

Effect of aging time, aging technique (dry- vs. wet-aging) and packaging on tenderness, pigment and lipid stability of Belgian Blue beef

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To evaluate the potential effect of aging technique (dry-aging vs. wet-aging), aging time (0, 21, 42 and 63 days) and packaging during display (vacuum vs. shrinkable film wrapping) on pH, tenderness, and pigment and lipid stability of Belgian Blue beef.

INTRODUCTION

Post-mortem aging is a process that occurs naturally in all muscle tissues, which improves palatability attributes of meat such as flavor and tenderness.

Wet-aging

Meat aged in sealed barrier vacuum packages at refrigerated temperatures.



Dry-aging

Unpacked wholesale cuts aged at controlled temperature and relative humidity.

- ⇒ unique flavor and superior quality
- ⇒ destined to high-end consumers



MATERIALS AND METHODS

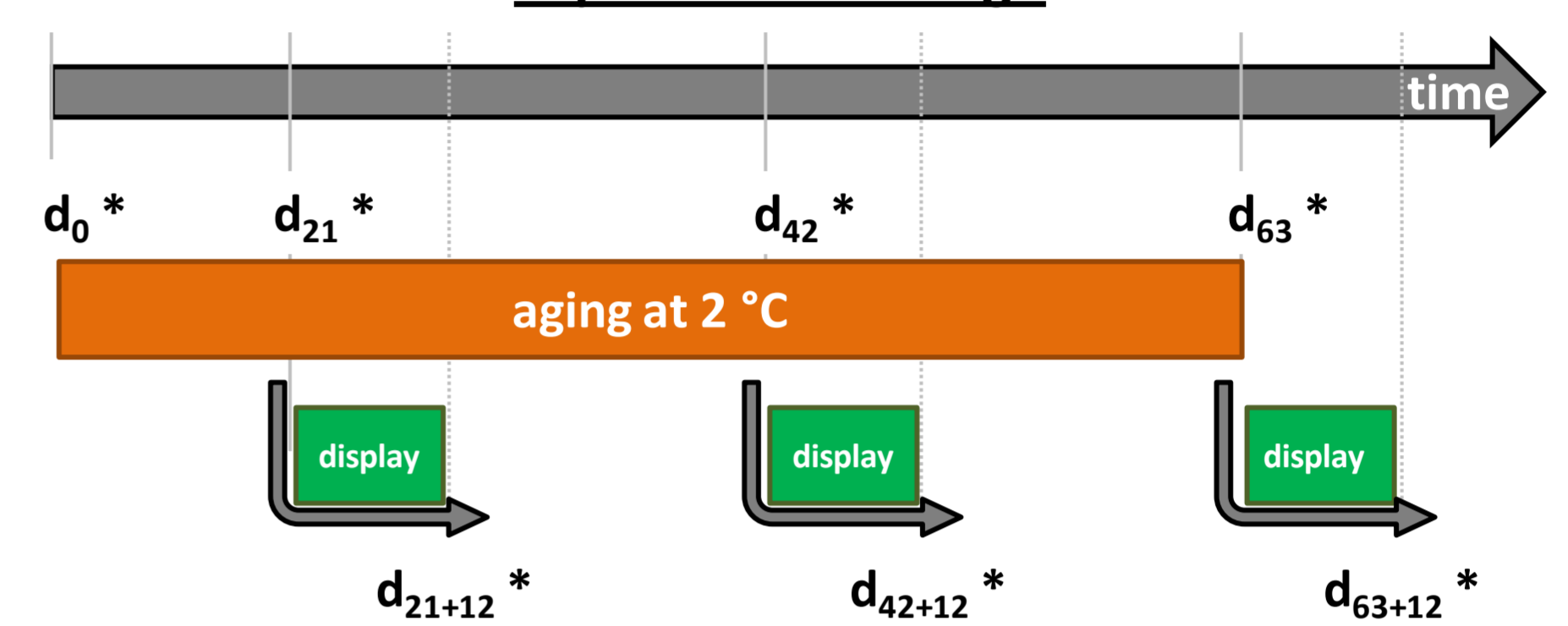
Samples

- 3 x ½ *longissimus dorsi* (wet aging)
- + 3 x ½ *longissimus dorsi* (dry aging)
- from 2 Belgian Blue cows

* Analysis

- pH
- Tenderness *only before display* (Warner-Bratzler shear force)
- Color (redness) (CIE L^ab*)
- Myoglobin oxidation (K/S₅₇₂:K/S₅₂₅ ratio)
- Lipid oxidation (TBARS: mg MDA-equivalent/kg)
- Statistics (ANOVA + Tukey test)

Experimental design



Aging conditions

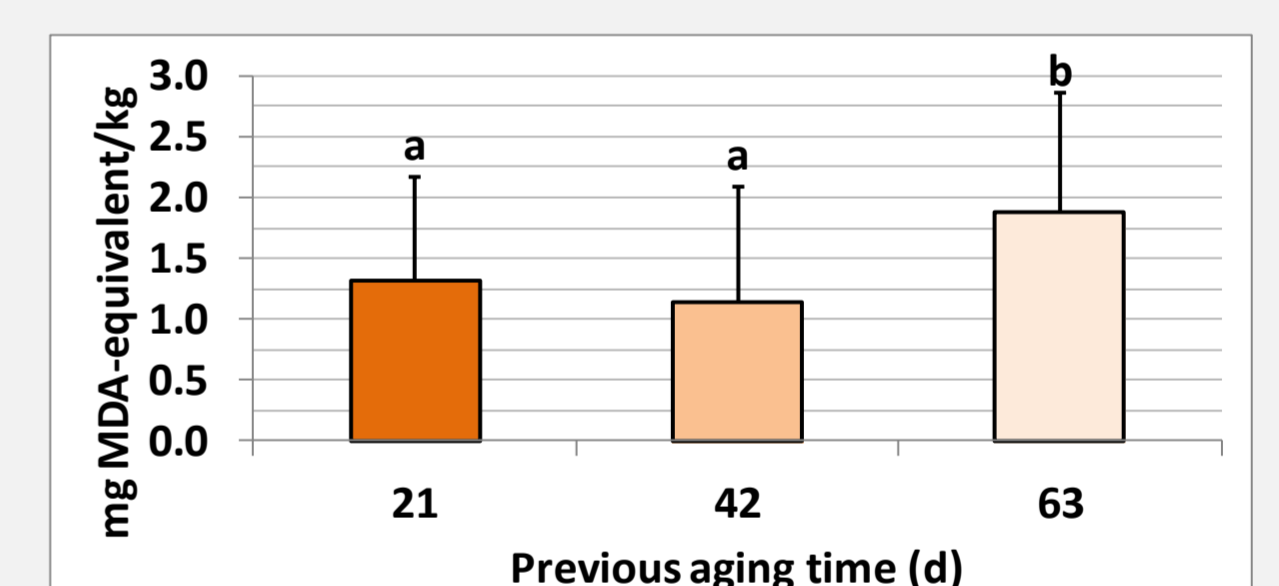
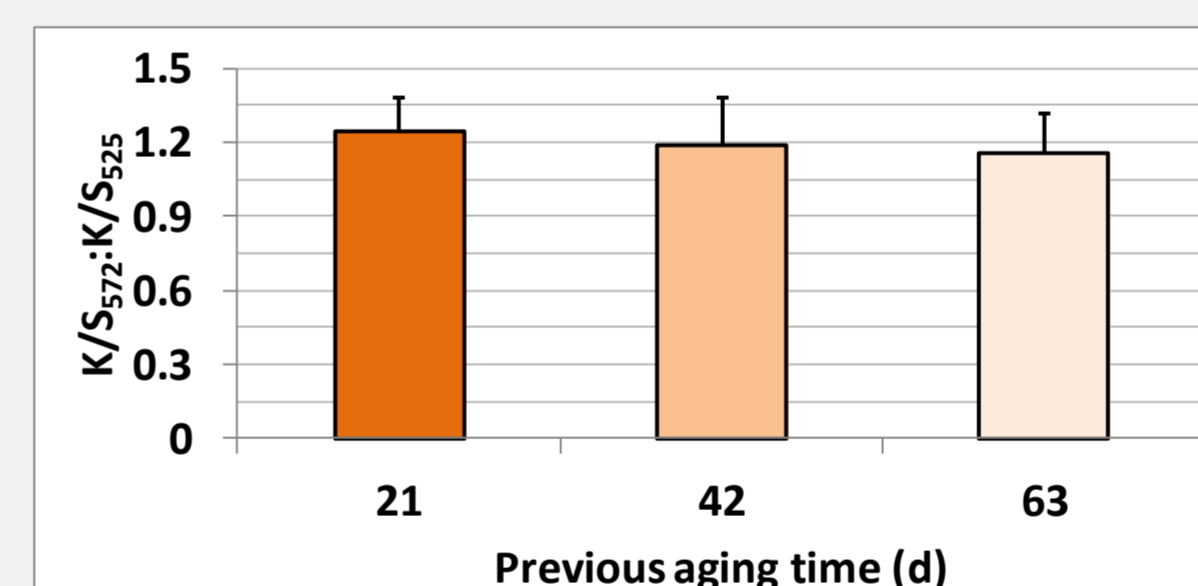
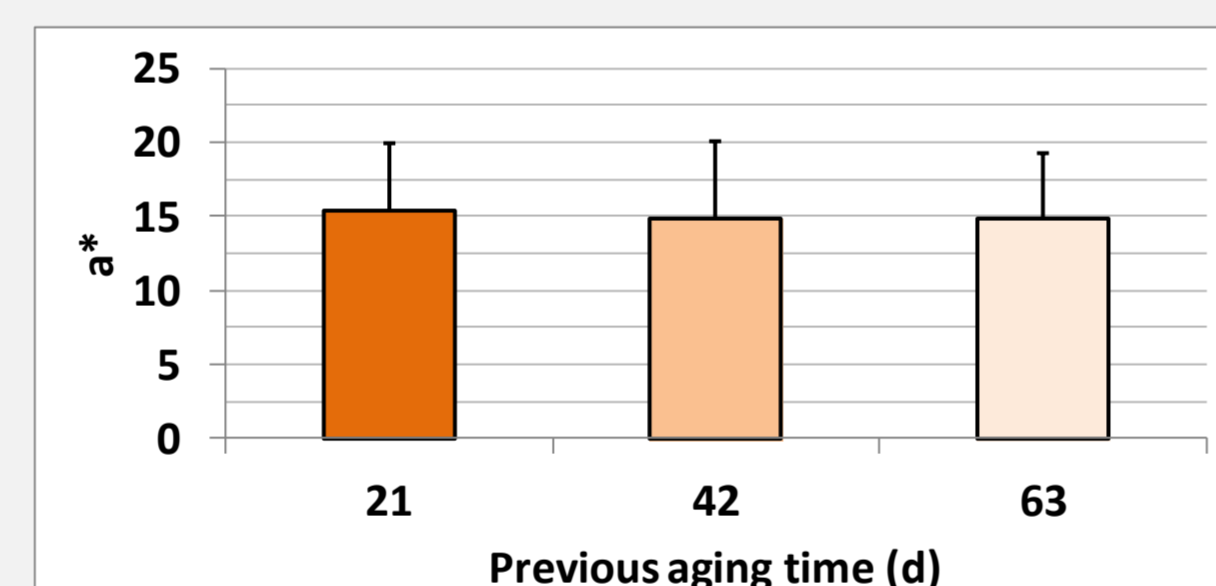
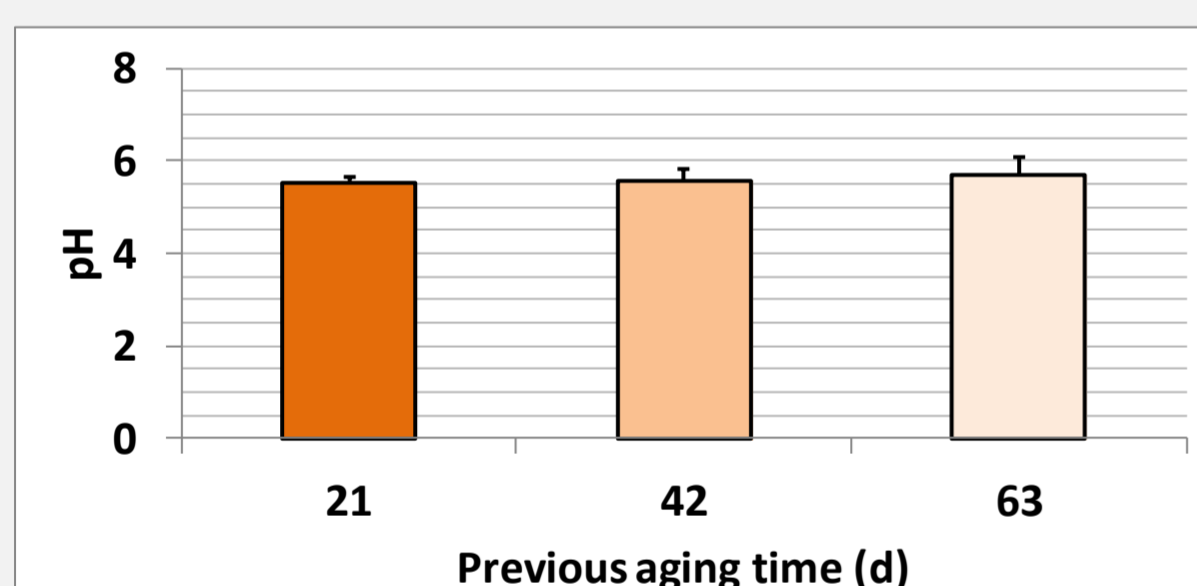
wet-aging (WA) or dry-aging (DA)

Display conditions

vacuum (VP) or shrinkable film wrapping (FW) (4 days at 4 °C + 8 days at 8 °C)

RESULTS AND DISCUSSION

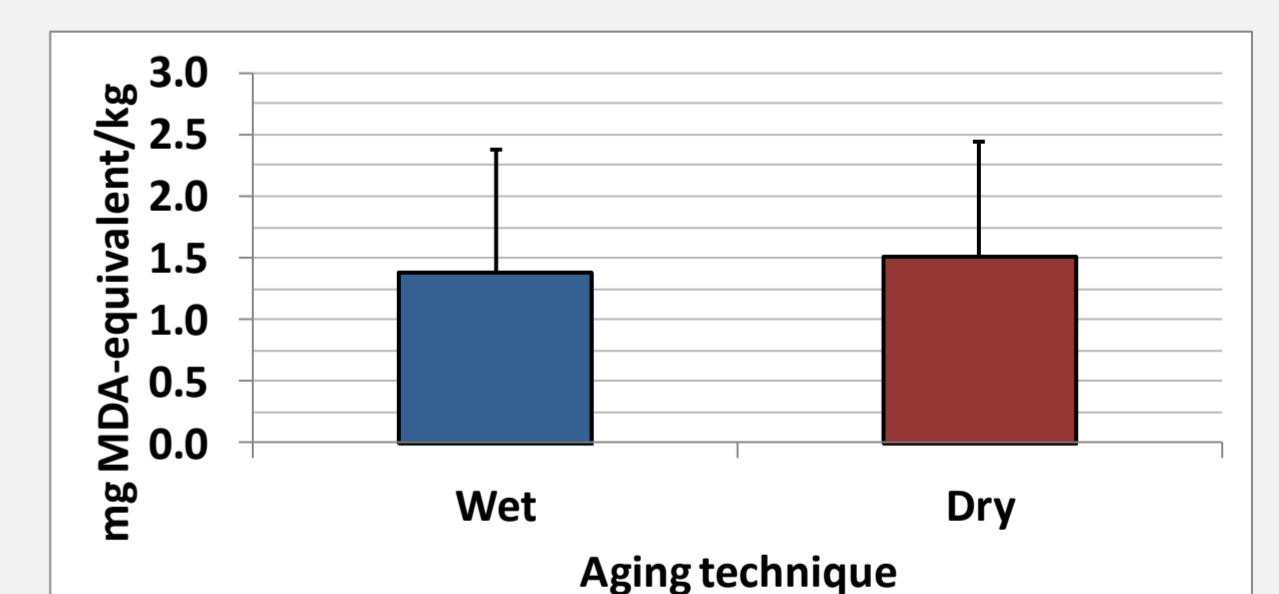
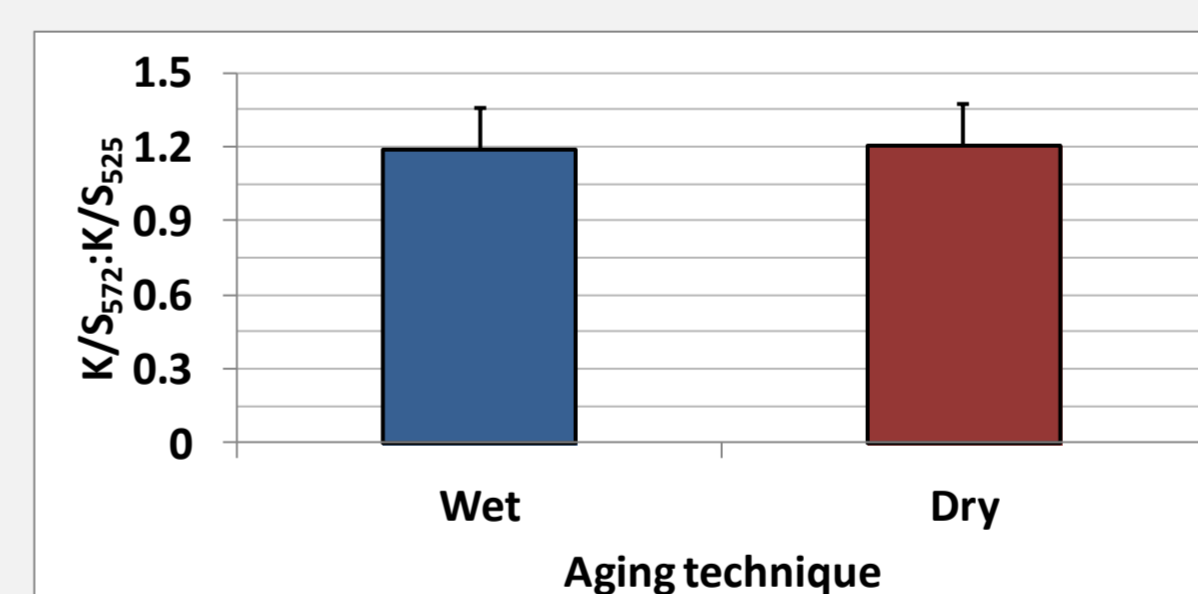
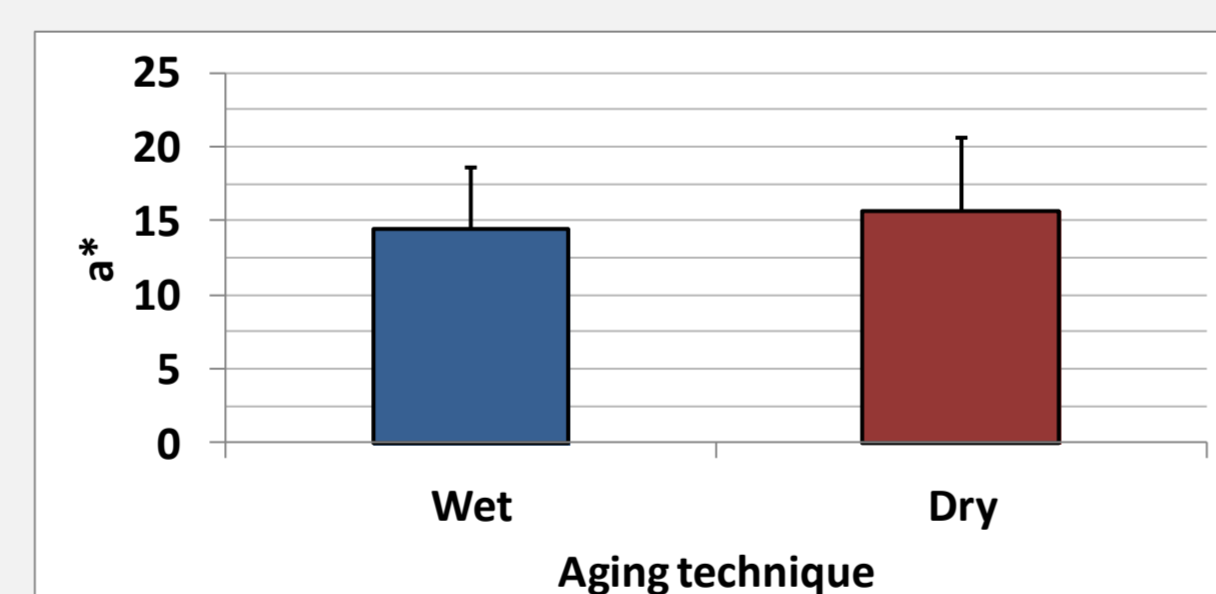
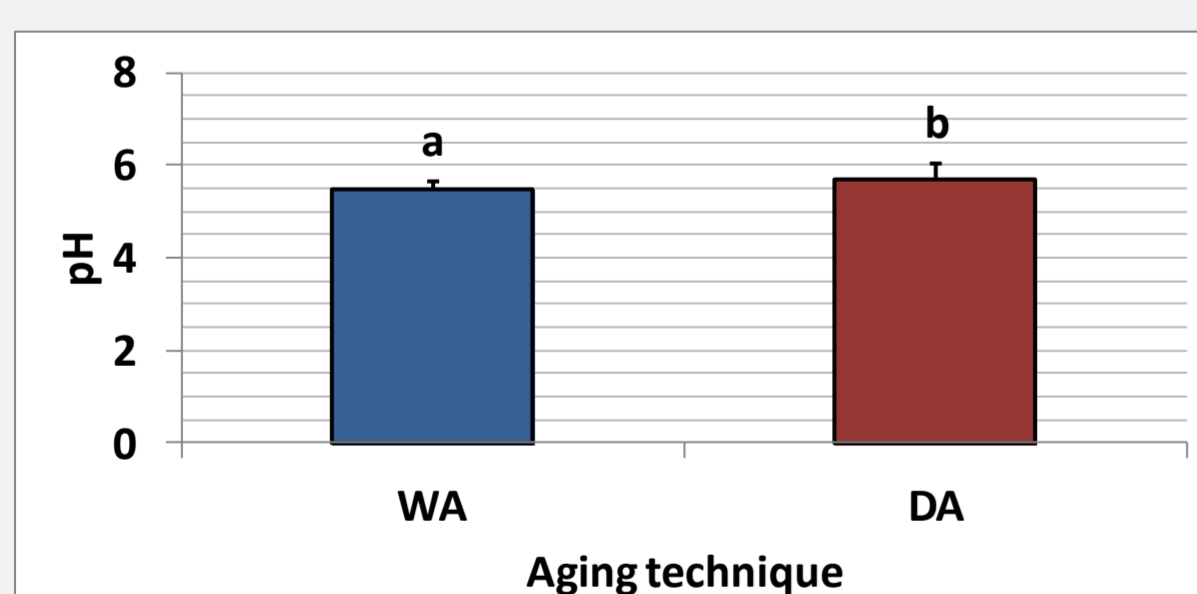
Effect of previous aging time



No effect of aging time on pH, redness (a*) and myoglobin oxidation (K/S₅₇₂:K/S₅₂₅ ratio) in samples after 21, 42 or 63 days of aging at 2 °C + 12 days of display (4 days at 4 °C + 8 days at 8 °C).

Previous aging time favored lipid oxidation (increase in mg MDA-equivalent/kg) during display.

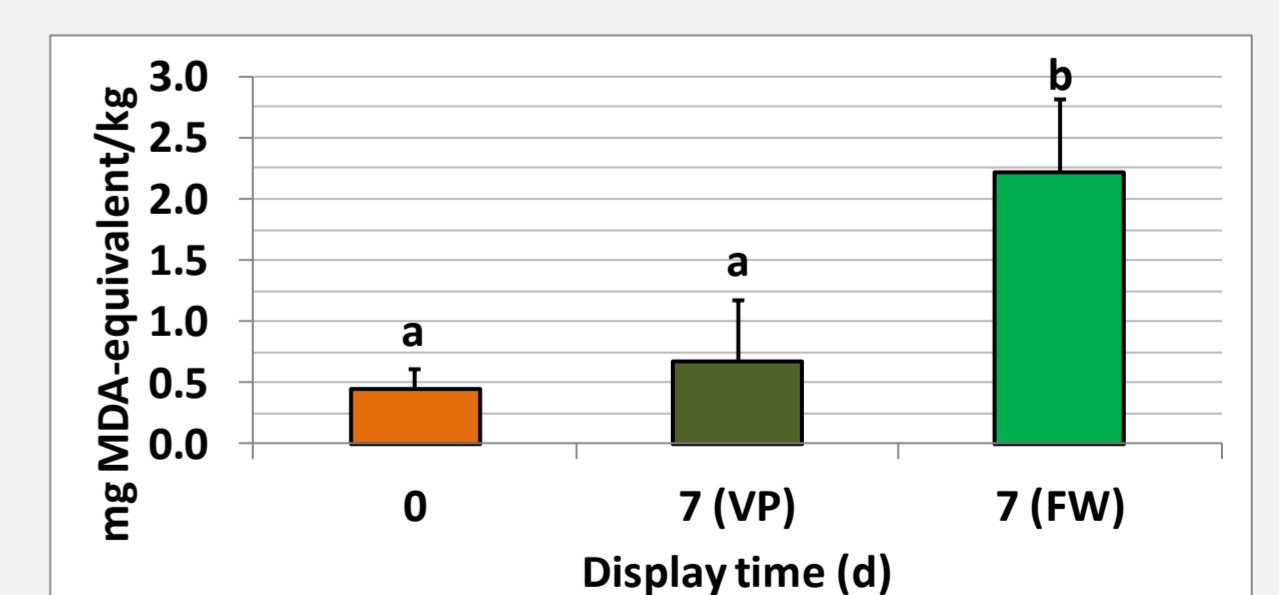
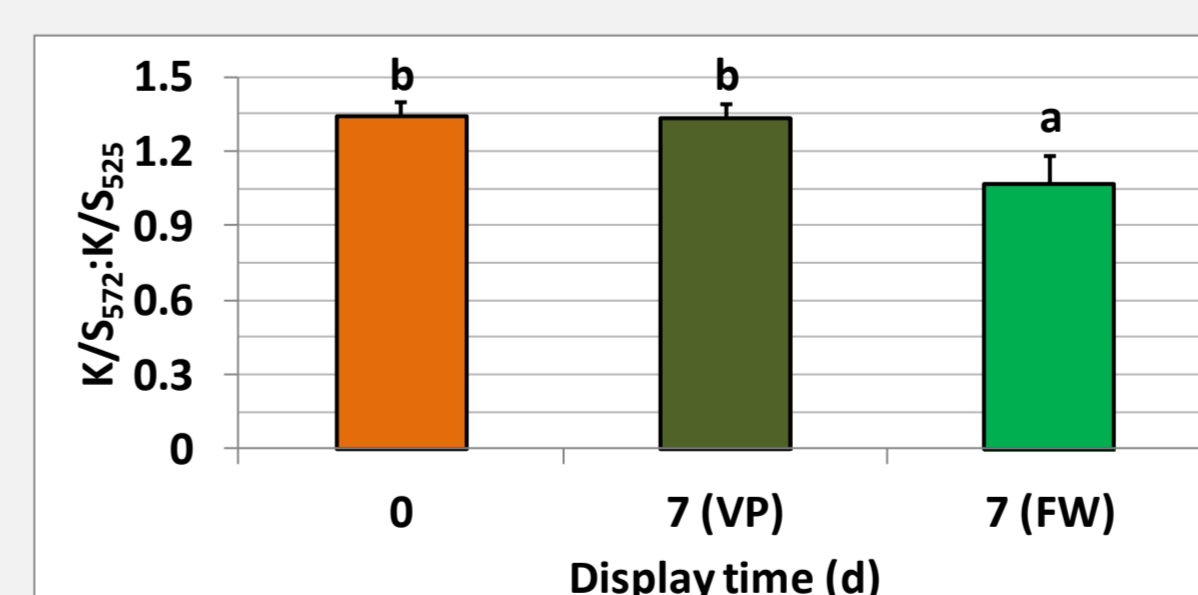
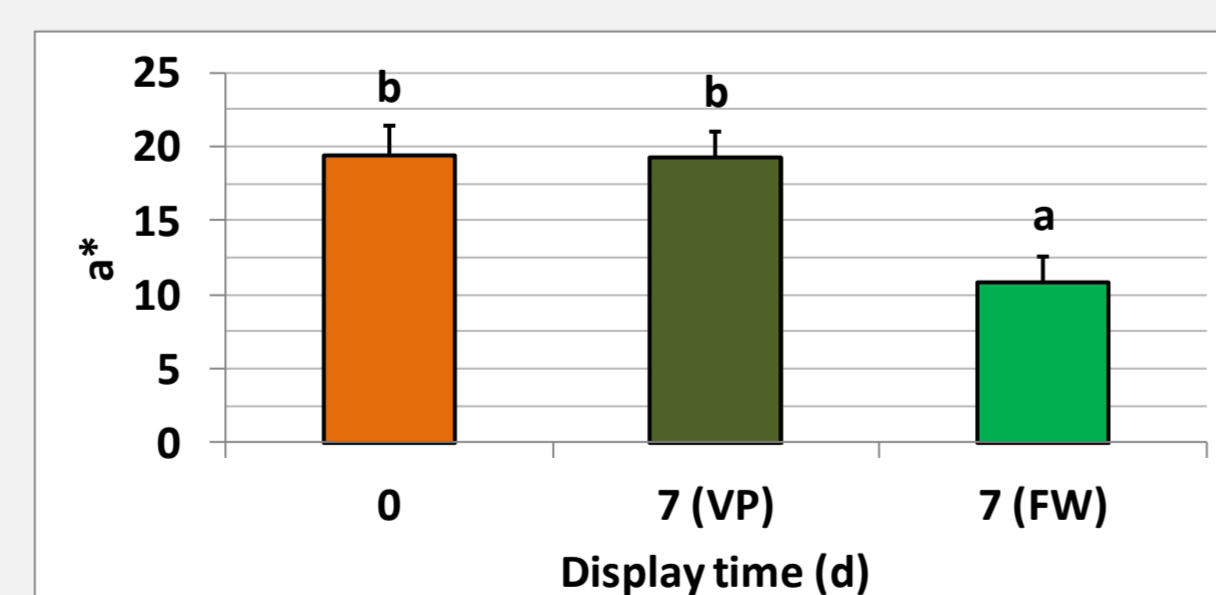
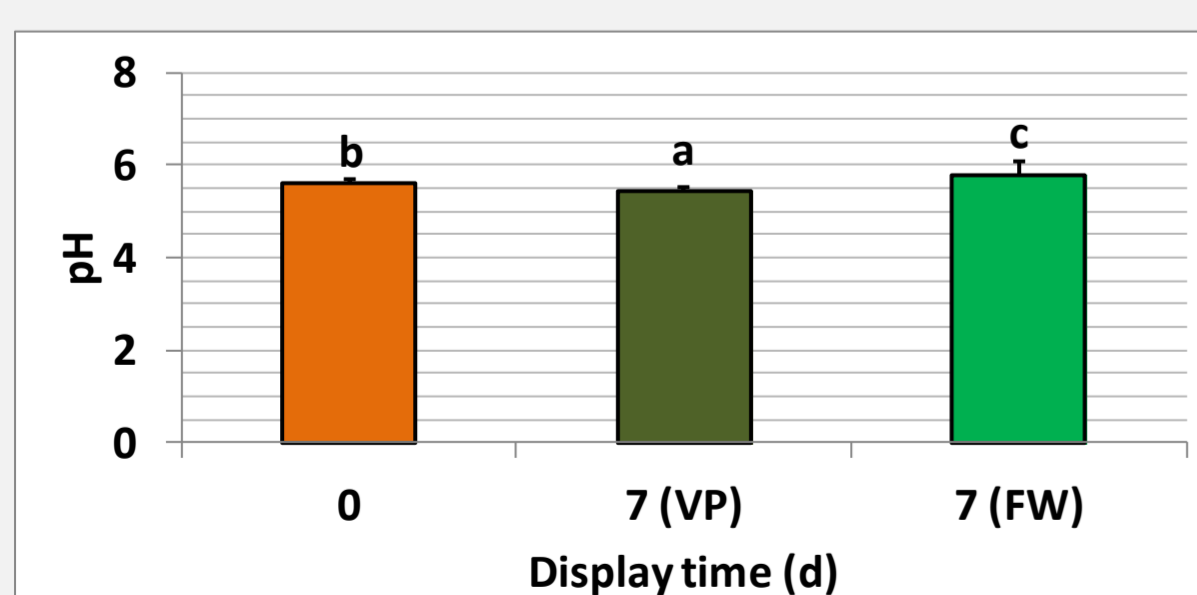
Effect of aging technique



The growth of proteolytic bacteria (e.g. *Pseudomonas*) in aerobic conditions and of lactic acid bacteria in low-oxygen conditions may have contributed to the difference in pH.

No effect of aging time on redness (a*), myoglobin oxidation (K/S₅₇₂:K/S₅₂₅ ratio) and lipid oxidation (mg MDA-equivalent/kg) in samples after 21, 42 or 63 days of aging at 2 °C + 12 days of display (4 days at 4 °C + 8 days at 8 °C).

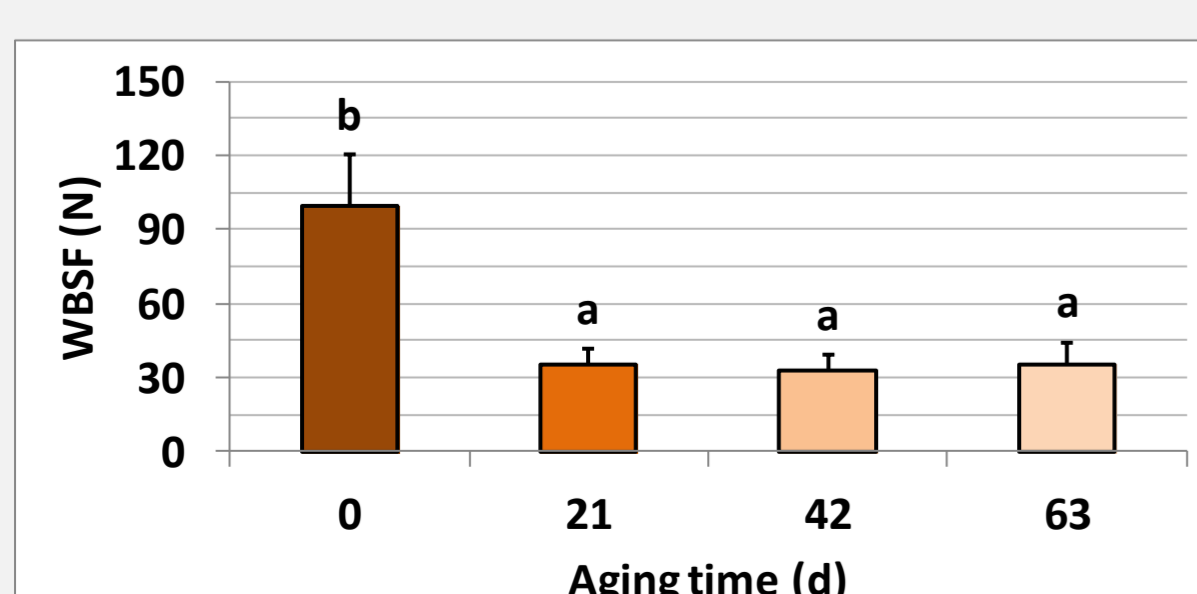
Effect of retail packaging



The growth of proteolytic bacteria (e.g. *Pseudomonas*) in aerobic conditions and of lactic acid bacteria in low-oxygen conditions may have contributed to the differences in pH.

Samples before display (orange bars) and samples after a seven-day display under vacuum conditions (dark green bars) presented no statistically significant differences in redness (a*), myoglobin oxidation (K/S₅₇₂:K/S₅₂₅ ratio) and lipid oxidation (mg MDA-equivalent/kg). The contact with atmospheric air in FW samples (display time = 7 days) may have contributed to meat discoloration (decrease in chromaticity a*), myoglobin oxidation (decrease in K/S₅₇₂:K/S₅₂₅ ratio) and lipid oxidation (increase in mg MDA-equivalent/kg).

Effect of aging on tenderness



Increase in tenderness (decrease in WBSF values) during the first 21 days of aging.

CONCLUSIONS

A higher sensitivity to oxidation was observed with shrink wrap packaging during display. Twenty-one days of aging allowed to achieve the maximum tenderness observed and would be compatible with a subsequent 12-day display under vacuum. Further research will be conducted to study the microbiological quality of these meats as well as their antioxidant capacity.