

CARCASS FAT AND INTRAMUSCULAR FAT DISTRIBUTION IN PIGS

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**Summary**  
The quality and quantity of intramuscular fat (IMF) were determined in 31 muscles from eight Duroc Jersey barrows of each of two lines, lean (L) and fat (F) at similar live-weight. Differences in IMF distribution were detected between lean and fat pigs. The total amount of IMF was almost identical (0.43 and 0.41 kg, respectively) in spite of big differences in total dissected fat (26.6 and 42.9 kg, respectively) or % total dissected fat (32.6 and 32.9%, respectively). The IMF% were higher in the fat pig muscles compared with the lean pig muscles except for Mm Vastus lateralis, Semimembranosus, Adductor, Sartorius, Obliquus internus abdominis and Transversus abdominis. The fatty acid composition of IMF shows a strong tendency for a higher percentage of linoleic fatty acid in muscles for the L pigs.

**Introduction**  
Intramuscular fat makes a positive contribution to the palatability characteristics of pork. Selection towards a maximum of muscularity is related to a strong reduction of the amounts of fatty tissues and intramuscular fat and has led to general decline in meat quality (Desmoulin, 1983). It might be possible that the amount of intramuscular and the quality of the fatty tissues become the limiting factors in breeding extremely lean pigs. In several studies low correlations have been found between the concentration of lipid in muscle and carcass fatness (Duniec et al., 1961; Hiner et al., 1965). This suggests that fat deposition in muscle is not part of the fattening process which occurs in the other fat depots of the body. This research was undertaken to determine the differences in the quality and quantity of intramuscular fat from Duroc Jersey pigs with different fat carcass levels but similar carcass weights.

**Materials and Methods**  
Eight barrows Duroc Jersey of each of the two lines, lean (L) and fat (F), reared at Pergamino Experimental Station of INTA were used as experimental animals. They were allocated to individual pens with water and a standard diet supplied ad libitum. Both lines at the end of the trial were similar in carcass weight (80.2+3.18 and 81.2+3.44 kg respectively for the L and F lines) but they were different in carcass fat % (32.9+2.95 and 32.9+4.50 respectively). The right side of the carcasses were kept at -15 C and 31 muscles dissected as soon as possible. Each muscle was weighted, minced and aliquot samples extracted with the method of Polch et al., (1957). Aliquot samples from the chloroform extract were used to determine the total weight of chemical fat (IMF) and for fatty acid analysis. TLC and GLC were used for the lipid separation and methyl ester analysis respectively as was previously reported by Garcia et al., 1986). The muscles studied were: Proximal pelvic limb (G1): Tensor fascia latae (TLF), Biceps femoris (BFE), Glutaeus medius, accessorius et profundus (GLU), Vastus lateralis (VLA), Vastus medialis (VME), Vastus intermedius (VIN), Rectus femoris (RFE), Sartorius (STE), Gracilis (GRA), Semimembranosus (SME), Adductor (ADD), Pectineus (PEC) and Sartorius (SAR). Distal pelvic limb (G2): Gastrocnemius et soleus (GAS), Fibularis digitorum superficialis et profundus & Fibularis longus (FPL) and Flexor digitorum longus (FDL). Spinal (G3): Psoas major (PMA) and Longissimus dorsi (LDO). Abdominal (G4): Obliquus externus abdominis (OEA), Obliquus internus abdominis (OIA), Transversus abdominis (TRA) and Rectus abdominis (REA). Proximal thoracic limb (G5): Deltoides (DEL), Infraspinatus (INF), Triceps brachii (TBR),

Teres major (TEM), Supraspinatus (SUP), Biceps brachii (BIB) and Subcapularis (SUB). Distal thoracic limb (G6): Extensor carpi radialis, Ulnaris lateralis, Abductor digiti I longus, Extensor digitorum communis et lateralis (ETL) and Flexor carpi radialis et Ulnaris. Proximal forelimb: Flexor digitorum profundus et superficialis (FTL).

The data were processed statistically by the NWASTATPAK Program Northwest Analytical Services, Inc. Portland Oregon USA.

### Results and Discussion

The individual IMF content as a percent of total IMF in each anatomical muscle group for L and F pigs are given in Figures 1, 2, 3 and 4. Significant differences in IMF distribution between the L and F pigs were detected. A significant accumulation ( $p < .05$ ) of IMF fat occurs in Mm BFE and GLU in F pigs and in GRA, SME and ADD in L pigs in the pelvic limb muscle group (Fig. 1). In spite of the big SD in the abdominal muscle group fat content shows the most important changes, specially in Mm OIA and REA ( $p < .05$ ). Differences in the other muscle groups were less significant. The total IMF fat weight in each anatomical muscle group as a percentage of total IMF in the L and F pigs is shown in Figure 5. Statistical differences ( $p < .05$ ) in the distribution of IMF between the groups are shown for groups 1, 2 and 3. The total amount of IMF fat was almost identical, 0.43 and 0.41 kg in spite of big differences in carcass fat. This results explain the correlation found between total body fat and IMF for many authors (Duniec et al., 1961; Hiner et al., 1965) and the difficulties in the estimation of the effects of variables like breed, sex, diet on the IMF content fat in a given muscle taking in account carcass fatness or any predictor indicator of carcass fatness as a back fat thickness. The increasing incidence of meat quality problems requires new methods of meat prediction (Kempster et al 1986).

The IMF% in the muscles from the two lines are given in Figure 1, 2, 3 and 4. The IMF% were higher in the F pigs compared with the L ones except for the Mm VLA, SME and TRA (Group 1); OIA and TRA (Group 4) and extensor muscles (Group 6).

The levels of linoleic acid (18:2) in the muscle lipids were always higher in the lean pigs than in the fat pigs (Fig.6). The values were from 6.3 to 13.4% in the lean pigs and from 4.9 to 10.9% in the fat pigs. The % of 18:2 was related negatively to the IMF % in the muscle but also at similar IMF% the F pigs have less 18:2 in total lipids than the L pigs. No differences in the average daily gain between the two lines were detected (0.970 vs 0.929 g for L and F pigs) and then the difference may be due to the different contribution of the lipid classes to the total muscular lipids. At similar diet the firmest fat was observed when growth rate and fat thickness were high (Wood, 1983).

### Conclusions

Differences in the distribution of intramuscular fat between lean and fat pigs were found. The total amount of intramuscular fat was similar in both lines. The fatty acid composition of intramuscular lipids showed a strong tendency for a higher percentage of linoleic acid in the lean pigs.

### References

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Fig. 1. IMF% and IMF weights as a % of total IMF in the anatomical muscle group.

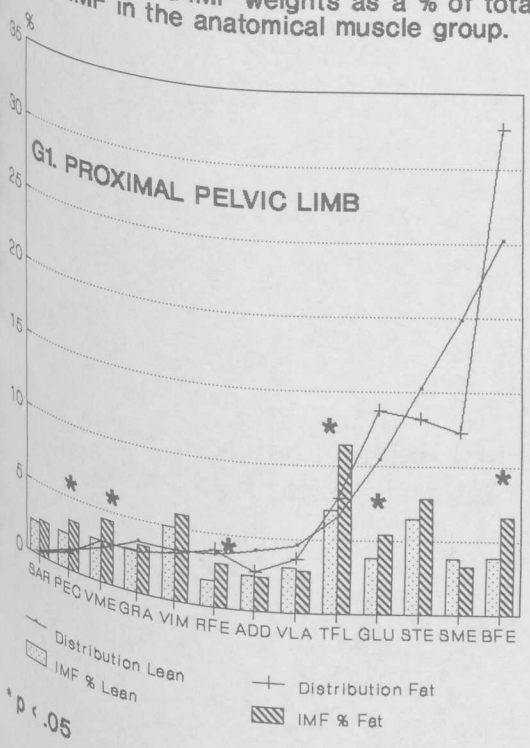


Fig. 2. IMF% and IMF weights as a % of total IMF in the anatomical group.

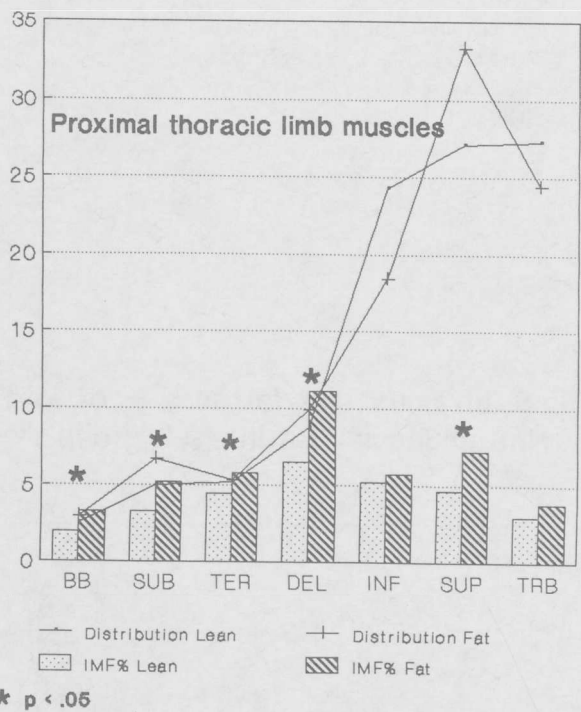
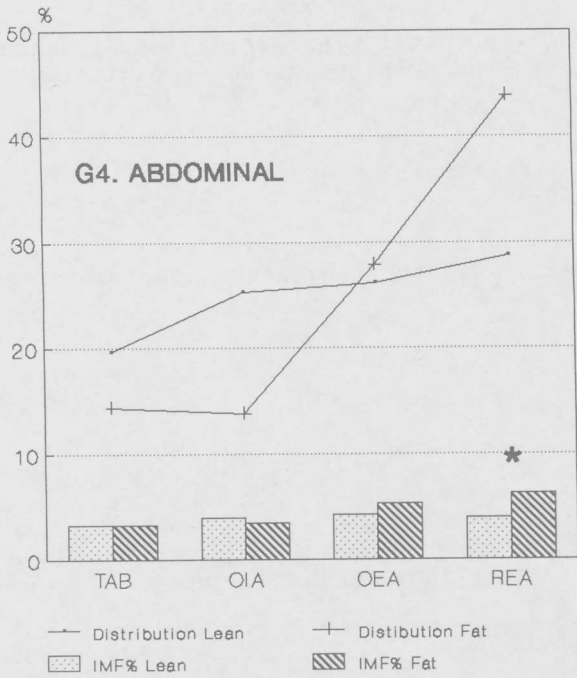
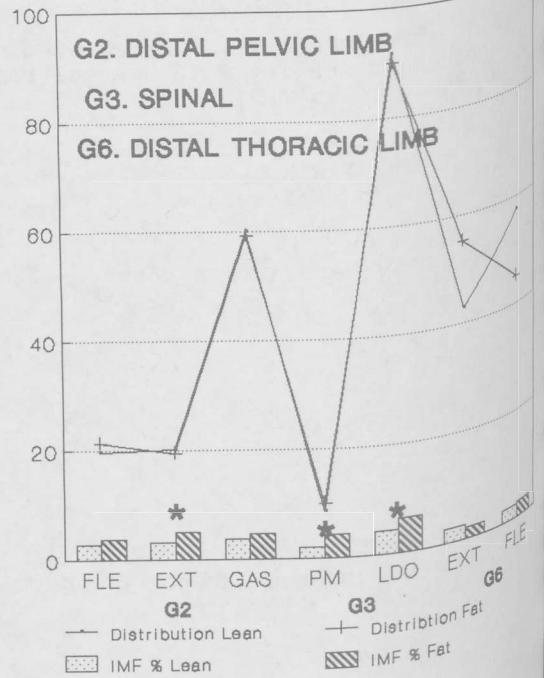


Fig. 3. IMF% and IMF weights as a % of total IMF in the anatomical group.



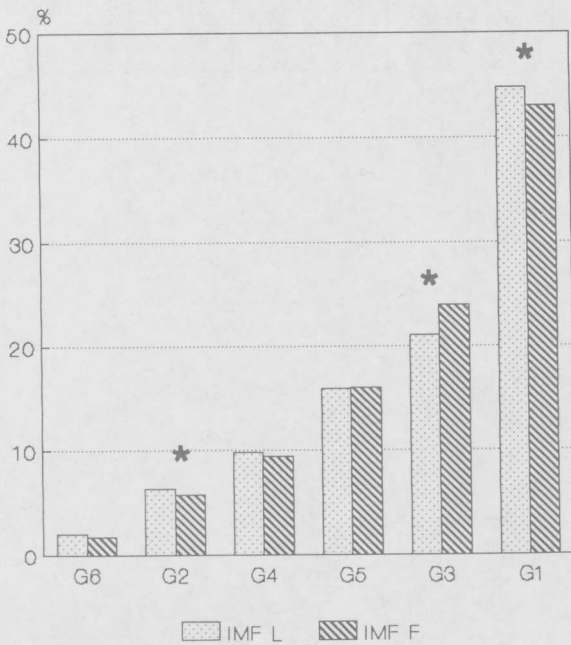
\* p < .05

Fig. 4. IMF% and IMF weights as a % of total IMF in the anatomical muscle group.



\* p < .05

Fig. 5. Intramuscular fat as a % of total intramuscular fat in each group.



\* p < .05.

Fig. 6. Relation between intramuscular fat and linoleic acid percentages.

