing temperatures used for PCR amplification are listed in Table 2. Magnesium chloride concentrations were 1.5 mM for all PCRs.

Nucleotide sequence accession number. The GenBank nucleotide sequence accession numbers for the sequences used in this study were as follows: CTX-M-1, X92506; CTX-M-2, X92507; CTX-M-3, AF550415; CTX-M-15, AY044436; CTX-M-4, Y14156; CTX-M-5, AF286192; CTX-M-6, AJ005044; CTX-M-7,

AJ005045; CTX-M-8, AF189721; CTX-M-9, AJ416345; CTX-M-10, AF255298; CTX-M-11, AJ310929; CTX-M-12, AF305837; CTX-M-13, AF252623; CTX-M-14, AF252622; CTX-M-16, AY029068; CTX-M-17, AF454633; CTX-M-18, AF325133; CTX-M-19, AF325134; CTX-M-20, AJ416344; CTX-M-21, AJ416346; CTX-M-22, AY080894; CTX-M-23, AF488377; CTX-M-24, AY143430; CTX-M-25, AF518567; CTX-M-26, AY455830; CTX-M-27, AY156923; CTX-M-28, AJ549244; CTX-M-29, AY267213; CTX-M-30, AY292654; CTX-M-31, AJ567482; CTX-M-32, AJ557142; CTX-M-33, AY238472; Toho-1, D37830; Toho-2, AF311345; Toho-3, AB038771; and FEC-1, AB098539.

RESULTS

Molecular detection of CTX-M genes. Currently, there are 40 gene sequences designated as *bla*_{CTX-M} in the National Center for Biotechnology Information GenBank database (3).

TABLE 2. Primers used for amplification

Target(s) ^a	Primer	Sequence ^b	Product size (bp)	Annealing temp (°C)	Nucleotide positions (bp) ^c	GenBank accession no. ^d
CTX-M group I	CTXM1-F3 CTXM1-R2	GAC GAT GTC ACT GGC TGA GC AGC CG C CGA CGC TAA TAC A	499	55	416–435 914–896	X92506
CTX-M group II	TOHO1-2F TOHO1-1R	GCG ACC TGG TTA ACT ACA ATC C CGG TAG TAT TGC CCT TAA GCC	351	55	313–334 663–643	X92507
CTX-M group III	CTXM825F CTXM825R	CGC TTT GCC ATG TGC AGC ACC GCT CAG TAC GAT CGA GCC	307	55	475–495 781–764	AF189721
CTX-M group IV	CTXM914F CTXM914R	GCT GGA GAA AAG CAG CGG AG GTA AGC TGA CGC AAC GTC TG	474	62	1857–1876 2330–2311	AF252622

^a Group I includes CTX-M-1, -3, -10 to -12, -15 (UOE-1), -22, -23, -28, -29, and -30. Group II includes CTX-M-2, -4 to -7, and -20 and Toho-1. Group III includes CTX-M-8. Group IV includes CTX-M-9, -13, -14, -16 to -19 and -21, and -27 and Toho-2. Finally, group V includes CTX-M-25 and -26.

^b Sequence of primer as synthesized 5' to 3'.

^c Nucleotide position in base pairs for the GenBank accession number sequence.

^d Accession number of the sequence used for primer design.

TABLE 3. Laboratory diagnosis of control strains producing known ESBLs

Strain	Enzyme	Result by: ^a									
		Screening ^b			Disk confirmation ^c		PCR primer pairs ^d				
		CPD	CAZ	CTX	CAZ/CLA	CTX/CLA	CTX-M-I group	CTX-M-II group	CTX-M-III group	CTX-M-IV group	
CF2	CTX-M-1	+	+	+	-	+	+	-	-	-	
Rio-4	CTX-M-2	+	-	+	-	+	-	+	-	-	
VER-1	CTX-M-3	+	+	+	-	+	+	-	-	-	
Cfr2525/96	CTX-M-3	+	+	+	-	+	+	-	-	-	
Eco3553/98	CTX-M-15	+	+	+	+	+	+	-	-	-	
34	CTX-M-5	+	+	+	-	+	-	+	-	-	
Rio-3	CTX-M-8	+	-	+	-	+	-	-	+	-	
785D	CTX-M-9	+	-	+	-	+	-	-	-	+	
EC97/38582	CTX-M-10	+	-	+	-	+	+	-	-	-	
EC984167	CTX-M-14	+	-	+	-	+	-	-	-	+	
CF1	CTX-M-14	+	-	+	-	+	-	-	-	+	
Rio-6	CTX-M-16	+	+	+	+	+	-	-	-	+	
BM4493	CTX-M-17	+	+	+	+	+	-	-	-	+	
ILT-2	CTX-M-18	+	+	+	-	+	-	-	-	+	
ILT-3	CTX-M-19	+	+	+	+	+	-	-	-	+	
CT1	Toho-1	+	-	+	-	+	-	+	-	-	
S1	SHV-2	+	+	+	+	+	-	-	-	-	
S2	SHV-7	+	+	+	+	+	-	-	-	-	
T1	TEM-3	+	+	+	+	+	-	-	-	-	
T4	TEM-10	+	+	-	+	+	-	-	-	-	
T5	TEM-50	+	+	+	+	+	-	-	-	-	

^a -, negative; +, positive.

^b NCCLS guidelines for screening for ESBL-producing bacteria. CPD, CAZ, and CTX were each used at 1 µg/ml.

^c NCCLS guidelines for ESBL disk confirmation tests.

^d PCR primers for CTX-M groups (refer to Materials and Methods for details). Group I includes CTX-M-1, -3, -10 to -12, -15 (UOE-1), -22, -23, -28, -29, and -30. Group II includes CTX-M-2, -4 to -7, and -20 and Toho-1. Group III includes CTX-M-8. Group IV includes CTX-M-9, -13, -14, -16 to -19, -21, and -27 and Toho-2. Finally, group V includes CTX-M-25 and -26.

Based on percent sequence similarities, *bla*_{CTX-M} genes can be clustered into five different groups (3). Nucleotide similarities between these genes were analyzed by using DNAsis for Windows 2.6 (Hitachi Software), and four primer sets were designed which would amplify family- or group-specific CTX-M genes. These groups were designated CTX-M groups I, II, III, and IV. These groups and the specific family members within the group are listed in Tables 1 and 2. The group III primer set was designed to amplify both *bla*_{CTX-M-8} and *bla*_{CTX-M-25}. This was done to simplify the number of primer pairs required to identify *bla*_{CTX} gene families. The forward primer of primer pair III (*bla*_{CTX-M-8}) has 100% identity to *bla*_{CTX-M-8} but has three mismatches at the 5' end for *bla*_{CTX-M-25}; the reverse primer is 100% identical to *bla*_{CTX-M-8} and has a 1-base mismatch at position 5 of the 18-base primer for *bla*_{CTX-M-25}. The accession numbers used for sequence alignments in this study are listed in Materials and Methods.

The designed primers were tested for specificity in separate PCRs using a DNA template prepared from control strains known to produce specific CTX-M β-lactamases or strains producing ESBLs other than CTX-M β-lactamases (Tables 1 and 3 and Fig. 1). PCR amplification of DNA template prepared from strains CF2, VER-1, Cfr2525/96, Eco3553/98, and EC97/38582 resulted in a single amplified product of 499 bp when CTX-M group I primers were used. No amplified product was identified with this primer set when DNA template from the rest of the control strains in Table 1 was used during PCR amplification. Group II primers amplified a single 351-bp fragment when DNA template was prepared from control strains Rio-4, 34, and CT1. All other control strain DNA tem-

plates resulted in no amplification product for this primer set. Group IV primers amplified a 474-bp product from DNA prepared from strains 785D, EC984167, CF1, Rio-6, BM4493, ILT-2, and ILT-3. This primer set was also very specific, re-

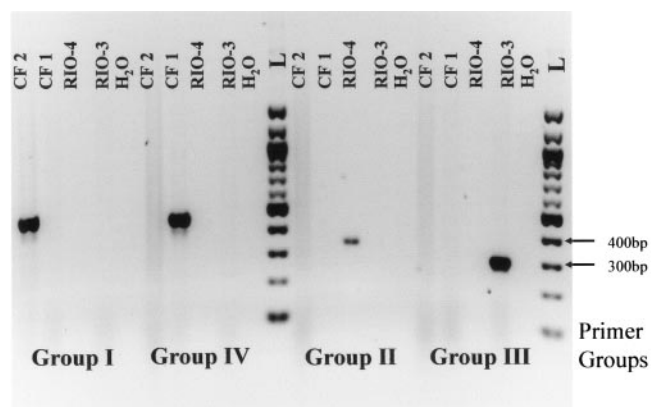


FIG. 1. CTX-M PCR. Four primer sets representing specific groups of *bla*_{CTXM} genes were evaluated for specificity with DNA prepared from strains known to produce specific CTX-M β-lactamases. The DNA templates were prepared from CF2 producing CTX-M-1, CF1 producing CTX-M-14, Rio-4 producing CTX-M-2, and Rio-3 producing CTX-M-8. H₂O represents the negative control, and L represents a 100-bp ladder (Invitrogen). Specific marker sizes in base pairs are shown on the right. Each primer pair was used in combination with both positive and negative templates. The primer groups used for each set of templates are listed below the figure and are defined in Table 2.

TABLE 4. Laboratory diagnosis of clinical strains producing ESBLs

Strains (n)	PCR for <i>bla</i> _{CTX-M} genes ^a	No. (%) of strains					
		Screening ^b			Disk confirmation ^c		
		CPD	CAZ	CTX	CAZ/CLA	CTX/CLA	CTX/CLA and CAZ/CLA
<i>E. coli</i> (51)	Negative	51 (100)	51 (100)	48 (94)	34 (67)	49 (96)	51 (100)
<i>K. pneumoniae</i> (5)	Negative	5 (100)	5 (100)	4 (80)	3 (60)	5 (100)	5 (100)
<i>E. coli</i> (24)	Positive for CTX-M-I group	24 (100)	24 (100)	24 (100)	21 (88)	24 (100)	24 (100)
<i>E. coli</i> (93)	Positive for CTX-M-IV group	93 (100)	71 (76)	93 (100)	3 (3)	93 (100)	93 (100)
<i>K. pneumoniae</i> (2)	Positive for CTX-M-IV group	2 (100)	0 (0)	2 (100)	0 (0)	2 (100)	2 (100)
Total (175)		175 (100)	151 (86)	171 (98)	61 (35)	173 (99)	175 (100)

^a PCR primers for CTX-M subgroups (see Materials and Methods for details). Group I includes CTX-M-1, -3, -10 to -12, -15 (UOE-1), -22, -23, -28, -29, and -30. Group II includes CTX-M-2, -4 to -7, and -20 and Toho-1. Group III includes CTX-M-8. Group IV includes CTX-M-9, -13, -14, -16 to -19, -21, and -27 and Toho-2. Finally, group V includes CTX-M-25 and -26.

^b NCCLS guidelines for screening ESBL-producing bacteria. CPD, CAZ, and CTX were each used at 1 µg/ml.

^c NCCLS guidelines for ESBL disk confirmation tests.

sulting in no amplification of DNA when template was prepared from the other control strains. Use of the group III primer set resulted in amplification of DNA prepared from only one strain, Rio-3, which produced CTX-M-8. An isolate producing CTX-M-25 was requested but not obtained. Therefore, we were unable to examine the ability of the group III primer set to amplify *bla*_{CTX-M-25}. A representative gel indicating the specificity of the primer pairs is shown in the Fig. 1. These data indicate a high level of specificity for these group-specific primer pairs.

Clinical bacterial strains. During the study period, 232 *E. coli* strains producing ESBLs were isolated from 168 patients and 11 *K. pneumoniae* strains were isolated from 7 patients. The overall frequencies of ESBL producers observed in this study were 1.3% for *E. coli* and 0.5% for *K. pneumoniae*. Of the 175 (168 plus 7) strains included in this study, 154 (88%) were isolated from urine cultures, 7 (4%) were isolated from blood, 5 (3%) were isolated from both the blood and urine of a single patient, 3 (2%) were isolated from wounds (purulent), 2 (1%) were isolated from sputum, 2 (1%) were isolated from both sputum and urine of a single patient, and 1 (1%) each was isolated from bronchoalveolar lavage and stool (associated with symptomatic diarrhea), respectively.

Antimicrobial susceptibility of clinical strains. All the ESBL-producing *E. coli* strains were resistant to CPD and PIP, 119 (71%) were resistant to CTX, 29 (17%) were resistant to CAZ, and 7 (4%) were resistant to TZP. Among the ESBL-producing *K. pneumoniae* strains, all were resistant to CPD and PIP, 3 (43%) were resistant to CTX, and 1 (14%) was resistant to CAZ. All of the ESBL-producing *K. pneumoniae* strains were susceptible to TZP.

Identification of *bla*_{CTX-M} genes in the clinical strains. All of the ESBL-producing *E. coli* and *Klebsiella* spp. were examined by PCR for the presence of *bla*_{CTX-M} genes. Of the 168 *E. coli* strains isolated during the study period, 24 (14%) were positive for *bla*_{CTX-M} genes from the CTX-M-I group, indicating CTX-M-1-like β-lactamases. Ninety-three (55%) were positive for *bla*_{CTX-M} genes from the CTX-M-IV group, indicating CTX-M-14-like β-lactamases, and the remainder (51 [31%]) were negative for *bla*_{CTX-M} genes (Table 4). Of the seven *K. pneumoniae* strains isolated during the study period, two (29%)

were positive for *bla*_{CTX-M} genes from the CTX-M-IV group and the remainder (five [71%]) were negative for *bla*_{CTX-M} genes (Table 4).

Screening and confirmation for ESBLs by NCCLS criteria.

Using the 1999 NCCLS criteria for detection of ESBLs in *E. coli* and *Klebsiella* spp., all of the control and clinical strains from the current study producing ESBLs were positive when screened with CPD (1 µg/ml) (Tables 3 and 4) (29). Because the CPD MICs for all organisms examined were greater than 4 µg/ml, the new guidelines set forth in 2003 by the NCCLS changing the CPD concentration to 4 µg/ml would not have affected the results described in this study. It is important to note that according to the references listed in Table 1, none of the control strains tested produced an ESBL other than the CTX-M β-lactamase identified. Almost every reference indicated the presence of TEM-1 in the control strains, and in the case of *K. pneumoniae* strains, a band with a pI of 7.6 was reported as SHV-1. The production of an AmpC β-lactamase was not indicated in any of the references describing these strains (see references in Table 1). All of the control strains expressing various ESBLs were detected with 1-µg/ml CTX except for TEM-10. In addition, 98% of the clinical strains from the current study were positive with 1-µg/ml CTX; the four exceptions were *bla*_{CTX-M}-negative strains. Using CAZ at 1 µg/ml did not detect control strains producing CTX-M-2, -8, -9, -10, and -14 and Toho-1 or 14% of the clinical strains from the current study, which were all positive for *bla*_{CTX-M-14-like} (Tables 3 and 4). The NCCLS disk confirmation test using CTX with and without CLA was positive for all of the control strains and 99% of the clinical strains from the present study (Table 3 and 4). The disk confirmation tests using CAZ with and without CLA were positive for the control strains producing CTX-M-15 (UOE-1), -16, -17, and -19. These β-lactamases are part of groups I and IV (Table 1 and 3). Sixty-five percent of the current clinical strains tested negative (35% positive) with CAZ with and without CLA (Table 4). These data reflect the molecular data, which indicated that only 3% of the current clinical strains with a *bla*_{CTX-M-14-like} gene tested positive by the CAZ and CLA disk confirmation test (Table 4). All of the strains (control and clinical) were positive for at least one of the confirmation tests, but by using both CTX/CLA and CAZ/

CLA disk confirmation tests, all strains producing an ESBL were detected (Table 4).

DISCUSSION

There has been a dramatic increase in the number of organisms reported in the literature that produce CTX-M- β -lactamases (3). This class of β -lactamases has been recognized worldwide as an important mechanism of resistance to oxyiminocephalosporins used by gram-negative pathogens (3). In most cases, organisms producing these enzymes display higher levels of resistance to CTX and ceftriaxone than CAZ (3). However, organisms producing some CTX-M variants, including CTX-M-16 and -19 and UOE-1, are resistant to CAZ (4, 33, 35).

Phenotypic differentiation of organisms producing CTX-M- β -lactamases from organisms producing other types of ESBLs can be difficult. The difficulty is due to overlapping phenotypes resulting in interference from other β -lactamases produced by the organism capable of hydrolyzing CAZ (2, 4, 35). Therefore, susceptibility testing which relies on identifying organisms that are resistant to CTX and/or ceftriaxone but susceptible to CAZ is not a reliable approach. In addition, the use of isoelectric focusing to identify β -lactamases is becoming obsolete because many isoelectric points of different β -lactamases overlap, including CTX-M- β -lactamases (3, 7). PCR amplification and sequencing of *bla*_{CTX-M} genes have been used to characterize organisms producing CTX-M- β -lactamases (6, 12, 14). Recently a molecular approach for screening ESBL-positive organisms for the presence of CTX-M genes was described by Edelstein et al. (19). Consensus primers, which recognize all the known variant genes of *bla*_{CTX-Ms} to date, were used to generate an amplified product of 544 bp. To identify the specific groups of *bla*_{CTX-Ms}, restriction fragment length polymorphism (RFLP) analysis was employed. The technique described in this study uses four specific primer sets to detect the various groups of CTX-M- β -lactamase genes, thus negating the need for RFLP analysis. Data generated using several control strains known to produce specific CTX-M- β -lactamases validated the specificity and sensitivity of these primer sets. A control strain producing a CTX-M-25 β -lactamase (belonging to group V) was not available, and although the primers designed to amplify *bla*_{CTX-M-8} (belonging to group III) contained one mismatch with respect to *bla*_{CTX-M-25} in the reverse primer and three mismatches in the 5' portion of the forward primer, these mismatches should still allow amplification from *bla*_{CTX-M-25} given the stringency with which the PCR is modified (unpublished data on mismatched primer pairs). Even though this PCR assay involves the use of four sets of primers, a single DNA fragment is amplified for each CTX-M group. Therefore, interpretation of results is simple and can be adapted in reference laboratories for screening multiple isolates for the presence of group-specific CTX-M- β -lactamase genes (Fig. 1).

The four-primer-pair PCR-based detection system was used to screen 168 ESBL-producing *E. coli* strains and 7 *K. pneumoniae* strains for the presence of genes encoding CTX-M- β -lactamases recovered from the Calgary Health Region (CHR) during 2000 to 2002. The CHR is a fully integrated, publicly funded health system that provides health care to the residents

of the cities of Calgary and Airdrie and approximately 20 nearby small towns, villages, and hamlets (overall population of 958,610 in 2001). In the CHR, Calgary Laboratory Services receives all clinical specimens submitted for bacteriologic testing, including those from all hospitals, nursing homes, physicians' offices, and community collection sites (15). The majority of ESBL-producing bacteria (68%) isolated in the CHR during 2000 to 2002 carried a CTX-M- β -lactamase gene (Table 4). To our knowledge, this is the first study that identified strains with *bla*_{CTX-M} genes as the predominant type of ESBL in a well-defined North American region, although a previous study has shown the presence of CTX-Ms in Canada (28).

The limitations of molecular analyses for resistance genes result from the presence of unknown mutations which might occur in the primer target region or the evolution of gene products which have not yet been identified at the genetic level. Therefore, any negative PCR result must be evaluated with this in mind. Thirty-five percent of the 175 ESBL-producing strains in this report were negative by PCR for *bla*_{CTX-M}. These data could indicate that the ESBL phenotype is due to production of ESBLs other than CTX-Ms. However, the negative PCR results in this report do not negate the possibility that modified *bla*_{CTX-Ms} were present in these isolates. Due to the increased complexity of β -lactam resistance in gram-negative organisms, the key to effective surveillance is the use of both phenotypic and genotypic analyses in concert.

The detection of organisms producing ESBLs remains a contentious issue. Proficiency testing studies performed by the Centers for Disease Control and Prevention and the College of American Pathologists have raised concerns about the current capacity of many laboratories to detect organisms producing ESBLs (22, 39–41). A study recently published showed that only 8% of microbiology laboratories from rural hospitals in the United States routinely screen for ESBL-producing organisms (38). Since the majority of patients infected with a strain of ESBL-producing *E. coli* or *Klebsiella* spp. identified in our study originated from the community (32), it is conceivable to predict that ESBL-producing organisms are present in the community in North America but are not being reported.

The NCCLS guidelines for ESBL detection in *E. coli* and *Klebsiella* spp. include an initial screening with either CPD, CTX, CAZ, ceftriaxone, or aztreonam, followed by a confirmation test using both CTX and CAZ in combination with clavulanate (30). A practice exists among some clinical laboratories of using CAZ as the initial screening drug and CAZ with CLA as the confirmation test (10, 16). Data from this study indicate that 14% of ESBL-producing strains will not be detected if CAZ is used as the initial screen. However, in this study, CPD detected all of the ESBL-positive strains (Table 4). The concern raised by this study is that only 35% of ESBL-producing strains (20% of CTX-M-producers) were reported as ESBL positive when CAZ with CLA was the only confirmation test (Table 4). Therefore, a more appropriate approach for initially screening organisms for the presence of ESBLs is to use CPD followed by disk tests using both CTX with CLA and CAZ with CLA to confirm the presence of ESBLs among *E. coli* and *Klebsiella* spp.

Organisms with genes encoding CTX-M-14-like β -lactamases were responsible for an outbreak among elderly patients in the Calgary community during 2000 (32). This outbreak was

only recognized after the PCR identification of CTX-M-14-like β -lactamase genes. If molecular detection assays, such as the one described in this paper are available at the time of an outbreak and performed by a reference laboratory, early recognition and the possible mechanism(s) by which resistance is spread can be identified in a timely manner. Extra efforts such as molecular procedures to identify resistance mechanisms can promote optimal patient care by early detection of antimicrobial-resistant organisms and implementation of appropriate infection control procedures.

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