TRANSCRIPTOMIC MARKERS OF BEEF MEAT TENDERNESS

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In Livestock species and more particularly in cattle, meat tenderness is a major factor to help producers to give value to their animals but also to satisfy the consumers. Although there is no genomic selection yet in beef cattle to meet these expectations, many researches have been conducted to get a better understanding of the physiological processes underlying meat tenderness and so to identify genes related to this trait.

OBJECTIVE: Identify key genes associated with beef tenderness by transcriptomic approaches

ANIMALS

25 Charolais young bulls 15 or 19 months of age

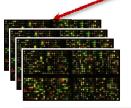


TRANSCRIPTOMIC ANALYSIS

Muscles:

- Longissimus thoracis (LT)
- Semitendinosus (ST)

Chips: 5 418 to 6 473 oligo



TENDERNESS EVALUATION

Shear force measurement: (Warner Brätzler-WB) 55°C grilled meat

COMPARISON OF TRANSCRIPTOME: according to meat tenderness

For each muscle:

Differentially expressed genes were identified by comparing groups of animals selected on the basis of their extreme WB value Correlations between gene expression level and WB value were determined using all animals data (n=25)

Up-regulated genes in the muscles of animals giving tender beef

LT: 20 ST: 195

194

CAPN3

CYP2E1

CAPN3: up-regulated ST muscle and negatively correlated with the WB value (r=-0,60)

Genes involved in apoptosis (DNAJA1, HSPA8) or genes belonging to the family of Heat Shock Protein (HSPCA, HSPD1...) were regulated in ST muscle giving tender beef but their expressions were not correlated with WB value

Down-regulated genes in the muscles of animals giving tender beef

LT: 23 ST: 204

DNAJA1 HSPA8 20 201 **HSPCA HSPD1** HSPB1, ATF3, U2AF1

HSPB1 and **ATF3**: positively correlated with the WB value in LT muscle (r=0,41 and 0,44)

In conclusion, the transcriptomic markers of beef meat tenderness are specific muscle. These genes are often involved in same pathway (apoptosis) and among them, DNAJA1 and HSPB1 were already shown to be related to beef sensory tenderness. These genes could be good genomic markers of tenderness evaluated by sensory or mechanical analyses.







