

A new method for detection of carbapenemase-producing Enterobacteriaceae carriage

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Poster **#P2333**

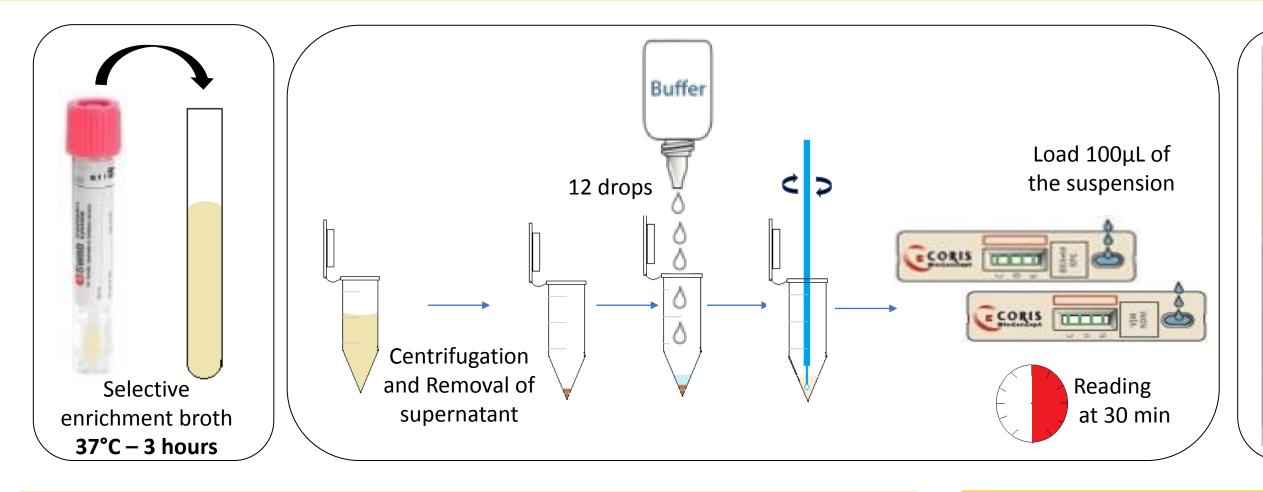
Introduction and purpose

Rapid detection of patients colonized with Carbapenemase-Producing Enterobacteriaceae (CPE) is fundamental for the control of their diffusion and limitation of the use of carbapenems. According to the latest epidemiological report, the main carbapenemases observed in France are OXA-48 and its variants followed by NDM and rarely VIM and KPC.

We have improved a method allowing early detection and identification of principal carbapenemase types directly on rectal eSwabs™, using O.K.N.V. K-SeT® (Coris Bioconcept).

Methods

- <u>Development step</u>: eSwabs[™] were spiked with CPE-negative stools and different concentration (10³, 10⁴, 10⁵, 10⁶ and 10⁷ CFU/mL) of 47 CPE strains (19 OXA-48, 10 NDM, 14 KPC, 3 VIM and 1 OXA-48 and NDM strain).
- Patient eSwabs™: Between 2017 and 2018, we have collected 94 rectal eSwabs™ (stored at -20°C) from patients in 3 hospitals. Among them, 21 were found positive for carbapenemase production by standard culture and confirmed by molecular typing (including 19 CPE, 1 NDM-producing *A. baumannii* strain and 1 VIM-producing *P. aeruginosa* strain).
- → After selective broth enrichment at 37°C for 3 hours, the protein extraction method of Coris Bioconcept is applied to the centrifugation pellet. Extract is tested with O.K.N.V K-SeT® kit according to the manufacturer's instructions. However, test result is read at 30 minutes.
- → Quantification of the CPE before and after selective enrichment is carried out on agar plates ChromID® CARBA SMART (bio-Mérieux).



Conclusion

Our new method of selective enrichment during 3 hours allow direct identification of low CPE carriage from rectal eSwab™ using O.K.N.V. K-SeT®.

Results

- <u>Development step</u>: Average yield of preanalytical enrichment of CPE is around 200 times. The detection limit (DL) is $\approx 10^4$ CFU/mL whatever the carbapenemase types.
- Patient eSwabs™: After selective enrichment, sensitivity and specificity were 100% and 100%, respectively.
- Of note, 1 NDM-producing *A. baumannii* strain and 1 VIM-producing *P. aeruginosa* strain were detected.

1 1/10				
1 VIM		Positive eSwabs™	Negative eSwabs™	n
5 NDM	Positive O.K.N.V.	21	0	21
15 OXA- 48	Negative O.K.N.V.	0	73	73
	Total	21	73	94
	Sensitivity (%)	100		
OXA-48 NDM	Specificity (%)		100	
■ KPC ■ VIM				

References

- *Dortet L. *et al.* Note technique : Détection des souches d'entérobactéries productrices d'une carbapénèmase. CNR associé de la résistance aux antibiotiques, « Entérobactéries productrices de carbapénèmases »
- *Wareham DW and Momin MHFA. Rapid Detection of Carbapenemases in Enterobacteriaceae: Evaluation of the RESIST-3 O.K.N (OXA-48, KPC, NDM) Multiplexed Lateral Flow Assay. J. Clin. Microbiol. Accepted manuscript posted online 1 February 2017
- *Bogaerts P. *et al.* Evaluation of a new lateral flow assay for the detection of VIM-producing bacteria. Poster 37th RICAI 2017