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Comparison of the SUPERCARBA, CHROMagar KPC, and Brilliance CRE screening media for detection of Enterobacteriaceae with reduced susceptibility to carbapenems $\frac{1}{2}, \frac{1}{2}, \frac{1}{2}$

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ABSTRACT

The recently developed SUPERCARBA medium was evaluated together with 2 commercially available selective culture media containing carbapenems: CHROMagar KPC (CHROMagar) and Brilliance CRE (Oxoid, Thermofisher Scientific). A total of 142 enterobacterial isolates were tested, including 131 isolates with reduced susceptibility to carbapenems. The SUPERCARBA medium has the highest sensitivity (96.5%) (detecting virtually all carbapenemase producers including OXA-48–like producers) as compared to Brilliance CRE (76.3%) and CHROMagar KPC (43%). The specificity of the screening media was similar, ranging from 57% to 68%.

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The spread of carbapenemase-producing Enterobacteriaceae is increasingly reported worldwide (Castanheira et al., 2011; Nordmann et al., 2011, 2012a; Poirel et al., 2012). Targeted surveillance of highrisk patients and screening are essential to prevent outbreaks of nosocomial infections by these organisms. The clinically significant carbapenemases in Enterobacteriaceae belong either to Ambler class A (KPC-type) hydrolysing all β -lactams except cephamycins; Ambler class B (NDM, VIM, and IMP), which are zinc-dependent metallo- β lactamases (MBL) hydrolysing all β -lactams except aztreonam; and Ambler class D enzymes (OXA-48-like) hydrolysing carbapenems but weakly (or not) broad-spectrum cephalosporins (Nordmann et al., 2011, 2012a). The level of resistance to carbapenems conferred by those carbapenemase producers may vary significantly, making their detection difficult when just based on their in vitro susceptibility profile (Landman et al., 2010). The SUPERCARBA medium has been specifically developed for the detection of carbapenemase producers and, in particular, of OXA-48 carbapenemase-producing organisms (Nordmann et al., 2012b). OXA-48 producers currently represent a worrisome threat in North African countries, the Middle East, Turkey, and the Indian subcontinent and Europe (Poirel et al., 2012). Moreover, the spread of MBL and KPC producers has created a real

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need for a reliable medium that is efficient for the detection of all types of carbapenemase-producing isolates (Adler et al., 2011; Vatopoulos, 2008). The present study was aimed to compare the performance of the recently developed SUPERCARBA medium (Nordmann et al., 2012b), which is supplemented with ertapenem, cloxacillin, and zinc sulfate, with commercially available selective media for the screening of carbapenemase producers.

One hundred and forty-two enterobacterial isolates were tested, including 131 isolates exhibiting reduced susceptibility or resistance to carbapenems. The B-lactamase content of all these isolates has been characterized at the molecular level (Table 1). The tested isolates were as follows: OXA-48–like producers (n = 43), KPC producers (n = 20), VIM producers (n = 18), IMP producers (n = 17), and NDM producers (n = 16), and noncarbapenemase producers with reduced susceptibility or resistance to carbapenems (AmpC overproducers, extendedspectrum β -lactamase [ESBL] producers combining porin deficiency) (n = 17), and carbapenem-susceptible isolates (n = 11). Strains with reduced susceptibility to ertapenem due to an overexpressed AmpC or to an ESBL, and/or porin deficiency had been previously characterized (inhibition of AmpC activity by using cloxacillin containing plates, polymerase chain reaction and sequencing of AmpC genes, measurement of carbapenem hydrolysis by UV spectrophotometry) (Nordmann et al., 2012b; Caroff et al., 2000).

MIC values of imipenem, ertapenem, and meropenem were determined by Etest and interpreted according to the updated 2012 Clinical and Laboratory Standards Institute (CLSI) guidelines (CLSI, 2012) (Table 1). The lowest detection limit of the carbapenemase

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Table 1

Sensitivity of detection of SUPERCARBA, Brilliance CRE, and CHROMagar KPC media for 142 carbapenemase- and/or ESBL/AmpC-producing enterobacterial isolates.

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Strains	β-Lactamase content	MIC (mg/L)		Lowest detection	on limit (CFU/mI	.)"
		IPM ^a	ETP	MEM	SUPER CARBA	Brilliance CRE	CHROMagar KPC
C. 1	tore (n. 12)						0
Carbapenemase OXA-48-	-type $(n = 43)$	0.5	2	0.5	1 101	1 101	5 106
K. pneumoniae BIC	0XA-48°	0.5	2	0.5	1×10^{1}	1×10^{1}	$\frac{5 \times 10^{\circ}}{1 \times 10^{5}}$
K. pneumoniae BEL	0XA-48	1	4	1	1×10^{1}	1×10^{1}	$\frac{1 \times 10^{\circ}}{1 \times 10^{\circ}}$
K. pneumoniae RAM	OXA-48	1	4	1	1×10^{1}	1×10^{4}	$\frac{1 \times 10^3}{1 \times 10^4}$
K. pneumoniae LIB	OXA-48	16	16	16	1×10^{1}	$>2 \times 10^{7}$	$\frac{5 \times 10^4}{2}$
K. pneumoniae SCO	OXA-48	0.5	0.75	0.25	1×10^{1}	$>2 \times 10^{7}$	$>2 \times 10^{7}$
K. pneumoniae LOU	OXA-48	4	16	0.5	1×10^{1}	$>6 \times 10^{7}$	$>6 \times 10^{7}$
K. pneumoniae TIK	OXA-48	0.75	2	0.38	1×10^{1}	1×10^{1}	$>1 \times 10^{\circ}$
K. pneumoniae OM14	OXA-48 + TEM-1	0.5	1	0.38	1×10^{1}	$>5 \times 10^{7}$	5×10^{7}
K. pneumoniae CHA	OXA-48 + TEM-1	0.38	1	0.5	1×10^{1}	$>3 \times 10^{7}$	$>3 \times 10^{7}$
K. pneumoniae BOU	OXA-48 + CTX-M-15	0.38	0.5	0.25	1×10^{1}	$>1 \times 10^{7}$	1×10^{8}
K. pneumoniae EGY	OXA-48 + CTX-M-15	2	3	2	1×10^{1}	1×10^{1}	1×10^{5}
K. pneumoniae ROU	OXA-48 + CTX-M-15	0.5	1.5	0.25	1×10^{1}	1×10^{1}	$>1 \times 10^{8}$
K. pneumoniae BEY	OXA-48 + CTX-M-15 + TEM-1	0.38	0.38	0.38	5×10^{2}	1×10^{1}	1×10^{8}
K. pneumoniae DAL	OXA-48 + CTX-M-15 + TEM-1	0.38	2	0.38	1×10^1	1×10^1	4×10^5
K. pneumoniae BAJ	OXA-48 + CTX-M-15 + SHV-28 + TEM-1	0.5	1.5	0.38	1×10^1	1×10^1	$>1 \times 10^{8}$
K. pneumoniae BEN	OXA-48 + CTX-M-15 + SHV-28 + TEM-1	0.38	1	0.25	1×10^1	1×10^1	$>1 \times 10^{8}$
K. pneumoniae DUW	OXA-48 + CTX-M-15 + TEM-1 + SHV-28	32	32	32	1×10^1	1×10^1	1×10^{1}
K. pneumoniae SIC	OXA-48 + CTX-M-15 + SHV-28	0.25	1	0.25	1×10^1	1×10^{1}	$>1 \times 10^{8}$
K. pneumoniae AEL	OXA-48 + CTX-M-15 + SHV-28 + OXA-1	0.5	6	0.38	1×10^{1}	1×10^{1}	5×10^2
K. pneumoniae AMS	OXA-48 + CTX-M-15 + TEM-1 + OXA-1	0.5	2	0.38	1×10^{1}	1×10^{1}	$>1 \times 10^{8}$
K. pneumoniae FLK	0XA-48 + CTX-M-15 + TEM-1 + SHV-11	0.5	3	0.38	1×10^{1}	1×10^{1}	$>1 \times 10^8$
K nneumoniae VFR	0XA-48 + CTX-M-15 + TFM-1 + SHV-11	0.38	2	0.30	1×10^{1}	1×10^{1}	$>1 \times 10^8$
K nneumoniae VSC	0XA-48 + (TX-M-15 + 0XA-1 + TFM-1)	0.75	2	0.75	1×10^{1}	1×10^{1}	$>1 \times 10^8$
K nnoumoniao UDA	$\mathbf{OYA}_{\mathbf{A}} = \mathbf{OYA}_{\mathbf{A}} + \mathbf{OYA}_{\mathbf{A}} = \mathbf{OYA}_{\mathbf{A}} + \mathbf{OYA}_{\mathbf{A}$	1.5	ູ່າ	12	1×10^{1}	1×10^{1}	$\frac{1}{1} \times 10^8$
K pneumonice OM11	$\mathbf{OXA} = \mathbf{O} + \mathbf{CIA} = \mathbf{I} + \mathbf{I} + \mathbf{I} = \mathbf{I} + \mathbf{I} + \mathbf{I} = \mathbf{I} + \mathbf{I} + \mathbf{I} = \mathbf{I} + \mathbf{I} = \mathbf{I} + \mathbf{I} = \mathbf{I} + \mathbf{I} = \mathbf{I} + I$	1.5	~32 0.75	12	1×10 1×10^{1}	1×10 1×10^{1}	5×10^7
K. prieumoniae UNITT	$\frac{1}{1000} + \frac{1}{100} + 1$	0.5	0.75	0.25	1×10 2×10^{1}	1×10	$\frac{J \times 10}{56 \times 10^7}$
E. COIL NOV		0.5	0.75	0.20	2×10^{-1}	$\sim 0 \times 10^{\circ}$	2×10^{-104}
E. COIL HAN	0XA - 48 + CIX - M - 15	3	16	1	5×10^{-1}	$>6 \times 10^{7}$	$\frac{3 \times 10^{\circ}}{4 \times 10^{\circ}}$
E. coli BOU	$\mathbf{OXA-48} + \mathbf{CIX}-\mathbf{M}-15$	0.5	0.75	0.12	2×10^{1}	$>4 \times 10^{\prime}$	$>4 \times 10^{7}$
E. coli OM3	0XA-48 + CIX-M-15 + TEM-1	0.5	1	0.38	1×10^{1}	1×10^{1}	$>1 \times 10^{8}$
E. coli OM22	OXA-48 + CTX-M-15 + TEM-1	0.5	1	0.25	1×10^{1}	1×10^{1}	$\geq 1 \times 10^{\circ}$
E. coli BER	OXA-48 + CTX-M-15 + TEM-1	0.38	1.5	0.19	5×10^{1}	$>1 \times 10^{7}$	$>1 \times 10^{7}$
E. coli AME	OXA-48 + CTX-M-24	0.25	0.5	0.19	2×10^{1}	$>1 \times 10^{8}$	$>1 \times 10^{8}$
E. coli ZAN	OXA-48 + TEM-1 + CTX-M-14	0.38	8	0.75	1×10^1	1×10^1	$>1 \times 10^{8}$
E. coli BON	OXA-48 + CTX-M-24 + TEM-1	0.38	0.5	0.19	1×10^2	1×10^2	$>1 \times 10^{8}$
E. coli BOK	OXA-48 + CTX-M-15	0.25	0.38	0.19	1×10^2	1×10^1	$>1 \times 10^{8}$
E. cloacae TUR	OXA-48 + SHV-5	0.5	0.5	0.5	1×10^1	1×10^1	1×10^7
E. cloacae 501	OXA-48 + CTX-M-15 + TEM-1	1	16	1.5	1×10^1	1×10^1	1×10^{1}
E. cloacae BEU	OXA-48 + CTX-M-15 + TEM-1 + SHV-12	0.5	8	0.5	1×10^2	1×10^{2}	1×10^4
C. koseri ROU	OXA-48	0.38	2	0.38	1×10^{1}	$>1 \times 10^{8}$	$\frac{1}{>1 \times 10^8}$
C. koseri VER	OXA-48	0.75	2	0.38	1×10^{1}	$>1 \times 10^8$	$> \frac{1}{1 \times 10^8}$
K pneumoniae HOL	0XA-181 + CTX-M-15	1	4	1	1×10^{1}	1×10^1	1×10^{1}
K pneumoniae KP3	0XA-181 + CTXM-15 + 0XA-1	0.5	2	0.5	1×10^{1}	1×10^{1}	$>1 \times 10^{8}$
P rettgeri RAP	0XA-181 + 0XA-1	8	1	2	5×10^{2}	1×10^{1}	1×10^1
VPC(n - 20)		0	1	2	$J \times 10$	1 × 10	1 × 10
KFC (n = 20)	VDC 2 \downarrow CUV 11	\ 22	\ 22	\ 22	1×10^{1}	$>6 \times 10^7$	1×10^{1}
K. pheumoniae 2303	VDC 2 + OVA 0 + TEM 1	>32	~32	<22 <22	1×10 1×10^{1}	$\frac{20 \times 10}{1 \times 10^1}$	1×10^{1}
K nneumoniae TOP	$KDC_2 + OXA_0 + TEM_1$	∕s∠ ∧	~52 17	~ 52 /	$1 \wedge 10$ $1 \vee 10^{1}$	1×10 1×10^{1}	1×10^{1}
K. pheumoniae TUO	$\mathbf{KFC} = 2 + \mathbf{O}\mathbf{X}\mathbf{A} + \mathbf{TEM} + 1$	4	12	4	1×10^{1}	1×10 $1 \cdots 10^{1}$	1 × 10
K. pneumonice 1HU	$\mathbf{VDC} = \mathbf{V} \mathbf{V} \mathbf{A} \mathbf{O} + \mathbf{T} \mathbf{E} \mathbf{W}^{-1}$	~52 6	≥52 0	∕ວ∠ ว	1×10 1×10^{1}	1×10 1×10^{1}	1×10^{1}
K. pheumoniae 20111	$\mathbf{NFC^2} + \mathbf{O}\mathbf{A} + \mathbf{IEW} = \mathbf{I}$	U \	0	2	1×10 1×10^{1}	1×10 1×10^{1}	1×10^{1}
K. prieumoniae IIF	$\mathbf{NPC-2} + \mathbf{OXA-9} + \mathbf{1EW-1}$	>32	>32	>32	1×10^{-1}	1×10^{1}	1 × 10 1 ··· 10 ¹
K. prieumoniae LIE	$\mathbf{NFC-2} + \mathbf{OXA-9} + \mathbf{1EWI-1}$	>32 24	>32 22	>32 10	5×10^{-1}	1×10^{1}	I × IU ¹
K. prieumoniae 588	NTC-2 + UXA-9 + STV-11 + 1EW-1	24 4	32	10	1 × 10°	1 × 10 ¹	$I \times IU^{1}$
к. pneumoniae YC	$\mathbf{NPC-2} + \mathbf{UXA-9} + \mathbf{SHV-11} + \mathbf{SHV-12} + \mathbf{IEM-1}$	4	24	2	1 × 10'	1×10^{1}	1 × 10 ¹
к. pneumoniae A28006	KPC-2 + CIX-M-2 + SHV-11 + TEM-1	16	24	32	2×10^{4}	$1 \times 10^{\circ}$	$I \times IU^{1}$
K. pneumoniae A33504	$\mathbf{KPC-2} + \mathbf{CTX} - \mathbf{M} - 2 + \mathbf{SHV} - 11 + \mathbf{OXA} - 9 + \mathbf{TEM} - 1$	>32	>32	>32	1×10^{1}	1×10^{1}	$1 \times 10^{\circ}$
K. pneumoniae COL	KPC-2 + TEM-1 + SHV-1 + CTXM-15	4	4	32	1×10^{1}	1×10^{1}	1×10^{1}
K. pneumoniae 475	KPC-2 + CTX-M-15 + SHV-11	16	>32	>32	1×10^{1}	1×10^{1}	1×10^{1}
K. pneumoniae KAM	KPC-3 + TEM-1	8	12	2	1×10^{1}	1×10^1	5×10^{3}
E. coli PSP	KPC-2 + TEM-1 + OXA-1	0.5	0.5	0.5	1×10^2	$1 imes 10^1$	1×10^4
E. coli COL	KPC-2 + CTX-M-9 + TEM-1	4	4	2	1×10^1	1×10^{1}	1×10^{3}
E. cloacae HPT2	KPC-2	1	1.5	0.75	$1 imes 10^1$	$>\!\!8 imes 10^7$	1×10^4
E. cloacae CFVL	KPC-2 + TEM-3	4	2	1	$1 imes 10^1$	$>6 \times 10^7$	5×10^{3}
E. cloacae HMG	KPC-2 + TEM-1	24	>32	16	1×10^2	1×10^{1}	1×10^{1}
E. cloacae HPTU	KPC-2 + SHV-11 + TEM-1	2	4	1.5	1×10^{1}	1×10^{1}	1×10^{3}
MBL $(n = 51)$		-	•	- 10			
K. pneumoniae HK	NDM-1 + CTX-M-15 + CMY-4 + $0XA-1$	>32	>32	>32	1×10^1	1×10^{1}	1×10^{1}
K nneumoniae 6647	NDM-1 + CTX-M-15 + $OXA-1 + OXA-10$	1	16	3	1×10^{1}	1×10^{1}	1×10^{1}
K nneumoniae 6750	NDM.1 + CTX-M-15 + TFM-1 + OYA-1 + OYA 0 + OYA 10 + CMV16	12	25	222	1×10^{1}	1×10^{1}	1×10^{1}
K pneumoniae 10MA	$NDM_1 + CTY_{-M-15} + TEM_1 + CHV 11 + CHV 20 + OVA 1 + OVA 0$	12 \22	<22 <22	<22 <22	1×10^{1}	1×10^{1}	1×10^{1}
K. pneumoniae TOWA	$\frac{1}{100} + \frac{1}{100} + \frac{1}$	∕⊃∠ 1 ⊑	~52 F	∕ວ∠ ว	1×10 1×10^{1}	1×10 1×10^{1}	1×10^{2}
K. prieumoniae 201VIA	$\mathbf{NDW} 1 + \mathbf{SHV} 1 + \mathbf{UAA} 1$	1.5	0	2	1×10^{-1}	1×10^{1}	1 × 10 1 ··· 10 ¹
к. pneumoniae AFR/	MUM-1 + CIX-M-15 + 1EM-1 + CMY-6 + 0XA-1 + 0XA-9	>32	>32	>32	1×10^{1}	1×10^{1}	I × IU'
к. pneumoniae IND	NDM-1 + CIX-M-15 + IEM-1 + SHV-28 + CMY-6 + OXA-1 + OXA-9	1	8	4	1×10^{1}	1×10^{4}	$1 \times 10^{\circ}$

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