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**Evaluation of morphological and functional** characteristics of Carnobacterium maltaromaticum isolated from vacuum-packaged beef with long shelf life



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# INTRODUCTION

A temperature near the freezing point of meat ( $\sim -2$  C), associated with vacuum packaging allows the preservation of this product up to several months, which makes possible the meat trade across the planet without resorting to freezing.

Carnobacterium maltaromaticum is a lactic acid bacterium (LAB), and many LAB are known for their bactericidal or bacteriostatic activity against other strains, species or genera.

## OBJECTIVE

To perform a morphological and functional characterization Carnobacterium of *maltaromaticum* with a potential bioprotective

In this way, the presence of certain lactic acid bacteria adapted to a low temperature on fresh meat could extend the shelf life and improve the microbial stability and safety of this product.

effect isolated from vacuum-packaged beef with long shelf life.

### MATERIALS AND METHODS



Longissimus dorsi Australian origin commercial shelf life = 140 days

> **Isolation of** Carnobacterium maltaromaticum

Morphological, biochemical and microscopic profiles and comparison to two reference strains: macroscopic and microscopic observations, Gram staining, catalase and oxidase tests, API 50 CH and API ZYM galleries.

Influence of different atmospheres on growth of *C. maltaromaticum* (on a sterile meat model)



irradiated

(sterile) minced

pork meat



inoculation

C. maltaromaticum

10<sup>5</sup> CFU/mL (1% v/w)



1) 100 % N<sub>2</sub>



storage (7 d)

1) +4 °C

2) +8 °C

3) +12 °C

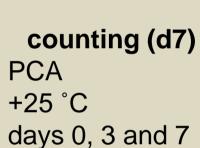












Microbiological stability of commercial beef inoculated with *C. maltaromaticum* 

Packaging (d0)

2) 70 % O<sub>2</sub> : 30 % CO<sub>2</sub>

3) 30 % O<sub>2</sub> : 70 % CO<sub>2</sub>







atmospheres









commercial	inoculation	vacuum-packaging	storage (7 d)	packaging (d7)	storage (7 d)	Counting (d14)
vacuum packed	C. maltaromaticum	(d0)	−1 °C	1) 100 % N <sub>2</sub>	+4 °C	total viable count (TVC)
psoas major	10 <sup>5</sup> CFU/mL (1% v/w)			2) 70 % $O_2$ : 30 % $CO_2$		lactic acid bacteria (LAB)
16 days after				,		Enterobacteriaceae (EB)
slaughter						Pseudomonas sp. (PS)
Slaughter						Brochothrix thermosphacta (BT)

# RESULTS Influence of different

9

log<sub>10</sub> CFU/g

Morphological, biochemical and enzymatic profiles

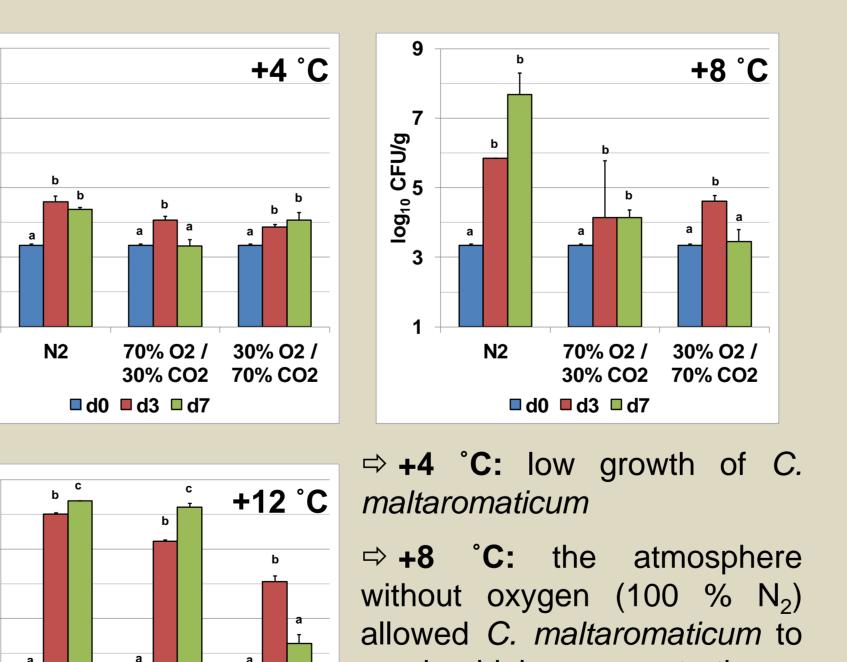
⇒ Similar profiles to two reference strains: LMG 11393 and LMG 22902

 $\Rightarrow$  **Colonies:** circular, convex, entire,  $\emptyset < 1$  mm, smooth, translucent, unpigmented and odorless

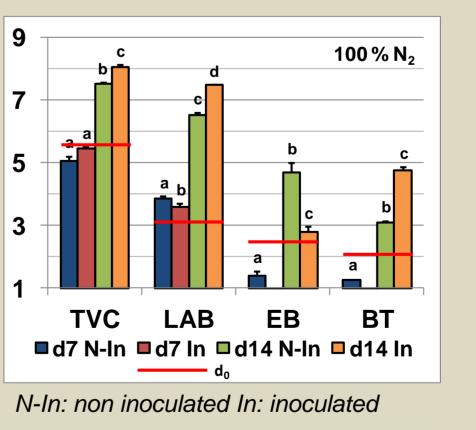
⇒ **Cells:** Gram positive bacillus arranged in pairs, catalase and oxidase negative

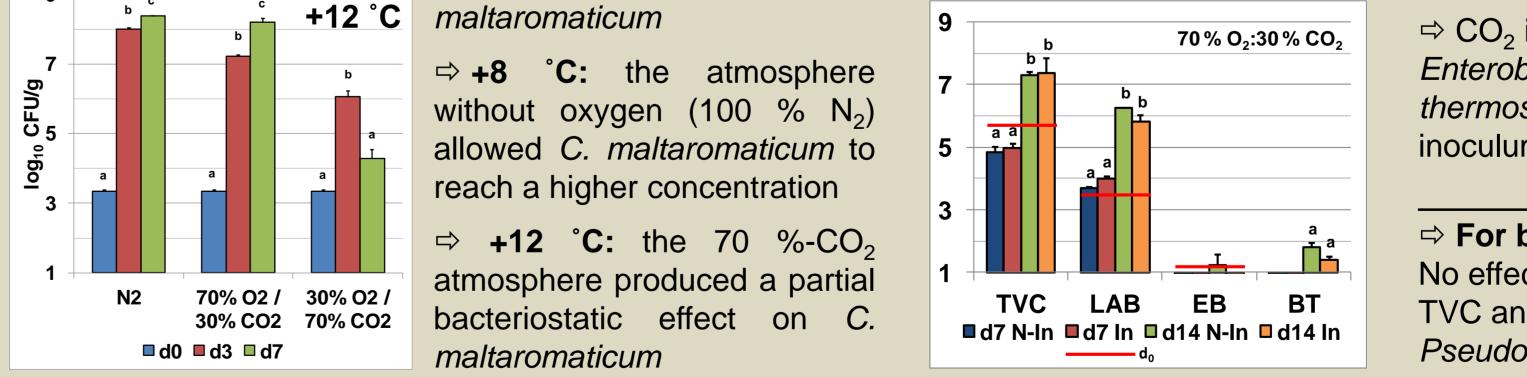
⇒ **Substrates:** glycerol, D-ribose, D-galactose, D-glucose, D-fructose, D-mannose, D-mannitol, methyl-α-Dmannopyranoside, methyl-α-D-glucopyranoside, Nacetylglucosamine, amygdalin, arbutin, esculin ferric citrate, salicin, D-cellobiose, D-maltose, D-lactose, Dmelibiose, D-saccharose, D-trehalose, gentiobiose, Dturanose and potassium gluconate

⇒ Enzymes: esterase (C4), esterase lipase (C8), valine arylamidase, acid phosphatase, naphthol-AS-BIphosphohydrolase and  $\beta$ -glucosidase



#### Microbiological stability of beef inoculated with C. maltaromaticum





 $\Rightarrow$  inoculum inhibited the growth of Enterobacteriaceae and favored the growth of B. thermosphacta

 $\Rightarrow$  CO<sub>2</sub> inhibited growth of Enterobacteriaceae and B. thermosphacta: no effect of inoculum

⇒ For both atmospheres: No effect of inoculum on TVC and LAB. Reduction of Pseudomonas sp. (data not shown)

## CONCLUSIONS

- ⇒ Morphological, biochemical and enzymatic profiles of the isolated strain similar to two reference strains
- $\Rightarrow$  Slower growth of *C. maltaromaticum* under 70 % O<sub>2</sub>:30 % CO<sub>2</sub> and 30% O<sub>2</sub>:70% CO<sub>2</sub>
- $\Rightarrow$  Antimicrobial effect of *C. maltaromaticum* against *Enterobacteriaceae* under N<sub>2</sub>
- ⇒ Perspectives: genotypic characterization of *C. maltaromaticum* and evaluation of its potential bioprotective effect

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