Targets and procedures for altering ruminant meat and milk lipids

D. Demeyer† and M. Doreau†

1Department of Animal Production and Department of Nutrition and Food Technology, University of Gent, Proefhoestraat 10, 9090 Melle, Belgium
2Ruminant Undernutrition Laboratory, Institut National de la Recherche Agronomique, Theix, 63122 Saint-Genès-Champanelle, France

Beef and dairy products suffer from a negative health image, related to the nature of their lipid fraction. Rumen lipid metabolism involves the presence of saturated lipids in ruminant tissues. Lipolysis, fatty acid biohydrogenation and formation of microbial fatty acids in the rumen and their effects on rumen outflow of fatty acids are discussed. Special emphasis is given to the formation of trans-fatty acids and the possibilities of decreasing biohydrogenation. Small differences in intestinal digestibilities of fatty acids are mentioned, followed by a discussion on transfer of absorbed fatty acids into milk and adipose tissue lipids. The preferential retention of polyunsaturated fatty acids as well as the balance between synthesis and incorporation of fatty acids in tissues is described. Dietary means for the modification of milk fat are listed, with special emphasis on the possibilities for enrichment in polyunsaturated fatty acids and the presence of conjugated linoleic acids. A description of the nature and development of fat depots in beef cattle is followed by a discussion of breed, conformation and feed effects on adipose tissue distribution and fatty acid composition. Special emphasis is given to the very lean Belgian Blue double-muscled breed. The review ends with a consideration of the limits to the modification of ruminant fats, involving considerations of consumer acceptance as well as animal welfare and environmental effects.

Résumé

Les produits laitiers et la viande bovine souffrent d’une image santé négative, liée à la nature de leur fraction lipidique. Le métabolisme ruminal des lipides est la cause de la présence de lipides saturés dans les tissus des ruminants. La lipolyse, la biohydrogénation des acides gras et la formation d’acides gras microbiens dans le rumen, ainsi que les effets sur le flux d’acides gras à la sortie du rumen sont discutés. Une attention particulière est portée à la formation d’acides gras trans et aux possibilités de limiter la biohydrogénation. De faibles différences de digestibilité entre acides gras sont mentionnées, suivies par une discussion sur le transfert des acides gras absorbés dans le lait et les tissus adipeux. La rétention préférentielle d’acides gras polyinsaturés et le bilan entre synthèse et incorporation d’acides gras sont décrits. Les moyens nutritionnels de modifier la matière grasse du lait sont répertoriés, en particulier en vue d’accroître la sécrétion d’acides gras polyinsaturés et d’acide linoléique conjugué. Une description de la nature et du développement des tissus adipeux chez le bovin à viande est accompagnée d’une discussion des effets de la race, de la conformation et de la ration sur la distribution et la composition en acides gras des tissus adipeux. Le cas de la race blanche bleue belge à forte muscularté et très maigre est spécialement analysé. En guise de conclusion, des considérations sur les limites à la modification des lipides des ruminants font appel aux notions d’acceptabilité par le consommateur, de bien-être animal et d’effets sur l’environnement.

Abbreviations: CLA, conjugated linoleic acid; FA, fatty acid; PUFA, polyunsaturated fatty acids.
*Corresponding author: Professor D. Demeyer, fax +32 9 264 9099, email Daniel.Demeyer@mug.ac.uk
For many years, consumers in industrialized countries have displayed an aversion to dietary fat (Claus, 1991) and, for health reasons, nutritional guidelines advocate a decrease in the consumption of (saturated) fat (Gormley et al. 1987). Nevertheless, total fat intake in western Europe continues to amount to about 40% of the dietary energy intake, equivalent to 130 g/d. Milk and milk products are the major sources of fat in the diets, accounting, on average, for about 30% of the total fat and 40% of the saturated fat intake in the UK (Mansbridge & Blake, 1997), Belgium (Demeyer et al. 1995a) and Germany (Flachowsky & Jahreis, 1997). Beef is estimated to contribute only about 5% of the total fat intake, but is regarded as a rich source of saturated fat. Milk and milk products can be made more nutritionally desirable through dairy technology, and can be upgraded by the incorporation of polyunsaturated fatty acids (PUFA), through, for example, directed or random interesterification or incorporation of conjugated linoleic acid (CLA) in meat and milk. The paper, therefore, will mainly discuss nutritional ways to improve the health value of milk and beef fat composition, particularly by increasing PUFA, adjusting the n-3:n-6 PUFA value and controlling trans-fatty acids (FA) and conjugated linoleic acid (CLA) in meat and milk. The paper summarizes information from other recent reviews (Demeyer et al. 1995b; Doreau et al. 1997; Doreau & Chilliard, 1997b; Wood & Enser, 1997), but emphasizes data obtained in Belgium, a country characterized by very intensive animal production and the use of extremely lean double-muscled cattle.

**Rumen metabolism and digestibility of lipids**

**Lipolysis**

Triacylglycerols, phospholipids and galactosyl lipids in ingested forage and concentrate feeds are rapidly hydrolysed in the rumen by extracellular lipases produced by a small number of bacteria. Some activity could be associated with the protozoal fraction (Harfoot & Hazlewood, 1997). No intermediate production of partial acylglycerols has been demonstrated, and free FA form the major lipid fraction in rumen and duodenal lipids, even when high amounts of fat are fed. Lipolytic activity has been shown to be inhibited by pH values below 6 (Van Nevel & Demeyer, 1996c) and by antibiotics such as ionophores (Van Nevel & Demeyer, 1996a). From results of incubation of rumen contents with free soyabean oil (Van Nevel & Demeyer, 1996a, c), it can be calculated that on average 67 and 80% of the linoleic and linolenic acids respectively were released from triacylglycerols, whereas 92% of the released FA were recovered after incubation, suggesting some specificity of lipolysis. At increasing rates of oil infusion into the rumen, the free FA : triacylglycerol and mono- : diunsaturated octadecenoic acid values in rumen total lipids and free FA respectively suggest that lipolytic activity is not limiting the rate of FA biohydrogenation in the rumen (Imming et al. 1993).

**Biohydrogenation**

Free FA liberated by lipolysis are adsorbed onto particles, where they are both hydrogenated and/or incorporated into the lipid fraction of the solid-associated bacteria. According to Harfoot & Hazlewood (1997), the balance between these two processes is one of the factors determining the extent of FA biohydrogenation in the rumen. A range of hydrogenating bacteria have been isolated and can be divided into groups A and B. Group A hydrogenate linoleic and α-linolenic acids mainly to the trans-11-octadecenoic acid (trans-11-18:1) or vaccenic acid, with smaller amounts of other positional and stereo-isomers of the same acid. The initial step in the biohydrogenation of linoleic acid involves the isomerization of the cis-9, cis-12-isomer to the cis-9, trans-11-isomer, which is followed by a preferential reduction of the cis-9 double bond to form the trans-11–18:1. Group A bacteria appear to be incapable of hydrogenating 18:1 acids, in contrast to the members of group B which are capable of hydrogenating a wide range of both cis- and trans-isomers of 18:1 to stearic acid. The trans-18:1 isoforms normally comprise about 70% of rumen 18:1 acids, and it was shown in a continuous culture of rumen micro-organisms, using specific i.r. detection techniques, that free FA may contain up to seven trans- and six cis-18:1 acids. The trans-11 isomer represented at least 80% of the total trans-18:1 acids (Fellner et al. 1995). Trans-18:1 and conjugated octadecadienoic acids or CLA constituted 13 and 0.5% of the total FA respectively, whereas the conjugated diene isomers were shown to contain 33% of the cis-9, trans-11 isomer (Fellner et al. 1997). Biohydrogenation only occurs with free FA, but the system is easily overloaded, especially in vitro, with inhibition of the process by the free acids and accumulation of trans-18:1 and CLA. Biohydrogenation of linoleic acid in a continuous culture of rumen micro-organisms was shown to be inhibited by ionophores, with accumulation of trans-18:1 and CLA (Fellner et al. 1997). However, CLA does not arise in linolenic acid biohydrogenation (Harfoot & Hazlewood, 1997). Inhibition of biohydrogenation (5%) by ionophores and low pH was also found by Van Nevel & Demeyer (1996a, c), but was less pronounced than inhibition of...
lipolysis (20 %). The inhibition of lipolysis and/or biohydrogenation by low pH may explain the increased degree of unsaturation in rumen and duodenal lipids (Kobayashi et al. 1992) and/or in body fat in animals fed on concentrate diets or treated with monensin. As shown by Gerson et al. (1985), however, other factors in addition to pH, probably related to changes in microbial populations, must be involved with concentrate diets. Biophosphatization of polyunsaturated C₂₀ and C₂₂ FA from fish oil has been shown in vivo by Doreau & Chilliard (1997a) and in vitro by Van Nevel et al. (1999). The latter authors found a slower lipolytic release of 18:1 acids from fish oil than from soyabean oil, and a significantly lower (P<0:05) but nevertheless significant extent of hydrogenation for eicosapentanoic (20:5n-3) and docosahexanoic (22:6n-3) acids compared with other PUFA.

**Formation of microbial fatty acids**

It is now clear that both bacteria and protozoa can synthesize and/or incorporate higher FA, the synthesis possibly including, under some conditions, linoleic acid (for review, see Demeyer, 1973; Demeyer & Hoozze, 1984). Incorporation includes linoleic acid being deposited in polar lipids or free FA. When high levels of free FA are available incorporation exceeds synthesis, whereas the opposite obviously occurs in the absence of added lipids (Demeyer et al. 1978; Bauchart et al. 1990). For the animal, bacterial lipids are more important than protozoal lipids, as protozoa are largely retained in the rumen. The bacterial lipids are characterized by a high proportion of branched-chain and uneven-numbered straight-C-chain FA, synthesized from propionate and/or amino acids.

The main lipid fractions are free FA and polar lipids, and the FA composition reflects incorporation of dietary FA, as well as biohydrogenation and bacterial synthesis (Table 1). Although considerable variability is apparent from the data reported, it is clear that a mixture of liquid- and particle-associated bacteria, when leaving the rumen, contains between 100 and 200 mg lipid/g DM, its FA containing between 5 and 15 mg trans-18:1 isomers/100 mg total FA, as well as 2–20 mg PUFA/100 mg total FA, the higher values being obtained with diets comprising PUFA-rich lipid supplements. This value is not to be ignored; per kg DM intake, a diet of maize silage contains about 400 g organic matter apparently digested in the rumen, equivalent to the production of about 120 g bacterial DM, representing about 12 g newly-synthesized lipid. Since, apart from hydrogenation, all dietary lipid (58 g/kg DM) leaves the rumen mainly unaltered, it can be estimated that duodenal lipids may contain up to 17 g bacterial lipids/100 g on such diets. Similar estimates have been reported previously (Jenkins, 1994; Demeyer & Van Nevel, 1995; Mansbridge & Blake, 1997).

**Rumen outflow of fatty acids**

Compared with rumen digesta, duodenal (Kobayashi et al. 1992) and abomasal (CI Van Nevel & DI Demeyer, unpublished results) digesta are slightly enriched in PUFA, a finding possibly related to the sloughing off of enterocytes. The duodenal flow of FA is generally higher than FA intake for diets not supplemented with lipids, but often lower for fat-supplemented diets. This finding probably reflects absorption or catabolism by rumen epithelial cells, and is apparent from relationships established between duodenal flow and intake of FA (Wu et al. 1991; Doreau & Chilliard, 1997b). The intercepts of such relationships indicate that between 6 and 10 g bacterial FA/kg DM intake are synthesized, a value comparable with 12 g/kg DM intake derived previously. FA degradation by rumen microbes *in vitro* is often minimal (Demeyer & Van Nevel, 1995), but has been clearly observed in continuous culture (Fellner et al. 1995). With concentrate diets, the extent of hydrogenation of C₁₈ PUFA is high; higher for linolenic acid (50–80 %) than for linoleic (35–60 %) acid and is generally lower for high-concentrate diets than for low-concentrate diets, as indicated in the literature (Doreau et al. 1997). This finding is in line with the suggested specificity of lipolysis, and is obviously related to variations in rumen pH and in microflora, as discussed previously.

**Biohydrogenation and trans-fatty acids**

A striking feature of duodenal lipid composition is its high content of *trans*-isomers, increasing with the completeness of rumen biohydrogenation of added unsaturated oils, this factor being also determined by the balance between biohydrogenation and outflow of oils (Table 2). Contents of stereo-isomers and positional isomers of 18:1 are reported, but it is clear that *cis*, *trans*-conjugated dienoic acids must also be present in duodenal contents. Indeed, Noble et al. (1974) have shown that sheep rumen micro-organisms

### Table 1. Fatty acid (FA) composition (mg/100 mg) of rumen bacterial lipids

<table>
<thead>
<tr>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Crude fat or total FA (mg/g DM)</td>
<td>50–90</td>
<td>66–81</td>
<td>98–237</td>
<td>146–163</td>
<td>60–214</td>
</tr>
<tr>
<td>SFA</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Branched-chain</td>
<td>12·0</td>
<td>5·3–11·4</td>
<td>11·4–17·6</td>
<td>–</td>
<td>3·0–14·7</td>
</tr>
<tr>
<td>Short-chain</td>
<td>73·0</td>
<td>51·8–54·6</td>
<td>61·2–72·2</td>
<td>65·2–67·8</td>
<td>39·8–64·7</td>
</tr>
<tr>
<td><em>trans</em>-Isomers</td>
<td>4·4–5·1</td>
<td>3·2–18·7*</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>MUFA</td>
<td>4·0</td>
<td>8·4–13·2</td>
<td>–</td>
<td>22·9–25·8</td>
<td>12·5–35·2</td>
</tr>
<tr>
<td>PUFA</td>
<td>4·0</td>
<td>1·3–0·8</td>
<td>3·9</td>
<td>5·6–5·5</td>
<td>6·3–19·9</td>
</tr>
</tbody>
</table>

* Mainly comprising the neutral-lipid fraction. SFA, saturated FA; MUFA, monounsaturated FA; PUFA, polyunsaturated FA.
produce trans-11-18:1 together with cis-9, trans-11-18:2 in vitro, and that the concentrations of both compounds reflect a product–precursor relationship when incubated with oils but not when incubated with free PUFA. Surprisingly, the feeding of high-oleate sunflower oil significantly increased the trans-18:1 content in total duodenal FA, although trans-isomers are not considered intermediates in the hydrogenation of oleic acid (cis-9-18:1). The trans-11:18:1 isomer makes up at least 80 % of the trans-18:1 isomers in accordance with the hydrogenation pathways established for rumen bacteria.

Techniques for decreasing biohydrogenation

A natural protection against hydrogenation is given by the coat of some oilseeds, but this protection is limited. Moreover, extruding oilseeds does not improve protection. Thus, for more than two decades, physical and chemical techniques for the protection of feedstuffs against hydrogenation have been evaluated. The first method to be developed was the coating of emulsified oils with formaldehyde-treated proteins. This technique is for the moment the most efficient method of increasing the amount of PUFA absorbed (Gulati et al. 1992; Van Nevel & Demeyer, 1996). Another form of physical treatment leads to fatty acyl amides, which partially resist hydrogenation. Techniques for the protection of feedstuffs against hydrogenation include the different isomers, whereas the low amounts of 18:3 may result in low accuracy of the estimates.

Absorption and incorporation into milk and adipose tissue lipids

Transport

The epithelial cells are the site for esterification into triacylglycerols and phospholipids, which are transported into the lymph as chylomicrons and VLDL, and further into the blood where these lipoproteins are found together with LDL and HDL. LDL, VLDL and chylomicrons contain approximately 10–15, 60 and 85 mg triacylglycerols/100 g respectively. Most preformed FA are delivered to the mammary gland by LDL and VLDL. Although HDL account for approximately 90 % of blood lipids they consist largely of phospholipids, cholesterol and cholesteryl esters, containing the major proportion of PUFA (Mansbridge & Blake, 1997). Lipoproteins transport FA mainly to the mammary gland in dairy cattle and mainly to adipose and muscle tissue in fattening animals.

Uptake by adipose tissue and by the mammary gland

Uptake of HDL-FA by the mammary gland is poor, which may explain the low PUFA content of milk fat. During passage into these tissues, triacylglycerols are largely or completely hydrolysed by a lipoprotein lipase (EC 3.1.1.34). There is little further modification of preformed FA within these tissues, except for mild to extensive desaturation of medium- and long-chain FA, particularly 18:0, by a Δ9-desaturase. The result is a lower 18:0/18:1 value in these tissues compared with duodenal digesta, a mechanism thought to ensure fluidity of tissue and milk. Adipocytes and

### Table 2. trans-Octadecenoic acid (trans-18:1) content of duodenal fatty acids (FA) in relation to oil feeding and biohydrogenation

<table>
<thead>
<tr>
<th>Reference</th>
<th>Dietary treatment</th>
<th>trans-18:1 (mg/100 mg total FA)</th>
<th>Biohydrogenation (%)*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Wu et al. (1991)</td>
<td>Diet: Alone</td>
<td>4-2</td>
<td>67-3</td>
</tr>
<tr>
<td></td>
<td>+ Ca salt palm oil (30 g/kg)</td>
<td>3-3</td>
<td>58-3</td>
</tr>
<tr>
<td></td>
<td>+ Ca salt palm oil (60 g/kg)</td>
<td>2-5</td>
<td>53-0</td>
</tr>
<tr>
<td>Enjalbert et al.</td>
<td>Diet: + soybean oil (SO; 57 g/kg)</td>
<td>37-0</td>
<td>89-5</td>
</tr>
<tr>
<td></td>
<td>+ emulsified SO (57 g/kg)</td>
<td>34-9</td>
<td>90-9</td>
</tr>
<tr>
<td></td>
<td>+ Ca salts SO (67 g/kg)</td>
<td>11-5</td>
<td>64-3</td>
</tr>
<tr>
<td>Kalscheur et al.</td>
<td>Diet: Alone</td>
<td>6-7</td>
<td>71-1</td>
</tr>
<tr>
<td></td>
<td>+ 18:1 sunflower oil (37 g/kg)</td>
<td>18-5</td>
<td>79-5</td>
</tr>
<tr>
<td></td>
<td>+ 18:2 sunflower oil (37 g/kg)</td>
<td>20-1</td>
<td>80-1</td>
</tr>
</tbody>
</table>

* Calculated as 100 – 100 (C18 unsaturated FA in duodenum/C18 unsaturated FA in diet), according to Wu et al. (1991).
mammary gland tissue incorporate chylomicron-derived FA into triacylglycerols, using glycerol formed from glucose. The specificity of FA distribution within the triacylglycerols is determined during the esterification of glycerol-3-phosphate with two FA, followed by the incorporation of a third FA into the dephosphorylated 1,2-diacyl-3-phosphate. Short-chain FA in milk triacylglycerols are, for example, esterified specifically at the 3 position. The FA distribution in adipose triacylglycerols conforms to the general rules formulated by Brockerhoff (1966) for mammalian depot triacylglycerols, i.e. saturated FA predominantly in the position sn-1, shorter-chain and unsaturated FA largely in position sn-2 and C_{18} and longer-chain FA in position sn-3. The branched-chain FA are mainly found in position sn-2, whereas the trans-18 : 1 isomers are distributed in a different manner from the cis-isomers (sn-2 position) and are mainly found in the sn-3 position (Christie, 1981). Phospholipids are formed through reactions between diacylglycerols and cytidine phosphate components, involving specificity of the acyltransferases involved.

**Synthesis de novo**

Biosynthesis of adipose and milk lipids uses diet-derived FA as well as FA synthesized de novo from acetate. The effect of the former is limited, however, as de novo synthesis and direct incorporation are regulated and balanced in relation to optimal fluidity of both adipocyte and milk lipids and membranes. Adipose tissue is the major site of FA synthesis in ruminants, except during lactation when the mammary gland becomes the predominant site. Synthesis of FA up to palmitic acid takes place in the cytoplasm from acetyl-CoA and β-hydroxybutyrate derived from mitochondrial oxidation. Mitochondria elongate palmitic acid to longer FA up to C_{22}, whereas microsomes are capable of elongation as well as desaturation of FA ≥ C_{18}. Ruminants, as all mammals, in contrast to plants, are incapable however of the introduction of double bonds at C-9, counted from the carboxyl group and the methyl end of the chain (Gurr & James, 1980). In the mammary gland, de novo synthesis is limited to short- and medium-chain FA (up to 16 : 0) that can be esterified only into the sn-2 and sn-3 positions of the glycerol backbone, thus the rate of FA synthesis in the mammary gland is dependent on the simultaneous supply of exogenous FA. Fatty acyl-CoA esters inhibit acetyl-CoA carboxylase (EC 6.4.1.2) in adipose tissue and mammary gland, and it has been shown with dispersed epithelial cells of both bovine and goat mammary glands that FA synthesized from acetate (C_{4}-C_{8}) were incorporated into triacylglycerols, depending on the type and amount of exogenous FA added; palmitic acid stimulated synthesis and incorporation, whereas stearic and linoleic acids were inhibitory (Hansen & Knudsen, 1987).

**Turnover and the balance between synthesis and incorporation**

As with all living tissues, the amounts of adipocyte lipids are the net result of both synthesis and degradation. Milk lipids, however, are secreted and thus subject to little turnover, this difference explaining the higher energetic efficiency of milk v. body tissue production. Turnover time of adipose tissue lipids varies with the level of feeding, physical activity and lactation, and may be up to hundreds of days. FA are liberated from adipose tissue for transportation as albumin-bound non-esterified FA to tissues requiring them for oxidative energy. Complex regulation of lipolytic activity occurs by both short- and long-term mechanisms, the former involving the action of a hormone-sensitive lipase, activated by adrenaline and inhibited by insulin. In the mammary gland, de novo synthesis is responsible for practically all FA up to C_{18} (short-chain FA) and for about half the medium-chain FA (12 : 0–16 : 0). The other half of the medium-chain FA and practically all long-chain FA (18 : 0 and longer) are derived from blood FA originating from the diet or from mobilization of body fat stores (Baer, 1996). About 30 % of milk fat is derived from acetate and perhaps 5 % from β-hydroxybutyrate (Freden, 1996).

**Preferential retention of polyunsaturated fatty acids**

Ruminants attempt to specifically retain the small amounts of PUFAs that escape hydrogenation by preferentially incorporating them into plasma cholesteryl esters via the lecithin-cholesterol acyl transferase (EC 2.3.1.43) pathway. It is also clear that they selectively incorporate polyenoic acids into membrane phospholipids. However, the ruminant has probably developed other mechanisms for the preferential retention of PUFAs. Vernon & Flint (1988) suggested limited oxidation of linoleic acid in ruminants, based on the finding that malonyl-CoA is much more effective in inhibiting carnitine acyltransferase when linoleyl-CoA is the substrate than when palmitoyl-CoA is the substrate in sheep liver, whereas the converse occurs in rat liver. Nevertheless, because of extensive rumen biohydrogenation, it has been suggested that ruminants can be delicately balanced in respect of their essential FA status (Ashes et al. 1995).

**Dietary methods for modifying milk fat**

Milk fat consists predominantly of triacylglycerol (98 mg/100 mg) with phospholipids and sterols comprising less than 1 and 0.5 mg/100 mg total lipid respectively. Over 500 FA are found esterified in the triacylglycerols. Their composition reflects the nature of dietary lipids, as well as the effects of rumen FA metabolism and FA synthesis in intestinal, mammary and adipose tissue. Saturated FA reaching the mammary gland are further desaturated by mammary desaturase activity, resulting in a lower 18 : 0/18 : 1 value in milk v. duodenal digesta, a mechanism used by the ruminant to preserve the fluidity of milk. The predominant FA in milk are the short- and medium-chain saturated FA (mainly 12 : 0, 14 : 0 and 16 : 0), accounting for about 50 mg/100 mg total FA, and the long-chain FA (mainly 18 : 0 and 18 : 1) for about 45 mg/100 mg total FA and including small amounts of PUFAs (2–3 mg, 18 : 2 and 18 : 3/100 mg total FA). The remainder consists of branched-chain and uneven-numbered saturated FA derived from bacterial lipids, as well as a series of trans- and positional isomers of 18 : 1 and 18 : 2. As discussed earlier, the distribution of FA over the three glycerol-C is not
random and triacylglycerols from C_{14} to C_{34} are present (Kennelly, 1996a; Mansbridge & Blake, 1997).

Such FA composition does not suggest a 'healthy' food, in view of the demonstrated serum LDL-raising properties of both saturated short- and medium-chain FA (Nicolisi, 1997) and of trans-unsaturated FA (Ip & Marshall, 1996). Furthermore, milk fat consumption cannot contribute to an improvement in the PUFA intake and the n-3:n-6 value in human nutrition, as recommended in a number of dietary guidelines (Truswell, 1995). The proposal of an 'ideal' nutritional fat by the Wisconsin Milk Marketing Board, enriched in monounsaturated FA (≥82 %) and low in saturated FA (<8 %; cited, for example, by Baer, 1996), and the new economical conditions of dairy production, especially in the EU, have resulted in pressure for a limitation of the production of milk fat and optimization of its FA composition.

Indeed, a number of countries outside and inside the EU value protein higher than fat with respect to milk payments, in line with the significant worldwide decline in the demand for milk fat, both as a result of substitution of vegetable spreads for butter and a consumer preference for milk with a reduced fat content (Kennelly, 1996a; Murphy & O’Mara, 1993). In contrast to milk fat content, milk fatty acid composition only shows nutritionally insignificant differences between cattle breeds (DePeters et al. 1995). Nutrition, and in particular fat supplementation, may alter milk fatty acid composition towards that advised in nutritional guidelines.

### Table 3. Effect of dietary unsaturated fat addition on milk fat yield and fatty acid (FA) composition (mg/100 g FA)

<table>
<thead>
<tr>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>P</td>
<td>FFS</td>
<td>GSB</td>
</tr>
<tr>
<td>4:0</td>
<td>2.5</td>
<td>2.5</td>
<td>3.3</td>
</tr>
<tr>
<td>6:0</td>
<td>1.9</td>
<td>1.6</td>
<td>2.5</td>
</tr>
<tr>
<td>8:0</td>
<td>1.2</td>
<td>0.9</td>
<td>1.6</td>
</tr>
<tr>
<td>10:0</td>
<td>2.6</td>
<td>1.8</td>
<td>4.1</td>
</tr>
<tr>
<td>12:0</td>
<td>3.2</td>
<td>2.0</td>
<td>3.7</td>
</tr>
<tr>
<td>14:0</td>
<td>11.1</td>
<td>7.4</td>
<td>10.3</td>
</tr>
<tr>
<td>14:1</td>
<td>1.9</td>
<td>1.2</td>
<td>0.8</td>
</tr>
<tr>
<td>16:0</td>
<td>23.8</td>
<td>19.7</td>
<td>23.8</td>
</tr>
<tr>
<td>16:1</td>
<td>2.5</td>
<td>2.2</td>
<td>1.8</td>
</tr>
<tr>
<td>SMCFAs</td>
<td>50.7</td>
<td>39.3</td>
<td>51.9</td>
</tr>
<tr>
<td>18:0</td>
<td>11.1</td>
<td>14.2</td>
<td>15.4</td>
</tr>
<tr>
<td>18:1, trans-11</td>
<td>2.3</td>
<td>3.6</td>
<td>–</td>
</tr>
<tr>
<td>18:1, cis-9</td>
<td>28.5</td>
<td>35.1</td>
<td>20.6</td>
</tr>
<tr>
<td>18:2</td>
<td>1.8</td>
<td>4.8</td>
<td>5.3</td>
</tr>
<tr>
<td>18:3</td>
<td>0.6</td>
<td>0.9</td>
<td>1.2</td>
</tr>
<tr>
<td>20:0</td>
<td>–</td>
<td>–</td>
<td>0.4</td>
</tr>
<tr>
<td>20:5</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>22:6</td>
<td>24.0</td>
<td>27.0</td>
<td>23.0</td>
</tr>
<tr>
<td>LCFA</td>
<td>42.0</td>
<td>55.0</td>
<td>45.2</td>
</tr>
</tbody>
</table>

SMCFA, short- and medium-chain FA (up to 16 : 0); LCFA, long-chain FA (from C_{18} on); P, pasture; FFS, toasted full-fat soybeans; GSB, RSB, TSB, ground, roasted and toasted soybeans respectively; MS + C, maize silage and concentrate-based diet; S + C, silage and concentrate-based diet; FO, fish oil.

* Holstein cows were fed on 3·2 kg FFS/d.
† Holstein cows were fed on a mixed ration of maize and silage with GSB or RSB at 175 g/kg DM in the total ration.
‡ Cows were fed on MS + C with or without TSB (85 g/kg DM in the total ration).
§ Also, Y Chilliard, JM Chardigny, J Chabrot, A Ollier, J-L Sebedio & M Doreau (unpublished results). Holstein cows were fed on S + C with or without FO (300 ml).

**Effect of dietary fat**

Fat supplements have been shown to influence more than the reduction in the supply of precursors for fatty acid synthesis in the mammary gland. The following effects have been documented:

(a) an inhibition of FA synthesis through inhibition of acetyl-CoA carboxylase in both mammary and adipose tissue (Vernon & Flint, 1988);
(b) an inefficient incorporation of PUFA, and particularly of the fish oil C_{20} and C_{22} FA and the trans-isomers derived from rumen biohydrogenation, into the 1-monoyacylglycerols required as acceptors for the newly-synthesized FA (Hansen & Knudsen, 1987);
(c) an inefficient incorporation of trans- v. cis-FA into small and large triacylglycerols, not mediated at the level of lipoprotein lipase activity (Gaynor et al. 1994);
(d) a specific reduction in insulin-dependent amino acid v. non-insulin-dependent glucose and FA uptake by the mammary gland, due to an induction of insulin resistance by FA, leading to a reduction in milk protein content (Palmquist & Moser, 1981). This reduction is brought about to the same extent by both cis- and trans-18 : 1 isomers (Gaynor et al. 1996).

Table 3 shows data illustrating a 23 % decrease in the short- and medium-chain FA content with a corresponding 30 % increase in the long-chain FA content as the result of supplementation with oil seeds. The latter increase mainly involves oleic acid. These effects are intensified by heat.
treatment of the soybeans (Chouinard et al. 1997). However, experiments in vitro indicated inhibition of biohydrogenation by roasting soybeans (Reddy et al. 1994). In contrast with soybean oil, fish oil supply modifies the proportion of short- and medium-chain FA to a small extent, suggesting that inhibition of FA incorporation and inhibition of de novo synthesis are both associated with the decrease in fat content.

Enrichment in linoleic acid and n-3 polyunsaturated fatty acids

Until now, the only way to increase the linoleic acid content in milk has been to use oils protected by protein coating (Kennelly, 1996b). Unprotected oils or seeds and Ca salts of PUFA do not give a linoleic acid content in milk higher than 5 mg/100 mg total FA (Doreau et al. 1997). The positive influence of n-3 PUFA on cardiovascular diseases (Kinsella et al. 1990) suggests it would be of benefit to increase such FA in milk.

Table 3 also illustrates the low transfer efficiency of 20 : 5n-3 (9.3 %) and 22 : 6n-3 (16.2 %) into milk-fat triacylglycerols. This finding contrasts with the values of 35–40 % obtained after post-rumen infusion (Hagemeister et al. 1988). A probable explanation relates to the partial, but significant, hydrogenation of the fish oil FA, demonstrated recently both in vivo and in vitro, as mentioned previously. Obviously, further research is needed to clarify the contradiction with negative findings in this respect (Ashes et al. 1992), as well as the surprising stimulatory effect of fish oil on rumen fibre digestion reported by Doreau & Chilliard (1997a). As described for linoleic acid, an enrichment of linolenic acid in milk is possible using efficiently protected linseed, whereas the intake of unprotected linseed generally does not result in an increase in milk linolenic acid (Kennelly, 1996b).

Trans-fatty acids

An increase in the 18 : 1 content of milk fat is associated with an increase in trans-18 : 1 isomer content, a change considered of importance in human nutrition (Wolff, 1995). Indeed, human consumption of trans-18 : 1 isomers has been shown to be related to the incidence of CHD (Willett et al. 1993), and dietary studies with human subjects have revealed their raising and lowering effects on LDL- and HDL-cholesterol respectively (Zock & Katan, 1997). Dietary guidelines often stipulate that trans-FA intake should not increase (Truswell, 1995). In line with this situation, trans-FA content in margarines has been shown to have decreased in recent years (Precht & Molkentin, 1997; Zock & Katan, 1997). Using combined argentation TLC and GC, it was recently established, however, that trans-18 : 1 isomers in milk fat predominantly comprise the trans-11 positional isomer, in line with the composition of rumen and duodenal contents (Precht & Molkentin, 1995), although the trans-11 isomer represents more than 80 mol/100 mol trans-monoenes in rumen lipids but only 40–60 mol/100 mol trans-monoenes in milk fat (Griiniari et al. 1997). In partially-hydrogenated oils, however, amounts of the trans-9, -10 and -11 isomers are almost equal. Also, evidence is available suggesting that the trans-FA present in milk and beef may not have the same negative characteristics as those present in partially-hydrogenated vegetable and fish oils (Willett et al. 1993; Zock & Katan, 1997).

The highest levels of trans-18 : 1 in milk are found with diets containing fish oil and oils rich in linoleic acid, which are biohydrogenated in the rumen, and diets containing partially-hydrogenated oils which contain different trans-18 : 1 isomers. Surprisingly, Kalscheur et al. (1997) have observed an increase in trans-18 : 1 after feeding high-oleic sunflower oil, and attribute this effect to a sufficient amount of dietary linoleic acid available for biohydrogenation. The response of trans-18 : 1 content in milk to increasing levels of dietary linoleic acid has not been established. It can be hypothesized, however, that combining a high-concentrate diet with a supply of fish oil or oils rich in linoleic acid may enhance the production of trans-18 : 1 due to a slower rate of hydrogenation in the rumen.

Trans-isomers of 18 : 1 are considered to have a major role in the mechanisms for milk fat reduction described earlier, and there is a patent describing their use for the regulation of milk fat content and milk yield (Erdman et al. 1995). However, their concentration in milk fat does not always indicate a reduction in milk fat (Kalscheur et al. 1997). Recent work, summarized by Griiniari et al. (1997), suggests that the production in the rumen of the trans-10-18 : 1 isomer and its transfer to milk lipid is specifically responsible for the reduction in milk fat. The isomer would not be prevalent in the trans-18 : 1 mixture present in partly-hydrogenated fats. The occurrence of this isomer would be specifically associated with concentrate feeding, but not with roughage feeding.

Increasing the conjugated linoleic acid content of milk

The potent anti-carcinogenic properties of CLA have been widely shown using experimental animals and in epidemiological studies (Belury, 1995; McGuire et al. 1997). Among human foods, dairy products and beef are especially rich in CLA (Kelly & Bauman, 1996). This specificity can be used to promote ruminant products, or at least limit the decrease in their consumption. Moreover, compared with milk, dairy products may have a higher content of CLA; processing may involve oxidation with free radical formation, followed by a reduction, yielding an increased CLA concentration (Lin et al. 1995). Four isomers have been found: cis-9-trans-11, cis-8-trans-10, cis-11-trans-13, cis-10-trans-12, the first isomer being quantitatively the most important. The comparative significance of these isomers for health remains to be established. Abomasal infusion of CLA caused an increase in both cis- and trans-isomers in milk CLA, showing a digestive origin (Chouinard et al. 1998b). However, a possible synthesis in the mammary gland from trans-11 by the Δ9-desaturase has been shown (Corl et al. 1998).

With winter diets not supplemented with lipids, the CLA content of milk is less than 1 mg/100 mg total FA. Pasture significantly increases the CLA content. However, the more marked effect is associated with supplementary FA (Table 4). From these findings it is clear that (1) saturated and monounsaturated FA do not increase CLA; (2) the maximal
increase in CLA is obtained with free oils rich in linoleic acid or salts of their FA, and in oils rich in 20 : 5 and 22 : 6; (3) a lower increase is obtained with oils rich in linolenic acid. A positive relationship between trans-18 : 1 and CLA was found by Lin et al. (1995) and Jiang et al. (1996). This finding is consistent with the highest values found with oils rich in linolenic acid and with fish oil, both of which induce large amounts of trans-18 : 1 FA. This finding may also explain why the effect of trans-18 : 1 is not always positively related to an increase in cardiovascular diseases (Wolff, 1995). However, it has been hypothesized that CLA may decrease butterfat content, independently of the positive relationship with trans-18 : 1 (Griinari et al. 1997).

Control of adipose and muscle tissue lipids

Nature and development of fat depots

In ruminants, lipids are mainly accumulated as triacylglycerols in adipocytes, located in subcutaneous, inter- and intramuscular adipose tissue, and abdominal, e.g. perirenal and omental fat depots. Total dissecatable fat in a beef carcass varies approximately between 100 and 400 g/kg, about half of which is subcutaneous and intramuscular fat. With increasing animal weight, the proportion of subcutaneous fat increases more than that of intramuscular tissue, i.e. it has a higher relative (allometric) growth rate. Different regions of subcutaneous fat also grow at different rates, and the rate of increase in intramuscular lipid is less than that in dissecatable fat (Wood, 1984). Young fat tissue has a high proportion of water and connective tissue and a low proportion of lipid, contained in small cells. When the animal gets older, fat depots, especially internal depots, increase mainly through an increase in cell size, with a decrease in the proportion of water and connective tissue in the tissue and a colour change from grey to more white. The fat cell diameter in adult animals may vary between 100 and 250 µm, indicating more than an eightfold difference in storage capacity (Lawrence & Fowler, 1997). Female animals deposit more fat than males, and castration increases fat deposition so that there is a great emphasis on the production of entire males, especially in southern and western Europe.

Triacylglycerols in intramuscular adipocytes, associated with the epi- and endomysium, determine the ‘marbling’ of meat, which increases with the age of the animal. Lipid is also present, as intracellular lipid droplets, in muscle fibres, and at higher concentrations in red muscle than in white muscle. In addition to triacylglycerols in adipocytes and in oil droplets, intramuscular lipid also comprises the membrane phospholipids of the muscle fibres. Whereas extraction of the triacylglycerols can be done using non-polar solvents such as diethyl ether, extraction of total lipids, including membrane phospholipids, requires the use of more polar solvents such as chloroform–methanol. Mean values calculated from data reported by Christie (1981), Larick & Turner (1989), Ashes et al. (1995) and Gandemer (1997) show that FA in phosphatidyl choline and phosphatidyl ethanolamine make up 48·1 (SE 0·7) and 25·3 (SE 2·0) mg/100 mg total membrane phospholipid-bound FA respectively; 20–30 % in each fraction being recovered as plasmalogens. It is claimed that total intramuscular lipid content is very important for sensory evaluation of beef, especially of texture and flavour. However, meat preparation and/or cultural differences have a considerable effect on sensory evaluation, since minimal intramuscular fat contents for optimal sensory quality vary between 15 and 30 g/kg for Denmark and the US respectively (Wood, 1990). With decreasing total intramuscular lipid content, the relative importance of phospholipids in total lipids increases. Indeed, as adipocyte size is lowered, or less adipocytes are present, amounts of muscle fibre membranes remain relatively constant.

Breed and conformation effects on adipose tissue distribution

There are important breed differences in the proportions of body fat in the various depots. Breeds noted for beef production have a higher value for dissecatable intramuscular fat than breeds noted for milk production, as shown in Table 5,

### Table 4. Effect of dietary lipids on concentration of conjugated linoleic acid (CLA) in milk (mg/100 mg total fatty acids)

<table>
<thead>
<tr>
<th>Reference</th>
<th>CLA &lt; 0·8</th>
<th>CLA 0·8–1·6</th>
<th>CLA &gt; 1·6</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dhiman et al. (1997)</td>
<td>Raw and roasted cracked soyabeans</td>
<td></td>
<td>Soyabean oil</td>
</tr>
<tr>
<td>Kelly et al. (1997)</td>
<td>Peanut oil</td>
<td>Linseed oil</td>
<td>Sunflower oil</td>
</tr>
<tr>
<td>Chilliard et al. (1999)</td>
<td>Untreated, roasted and micronized soyabeans</td>
<td></td>
<td>Fish oil</td>
</tr>
<tr>
<td>Chouinard et al. (1998a)</td>
<td>High-oil maize</td>
<td>Extruded soyabean</td>
<td>Soyabean Ca salts</td>
</tr>
<tr>
<td>Dhiman et al. (1998)</td>
<td>Tallow</td>
<td>Rapeseed Ca salts</td>
<td>Linseed Ca salts</td>
</tr>
<tr>
<td>Grinari et al. (1998)</td>
<td>Extruded soyabean</td>
<td></td>
<td>Fish oil</td>
</tr>
<tr>
<td></td>
<td>Extruded cottonseeds</td>
<td></td>
<td>Sunflower oil</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Rapeseed Ca salts</td>
</tr>
</tbody>
</table>
also illustrating the low intramuscular fat content of double-muscled Belgian Blue cattle compared with Holstein cattle.

Furthermore, the subcutaneous fat: dissectable fat value increases with meat yield. For the same degree of subjective EU carcass fatness classification (e.g. 2; EEC, 1981, 1991), based to a large extent on visual inspection of the subcutaneous fat cover, the carcass dissectable fat content, estimated by dissection of a one-rib cut, decreased with increasing conformation, the EU classification of muscularity (EEC, 1981, 1991), from 200 to 123 g/kg. This finding suggests that the relative importance of subcutaneous tissue in total adipose tissue increases with increasing muscularity (G Van de Voorde, S De Smet and DI Demeyer, unpublished results).

**Fatty acid composition**

Beef lipid composition, as well as that of milk fat, reflects rumen metabolism of dietary lipids. Relative amounts of ‘unusual’ odd-chain, branched-chain and trans-monoenoic acids of 3-3, 2-7 and 4-2 mol/100 mol total FA respectively have been reported. The multi-branched FA pristanic and phytanic acids are present, whereas the trans-18:1 series is dominated by the trans-11 isomer (Christie, 1981). Beef meat and beef tallow display slightly higher relative proportions of trans-10- and trans-11-18:1 (approximately 65 mol/100 mol total trans-18:1 isomers, which represent 4-9 (SD 0-9) mol/100 mol total FA) compared with butterfat (55–60 mol/100 mol total trans-18:1 isomers), the opposite being found for trans-13- and trans-16-isomers. This finding would indicate that some subtle differences exist in the metabolic selectivity for individual trans-18:1 acids between the mammary gland and muscles or adipose tissue (Bayard & Wolff, 1996). In earlier work, the same group reported higher trans-18:1 contents in total FA for beef tallow (4-6 mol/100 mol) than for meat (1-95 mol/100 mol). Intramuscular trans-18:1 was concentrated in triacylglycerol-FA (2-53 mol/100 mol), and much less in polar lipid FA (0-76 mol/100 mol; Wolff, 1995).

Saturated and unsaturated aldehydes, possibly derived metabolically from FA, have been detected in adipose tissue from ruminants, and some of these may be important in flavour determination of meat and meat products (Christie, 1981). It is generally recognized that beef fat has a more saturated nature than pork and poultry fat, because of the biohydrogenation processes in the rumen (Demeyer et al. 1995a). This statement cannot be generalized, however. There are, for example, clear indications of specific deposition and/or metabolism (desaturation) of absorbed FA in the various triacylglycerol depots. As in all large animals, internal fat depots in mature animals are characterized by high proportions of saturated FA. Saturation, characterized by the 18:0/18:1 value, decreases with decreasing distance from the animal’s exterior. Effects of temperature differences between these sites on desaturase activity are held partly responsible for this finding. In contrast to pigs, the concentration of saturated acids in fat depots decreases with age and body fat content in cattle (Wood, 1984). This finding seems to be limited to domesticated cattle, however, as adipose tissue triacylglycerols of lean African wild ruminants were found to contain higher proportions of unsaturated FA, possibly because of dietary differences (Sinclair & O’Dea, 1990). Effects of breed are illustrated by the more saturated character of Hereford cow subcutaneous adipose tissue compared with that of Brahman cows (Huerta-Leidenz et al. 1993). The effect of site is illustrated for double-muscled animals in Table 6.

The data in Table 6 also illustrate that the triacylglycerol fraction of intramuscular lipids shows a FA composition similar to that of subcutaneous fat. The phospholipid fraction on the other hand is probably fairly representative of the membrane structure and is characterized by high levels of PUFA, particularly linoleic and arachidonic acids. In fact, skeletal muscle appears to be a major repository for these essential FA in ruminants (Christie, 1981). Concentrations of total FA in the M. longissimus thoracis increased with increasing carcass fat content (r 0-41), the increase being limited to the triacylglycerol fraction, a finding also illustrating the distinctive nature of the polar lipid fraction. The concentration of FA in the latter amounts to 3-1 mg/g wet tissue, a value similar to those reported by Larick & Turner (1989) and Gandemer (1997). The PUFA

### Table 5. Carcass and intramuscular fat content of Holstein, double-muscled (DM) and dual-purpose (M) Belgian Blue (BB) cattle fed on different diets

<table>
<thead>
<tr>
<th>Reference</th>
<th>Animals and diets</th>
<th>Dissectable fat in carcass (g/kg)</th>
<th>Intramuscular fat in longissimus thoracis* (g/kg)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fiems et al. (1998)</td>
<td>DM BB bulls of about 700 kg</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>High-energy</td>
<td>131</td>
<td>9.3</td>
</tr>
<tr>
<td></td>
<td>Low-energy</td>
<td>123</td>
<td>8.6</td>
</tr>
<tr>
<td>Minet et al. (1996)</td>
<td>Bulls of about 500–550 kg</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Holstein</td>
<td>252</td>
<td>46.4</td>
</tr>
<tr>
<td></td>
<td>MBB</td>
<td>242</td>
<td>22.0</td>
</tr>
<tr>
<td></td>
<td>DM BB</td>
<td>116</td>
<td>7.5</td>
</tr>
<tr>
<td></td>
<td>DM BB bulls of about 600 kg</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Control: Alone</td>
<td>134</td>
<td>6.8</td>
</tr>
<tr>
<td></td>
<td>+ Milk fat</td>
<td>137</td>
<td>8.8</td>
</tr>
<tr>
<td></td>
<td>+ Vegetable fat</td>
<td>149</td>
<td>11.2</td>
</tr>
<tr>
<td>Webb et al. (1998)</td>
<td>DM BB cows of about 700 kg</td>
<td>126</td>
<td>11.2</td>
</tr>
</tbody>
</table>

* Extracted with chloroform–methanol in Webb et al. (1998); determined by near i.r. reflectance spectroscopy in Fiems et al. (1998); procedure not mentioned in Minet et al. (1996).
Table 6. Fatty acid (FA) composition (mg/100 mg FA) of muscle and adipose tissue lipids of double-muscled Belgian Blue cows (Webb et al. 1998)

<table>
<thead>
<tr>
<th>FA</th>
<th>Intramuscular fat</th>
<th>Intermuscular fat</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>PL (3-1)*</td>
<td>TAG (8-1)*</td>
</tr>
<tr>
<td>14:0</td>
<td>0·25</td>
<td>2·15</td>
</tr>
<tr>
<td>16:0</td>
<td>14·61</td>
<td>27·19</td>
</tr>
<tr>
<td>16:1</td>
<td>2·87</td>
<td>3·35</td>
</tr>
<tr>
<td>18:0</td>
<td>12·58</td>
<td>16·12</td>
</tr>
<tr>
<td>18:1</td>
<td>19·11</td>
<td>43·10</td>
</tr>
<tr>
<td>18:2</td>
<td>23·10</td>
<td>2·44</td>
</tr>
<tr>
<td>18:3</td>
<td>3·57</td>
<td>0·63</td>
</tr>
<tr>
<td>20:4</td>
<td>5·88</td>
<td>0·00</td>
</tr>
<tr>
<td>Other</td>
<td>18·03</td>
<td>5·02</td>
</tr>
</tbody>
</table>

*M g FA/g wet tissue; PL, polar lipids; TAG, triacylglycerols.

content in the polar lipids (32·6 mg/100 mg) is similar to recent values reported for beef (21.5–38·7 mg/100 mg), as well as for pork (29·9–38·7 mg/100 mg; Demeyer et al. 1995a).

Because of the low content of total intramuscular lipid, its FA composition is determined to a large extent by that of the polar lipid fraction and is similar to that of pork intramuscular lipids, characterized by high and low contents of PUFA and trans-FA respectively, an important finding from a nutritional point of view. In summary, dissectable fat of lean animals is more saturated than that of fat animals, whereas the opposite is true for intramuscular fat.

Effects of fat feeding

Unlike milk production, lean beef production is not limited by energy intake. However, fats may be cheaper alternatives in feed formulation, and their use allows modification of beef FA composition, providing other incentives for the increasing practice of fat supplementation in beef production. From a literature survey, Clinquart et al. (1995) conclude that, in general, fat addition to a finishing diet results in improved weight gain at lower intakes associated with a 5–15 % higher fat content of the carcasses. At incorporation levels higher than 5 %, however, growth may be impaired, possibly related to impairments in both digestion and intake; the authors cite two non-published experiments from their laboratory, however, in which decreases in carcass fat content of up to 28 % were observed. This finding might be explained by inhibitions of de novo FA synthesis, as reported with dairy cattle. From their own results, summarized in Table 7, it is clear that unsaturated oils provoke effects comparable with those observed in milk fat: a decrease in short- and medium-chain FA content, and an increase in long-chain FA, mainly 18:1. These effects are less pronounced in intramuscular lipids and with heat-treated beans vs. free oils. With free oil 18:2 is preferentially retained in intramuscular lipids, again reflecting the separate metabolic importance of muscle polar lipids. Heat treatment of soyabean seems to protect 18:2 from biohydrogenation, as discussed earlier. As with dairy cattle, evidence is available suggesting that fat supplementation induces effects beyond changes in the proportion of dietary and newly-synthesized FA as substrates for fat synthesis. Van Eenaeme et al. (1991) have shown, for example, that fat supplementation increased plasma insulin levels with a reduction in blood glucose levels.

It is more difficult to justify dietary interventions to change the FA composition of beef fat than it is for milk fat. Indeed, beef dissectable fat is not consumed directly and is used as tallow in food and non-food industries. Furthermore, intramuscular FA composition has been shown to display effects comparable with those observed in milk fat: a decrease in short- and medium-chain FA, and with heat-treated beans a decrease in 18:2.

Table 7. Percentage change in fatty acid (FA) composition of perirenal and intramuscular (longissimus thoracis) fat by dietary addition of soyabean oil (33 g/kg) or heat-treated soyabean (150 g/kg) (Clinquart et al. 1995)

<table>
<thead>
<tr>
<th>FA</th>
<th>Soyabean oil</th>
<th>Heat-treated soyabean</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Perirenal</td>
<td>Intramuscular</td>
</tr>
<tr>
<td>SMCAFA</td>
<td>−21.2</td>
<td>−10.8</td>
</tr>
<tr>
<td>18:0</td>
<td>+2.0</td>
<td>+3.8</td>
</tr>
<tr>
<td>18:1</td>
<td>+34.1</td>
<td>−2.3</td>
</tr>
<tr>
<td>18:2</td>
<td>−8.7</td>
<td>+7.1</td>
</tr>
</tbody>
</table>

SMCAFA, short- and medium-chain FA.
only, in the phospholipid fraction, whereas little or no n-3 PUFA are retained in the triacylglycerols, in contrast to pigs (Wood & Enser, 1997). Chain elongation occurs since dietary linseed oil also increases 20:5 n-3 in intramuscular total lipids and phospholipids (Vatansever et al. 1998). On the other hand, it has been clearly shown that linseed oil supplementation increases 18:3n-3 in adipose tissues (Climquart et al. 1991). In more detailed work, Ashes et al. (1995) discuss the specific substitution of 20:5 for 20:4 in the phosphatidyl inositol of membrane phospholipids. They refer to the associated changes to be expected in the nature and effects of phosphatidyl inositol-derived secondary messenger molecules. In a more general approach, these effects include possible changes in gene expression (Jump et al. 1996) and immune reactions (Miles & Calder, 1998), mediated through altered prostaglandins and leukotrienes, when muscle membrane PUFA are modified. As discussed for milk production, such considerations, also related to animal welfare, should be taken into account when evaluating methods for dietary manipulation of muscle lipids. They should be of similar importance to considerations of the greater susceptibility to oxidation and flavour defects of PUFA-enriched meat and meat products (Gandemer, 1997). Similar considerations should indeed apply for further selection and breeding of double-muscled cattle (Demeyer et al. 1995b).

The use of tissue active growth promoters as well as direct immunological depression of body fat (Flint et al. 1994) will not be discussed here. Indeed, regulatory and/or consumer attitudes do not (yet) allow such technologies. Also, they may adversely affect sensory meat quality (Fiems et al. 1990).

Finally, it should be realized that, as was observed for milk production, simple feeding practices may go a long way towards optimizing beef FA composition. From data presented by Wood & Enser (1997), n-3:n-6 values for PUFA in intramuscular lipids of 0-12 to 0-77 can be calculated for concentrate- and grass-fed cattle respectively. This finding suggests that the higher level of 18:3n-3 normally found in meat compared with milk is due to a more efficient incorporation into intramuscular lipids than into milk lipids rather than to the concentrate feeding of fattening animals. The difference, however, was partly due to a lower 18:2 meat content for the concentrate-fed animals (Enser et al. 1998)

**Limits to modification of ruminant fat**

Processing of feed for fat protection may not be acceptable to consumers or governments, as illustrated by the refusal to accept formaldehyde treatment, in spite of extensive experimentation (McDonald & Scott, 1977). The same argument may hold for newer technologies involving the preparation of fatty acyl amides. The use of feed additives aimed at modification of milk production is, at least for the EU, out of the question. A good example is monensin, shown to increase duodenal flow of unsaturated FA (Kobayashi et al. 1992). A definite ban of its use could be supported by the finding that it is absorbed from the intestinal tract and elicits a series of metabolic effects in the animal when infused intravenously (Armstrong & Spears, 1988). Consumer and governmental aversion is even greater against the use of production modifiers active at the animal tissue level, although reduction of short- and medium-chain FA and increases in long-chain FA in milk fat have been achieved in cows treated with bovine somatotropin (Baer, 1996). Direct interference with mammary tissue metabolism through genetic manipulation has been suggested for reduction of acetyl-CoA carboxylase, the enzyme regulating de novo synthesis of fat (Chalupa et al. 1996). A detailed proposal for the development of a recombinant bovine stearoyl-CoA desaturase (EC 1.14.99.5) for use in selection and/or treatment of milk subsequent to its removal from the cow is available (McFadden & Trumble, 1997). Again, consumer acceptance of such technologies is questionable. The limitation on the use of tallows in the UK because of the risk of transmission of factors linked to bovine spongiform encephalopathy is perhaps the best example of the care that should be taken when proposing dietary or other interventions. In an excellent paper that should inspire animal production as a whole, Fredeen (1996) states that modification of milk composition via nutrition should not be approached simply from the point of view of economic advantage. Environmental, welfare and moral considerations should also be considered, analogous to a general quality concept for animal production (Demeyer, 1997). For example, dietary supplements should also be evaluated for environmental effects related to, for example, methanogenesis (Dong et al. 1997), whereas the role of pasture v. concentrate feeding in providing more acceptable unsaturation of milk fat (Murphy et al. 1995) and improvement of PUFA-derived flavour compounds in dairy products (Urbach, 1990) should perhaps receive more attention, related also to the need for extensification. Finally, the greater susceptibility to oxidation and flavour defects of dairy products enriched in PUFA should be considered (Doreau & Chilliard, 1997b; Doreau et al. 1997).

**Acknowledgements**

The authors thank D. Baevens, J. Chabrot, A. Christophe, W. De Greyt, S. De Smet, R. De Schrijver, L. Fiems, V. Fievez, J.L. Hornick, L. Istasse, E. Teller, J. Van Camp and C. Van Nevel for help with the literature search, interpretation of data and preparation of the manuscript. Part of the work reported by D. D. was supported by the IWONL, Ministry of Agriculture, Brussels.

**References**


Enjalbert F, Nicot MC, Vernay M, Moncoulou R & Griess D (1994) Effect of different forms of polysaturated fatty acids on


Kelly ML, Berry JR, Dwyer DA, Bauman DE, Van Amburgh ME & Grinnar JM (1997) Effect of dietary fatty acid sources on...
conjugated linoleic acid (CLA) levels in milk from lactating dairy cows. *Journal of Dairy Science* 80, Suppl. 1, 243.

Kennelly JJ (1996a) Producing milk with 2.5% fat – the biology and health implications for dairy cows. *Animal Feed Science and Technology* 60, 161–180.

Kennelly JJ (1996b) The fatty acid composition of milk fat as influenced by feeding oilseeds. *Animal Feed Science and Technology* 60, 137–152.


Willett WC, Stampfer MJ, Manson JE, Colditz GA,Speizer FE, Rosner BA, Sampson LA & Hennekens CH (1993) Intake of


© Nutrition Society 1999