

Evaluation of a new chromogenic medium, chromID<sup>TM</sup> CARBA, for the detection of carbapenemase-producing *Enterobacteriaceae* N. Bereksi<sup>1</sup>, D. Giraud<sup>1</sup>, F. Joyeux<sup>1</sup>, L. Barbaux<sup>2</sup>, S. Ghirardi<sup>2</sup>, S. Orenga<sup>2</sup>, F. Villeval<sup>1</sup>, G. Zambardi<sup>2</sup> <sup>1</sup>bioMérieux 69290 Craponne, France. <sup>2</sup> bioMérieux, La Balme les Grottes, 38390 France

#### **INTRODUCTION AND PURPOSE**

Carbapenemase Producing *Enterobacteriaceae* (CPE) are multi-resistant emerging bacteria which can be responsible for hospital acquired infections and outbreaks. Detection of CPE carriers is of particular importance for prevention and epidemiological monitoring of these infections. In this context, chromogenic media for CPE should make screening easier due to their selectivity and the use of different colours to discriminate targeted species. The aim of this study was to evaluate the performance of two chromogenic media for CPE detection, namely chromID<sup>TM</sup> CARBA (a prototype medium from bioMérieux based on the same principles than the ones – ID Carba – previously tested by Perry *et al.*, 2011) and CHROMagar™ KPC (CHROMagar). A commonly used selective home-brewed medium, MacConkey supplemented with 1 mg/L imipenem (McC+I), was also included in the study.

### **METHODS**

A total of 194 isolates was tested, including 127 CPE with different types of carbapenemases (53 KPC, 44 NDM, 13 VIM, 11 IMP, 6 OXA-48) and 67 isolates not producing a carbapenemase. All microorganisms were inoculated directly onto each medium with 10  $\mu$ L of a 0.5 McFarland calibrated suspension before 24 h includation at 34-38°C. Two batches of chromID<sup>TM</sup> CARBA were used: one freshly prepared (B1) and one close to the expiry date (B2).

## RESULTS

Sensitivity and specificity results for CPE detection are shown in the table below. Typical colonies of *Klebsiella pneumoniae* (coloured in green) and *Escherichia coli* (coloured in pink) are presented for each medium tested. For chromID<sup>TM</sup> CARBA, the sensitivity for CPE detection varied from 89.8% (B1) to 96.1% (B2). By comparison, the sensitivity was 89.0% for CHROMagar<sup>TM</sup> KPC and only 68.5% for McC+I. Variation in sensitivity of detection by the two batches of chromID<sup>TM</sup> CARBA was similar and usually higher when compared to CHROMagar<sup>TM</sup> KPC and McC+I. However, detection of IMP and OXA-48 could be more difficult with a fresh medium than with a medium close to the expiry date. Nonetheless, the specificity of chromID<sup>TM</sup> CARBA (B2) remained stable over time.

### chromID CARBA



chromID CARBA



Туре	Total	Microorganism (nb of strain)	chromID Carba (Bl)	chromID Carba (B2)	CHROMagar KP C	Mac Conkey (McC+l)
Class A KPC	53	<i>E. coli</i> (6)	96.2%	100%	96.2%	56.6%
		Klebsiella (43)				
		Enterobacter (4)				
Class B NDM	44	<i>E. coli</i> (31)	97.7%	97.7%	90.9%	77.8%
		Klebsiella (5)				
		Enterobacter (7)				
		Citrobacter*(1)				
Class B VIM	13	E. coli (5)	84.6%	92.3%	61.5%	69.2%
		Klebsiella (2)				
		Enterobacter (5)				
		Serratia (1)				
Class B IMP	11	E. coli (2)	45.5%	72.7%	81.8%	81.8%
		Klebsiella (1)				
		Enterobacter ** (7)				
		Serratia (1)				
Class D OXA-48	6	E. coli (1)	66.7%	100%	83.3%	83.3%
		Klebsiella (5)				
Sensitivity						
(True positive/ (True positive +			89.8%	<b>96.1%</b>	89%	68.5%
false ne	gative)	x 100				
Specificity						
(True negative / (True negative + 97% 94% 94% 96						96%
talse positive+) x 100						

\* A strain of *Citrobacter freundii* expressing both ß-galactosidase and ß-glucosidase produce violet colonies on **chromID™ CARBA** and a mixture of pink and blue colonies on **CHROMagar™ KPC**.

\*\* 3 of 7 strains of IMP producing *Enterobacter* that grow on **chromID™ CARBA** produced violet colonies associated to positivity of both ß-glucuronidase and ß-glucosidase activities.

# **CONCLUSION**

This study highlights the superior sensitivity of both chromogenic media over the **imipenem supplemented MacConkey**. In comparison to **CHROMagar™ KPC**, **chromID™ CARBA** presents three advantages:

- (i) trend to higher sensitivity,
- (ii) ready to use plates and
- (iii) extended shelf life.

As such, it has the potential of being a very useful tool for the screening of patients who carry the widespread KPCand NDM-producing *Enterobacteriaceae*. These results should be confirmed with clinical samples such as rectal swabs.