

Impact of the isolation medium for the detection of OXA-48 and KPC-producing Gram negative bacteria by immunochromatographic assays

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Introduction and Purpose

The development of new rapid diagnostic tests to track antimicrobial resistance constitutes one priority core action in the current context of multidrug resistance. We recently developed two lateral flow **immunochromatographic assays (ICAs, OXA-48 K-Set[®] and KPC K-Set[®], Coris Bioconcept, Gembloux, Belgium)** for direct confirmation of OXA-48-like and KPC-like carbapenemase based on monoclonal antibodies that were generated by immunization in mice. The evaluation of these two ICAs was already performed showing **100 % sensitivity and specificity (1, 2, 3)** with colonies grown on TSA blood agar. Here we assessed the impact of different culture isolation media and also the **impact of aged cultures** on the performance of these IC tests.

Methods

Immunochromatographic assays (ICAs):

For OXA-48 K-Set, 22 carbapenemase-producing *Enterobacteriaceae* (18 OXA-48-like, one of each VIM-1, KPC-2, NDM-1, IMP-4) and one carbapenem-non-susceptible, non carbapenemase (CTX-M-15-producing *E. coli*) were tested.

For KPC K-Set, 10 carbapenemase-producing strains (7 KPC, one of each OXA-48, IMP, NDM) and one carbapenem-non-susceptible, non carbapenemase (CTX-M-15-producing *E. coli*) were tested. ICAs were performed and results interpreted after 15 min as per manufacturers' instructions.

Culture isolation media:

All strains were grown at 37°C for 24h prior testing on the following media:

Non-chromogenic and non-selective media (TSA blood agar, Mueller-Hinton), differential media (McConkey, Drigalski), chromogenic non-selective media (Uriselect4, CHROMagar orientation medium) and chromogenic-selective media (CHROMID agar, KPC colorex).

Aged cultures:

OXA-48 and OXA-181-positive bacteria were grown on three different media (TSA, chromID BLSE and chromID OXA-48) at 37°C for 24 h. The plates were then left at room temperature for 7 days and 7 additional days at 4°C. Colonies of the same tested once a day during 15 days to evaluate the performance of the test on aged cultures.

Molecular testing:

All strains characterized by in house multiplex PCR targeting carbapenemases, ESBLs and by PCR-sequencing. A subset of strains was typed by MLST.

Results

A) OXA-48 K-Set : 23 *Enterobacteriaceae* on 18 different culture media

Species	Carbapenemase	β-lactamase genes		MICs (ng/ml)		Non chromogenic - non selective media		Differential media		Chromogenic - non selective media		Chromogenic - selective media	
		β-lactamase-PhoP/K	β-lactamase-PhoP/K	MEM	ERT	TEM	TSA	Colombia sheep blood agar	Uriselect 4	McConkey	Drigalski	Uriselect 4	CHROMID BLSE
<i>K. pneumoniae</i> PEP136	OXA-48	SHV-11	TEM-1, CTX-M-27	<1	0.5	<1	+	+	+	+	+	+	+
<i>E. coli</i> PEP141	OXA-48	SHV-12, CTX-M-9	TEM-1, CTX-M-27	0.5	4	<1	+	+	+	+	+	+	+
<i>E. cloacae</i> PEP143	OXA-48	SHV-12, CTX-M-9	TEM-1, CTX-M-27	4	>256	>256	+	+	+	+	+	+	+
<i>C. freundii</i> PEP157	OXA-48	TEM-1, OXA-9, OXA-18, CTX-M-9	TEM-1, CTX-M-27	1	4	>256	+	+	+	+	+	+	+
<i>K. oxytoca</i> PEP158	OXA-48	None	TEM-1, CTX-M-9	1	4	>256	+	+	+	+	+	+	+
<i>E. aerogenes</i> CNR20140467	OXA-48	SHV, CTX-M-9	TEM-1, CTX-M-9	2	16	>256	+	+	+	+	+	+	+
<i>E. aerogenes</i> CNR20140620	OXA-48	TEM	TEM-1, CTX-M-9	1	8	>256	+	+	+	+	+	+	+
<i>C. baileyi</i> CNR20140667	OXA-48	OXA-1	TEM-1, CTX-M-9	2	16	>256	+	+	+	+	+	+	+
<i>E. coli</i> CNR20140687	OXA-48	SHV	TEM-1, CTX-M-9	2	16	>256	+	+	+	+	+	+	+
<i>S. marcescens</i> CNR20130553	OXA-48	None	TEM-1, CTX-M-9	8	>256	>256	+	+	+	+	+	+	+
<i>E. coli</i> CNR20140388	OXA-162	CTX-M-1	TEM-1, CTX-M-9	<0.25	0.5	>256	+	+	+	+	+	+	+
<i>E. coli</i> CNR20140774	OXA-181	TEM, OXA-1, CTX-M-1	TEM-1, CTX-M-9	<0.25	2	>256	+	+	+	+	+	+	+
<i>E. coli</i> CNR20130516	OXA-204	TEM, CMY-2	TEM-1, CTX-M-9	<0.25	2	ND	+	+	+	+	+	+	+
<i>K. pneumoniae</i> CNR20130877	OXA-232	TEM, SHV, CTX-M-1	TEM-1, CTX-M-9	>32	ND	ND	+	+	+	+	+	+	+
<i>E. coli</i> CNR20130258	OXA-244	None	TEM-1, CTX-M-9	0.5	2	64	+	+	+	+	+	+	+
<i>K. pneumoniae</i> CNR20140689	KPC-2	SHV, OXA-1, CTX-M-1	TEM-1, CTX-M-9	>32	>32	>256	+	+	+	+	+	+	+
<i>K. pneumoniae</i> NDM-1 + OXA-232	SHV-28, OXA-1	SHV-28, OXA-1	TEM-1, CTX-M-9	>32	>32	ND	+	+	+	+	+	+	+
<i>C. freundii</i> CNR20120598	VIM-4 + OXA-48	TEM, OXA-1, CTX-M-1	TEM-1, CTX-M-9	8	16	>256	+	+	+	+	+	+	+
<i>K. pneumoniae</i> COL20140006	IMP-4	TEM, SHV	TEM-1, CTX-M-9	4	4	64	+	+	+	+	+	+	+
<i>K. pneumoniae</i> PEP075	KPC-2	TEM-1, SHV-11, SHV-12, OXA-9, OXA-1	TEM-1, CTX-M-9	8	32	64	+	+	+	+	+	+	+
<i>K. pneumoniae</i> PEP134	NDM-1	TEM-1, SHV-12, OXA-1, OXA-9, CTX-M-15	TEM-1, CTX-M-9	16	>32	256	+	+	+	+	+	+	+
<i>K. pneumoniae</i> PEP070	VIM-1	TEM-1, SHV-28, OXA-1, CTX-M-15	TEM-1, CTX-M-9	4	2	>256	+	+	+	+	+	+	+
<i>K. pneumoniae</i> PEP108	None	TEM-1, SHV-28, OXA-1, CTX-M-15	TEM-1, CTX-M-9	<0.25	<0.25	8	+	+	+	+	+	+	+

Table 1. Detection of OXA-48-like carbapenemase producers from strains grown on 18 media using the ICA OXA-48 K-Set (+, positive result; -, negative result; ND, not done; NG, no growth; Meropenem (MEM), Ertapenem (ERT), Temocillin (TEM)

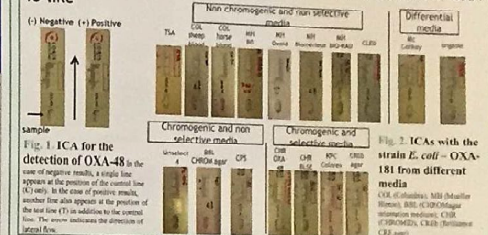
D) KPC K-Set: 10 *Enterobacteriaceae* and 1 *P. aeruginosa* on 11 different culture media

Species	Carbapenemase	β-lactamase genes		MICs (ng/ml)		Non chromogenic - non selective media		Differential media		Chromogenic - non selective media		Chromogenic - selective media	
		β-lactamase-PhoP/K	β-lactamase-PhoP/K	MEM	ERT	TEM	TSA	Colombia sheep blood agar	Uriselect 4	McConkey	Drigalski	Uriselect 4	CHROMID BLSE
<i>K. pneumoniae</i> CNR2012580	KPC-2	TEM, SHV, SHV-4-like	TEM-1, CTX-M-27	258	4	16	ND	+	+	+	+	+	+
<i>K. pneumoniae</i> CNR2012497	KPC-3	TEM, SHV	TEM-1, CTX-M-27	512	>32	>32	ND	+	+	+	+	+	+
<i>C. baileyi</i> CNR20140729	KPC-3	SHV	TEM-1, CTX-M-27	2	8	16	+	+	+	+	+	+	+
<i>E. coli</i> CNR20140325	KPC-2	SHV	TEM-1, CTX-M-27	2	2	64	+	+	+	+	+	+	+
<i>P. aeruginosa</i> PVO KPC-2	KPC-2	None	TEM-1, CTX-M-27	ND	ND	ND	+	+	+	+	+	+	+
<i>K. pneumoniae</i> PEP241	KPC-3 + OXA-48	TEM, SHV, OXA-9	TEM-1, CTX-M-27	512	>32	>32	>256	+	+	+	+	+	+
<i>E. coli</i> PEP141	OXA-48	TEM, SHV	TEM-1, CTX-M-27	4	4	64	+	+	+	+	+	+	+
<i>K. pneumoniae</i> COL20140006	IMP-4	TEM-1, OXA-10, CMY-16	TEM-1, CTX-M-27	8	4	128	+	+	+	+	+	+	+
<i>E. coli</i> PEP135	NDM-1	TEM-1, SHV-5, OXA-10	TEM-1, CTX-M-27	4	2	>256	+	+	+	+	+	+	+
<i>K. pneumoniae</i> PEP070	VIM-1	TEM-1, SHV-28, OXA-1, CTX-M-15	TEM-1, CTX-M-27	<0.25	<0.25	8	+	+	+	+	+	+	+
<i>K. pneumoniae</i> PEP108	No Carbapenemase	TEM-1, SHV-28, OXA-1, CTX-M-15	TEM-1, CTX-M-27	<0.25	<0.25	8	+	+	+	+	+	+	+

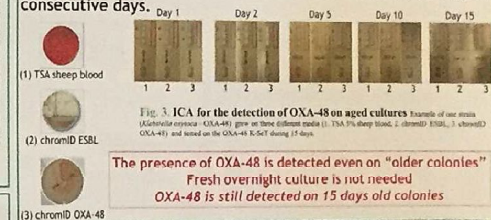
Table 2. Detection of KPC-like carbapenemase producers from strains grown on 11 media using the ICA KPC K-Set (+, positive result; -, negative result; ND, not done; Meropenem (MEM), Ertapenem (ERT), Temocillin (TEM)

All culture media tested (including Drigalski and McConkey) are compatible with the use of OXA-48 and KPC-K-Set (specificity and sensitivity of 100%)

B) Immunochromatographic assay (ICA) for the detection of OXA-48-like



C) Aged cultures : 1 strain OXA-48 and 1 strain OXA-181 were grown 3 different media and tested with OXA-48 K-Set during 15 consecutive days.



Conclusions

- OXA-48 and KPC K-Set are very robust assays:
- Various isolation media are compatible with these tests including Drigalski and McConkey
- Colonies do not need to be freshly cultured
- Simple and easy to implement as first line testing for rapid confirmation OXA-48 and KPC.

References

- Glupczynski Y and al. Evaluation of two new commercial immunochromatographic assays for the rapid detection of OXA-48 and KPC carbapenemases from cultured bacteria. J Antimicrob Chemother. 2016 Jun 28;doi:10.1093/ac/ckw147.
- Dozen M and al. Prospective evaluation of the OXA-48 K-Set assay, an immunochromatographic test for the rapid detection of OXA-48 type carbapenemase. J Antimicrob Chemother. 2016 Jun 10;doi:10.1093/ac/ckw147.
- Wermans D and al. Evaluation of an immunochromatographic lateral flow assay (OXA-48 K-Set) for rapid detection of OXA-48-like carbapenemase in Enterobacteriaceae. J Clin Microbiol. 2016 Feb;doi:10.1128/JCM.01131-15.

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