



# **Influence of castration and postnatal energy restriction on the contractile and metabolic characteristics of bovine muscle**

B Picard, J Robelin, Y Geay

## **► To cite this version:**

B Picard, J Robelin, Y Geay. Influence of castration and postnatal energy restriction on the contractile and metabolic characteristics of bovine muscle. *Annales de zootechnie*, 1995, 44 (4), pp.347-357. <hal-00889189>

**HAL Id: hal-00889189**

**<https://hal.science/hal-00889189v1>**

Submitted on 11 May 2020

**HAL** is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



HAL Authorization

Original article

## Influence of castration and postnatal energy restriction on the contractile and metabolic characteristics of bovine muscle

B Picard, J Robelin, Y Geay

INRA-Theix, laboratoire croissance et métabolismes des herbivores,  
63122 Saint-Genès-Champagnelle, France

(Received 25 July 1994; accepted 5 January 1995)

**Summary** — Twenty-four Montbeliard calves were raised with milk replacer to achieve a high or low rate of gain, 1 167 and 658 g/d, respectively. At 146 kg, the animals were weaned and 6 calves from each group were castrated. Thereafter, and until slaughter, animals of the same weight received the same diet in both groups. Biopsy samples of *Semitendinosus* (ST) muscle were taken in all animals at 140 kg, 305 kg, and at slaughter, 541 kg. The biochemical, contractile and metabolic characteristics of the ST samples were analyzed. The results showed that castration at the age of 4 months had a significant effect only on muscle characteristics measured at slaughter, 13 months after castration. The muscles of the steers contained smaller fibers, fewer type IIA fibers and more type IIB fibers than those of the bulls. In contrast, feed restriction before weaning resulted in changes to fiber characteristics after weaning at 305 kg. Restricted animals had smaller muscle fibers, a lower percentage of type I fibers and a higher percentage of type IIB fibers than animals fed *ad libitum*. These changes were no longer observed at slaughter.

**muscle / fiber type / castration / feed restriction**

**Résumé** — Influence de la castration et de la restriction énergétique postnatale sur les caractéristiques contractiles et métaboliques des muscles de bovins. Vingt-quatre veaux Montbéliards âgés d'environ 3 sem ont été répartis en 2 groupes. Le premier a reçu une ration permettant un gain moyen quotidien (GMQ) de 1 167 g/j, le second a été limité à un GMQ de 658 g/j jusqu'à la fin du sevrage, à l'âge de 4 mois en moyenne. Ensuite, jusqu'à l'abattage, les 2 groupes ont reçu, à même poids, le même régime. La moitié des animaux de chaque groupe a été castrée à 4 mois. Des échantillons du muscle *Semitendinosus* (ST) ont été prélevés par biopsie sur l'ensemble des animaux, à 2 stades correspondant respectivement à un poids moyen de 140 kg (à la fin du sevrage), 305 kg (9 mois environ) et à l'abattage à un poids moyen de 541 kg (à 17 mois en moyenne). Sur ces échantillons les caractéristiques biochimiques, contractiles et métaboliques du muscle ST ont été analysées. La castration à 4 mois n'a pas d'effet significatif sur les caractéristiques musculaires mesurées à 9 mois. En revanche, à l'abattage, 13 mois après castration, les animaux castrés par rapport aux animaux entiers, renferment moins de fibres de type IIA et plus de fibres de type IIB ; enfin, ils présentent des fibres mus-

*culaires de plus petite section. La restriction alimentaire avant le sevrage entraîne des modifications des caractéristiques des fibres qui sont encore mesurables 5 mois après le sevrage : les animaux restreints ont des fibres de plus petite section, un pourcentage de fibres IIA plus faible, et un pourcentage de fibres IIB supérieur à celui des animaux alimentés ad libitum. Ces modifications n'apparaissent plus à l'abattage, 8 mois plus tard. Une restriction énergétique appliquée avant le sevrage a donc des conséquences réversibles sur les caractéristiques musculaires.*

***muscle / type de fibre / castration / restriction énergétique / alimentation***

## INTRODUCTION

The commercial value of cattle depends on the muscle mass of the carcass and its adipose deposit content. The palatability of meat is partly governed by the contractile and metabolic characteristics of the muscles. Diet and castration can modify the growth rate of the different tissues. Steroid hormones affect the development of muscles and their biological characteristics and hence the quality of the meat. The meat of cows is more tender than that of steers and bulls (Field, 1971; Seideman *et al*, 1982). However, these effects depend on the muscle: the sensitivity of the neck, shoulder, cheek and thigh muscles to androgens decreases in that order (Butterfield, 1988). Certain muscles have sexual dimorphism, in particular the *Splenius* in cattle (Young and Bass, 1984), the *Temporalis* in guinea pigs (Bass *et al*, 1971; Lyons *et al*, 1986), the larynx in *Xenopus laevis* (Sassoon *et al*, 1987; Kelley *et al*, 1989) and the *Levator ani* in rats (D'Albis *et al*, 1991). Late castration is often performed in cattle to achieve rapid growth in the young animal and greater meat tenderness at slaughter.

Feed intake is another factor in cattle rearing that can modify growth rate, carcass composition and muscle characteristics. The effects on muscle characteristics vary depending on whether feed restriction is implemented before or after weaning (Haltia *et al*, 1978; Goldspink and Ward, 1979; Beverly *et al*, 1991; Yambayamba and Price, 1991). There is little available evidence on

postnatal energy restriction in cattle. In rats, postnatal restriction had irreversible effects (Bedi *et al*, 1982; Beverly *et al*, 1991). In contrast, Nordly *et al* (1987) showed that a restricted energy diet in gestating ewes had no effect on the muscle fiber characteristics of lambs observed at slaughter.

The aim of this study, therefore, was to determine the influence of castration and energy restriction in the early postnatal period on the contractile and metabolic characteristics of bovine muscle during the growth phase and at slaughter, which was performed, on average, at 17 months of age.

## MATERIALS AND METHODS

### ***Animal management and muscle sampling***

Twenty-four newborn Montbeliard calves were allocated, by age, weight and birth weight to 2 groups (I and II). At an average age of 4 months, the animals of the 2 groups were weaned when they attained 146 kg. Six calves from each group were castrated.

During the milk feeding period, group I received *ad libitum* a diet of reconstituted milk to achieve a high rate of gain (1 167 g/d), whereas the animals of group II received a restricted diet of reconstituted milk to achieve a low rate of gain (658 g/d). In the postweaning period, animals of the same weight were fed the same diet of 70% corn silage and 30% concentrate, calculated for a predetermined body weight gain of 1 100 g/d.

During these 2 periods, all of the animals were weighed weekly.

The circulating testosterone concentration was measured by radioimmunoassay at 4 months, just before castration, after castration and at 9 months.

Biopsy samples of *Semitendinosus* (ST) muscle were taken in animals of both groups at 3 stages (fig 1):

*Stage 1:* during the weaning period at the end of feed restriction (mean weight 146 kg),

*Stage 2:* 5 months after castration and the end of restriction (mean weight 305 kg),

*Stage 3:* at slaughter (mean weight 541 kg).

The effect of castration was studied on samples taken during stages 2 and 3 and that of diet on samples from stages 1, 2 and 3. The samples were frozen in isopentane and then in liquid nitrogen and stored at  $-80^{\circ}\text{C}$ .

## Muscle analysis

### Determination of protein/DNA ratio

Total muscle protein content, expressed in mg/g of muscle, was determined according to the method of Lowry *et al* (1951) with a preparation of 1 mg/ml of bovine albumin serum as standard. The DNA content, expressed in  $\mu\text{g/g}$  of muscle, of the samples taken at slaughter was measured by the method of Labarca and Paigen (1980).

The protein/DNA ratio was expressed in mg of protein per mg of DNA.

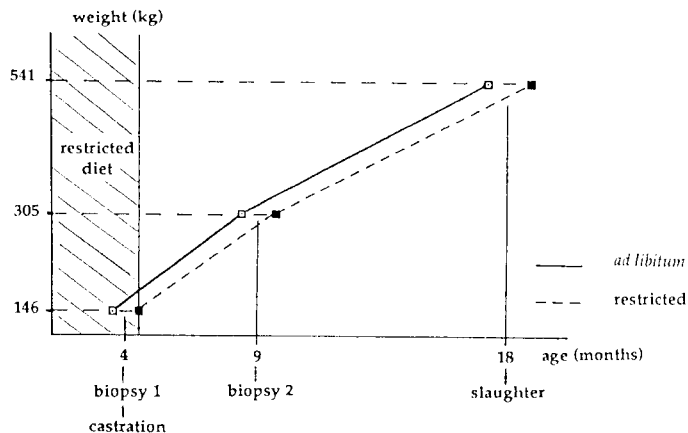
### Determination of contractile type of muscle fibers

#### Histological, immunohistochemical and histochemical studies

Serial sections 10  $\mu\text{m}$  thick were cut perpendicular to the muscle fibers with a cryotome at  $-25^{\circ}\text{C}$ . Three types of staining were performed on the sections to determine the size of the fibers, their contractile activity and their metabolic activity.

The muscle fibers were treated with azorubine, which stains all fibers irrespective of contractile or metabolic type. It shows the general histological architecture of the muscle and thereby allows determination of the number and diameter of the fibers. Slow (I) and fast (II) contractile types were determined by immunohistochemistry according to the immunofluorescence technique described by Pons *et al* (1986), who used monoclonal antibodies specific to myosin heavy chains (MHC). Their production, purification and characterization have been described by Léger *et al* (1985). Their reactivity in cattle has been tested by Robelin *et al* (1993). Type IIA (oxidoglycolytic) fibers were distinguished from type IIB (glycolytic) by succinate dehydrogenase (SDH) activity according to the method of Pearse (1968). This enzyme, which is present in the cell wall of mitochondria, is characteristic of an oxidative metabolism. Fibers of this metabolic type (I, IIA) stained dark blue in the presence of nitro-

**Fig 1.** Experimental design. Stage 1 corresponds to the end of the period of energy restriction when biopsy samples were taken in the 2 groups of 6 animals at a mean age of 4 months and a mean weight of 146 kg. Stage 2 corresponds to the second biopsy when animals had a mean age of 9 months and a mean weight of 305 kg. Stage 3 corresponds to slaughter, performed at a mean age of 17 months for a mean weight of 541 kg.



bleuetetrazolium, which is reduced by the action of the enzyme, whereas glycolytic fibers (IIB) stained pale blue. A fourth type of fiber, IIC, was evidenced. The cells were recognized by both anti-MHC1 (slow) and anti-MHC2 (fast) antibodies and had an oxidoglycolytic metabolism.

The proportion of each type of fiber and their average area were determined in 2 different sites on each serial section. Between 100 and 200 fibers were analyzed by an image-analysis computer program (Visilog).

#### *ELISA: enzyme-linked immunosorbent assay*

This technique was used to quantify the proportion of slow isoform MHC1. It was based on the method of Winkelmann *et al* (1983) modified by Picard *et al* (1994).

#### **Determination of muscle metabolic type**

The metabolic type of the muscles studied was determined by measuring the enzyme activities of isocitrate dehydrogenase (ICDH) according to the method of Briand *et al* (1981) and of lactate dehydrogenase (LDH) by the method of Ansary (1974). ICDH activity was measured only on the samples taken at slaughter since the biopsy specimens were not large enough to permit the performance of all the assays.

#### **Analysis of results**

The results were subjected to analysis of variance with the GLM module of the SAS package (1985), to study the effect of castration and diet.

### **RESULTS**

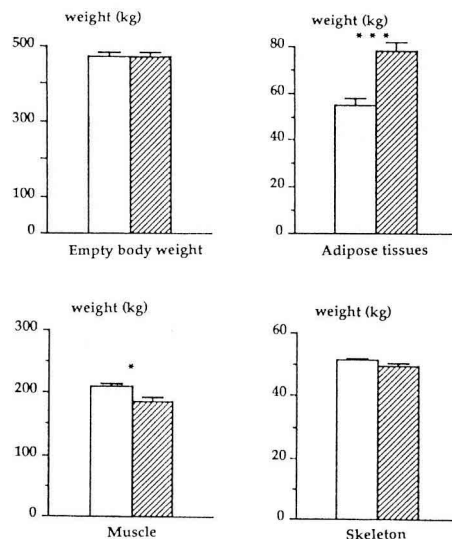
#### **Effects of castration**

At slaughter, the castrated animals had a lower muscle weight and a greater overall amount of adipose deposit content than the bulls (fig 2).

There was no significant difference in muscle characteristics between the bulls and the animals castrated at stage 2, 5 months after castration. In contrast, 13 months after castration, the steers had a lower oxidative metabolism and a smaller proportion of MHC1 than the bulls (fig 3). Their muscles contained significantly lower proportions of type IIA fibers and a significantly higher percentage of type IIB (fig 4), and the surface of all their muscle fibers was smaller (fig 5).

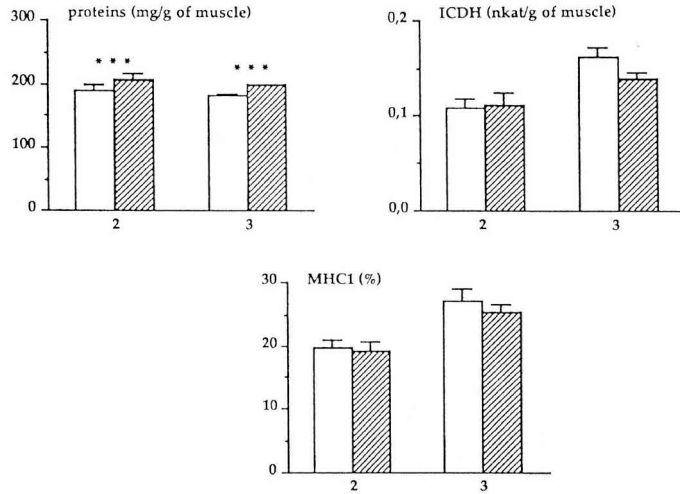
#### **Effects of energy restriction**

There was no significant difference at slaughter between restricted animals and those fed *ad libitum* in total adipose tissue content, muscle and skeleton weight (fig 6). However, significant differences were found at stage 2, 5 months after diet restriction, but at slaughter were no longer observed.



**Fig 2.** Weight and body composition: empty body weight; adipose deposit content; muscle weight; skeleton weight of bulls □ and steers ▨ at slaughter. \*  $P < 0.05$ , \*\*  $P < 0.01$ , \*\*\*  $P < 0.001$ .

**Fig 3.** Biochemical characteristics of muscles: total muscle protein content; isocitrate dehydrogenase (ICDH) activity (oxidative); percentage of slow myosin heavy chains (MHC1) of bulls  $\square$  and steers  $\text{▨}$  measured at 9 (stage 2) and 17 months (stage 3).



At stage 1, the only dissimilarity between animals was the higher proportion of protein in the muscles of those fed a restricted diet. Conversely, at stage 2, restricted animals had less muscle protein than those fed *ad libitum* (fig 7). At this stage, significant differences were observed in the proportion of the different types of fiber. Restricted animals had fewer type I fibers and more type IIB fibers, but the same proportion of type IIA fibers as animals fed *ad libitum* (fig 8). While there was no difference in the surface of the latter 2 types of fibers, the surface of type I fibers was significantly smaller in restricted animals (fig 9). At stage 3, although no significant difference was observed between the 2 groups, the animals on a restricted diet tended to have

fewer type IIB fibers and more type IIA than those fed *ad libitum* (fig 8).

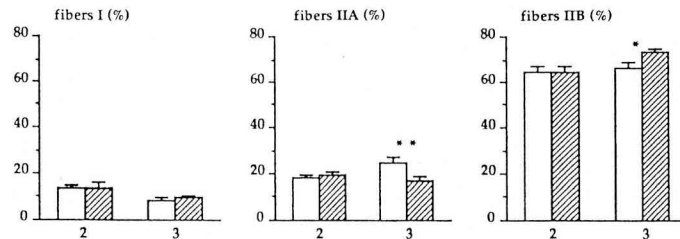
At all 3 stages studied, the ICDH activity tended to be lower in restricted animals, but the differences were not significant (fig 7). Likewise, there was no significant difference in the percentage of MHC1 between the animals at any of the 3 stages (fig 7).

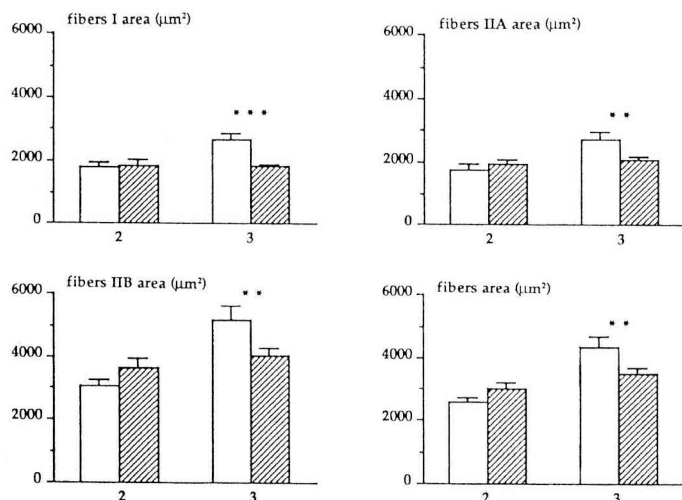
## DISCUSSION

### Effects of castration

The effects of castration on carcass composition observed in this study were consistent with previous findings. Castration

**Fig 4.** Proportions of the different fiber types: type I fibers (slow, oxidative); type IIA fibers (fast, oxidoglycolytic); type IIB fibers (fast, glycolytic) in bulls  $\square$  and steers  $\text{▨}$  measured at 9 (stage 2) and 17 months (stage 3).



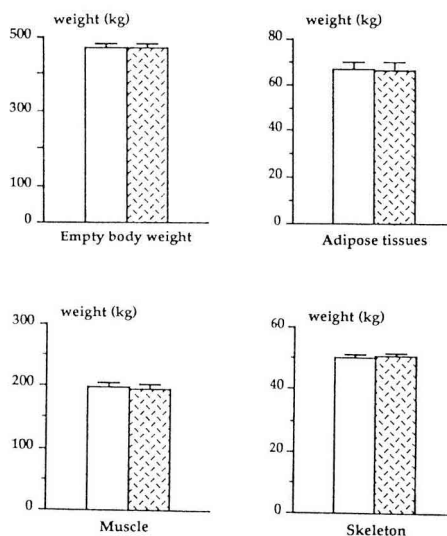


**Fig 5.** Surface of the different fiber types: type I fibers (slow, oxidative); type IIA fibers (fast, oxidoglycolytic); type IIB fibers (fast, glycolytic); mean surfaces in bulls □ and steers ▨ measured at 9 (stage 2) and 17 months (stage 3).

decreases muscle growth and increases adipose tissue content (Field, 1971; Raju, 1975; Seidman *et al*, 1982) as a result of the effect of testosterone on muscle protein anabolism. In several species, testosterone has been observed to enhance protein synthesis and hence to increase the body mass and size of certain muscles (Kochakian and Stettner, 1949; Field, 1972; Kochakian, 1975; Martinez *et al*, 1984; Griggs *et al*, 1989). In light of this evidence, it is surprising that in our experiment the protein content of ST muscles was significantly greater in the steers. It should be added, however, that the protein content measured in *Maseter*, *Rectus abdominis* and *Tensor fascia lata* muscles did not vary significantly between bulls and steers at the 2 stages studied (unpublished results).

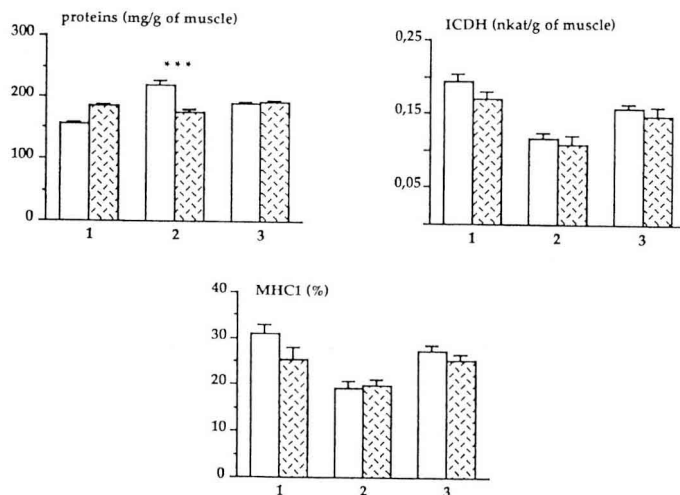
The original aspect of our results was to show that changes in muscle characteristics occurred between 5 and 13 months after castration. Five months after castration, no effect on muscle characteristics was observed, whereas after 13 months, significant differences were noted between bulls and steers. The principal differences were in the size and proportion of fiber types, but all features measured varied consistently.

Between birth and 4 months of life, testosterone plasma levels in calves are at very low levels, between 0 and 1 ng/ml (Lacroix *et al*, 1977). However, between 4 and 6 months, high peaks may be observed

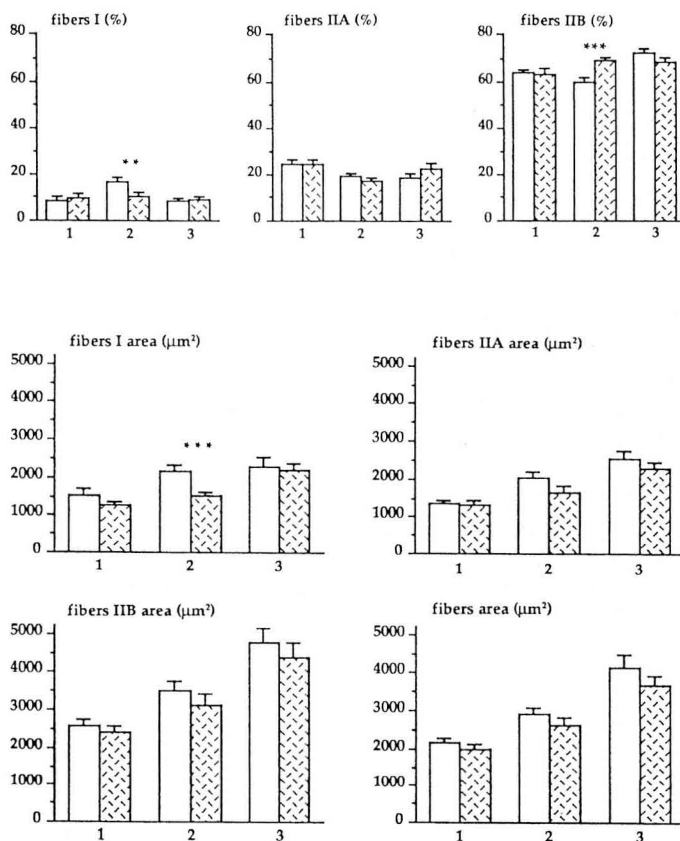


**Fig 6.** Weight and body composition: empty body weight; total adipose tissue content; muscle weight; skeleton weight of animals fed a restricted diet □ or *ad libitum* ▨, measured at slaughter.

**Fig 7.** Biochemical characteristics: total muscle protein content; isocitrate dehydrogenase (ICDH) activity (oxidative); percentage of slow myosin heavy chains (MHC1) of the muscles of animals fed a restricted diet  $\square$  or *ad libitum*  $\boxtimes$ , measured at 4 (stage 1), 9 (stage 2) and 17 months (stage 3).



**Fig 8.** Proportions of the different types of muscle fibers: type I fibers (slow, oxidative); type IIA fibers (fast, oxidoglycolytic); type IIB fibers (fast, glycolytic) of animals fed a restricted diet  $\boxtimes$  or *ad libitum*  $\square$ , measured at 4 (stage 1), 9 (stage 2) and 17 months (stage 3).



**Fig 9.** Surfaces of the different types of muscle fibers: type I fibers (slow, oxidative); type IIA fibers (fast, oxidoglycolytic); type IIB fibers (fast, glycolytic); mean surfaces of animals fed a restricted diet  $\boxtimes$  or *ad libitum*  $\square$ , measured at 4 (stage 1), 9 (stage 2) and 17 months (stage 3).



whereas between 6 and 12 months, testosterone levels return to very low levels and then increase after 12 months (Rawlings *et al*, 1972; Karg *et al*, 1975; Lacroix *et al*, 1977). The period from 4 to 12 months of life corresponds to the onset of puberty, which is generally completed by the age of 1 year or thereabouts (Thibier, 1975; Lacroix *et al*, 1977). It is likely that testosterone is present before the end of puberty but that its muscle receptor is not yet active. The hormone-receptor system may then become functional once puberty has begun. It can therefore be supposed that the differences in muscle characteristics between bulls and steers begin to occur after puberty.

The effects of castration on muscle characteristics that we observed at the age of 17 months were comparable to those reported previously in animals of various species, including cattle, but which were not castrated at the same age as in our experiment. All cell surfaces were found to be smaller in castrated animals. This difference seems to be the first effect of castration. For example, Tobin and Pécot-Dechavanne (1982) showed that 1 month after castration, the *Levator ani* muscle in rats were severely atrophied, not as a result of the number of fibers but of their diameter. The same effect has been observed in cattle, in different muscles and breeds (Dreyer *et al*, 1977; Ockerman *et al*, 1984; Young and Bass, 1984; Seideman and Crouse, 1986; Clancy *et al*, 1986). In frogs, Regnier and Herrera (1993) reported a positive correlation between fiber size and testosterone plasma levels. In contrast, Laflamme *et al* (1973) observed no effect of castration on fiber size, DNA and protein contents in *Longissimus dorsi* muscles from 10 pairs of cattle twins.

The bulls had more type IIA fibers and fewer type IIB fibers than the steers. These findings are in agreement with the results observed in different studies on cattle cas-

trated at a later age (Dreyer *et al*, 1977; Ockerman *et al*, 1984; Young and Bass, 1984; Seideman and Crouse, 1986; Clancy *et al*, 1986). They lend weight to the hypothesis that androgens slow down the effect of age on the transformation of type IIA fibers into type IIB fibers and suggest that steroids have a modulating effect on the expression of genes encoding for myosin isoforms. Lyons *et al* (1986) showed that testosterone triggered the appearance of fast type 2a MHC in the *Temporalis* muscle of male rats. Likewise, in rat cardiac muscles, Morano *et al* (1990) showed that castration favored the expression of MHC2b, whereas the administration of testosterone resulted in the appearance of MHC2a. These authors demonstrated that testosterone regulates the expression of cardiac MHC at a pretranslational level. Differences in the expression of MHC2a and 2b have also been described by Dalla Libera *et al* (1980) in guinea pigs. Our results on the expression of MHC1 showed no significant differences between bulls and steers. However, in steers the muscles tended to contain lower percentages of MHC1.

The oxidative mechanism of ST muscle was generally lower in the steers, but the difference was not significant. This finding is consistent with the differences observed in the proportions of the different fiber types. It is also in agreement with the well-documented observation that the muscles of bulls are redder than those of steers (Seideman and Crouse, 1986).

### **Effects of energy restriction**

Energy restriction was observed to have an effect on muscle characteristics 5 months after it was stopped. However, at slaughter (13 months after the treatment), no significant difference was found between the 2 groups of animals. This result is at variance with different published reports

which state that a period of undernutrition before weaning has irreversible effects on muscle characteristics (Dobbin, 1974; Bedi *et al*, 1982).

Results from the literature show that the effect of energy restriction on muscle characteristics differs according to whether it is implemented before or after weaning. Haltia *et al* (1978) and Beverly *et al* (1991) reported a decrease in the proportion of type I fibers accompanied by an increase in that of type IIB fibers in rats which had undergone energy restriction during the perinatal period. As rats are less developed at birth than cattle, it is difficult to compare the 2 sets of results. However, various authors have shown that energy restriction after weaning decreases the proportion of type IIB fibers and increases that of type IIA fibers in different species (Goldspink and Ward, 1979, in rats; Moody *et al*, 1980; and Solomon and Lynch, 1988, in sheep; and Seideman and Crouse, 1986, in cattle). These changes are commonly explained as a consequence of the hypothyroidism that occurs during a period of undernutrition (Beverly *et al*, 1991). It is known that thyroid hormones regulate the expression of type IIB myosin isoform (MHC2b) (Bacou and Vigneron, 1988). The changes are accompanied by a decrease in the oxidative activity of the muscles of animals on a restricted diet (Raju, 1975; Howells and Jordan, 1978).

The size of the different fiber types is, in contrast, always decreased after a period of energy restriction, whether it be before or after weaning (Moss, 1971; Haltia *et al*, 1978; Goldspink and Ward, 1979; Bedi *et al*, 1982; Burwin *et al*, 1988; Lanz *et al*, 1992). This diminution is due to a decrease in muscle DNA and protein content (Moss, 1971; Ward and Stickland, 1993), as observed in our experiment at stage 2 (9 months, 305 kg).

The differences noted in muscle characteristics at stage 2 are in agreement with the results of Haltia *et al* (1978) and of Be-

erly *et al* (1991) in rats. The fact that by the time of slaughter there were no longer any effects of energy restriction observed, either on body composition or muscle characteristics, is in agreement with the findings of Morgan (1972), who stated that undernutrition at the beginning of postnatal life had no effect on the quality of meat, as had been suggested in an early study by Winchester and Ellis (1956).

## CONCLUSION

This study demonstrates the opposing effects of castration and energy restriction implemented at an early age on muscle characteristics. The effect of castration increased with age whereas restriction had only a temporary impact, which disappeared as the animal matured. It therefore yields novel results on the effects of 2 factors involved in cattle rearing — androgens and feed intake — about the contractile and metabolic characteristics of muscles. The original result of this study is that the differences in the muscle characteristics due to early castration were not observed until after puberty, which suggests that the testosterone-receptor system of the muscles was not operational until that time. The principal differences observed at slaughter were in the size and distribution of the different fiber types. In particular, steers had smaller fibers than the bulls, and their muscles contained more type IIB fibers and fewer type IIA fibers.

Undernutrition between birth and weaning had temporary effects on the distribution of type I and type IIB fibers. One of the most salient points to emerge from this study is that the energy restriction during the perinatal period has no effect on the carcass composition or on the contractile and metabolic characteristics of muscles at the standard age for slaughter of about 16 months.

## ACKNOWLEDGMENTS

The authors thank Rt Jailler and G Cuyllé for animal management and slaughtering, and C Barboiron for chemical analysis and Rd Jailler for statistical analysis.

## REFERENCES

- Ansary (1974) Individualité musculaire chez le bovin : étude de l'équipement enzymatique de quelques muscles. *Ann Biol Anim Biochim Biophys* 14, 471-486
- Bacou F, Vigneron P (1988) Propriétés des fibres musculaires squelettiques. I. Influence de l'innervation motrice. *Reprod Nutr Dev* 28, 1387-1453
- Bass A, Gutmann E, Hanzlikova V, Syrový L (1971) Sexual differentiation of enzyme pattern and its conversion by testosterone in the temporal muscle of the guinea pig. *Physiol Biochem* 20, 423-431
- Bedi KS, Birzgalis AR, Mahon M, Smart JL, Wareham AC (1982) Early life undernutrition in rats. I. Quantitative histology of skeletal muscles from underfed young and refed adult animals. *Br J Nutr* 47, 417-431
- Beverly SB, Daoud MJ, Laframboise WA, Watchko JF, Foley TP, Butler-Browne GS, Whalen RG, Guthrie RD, Ontell M (1991) Effects of perinatal undernutrition on elimination of immature myosin isoforms in the rat diaphragm. *Lung Cell Mol Physiol* 5, 49-54
- Briand M, Talmant A, Briand Y, Monin G, Durand R (1981) Metabolic types of muscle in sheep. II. Lactate dehydrogenase activity and LDH isozyme distribution. *Eur J Appl Physiol* 46, 359-365
- Burwin DG, Ferrell CL, Britton RA (1988) Effect of feed intake of lambs on visceral organ growth and metabolism. In: *Energy metabolism of animals. Proc 11th Symposium INRA*, 103-106
- Butterfield RM (1988) *New concept of sheep growth*. Dept of Vet Anatomy, Univ Sidney, Epping, Australia
- Clancy MJ, Janet M, Lester, Roche JF (1986) The effects of anabolic agents on the fibers of the 1 dorsi muscle of male cattle. *J Anim Sci* 100-111
- D'Albis A, Janmot C, Mira JC, Couteaux R (1991) Characterization of a ventricular V1 myosin isoform in rabbit masticatory muscles. Development and neural regulation. *BAM* 1, 23-24
- Dalla Libera L, Sartore S, Pierobon-Bormioli S, Schiaffino S (1980) Fast-white and fast-red isomyosins in guinea pig muscles. *Biochem Biophys Res Com* 96, 1662-1670
- Dobbing J (1974) *Scientific fundations of paediatrics* (JA Davis, J Dobbing, eds), Heinemann, London, Philadelphia, 565 p
- Dreyer JH, Naude RT, Henning JNN, Rossow E (1977) The influence of breed, castration and age on muscle fibre type and diameter in Friesland and Africaner cattle. *S Afr J Anim Sci* 7, 171-180
- Field RA (1971) Effect of castration on meat quality and quantity. *J Anim Sci* 32, 849-858
- Goldspink G, Ward PS (1979) Changes in rodent muscle fiber types during post-natal growth, undernutrition and exercise. *J Physiol* 296, 453-469
- Griggs RC, Kingston W, Jozefowics RF, Herr BE, Forbes G, Halliday D (1989) Effect of testosterone on muscle mass and muscle protein synthesis. *J Appl Physiol* 66, 498-503
- Haltia M, Berlin O, Schucht H, Sourander P (1978) Post-natal differentiation and growth of skeletal muscle fibers in normal and undernourished rats. A histochemical and morphometric study. *J Neurol Sci* 36, 25-39
- Howells KF, Jordan TC (1978) The effects of pre- and perinatal undernutrition on the succinic dehydrogenase content of muscle fibers from fast and slow rat muscles. *Histochemistry* 58, 97-102
- Karg H, Gimenez T, Hartl M, Hoffman B, Schallenberger E, Schams D (1975) Testosterone, luteinizing hormone (LH) and follicle stimulating hormone (FSH) in peripheral plasma bulls: levels from birth through puberty and short-term variations. *Zentralbl Veterinarmed [A]* 23, 793-803
- Kelley DB, Sassoon D, Segil N, Scudder M (1989) Development and hormone regulation of androgen receptor levels in the sexually dimorphic larynx of *Xenopus laevis*. *Dev Biol* 131, 111-118
- Kochakian CD (1975) Definition of androgen and protein anabolic steroids. *Pharmacol Ther* 1, 149-177
- Kochakian CD, Stettner (1949) Renotropic-androgenic and somatotrophic properties of further steroids. *Am J Physiol* 158, 51-56
- Labarca C, Paigen K (1980) A simple, rapid, sensitive DNA assay procedure. *Anal Biochem* 102, 344-352
- Lacroix A, Garnier DH, Pelletier J (1977) Temporal fluctuations of plasma LH and testosterone in Charolais bull calves during the first year of life. *Ann Biol Anim Biochim Biophys* 17, 1013-1019
- Laflamme LF, Trenkle A, Topel DG (1973) Effect of castration or breed type on growth of the *Longissimus* muscle in male cattle. *Growth* 37, 249-256
- Lanz JK, Donahoe M, Rogers RM, Ontell M (1992) Effects of growth hormone on diaphragmatic recovery from malnutrition. *Am Physiol Soc* 161, 801-805
- Léger JOC, Bouvagnet P, Pau B, Roncucci R, Léger JJ (1985) Levels of ventricular myosin fragments in human sera after myocardial infarction, determined with monoclonal antibodies to myosin heavy chains. *Eur J Clin Invest* 15, 422-429
- Lowry OH, Rosebrough NJ, Farr AL, Randall RJ (1951) Protein measurement with the Folin phenol reagent. *J Biol Chem* 193, 265

- Lyons G, Kelley A, Rubinstein W (1986) Testosterone-induced changes in contractile protein isoforms in the sexually dimorphic temporalis muscle of the guinea pig. *J Biol Chem* 26, 13278-13284
- Martinez JA, Buttery PJ, Pearson JT (1984) Mode of action of anabolic agents. Effects of testosterone on muscle protein turn-over in the female rat. *Br J Nutr* 52, 515-521
- Moody WG, Kemp JD, Mahyuddin M, Johnston DM, Ely DG (1980) Effect of feeding systems, slaughter weight and sex on histological properties of lamb carcasses. *J Anim Sci* 50, 249-256
- Morano I, Gerstner J, Caspar Rüegg J, Ganter U, Ganter D, P Vosberg H (1990) Regulation of myosin heavy chain expression in the hearts of hypertensive rats by testosterone. *Circ Res* 66, 1585-1590
- Morgan JHL (1972) Effect of plane of nutrition in early life on subsequent live-weight gain, carcass and muscle characteristics and eating quality of meat in cattle. *J Agric Sci (Camb)* 78, 417
- Moss FP (1971) The relationship between the dimensions of the fibers and the numbers of nuclei during restricted growth, regrowth and compensatory growth of skeletal muscles. *Am J Anat* 122, 565
- Nordly DJ, Field RA, Riley ML, Kercher CJ (1987) Effects of maternal undernutrition during early pregnancy on growth, muscle cellularity, fiber type and carcass composition in lambs. *J Anim Sci* 64, 1419-1427
- Ockerman HW, Jaworek D, Van Stavern B, Parrett N, Pierson CJ (1984) Castration and sire effects on carcass traits, meat palatability and muscle fibre characteristics in Angus cattle. *J Anim Sci* 59, 981-990
- Pearse AGE (1968) *Histochemistry theoretical and applied*, 2d ed, vol 2 (JA Churchill, ed), London, 948 p
- Picard B, Léger JOC, Robelin J (1994) Quantitative determination of type IMHC in bovine muscle with monoclonal antibodies. *Meat Sci* 36, 333-345
- Pons F, Léger JOC, Chevallay M, Tome FMS, Fardeau M, Léger JJ (1986) Immunocytochemical analysis of myosin heavy chains in human fetal skeletal muscle. *J Neurol Sci* 76, 151-163
- Raju NV (1975) Effect of early malnutrition on muscle function and metabolism in rats. *Life Sci* 15, 949-960
- Rawlings NC, Hafs HD, Swason LV (1972) Testicular and blood plasma androgens in Holstein bulls from birth through puberty. *J Anim Sci* 34, 435-440
- Regnier M, Herrera AA (1993) Differential sensitivity to androgens within a sexually dimorphic muscle of male frogs (*Xenopus laevis*). *J Neurobiol* 24, 1215-1228
- Robelin J, Picard B, Listrat A, Jurie C, Barboiron C, Pons F, Geay Y (1993) Myosin expression in *Semitenid-nosus* muscle during fetal development of cattle: immunocytochemical and electrophoretic analysis. *Reprod Nutr Dev* 33, 25-41
- SAS (1985) *SAS user's guide*, Version 6 Edition, SAS Institute Inc, Cary, NC, USA
- Sassoon D, Gray G, Kelley DB (1987) Androgen regulation of muscle fiber type in the sexually dimorphic larynx of *Xenopus laevis*. *J Neurosci* 7, 3198-3206
- Seideman SC, Crouse JD (1986) The effects of sex condition, genotype and diet on bovine muscle fiber characteristics. *Meat Sci* 17, 55-72
- Seideman SC, Cross HR, Oltjen RR, Schanbacher BD (1982) Utilization of the intact male for red meat production: a review. *J Anim Sci* 55, 826-840
- Solomon MB, Lynch GP (1988) Biochemical histochemical and palatability characteristics of young ram lambs as affected by diet and electrical stimulation. *J Anim Sci* 66, 1955
- Thibier M (1975) Variations circadiennes de la LH plasmatique chez le jeune taurillon. *Ann Endocrinol (Paris)* 36, 205-210
- Tobin C, Pécot-Dechavassine M (1982) Effect of castration on the morphology of the motor end-plates of the rat levator ani muscle. *Eur J Cell Biol* 26, 284-288
- Ward SS, Stickland NC (1993) The effect of undernutrition in the early post-natal period on skeletal muscle tissue. *Br J Nutr* 69, 141-150
- Winchester CF, Ellis NR (1956) Delayed growth of beef cattle. *USDA Tech Bull* 1159
- Winkelmann DA, Lowey S, Press JL (1983) Monoclonal antibodies localize changes on myosin heavy chain isozymes during avian myogenesis. *Cell* 34, 295-306
- Yambayamba E, Price MA (1991) Fiber type proportions and diameters in the *Longissimus* muscle of beef heifers undergoing catch-up (compensatory) growth. *Can J Anim Sci* 71, 1031-1035
- Young OA, Bass JJ (1984) Effect of castration on bovine muscle composition. *Meat Sci* 11, 139-156