











# Characterisation of adipose tissue in immunocastrated pigs

Poklukar, K.<sup>1</sup>, Čandek-Potokar, M.<sup>1,2</sup>, Batorek Lukač, N.<sup>1</sup>, Vrecl Fazarinc, M.<sup>3</sup>, Fazarinc, G.<sup>3</sup>, Kress, K.<sup>4</sup>, Weiler, U.<sup>4</sup>, Stefanski, V.<sup>4</sup>, Škrlep, M.<sup>1</sup>

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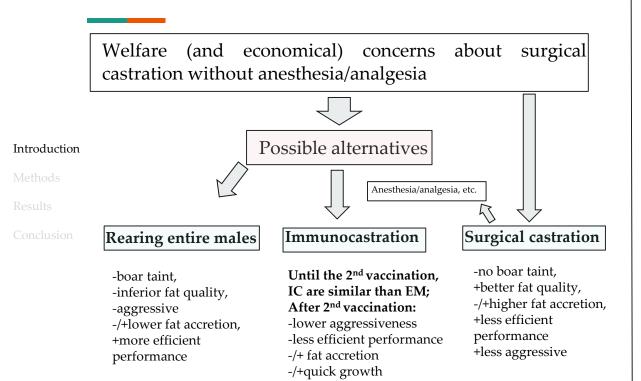
<sup>&</sup>lt;sup>1</sup> KIS, Hacquetova 17, 1000 Ljubljana

<sup>&</sup>lt;sup>2</sup> UM FKBV, Pivola 10, 2311 Hoče, Slovenia

<sup>&</sup>lt;sup>3</sup> UL VF, Gerbičeva 60, 1000 Ljubljana, Slovenia

<sup>&</sup>lt;sup>4</sup> UHOH, Garbenstr. 17, 70599 Stuttgart, Germany

# Introduction



# Objective of the study

The objective of the study was to evaluate the **effect of immunocastration** on **fat accretion** from morphological, histomorphological and biochemical perspective by comparing **immunocastrated pigs**(IC) with **entire male** (EM) and **surgically castrated** (SC) pigs.

#### Animals:

- 36 pigs, Landrace x Pietrain crossbreed
- 12 EM, 12 IC, 12 SC
  - Vaccination with Improvac® at the age of 12 and 21 weeks
- Slaughtered at the age of 26 weeks, and live weight  $121.7 \pm 1.6$  kg.

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#### Measurements:

- Backfat thickness at the level of the last rib (mm), on withers (mm) and above *Gluteus medius* muscle (mm)
- Belly lean meat (%)
- Ratio between fat and meat area



Sampling of the inner and outer back fat layer on withers. Determination of:

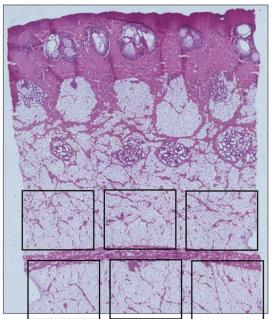
## • **Histomorphological** analysis:

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- → Backfat on withers was dissected, fixed, embedded in paraffin and sectioned.
- → Sections were stained with hematoxylin and eosin.
- → Digital images were acquired with light microscope.
- → Image J program was used for determination of:
  - dermis thickness (μm),
  - number of adipocytes per fascicle,
  - fascicle surface area (μm²),
  - adipocyte surface area (μm²).





Six randomly chosen fields (3 in inner and 3 in outer layer) exhibiting the best membrane integrity were used for cellularity analysis on each slide.

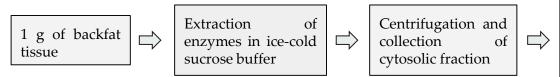
Sampling of the inner and outer back fat layer on withers. Determination of:

• **Fatty acid composition** of inner back fat layer using gas chromatography.

Fatty acids were extracted and trans-methylated as described in Park and Goins, 1994; fatty acid methyl esters were identified with Hewlett Packard 6890 FID System. Results are expressed as percentage of total FA.

• **Lipogenic enzyme activities** (in nmol/min per g of tissue) determined in aliquots of inner backfat layer:

- fatty acid synthase activity (FAS),
- o malic enzyme activity (ME),
- o glucose-6-phosphate dehydrogenase activity (G6PDH),
- o citrate synthase activity (CS).



The data were analysed by one-way **ANOVA** using R software.

After extraction of cytosolic fraction, activities of ME, G6PDH, FAS and CS were assayed (as described in Bazin and Ferre, 2001) by **absorbance at 340 nm** to assess the appearance (for ME and G6PDH) or the decrease (for FAS, CS) of NADPH.

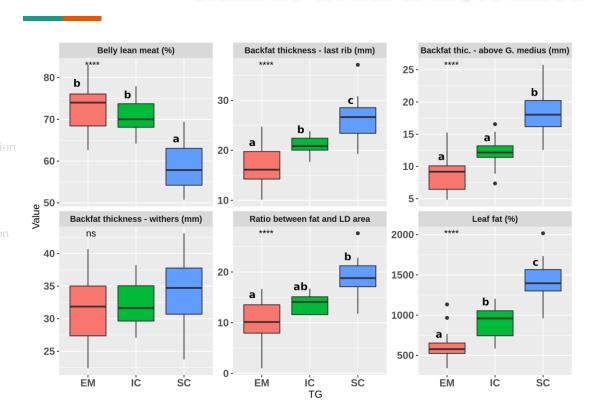
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# CARCASS TRAITS of EM, IC and SC

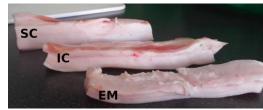


### IC compared to EM exhibited:

- → Î values for backfat thicknesses at the level of the last rib,
- → 1 leaf fat percentage.

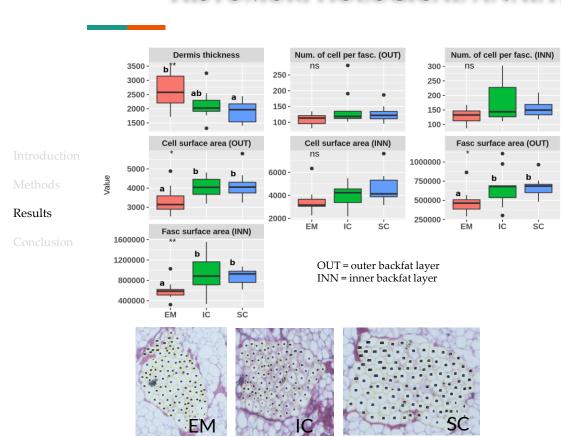
#### IC compared to SC exhibited:

- → Î belly lean meat percentage,
- → packfat thicknesses at the level of the last rib and above *Gluteus medius*,
- → **1** leaf fat percentage.



# Results

### HISTOMORPHOLOGICAL ANALYSIS of BACKFAT



## IC compared to EM:

- → Increased adipocyte surface area in outer backfat layer.
- Increased fascicle surface area in outer and inner backfat layer.
- → Tendency (p=0.0501) of increased **adipocyte proliferation** in inner backfat layer.

### EM compared to SC:

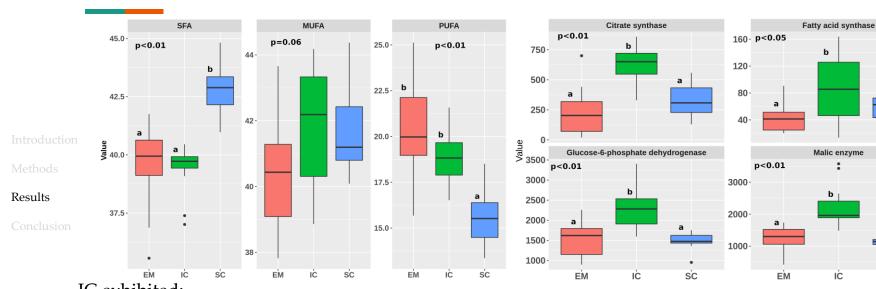
- → tendency (p=0.0537) of decreased adipocyte surface area in inner backfat layer.
- → exhibited thicker dermis.



Probably higher collagen content in EM

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## FATTY ACID COMPOSITION and LIPOGENIC ENZYME ACTIVITIES



### IC exhibited:

- ↓ SFA content (especially palmitic and stearic acid) and ÎPUFA content (especially linoleic acid) compared to SC.
- Immunocastration caused varied response of animals in MUFA deposition!

#### IC have:

1.4 to 2.7-fold ↑ lipogenic enzyme activities than EM and SC.

b

ΙĊ

SC

<sup>\*</sup>Enzyme activities are in nmol/min per g of tissue

### Compared to EM, immunocastration caused:

#### On carcass level:

→ enlargement of adipose tissue (leaf fat percentage, backfat thicknesses at the level of the last rib).

#### On the cellular level:

- → expansion of adipocytes in outer back fat layer, as well as expansion of fascicle surface area in outer and inner back fat layer,
- → tendency for adipocyte proliferation in inner back fat layer,
- → outer backfat layer compared to inner seems to be more responsive to immunocastration.

#### On biochemical level:

- $\Rightarrow$  lipogenic enzyme activity was  $\hat{1} \Longrightarrow$  showing the change in lipid metabolism ( $\hat{1}$  fatty acid synthesis),
- → SFA and PUFA content in backfat of IC was similar than in EM. IC animals had different response to immunocastration (high variability, tendency to elevated MUFA content).

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# Compared to SC, IC exhibited:

#### On carcass level:

→ \$\psi\$ smaller backfat thickness and leaf fat percentage.

#### On biochemical level:

- → 1 lipogenic enzyme activities,
- → \$\sigma \sigma \sigma \text{TUFA content.}

# EM compared to SC exhibited:

→ thicker dermis (probably î collagen content).

# Thank you for your attention!

These results were accomplished within project SuSI, co-financed by Susan EraNet and Slovenian Ministry of Agriculture, Forestry and Food. Collaboration within COST action CA15215 IPEMA and core financing of Slovenian Research Agency (grant P4-0133, PhD scholarship for KP) are also acknowledged.











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