The combined use of triacylglycerols containing medium-chain fatty acids and exogenous lipolytic enzymes as an alternative to in-feed antibiotics in piglets: concept, possibilities and limitations. An overview

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In the search for alternatives to banned in-feed antibiotics, a concept was developed based on studies with medium-chain fatty acid-containing triacylglycerols (MCTAG) and selected lipases for in situ generation of diacylglycerols, monoacylglycerols and medium-chain fatty acids (MCFA) in the stomach and proximal gut of piglets. MCFA are known to have strong antibacterial properties but can hardly be used as such because of their repellent odour and taste. Those problems could be overcome by the generation of MCFA in situ. The concept was tested in vitro and validated in vivo with gastric-cannulated piglets and under field conditions, including effects on zootechnical performance, with classical antibacterial growth promoters or organic acids acting as positive controls. Furthermore, the metabolic and dietary constraints on the nutritional and nutritive use of MCTAG and/or MCFA (for example, the effects on digestive physiology, gut flora, feed intake, performance, carcass composition) are reviewed. The role of natural pre-duodenal lipase activity, the presence of endogenous plant lipase activity in raw materials and the feasibility for exogenous lipase addition to the feed are discussed, in order to optimize the concept. The present review illustrates the similarity of the action of MCFA and commonly used antimicrobials on the flora (total flora, Gram-positive flora, Gram-negative flora, potential pathogens) and epithelial morphology and histology in the foregut. These observations are believed to be the basis for obtaining optimal growth performances. In addition, these naturally occurring antimicrobial agents have little or no human or animal toxicity and induce no problems of residues and cross-resistance induction. They are proposed as a valuable alternative to in-feed antibiotics, used for growth promotion, and even for the preventive and curative treatment of gastrointestinal diseases.

Animal feeds: Antimicrobials: Gut flora: Medium-chain fatty acids: Lipase

Introduction

As a general ban of all in-feed antibiotics in the European Union is foreseen for 2005, animal nutritionists are highly interested in active alternatives. Because there is a general consensus that the growth-promoting effect is mediated through the antibiotics’ regulating influence on the gut flora and altering directly or indirectly epithelial functions in the small intestine (Thomke & Elwinger, 1998; Anderson et al., 1999), a number of alternatives with comparable effects have been proposed, although a clear-cut validation is lacking for most of them. These are: enzymes, probiotics and prebiotics, certain non-digestible oligosaccharides, fermented liquid feeds, blood plasma proteins, Zn, dietary acidifiers, herbs and plant extracts, antimicrobial peptides and lytic phages (Roth & Kirchgessner, 1998; Thomke & Elwinger, 1998; Cowan, 1999; Verstegen & Schaafsma, 1999; Jensen et al., 2003; Joerger, 2003; Lis-Balchin, 2003; Mroz, 2003). However, at present, none of these compounds can completely replace antibiotics in the diet of piglets. Non-nutritional strategies (weaning age, management, environmental control, genetics), although very important, are not considered in the present review.

The most promising compounds for the feeding of weaner piglets, growers and finishers, seem to be the

Abbreviations: IEL, intra-epithelial lymphocytes; LCTAG, long-chain fatty acid-containing triacylglycerols; MCFA, medium-chain fatty acids; MCTAG; medium-chain fatty acid-containing triacylglycerols; NEFA, non-esterified fatty acids; SCFA, short-chain fatty acids; TAG, triacylglycerols.

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organic short-chain fatty acids (SCFA); formic, acetic, pro- 
pionic and butyric acids, commonly known as volatile fatty acids; and lactic, sorbic, fumaric, malic, tartaric and citric 
acids (Partanen & Mroz, 1999). However the health and 
performance effects of these organic acids are not always 
consistent (Jensen, 1998). The considerable variation in the 
efficacy of different organic acids seems to be related to 
differences in dietary composition and the physico-chemi- 
cal characteristics of the ingredients (for example, acid-
binding capacity), animals (for example, age) and type and 
level of acid(s) or their salts.

**Background of the concept**

In most mammals the alimentary tract becomes heavily 
populated with bacteria within a few hours after birth. 
Afterwards, a stabilized population persists throughout life, 
greatly depending on the relationship of age with diet. Baby 
rabbits are exceptional in this respect, because during the 
sucking period their stomach and small intestine are virtu-
ally sterile.

Further studies revealed that the fat present in rabbit milk 
was transformed into antimicrobial substances by lipolytic 
enzymes present in the stomach wall of the sucking rabbit 
(Canas-Rodriguez & Smith, 1966; Smith, 1966). Indeed, 
antimicrobial activity was not found in the rabbit milk itself 
and neither triacylglycerols (TAG) nor diacylglycerols 
inactivate microbes (Isaacs, 2001).

SCFA and medium-chain fatty acids (MCFA) are fatty 
acids made up of 1–5 and 6–12 C atoms, respectively. The 
antimicrobial activity of SCFA and MCFA or their deriv-
atives (for example, monoacylglycerols) has been known for 
a long time and is summarized in the work of Kabara 
(1972). They were further classified as food-grade germici-
dal agents, pharmaceutical preservatives, silage additives or 
feed preservatives. Meeus (1994) and Decuypere & Meeus 
(1995) tested the antimicrobial activity of MCFA 
in vitro 
against the components of the small-intestinal microflora of 
weanling piglets. They suggested that, in combination with 
a proper lipase, MCFA could be an interesting alternative to 
antibiotic growth promoters. By substituting part of the tallow 
in the milk replacer with tricaprin or tricaprylin, the 
growth of preruminant calves was increased by 40 % 
(Aurousseau et al. 1984). The authors explained this phe-

omenon by the high antimicrobial efficacy of MCFA 
released by the enzymes in the gut lumen and by their 
fuelling action for the enterocytes. In the stomach contents 
of babies, inhibitory fatty acids or their derivatives could be 
generated by endogenous lipases, as noted by Hamosh et 
al. (1981, 1989), Isaacs et al. (1990, 1992, 1995) and 
Hamosh (1997).

### Occurrence of medium-chain fatty acid-containing 
triacylglycerols in man and animals

The occurrence of medium-chain fatty acid-containing 
TAG (MCTAG) in the milk of women and different ani-
mals is variable and the causes of this are not well under-
stood. There is some evidence that their presence and 
concentration are related to the level of immaturity of the 
young at birth (Smith, 1980). A more in-depth literature

<table>
<thead>
<tr>
<th>Fatty acid</th>
<th>MCTAG oil</th>
<th>Coconut oil</th>
<th>Palm-kernel oil</th>
<th>Human milk</th>
<th>Goats milk</th>
<th>Horse milk</th>
<th>Cat milk</th>
<th>Sheep milk</th>
<th>Rabbit milk</th>
<th>Sows’ milk</th>
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<tr>
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<td>8.0</td>
<td>16.0</td>
<td>24.0</td>
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<td>40.0</td>
<td>48.0</td>
<td>56.0</td>
<td>64.0</td>
<td>72.0</td>
</tr>
<tr>
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<td>4.0</td>
<td>8.0</td>
<td>16.0</td>
<td>24.0</td>
<td>32.0</td>
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<td>56.0</td>
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<tr>
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<td>4.0</td>
<td>8.0</td>
<td>16.0</td>
<td>24.0</td>
<td>32.0</td>
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<td>48.0</td>
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<td>64.0</td>
<td>72.0</td>
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<tr>
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<td>8.0</td>
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<td>48.0</td>
<td>56.0</td>
<td>64.0</td>
<td>72.0</td>
</tr>
</tbody>
</table>

MCTAG, medium-chain fatty acid-containing triacylglycerols.
Alternative to in-feed antibiotics in piglets

Table 2. Presence and anatomical site of mammalian and poultry preduodenal lipase activities* (mainly after Moreau et al. 1988)

<table>
<thead>
<tr>
<th>Anatomical site</th>
<th>Rat</th>
<th>Mouse</th>
<th>Rabbit</th>
<th>Dog</th>
<th>Horse</th>
<th>Pig</th>
<th>Calf and Sheep</th>
<th>Poultry</th>
<th>Man</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tongue†</td>
<td>xxx</td>
<td>x</td>
<td>x</td>
<td>x</td>
<td>x</td>
<td>x</td>
<td>xx</td>
<td>(x)</td>
<td></td>
</tr>
<tr>
<td>Pharynx‡</td>
<td></td>
<td>x</td>
<td>xxxx</td>
<td>xxx</td>
<td>x</td>
<td>x</td>
<td>xx</td>
<td>(x)</td>
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<tr>
<td>Gizzard and proventriculus</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Gastric cardia</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Gastric fundus§</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

x, Activity < 10 U/g tissue; xx, activity of 10–100 U/g tissue; xxx, activity of 100–200 U/g tissue; xxxx, activity > 200 U/g tissue; (x), activity present but no exact data available.

* Lipase activity is expressed in U/g fresh tissue measured under optimal conditions, 1 U being the amount of enzyme releasing 1 µmol fatty acid/min (pH 6.5, 30°C) from an olive-oil emulsion in a pH stat apparatus (Committee on Food Chemicals Codex, 1981).
† Chief cells.
‡ Root of tongue and pharynx-glosso-epiglottic area.
§ Von Ebner’s glands.

search supports the relatively important contribution of MCFA in the milk-lipid of certain mammals including the rabbit, goat, mare, rat, mouse and elephant, while in other species (cow, sheep, man) the concentrations are rather low or even negligible; for example, in the milk of the sow, camel or guinea-pig (Table 1).

Occurrence of preduodenal lipases in man and animals
In man and most mammals there is a more or less developed preduodenal lipase. The origin is different; lingual and/or pharyngeal or gastric, and for both the term preduodenal lipases (to differentiate from pancreatic lipases) is used. Common properties of the preduodenal lipases are that they are active over a broad and rather acid pH range (except in the pig) (Höller, 1970; Newport & Howarth, 1985; Moreau et al. 1988). These preduodenal lipases have a strong preference for MCFA in milk fat. High activities are found in human subjects, preruminant calves, young rabbits and dogs, while activities are moderate in piglets and low in carnivorous birds. In poultry and other birds they are absent (Table 2).

Natural sources of medium-chain fatty acid-containing triacylglycerols other than milk
Although the milk of certain mammals is certainly the most abundant source of MCFA, they also occur in the fat of some seeds and plants (for example, Cuphea). They have been commercialized by many firms, and have slightly variable composition and are under different brands as tailored lipid sources; for example, Captex (Capital City Products, Columbus, OH, USA) and Neobee (Stepan Co., Maywood, NJ, USA). These MCTAG sources are almost pure and are prepared industrially from coconut and palm-kernel oils by the enzymic or chemical esterification of glycerol with octanoic and decanoic acids.

Possibilities, uses and side effects of medium-chain fatty acid-containing triacylglycerols and medium-chain fatty acids in nutrition
Intact MCTAG have been used in human nutrition as an energy source, especially in clinical settings and parenteral feeding (premature infants, fat malabsorption syndromes, severe surgery, cancer), for approximately 50 years because of their unique properties. Their most important properties include the rapid and complete hydrolysis by lingual, gastric and pancreatic lipases and their beneficial effect on the intestinal mucosa. Other important properties are: their direct transport via the portal blood to the liver (without chylomicron formation or re-esterification); their preferential oxidation in the mitochondria to CO2 and ketone bodies (less dependence on carnitine) providing a rapidly available energy source (Velasquez et al. 1996).

However, in animals, an excess of non-esterified MCFA can have serious unwanted side effects, especially when given in high doses over a short time (for example, as a force-fed energy booster in the form of a lipid bolus). In neonatal piglets this can be ketogenic and narcotic (Samson et al. 1956; Lin et al. 1995). Also in man, the ingestion of > 30 g MCTAG in a short period of time induces nausea and gastrointestinal discomfort (Brouns & Van der Vusse, 1998). Moreover, MCFA may be a stimulus to the secretion of cholecystokinin, and perhaps other intestinal hormones, resulting in a pronounced satiety action that could interfere with gastric emptying and feed intake (Mabayo et al. 1992, 1994). However, recent research has revealed that MCTAG have only minor effects on cholecystokinin release (Symersky et al. 2002). The strong goat-like odour (Molimard et al. 1997) and repellent taste of non-esterified MCFA (Cera et al. 1989b; Timmermann, 1993) can also be a cause of a lower feed intake. Salts of MCFA, on the other hand, may disturb the acid–base balance in the animal.

The authors of the present study thought that the in situ generation (for example, directly in the stomach) of diacylglycerols, monoacylglycerols and MCFA from intact MCTAG could avoid these side effects and so should always be highly preferred above the direct supplementation of the diet with non-esterified MCFA or their salts. Also because of the more prolonged retention time in the stomach and a slower absorption rate, a stronger antimicrobial efficacy of monoacylglycerols above non-esterified MCFA can be expected (Kabara et al. 1972; Kabara, 1984; Isaacs et al. 1990, 1995).

This was the origin of the authors’ studies on the influences of the combined feeding of MCTAG together with an appropriate lipase, of which the origin can be various, but preferably easily commercially available. These commercial lipases are quite different in origin: plants (wheat, cas-
tor bean, rapeseed, mustard); animals (pre-gastric esterase and rennet paste from calf, kid and lamb); micro-organisms (Candida, Rhizopus, Penicillium, Pseudomonas) (Pandey et al. 1999).

The present paper summarizes the concept, possibilities and limitations of applying TAG containing MCFA in combination with lipases as an alternative to in-feed antibiotics in piglets. The results presented here concerning the development of the concept are mostly based on the authors’ research, recently published in detail elsewhere (Dierick & Decuyper, 2002; Dierick et al. 2002a,b, 2003), but more emphasis is put in the present review on the possibilities and limitations. To the authors’ knowledge, no data in the literature are available regarding the simultaneous use of specific intact MCTAG and lipolytic enzymes in animal nutrition for the purposes of growth promotion.

Although the energetic evaluation of the MCFA was not part of the authors’ experiments, the metabolic and dietary aspects of the use of MCTAG and MCFA (effects on digestive physiology, gut flora, feed intake, performance, carcass composition) are also highlighted in the present overview. In addition, the role of preduodenal lipase activity, the presence of endogenous plant and microbial lipase activity in feedstuffs and the potential for exogenous lipase addition to the feed are discussed, leading to an optimal application of the concept.

Development and validation of the concept

Efficacy in vitro

An in vitro screening was carried out with a selection of commercial lipases for studying their activity on different MCFA containing natural or synthetic substrates (Dierick et al. 2002a). The conditions prevailing in vivo in the stomach of piglets (for example, an acid pH range of 3 to 6, a coarse emulsified state of the fat source, the presence of inhibiting components in the diet, the presence of pepsin and gastric flora, a mean retention time of 3 h) were simulated as much as possible. This was to avoid bias induced by the use of unnatural standard procedures. In the first set of experiments, lipolysis was studied with four selected MCFA-containing fat sources. These were: coconut oil; MCTAG1 (Aldo; Lonza Inc., Fair Lawn, NJ, USA); MCTAG2 (Stabilox; Lodders Croklaan B.V., Wormerveer, The Netherlands); butter oil. Six lipases of different origin were used (L1–L6; microbial, porcine and calf pancreas). Depending on the conditions, up to 20 % of the MCFA could be enzymically released in the medium. From these studies it is clear that some of the lipases tested were acid- and pepsin-resistant and that appropriate amounts of MCFA could be liberated to control the bacterial population, indicating that they could be an alternative to in-feed antibiotics.

In a second set of experiments (Dierick et al. 2002a), the generation and antimicrobial effects of MCFA from three selected MCFA-containing fat sources (coconut oil, MCTAG1, and MCTAG2) and one selected effective microbial lipase (L5), applied under different combinations and concentrations, were studied. A minimal concentration of 0·35 g/100 g incubation fluid or 0·025 m-MCFA in the medium (for example, stomach, proximal gut) seems necessary in order to obtain a significant (>10-fold) suppression of the flora (total anaerobic count, Escherichia coli), corresponding to 0·025 m-non-esterified MCFA in the medium. This amount can be obtained by selecting an appropriate combination of sources and doses of fat and lipases (Table 3). The inhibitory effect on the flora of the fat sources used decreases further (MCTAG1 oil > MCTAG2 oil > coconut oil) with an increase in the molecular weight of the most important MCFA present in the TAG used. There was also a correlation between microbial growth inhibition and the lipophilic character of the fatty acids. This was also reported by Freese et al. (1973) and Sheu et al. (1975); 5–10 mM-C6 : 0 and -C8 : 0 were needed for 50 % growth inhibition of E. coli, while for C10 : 0 and C12 : 0 the minimum inhibitory concentration value was >10 mM. Because the pKa (acid strength) of MCFA is about 4·9 and the pH in the piglet stomach (fed ad libitum) ranges between 3 and 6 (N Dierick and J Decuyper, unpublished results), it can be accepted that most of the MCFA will be undissociated. In the more fat-soluble (undissociated) form, they can freely penetrate through the semi-permeable peptidoglycan (Gram-positive) or phospholipid (Gram-negative) membrane of the micro-organisms into the cytoplasm by passive diffusion, as originally formulated by Jacobs (1940). In the cell, dissociation into proton and anion, due to the alkaline pH in the cytoplasm, will lower pH, suppress cytoplasmic enzymes and nutrient transport systems and uncouple ATP-driven pumps, leading to cellular death (Freese et al. 1973; Hsiao & Siebert, 1999). The inhibitory doses found in the present experiment agree well with the concentrations (25–30 mM) of acids proposed for inhibiting the growth of Gram-positive and Gram-negative microbes in food (Freese et al. 1973; Ostling & Lindgren, 1993), pharmaceutical preparations (Kabara, 1984) and silages (Woollford, 1975). Nevertheless, those bacterial species are very different from the normal dominant species in the pig gut. Using a mathematical model based on principal component analysis, Hsiao & Siebert (1999) obtained minimum inhibitory concentration values for caproic, caprylic and capric acids of 2·2, 3·4 and 19·3 g/l for E. coli in food. Their experiments confirm very well our findings and the importance of the pKa, molecular weight and the polarity of the MCFA.

Efficacy in vivo

A first in vivo experiment (Dierick et al. 2002b) was set up to verify whether the results obtained in vitro could also be obtained in situ in gastric-cannulated piglets. Indeed, the addition (5 %) of MCTAG (coconut oil, MCTAG1 oil, butter oil) to piglet diets in combination with selected lipolytic enzymes (1000 parts per million, in the feed) clearly regulated and stabilized the gastrointestinal flora in the stomach. It was striking that the extent of release of MCFA in the stomach corresponded closely to the degree of suppression of the bacterial load in the stomach (Table 4). The most pronounced reduction in total bacterial load in the stomach occurred with 1·01 g non-esterified fatty acids (NEFA)/100 g fresh contents (equivalent to 60 % hydrolysis of the fat) or 0·68 g MCFA/100 g obtained with MCTAG1 oil + L5, followed by coconut oil + L5, by which 0·82 g NEFA or
Table 3. In vitro release of medium-chain fatty acids (MCFA) from different fat sources under different incubation conditions and their effect on bacterial growth in gastric-simulating conditions (Dierick et al. 2002a)

<table>
<thead>
<tr>
<th>Incubations*</th>
<th>Start (blank)</th>
<th>0/0 (control)</th>
<th>2.5/10000</th>
<th>5/10000</th>
<th>10/10000</th>
<th>2.5/1000</th>
<th>5/1000</th>
<th>10/1000</th>
<th>2.5/100</th>
<th>5/100</th>
<th>10/100</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fat source</td>
<td></td>
<td></td>
<td></td>
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<td></td>
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<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Coconut oil</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sum MCFA†</td>
<td>0.00</td>
<td>0.00a</td>
<td>0.04a</td>
<td>0.07b</td>
<td>0.18b</td>
<td>0.03a</td>
<td>0.06p</td>
<td>0.10p</td>
<td>0.02a</td>
<td>0.03a</td>
<td>0.06p</td>
</tr>
<tr>
<td>Total anaerobic count‡</td>
<td>6.3</td>
<td>7.1a</td>
<td>7.2a</td>
<td>6.2b</td>
<td>6.2b</td>
<td>6.4a</td>
<td>6.5a</td>
<td>6.4a</td>
<td>6.9a</td>
<td>7.0a</td>
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</tr>
<tr>
<td>Escherichia coli§</td>
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<td>2.0a</td>
<td>0.0a</td>
<td>0.0a</td>
<td>0.0a</td>
<td>2.0a</td>
<td>1.8a</td>
<td>1.6a</td>
<td>1.9a</td>
<td>0.0a</td>
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<tr>
<td>MCTAG1 oil</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sum MCFA†</td>
<td>0.00</td>
<td>0.00a</td>
<td>0.15b</td>
<td>0.33b</td>
<td>0.61b</td>
<td>0.09b</td>
<td>0.18b</td>
<td>0.37b</td>
<td>0.07b</td>
<td>0.11b</td>
<td>0.21b</td>
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<td>6.8b</td>
<td>5.9a</td>
<td>0.0a</td>
<td>0.0b</td>
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<td>4.8b</td>
<td>3.8b</td>
<td>6.5a</td>
<td>6.5a</td>
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<td>Escherichia coli§</td>
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<td>2.5a</td>
<td>2.1a</td>
<td>0.0a</td>
<td>0.0a</td>
<td>3.1a</td>
<td>2.5a</td>
<td>1.8a</td>
<td>2.9a</td>
<td>2.8a</td>
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<tr>
<td>MCTAG2 oil</td>
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</tr>
<tr>
<td>Sum MCFA†</td>
<td>0.00</td>
<td>0.00a</td>
<td>0.15b</td>
<td>0.28b</td>
<td>0.56b</td>
<td>0.11b</td>
<td>0.19b</td>
<td>0.34b</td>
<td>0.09b</td>
<td>0.14a</td>
<td>0.22b</td>
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<td>7.0a</td>
<td>5.5b</td>
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<td>1.8b</td>
<td>6.3b</td>
<td>6.3b</td>
<td>5.6b</td>
<td>6.5b</td>
<td>6.6b</td>
<td>6.7a</td>
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<tr>
<td>Escherichia coli§</td>
<td>3.1</td>
<td>3.3a</td>
<td>0.0b</td>
<td>0.0b</td>
<td>0.0b</td>
<td>3.1a</td>
<td>1.8b</td>
<td>1.6b</td>
<td>3.3a</td>
<td>3.2a</td>
<td>3.3a</td>
</tr>
</tbody>
</table>

MCTAG, medium-chain fatty acid-containing triacylglycerols.

a,b Mean values within a row with unlike superscript letters were significantly different from control (condition 0/0) (P<0.05).

* Incubation time (3 h) and conditions (percentage fat content in medium and lipase dose (parts per million of fat)). The lipase used was ‘L5’, of microbial origin (6563 U/g; Kemin Europa, Herentals, Belgium).

† Release of MCFA in the medium (g/100 g incubation fluid).

‡ Reinforced Clostridial agar medium for total anaerobic count (colony-forming units log10/ml incubation fluid).

§ Eosin Methylene Blue agar for E. coli count (colony-forming units log10/ml incubation fluid).

II MCTAG1 contained (g/100 g fat): C4:0, 0.00; C6:0, 2.70; C8:0, 67.20; C10:0, 26.90; C12:0, 0.40. MCTAG2 contained (g/100 g fat): C4:0, 0.00; C6:0, 0.16; C8:0, 53.69; C10:0, 39.50; C12:0, 0.29.

0·30 g MCFA/100 g was released. Butter oil + L5 had the least activity with 0·73 g NEFA or 0·06 g MCFA/100 g gastric contents.

Confirmation of the in vivo results

It is clear that in the in vivo experiment of Dierick et al. (2002b), the results obtained in the in vitro experiment were confirmed. However an additional validation of the results in more commercial settings was required. Therefore a combined growth trial and slaughter experiment with newly weaned piglets was set up (Dierick et al. 2002b). Four diets (A (negative control), 2·5 % soyabean oil; B, 2·5 % MCTAG2 oil; C, 2·5 % MCTAG2 oil + 1000 parts per million lipase L5; D, 2·5 % soyabean oil + 1·5 % organic acids) were used. The last experimental group served as a positive control and was fed a combination of oils, which are rather expensive and industrially produced, as a natural source of MCFA (75 % C10:0 in the fat seeds, as a natural source of MCFA (75 % C10:0 in the fat content), together with an exogenous lipase (500 parts per tence of a correlation between the amount of released MCFA in the stomach and the inhibitory effect on the flora. The correlations were r 0·86 for MCTAG1 (P < 0·01); r 0·83 for MCTAG2 (P < 0·01); r 0·50 for coconut oil (P < 0·05). This means that the higher the C8:0 + C10:0 and the lower the C12:0 content in the MCFA profile in the fat source, the higher was the correlation.

Efficacy in the field

The in vivo experiment described earlier (Dierick et al. 2002b) was carried out in a commercial setting. The rest of the early weaned piglets were involved in a growth trial in which it was clearly demonstrated that the manipulation of the gut ecosystem by the enzymic in situ release of MCFA in the stomach and duodenum resulted in improved performance (Dierick et al. 2002b, 2003; Tables 5 and 6). An increase in daily gain of more than 10 %, combined with a 3 % better feed conversion with the diets containing MCTAG2 oil or MCTAG2 oil + lipase was obtained, exceeding the performance observed with the control diets based on soyabean oil, supplemented or not with organic acids (Table 6).

In a search for an alternative to commercial MCTAG oils, which are rather expensive and industrially produced, the effects of adding a combination of milled Cuphea seeds, as a natural source of MCFA (75 % C10:0 in the fat content), together with an exogenous lipase (500 parts per...
Table 4. Content of non-esterified and total fatty acids (FA), overall degree of hydrolysis (DH) and microbiota in gastric contents of cannulated piglets as influenced by the different dietary treatments (four piglets per treatment; Expt I) (Dierick et al., 2002b)

<table>
<thead>
<tr>
<th>D1</th>
<th>D2</th>
<th>D3</th>
<th>D4</th>
<th>D5</th>
<th>D6</th>
<th>D7</th>
<th>D8</th>
<th>D9</th>
</tr>
</thead>
<tbody>
<tr>
<td>Oil*</td>
<td>Coconut</td>
<td>Coconut</td>
<td>Coconut</td>
<td>MCTAG1</td>
<td>MCTAG1</td>
<td>MCTAG1</td>
<td>Butter oil</td>
<td>Butter oil</td>
</tr>
<tr>
<td>Enzyme†</td>
<td>0</td>
<td>L2</td>
<td>L5</td>
<td>0</td>
<td>L2</td>
<td>L5</td>
<td>0</td>
<td>L2</td>
</tr>
<tr>
<td>Non-esterified MCFA (g/100 g fresh contents)</td>
<td>4:0</td>
<td>0.00&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.01&lt;sup&gt;b&lt;/sup&gt;</td>
<td>0.01&lt;sup&gt;b&lt;/sup&gt;</td>
<td>0.00&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.01&lt;sup&gt;b&lt;/sup&gt;</td>
<td>0.01&lt;sup&gt;b&lt;/sup&gt;</td>
<td>0.00&lt;sup&gt;ab&lt;/sup&gt;</td>
</tr>
<tr>
<td>6:0</td>
<td>0.00&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.00&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.01&lt;sup&gt;bc&lt;/sup&gt;</td>
<td>0.00&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.02&lt;sup&gt;e&lt;/sup&gt;</td>
<td>0.02&lt;sup&gt;e&lt;/sup&gt;</td>
<td>0.00&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.01&lt;sup&gt;c&lt;/sup&gt;</td>
</tr>
<tr>
<td>8:0</td>
<td>0.02&lt;sup&gt;ab&lt;/sup&gt;</td>
<td>0.05&lt;sup&gt;b&lt;/sup&gt;</td>
<td>0.05&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.09&lt;sup&gt;c&lt;/sup&gt;</td>
<td>0.43&lt;sup&gt;d&lt;/sup&gt;</td>
<td>0.47&lt;sup&gt;d&lt;/sup&gt;</td>
<td>0.00&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.01&lt;sup&gt;bc&lt;/sup&gt;</td>
</tr>
<tr>
<td>10:0</td>
<td>0.01&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.03&lt;sup&gt;c&lt;/sup&gt;</td>
<td>0.03&lt;sup&gt;c&lt;/sup&gt;</td>
<td>0.03&lt;sup&gt;c&lt;/sup&gt;</td>
<td>0.16&lt;sup&gt;f&lt;/sup&gt;</td>
<td>0.17&lt;sup&gt;f&lt;/sup&gt;</td>
<td>0.00&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.01&lt;sup&gt;bc&lt;/sup&gt;</td>
</tr>
<tr>
<td>12:0</td>
<td>0.05&lt;sup&gt;b&lt;/sup&gt;</td>
<td>0.19&lt;sup&gt;c&lt;/sup&gt;</td>
<td>0.21&lt;sup&gt;c&lt;/sup&gt;</td>
<td>0.00&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.01&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.01&lt;sup&gt;b&lt;/sup&gt;</td>
<td>0.00&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.01&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
<tr>
<td>Sum of non-esterified MCFA (g/100 g fresh contents)</td>
<td>0.07&lt;sup&gt;ab&lt;/sup&gt;</td>
<td>0.28&lt;sup&gt;c&lt;/sup&gt;</td>
<td>0.30&lt;sup&gt;c&lt;/sup&gt;</td>
<td>0.12&lt;sup&gt;c&lt;/sup&gt;</td>
<td>0.61&lt;sup&gt;d&lt;/sup&gt;</td>
<td>0.68&lt;sup&gt;d&lt;/sup&gt;</td>
<td>0.02&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.01&lt;sup&gt;bc&lt;/sup&gt;</td>
</tr>
<tr>
<td>Sum of total non-esterified FA (g/100 g fresh contents)</td>
<td>0.32&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.79&lt;sup&gt;bc&lt;/sup&gt;</td>
<td>0.82&lt;sup&gt;bc&lt;/sup&gt;</td>
<td>0.29&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.91&lt;sup&gt;cd&lt;/sup&gt;</td>
<td>1.01&lt;sup&gt;d&lt;/sup&gt;</td>
<td>0.20&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.68&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
<tr>
<td>Total FA (g/100 g fresh contents)</td>
<td>1.93&lt;sup&gt;e&lt;/sup&gt;</td>
<td>1.83&lt;sup&gt;cd&lt;/sup&gt;</td>
<td>1.83&lt;sup&gt;cd&lt;/sup&gt;</td>
<td>1.54&lt;sup&gt;cd&lt;/sup&gt;</td>
<td>1.56&lt;sup&gt;d&lt;/sup&gt;</td>
<td>1.66&lt;sup&gt;d&lt;/sup&gt;</td>
<td>1.21&lt;sup&gt;a&lt;/sup&gt;</td>
<td>1.46&lt;sup&gt;ab&lt;/sup&gt;</td>
</tr>
<tr>
<td>DH ((non-esterified FA/total FA) × 100)</td>
<td>16.6&lt;sup&gt;a&lt;/sup&gt;</td>
<td>42.8&lt;sup&gt;b&lt;/sup&gt;</td>
<td>44.5&lt;sup&gt;b&lt;/sup&gt;</td>
<td>18.9&lt;sup&gt;a&lt;/sup&gt;</td>
<td>56.6&lt;sup&gt;c&lt;/sup&gt;</td>
<td>61.0&lt;sup&gt;d&lt;/sup&gt;</td>
<td>16.7&lt;sup&gt;a&lt;/sup&gt;</td>
<td>46.1&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

Flora components (log<sub>10</sub> CFU/g fresh contents)

<table>
<thead>
<tr>
<th>Total count</th>
<th>Lactobacilli</th>
<th>Streptococci</th>
<th>Escherichia coli</th>
</tr>
</thead>
<tbody>
<tr>
<td>7.1&lt;sup&gt;a&lt;/sup&gt;</td>
<td>6.6&lt;sup&gt;a&lt;/sup&gt;</td>
<td>5.1&lt;sup&gt;a&lt;/sup&gt;</td>
<td>2.8&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>5.3&lt;sup&gt;b&lt;/sup&gt;</td>
<td>5.1&lt;sup&gt;a&lt;/sup&gt;</td>
<td>4.5&lt;sup&gt;b&lt;/sup&gt;</td>
<td>3.0&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>5.6&lt;sup&gt;b&lt;/sup&gt;</td>
<td>5.3&lt;sup&gt;a&lt;/sup&gt;</td>
<td>3.6&lt;sup&gt;b&lt;/sup&gt;</td>
<td>3.9&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>6.1&lt;sup&gt;a&lt;/sup&gt;</td>
<td>5.9&lt;sup&gt;a&lt;/sup&gt;</td>
<td>6.3&lt;sup&gt;a&lt;/sup&gt;</td>
<td>3.8&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>4.4&lt;sup&gt;b&lt;/sup&gt;</td>
<td>3.9&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.0&lt;sup&gt;b&lt;/sup&gt;</td>
<td>2.6&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
<tr>
<td>4.3&lt;sup&gt;b&lt;/sup&gt;</td>
<td>3.4&lt;sup&gt;a&lt;/sup&gt;</td>
<td>2.1&lt;sup&gt;b&lt;/sup&gt;</td>
<td>2.0&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
<tr>
<td>6.5&lt;sup&gt;a&lt;/sup&gt;</td>
<td>6.1&lt;sup&gt;a&lt;/sup&gt;</td>
<td>5.3&lt;sup&gt;a&lt;/sup&gt;</td>
<td>4.3&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>6.0&lt;sup&gt;b&lt;/sup&gt;</td>
<td>5.6&lt;sup&gt;a&lt;/sup&gt;</td>
<td>4.4&lt;sup&gt;b&lt;/sup&gt;</td>
<td>3.4&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

MCTAG, medium-chain fatty acid-containing triacylglycerols; MCFA, medium-chain fatty acids; CFU, colony-forming units.

<sup>a</sup> Mean values within a row with unlike superscript letters were significantly different (P < 0.05).

* 5% oil in diets. For details of MCTAG1, see Table 3.

† Lipase 'L2' was of microbial origin (9362 U/g; Kemin Europa, Herentals, Belgium); lipase 'L5' was of microbial origin (6563 U/g; Kemin Europa). Concentration of lipases was 1000 parts per million in diets.
### Table 5. Effect of the diet on the content of non-esterified and total fatty acids (FA) and overall degree of hydrolysis (DH) in gastric contents of slaughtered weaning piglets; effects on the gastric and duodenal flora (five piglets per treatment) (Expt II) (Dierick et al. 2002b)

<table>
<thead>
<tr>
<th>Diet</th>
<th>A</th>
<th>B</th>
<th>C</th>
<th>D</th>
</tr>
</thead>
<tbody>
<tr>
<td>Oil*</td>
<td>Soya</td>
<td>MCTAG2</td>
<td>MCTAG2</td>
<td>Soya</td>
</tr>
<tr>
<td>Lipase†</td>
<td>0</td>
<td>0</td>
<td>L5</td>
<td>0</td>
</tr>
<tr>
<td>Organic acids‡</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>Mixture</td>
</tr>
</tbody>
</table>

**Non-esterified MCFA (g/100 g fresh contents)**

<table>
<thead>
<tr>
<th>Chain length</th>
<th>A</th>
<th>B</th>
<th>C</th>
<th>D</th>
</tr>
</thead>
<tbody>
<tr>
<td>6 : 0</td>
<td>0·00</td>
<td>0·00</td>
<td>0·00</td>
<td>0·00</td>
</tr>
<tr>
<td>8 : 0</td>
<td>0·00a</td>
<td>0·13b</td>
<td>0·25c</td>
<td>0·00a</td>
</tr>
<tr>
<td>10 : 0</td>
<td>0·01a</td>
<td>0·07b</td>
<td>0·17c</td>
<td>0·00a</td>
</tr>
<tr>
<td>12 : 0</td>
<td>0·01</td>
<td>0·01</td>
<td>0·02</td>
<td>0·01</td>
</tr>
</tbody>
</table>

**Sum of non-esterified MCFA (g/100 g fresh contents)**

<table>
<thead>
<tr>
<th>Chain length</th>
<th>A</th>
<th>B</th>
<th>C</th>
<th>D</th>
</tr>
</thead>
<tbody>
<tr>
<td>0·02a</td>
<td>0·22b</td>
<td>0·45c</td>
<td>0·01a</td>
<td></td>
</tr>
</tbody>
</table>

**Sum of total non-esterified FA (g/100 g fresh contents)**

<table>
<thead>
<tr>
<th>Chain length</th>
<th>A</th>
<th>B</th>
<th>C</th>
<th>D</th>
</tr>
</thead>
<tbody>
<tr>
<td>1·05a</td>
<td>1·25ab</td>
<td>1·35b</td>
<td>1·07a</td>
<td></td>
</tr>
</tbody>
</table>

**DH ((non-esterified FA/total FA) \times 100)**

<table>
<thead>
<tr>
<th>DH</th>
<th>A</th>
<th>B</th>
<th>C</th>
<th>D</th>
</tr>
</thead>
<tbody>
<tr>
<td>26·7a</td>
<td>35·2a</td>
<td>70·4b</td>
<td>28·9a</td>
<td></td>
</tr>
</tbody>
</table>

**Flora components (log_{10} CFU/g fresh contents)**

**Stomach**

- **Total count**
  - A: 7·0a
  - B: 7·0ac
  - C: 5·9b
  - D: 6·9ac
- **Lactobacilli**
  - A: 7·2ac
  - B: 7·6a
  - C: 6·6bc
  - D: 7·3a
- **Streptococci**
  - A: 4·2a
  - B: 0·6b
  - C: 5·3a
  - D: 5·1a
- **Escherichia coli**
  - A: 4·6a
  - B: 0·8bc
  - C: 2·0b
  - D: 0·0c

**Duodenum**

- **Total count**
  - A: 6·4a
  - B: 6·1a
  - C: 5·6a
  - D: 5·9a
- **Lactobacilli**
  - A: 6·9
  - B: 6·8
  - C: 5·9
  - D: 6·4
- **Streptococci**
  - A: 1·6a
  - B: 0·0a
  - C: 4·7b
  - D: 4·7b
- **E. coli**
  - A: 4·9a
  - B: 4·8a
  - C: 1·8b
  - D: 1·8b

**MCTAG, medium-chain fatty acid-containing triacylglycerols; MCFA, medium-chain fatty acids; CFU, colony-forming units.**

* 2·5 % oil in diets. For details of MCTAG2, see Table 3.
† Lipase ‘L5’ was of microbial origin (6563 U/g; Kemin Europa, Herentals, Belgium). Concentration was 500 parts per million in diets.
‡ 1·5 % mixture containing 25 % citric acid, 75 % fumaric acid, 50 % calcium formate.

### Table 6. Effect of diet on feed intake, growth rate and feed conversion ratio (FCR) of early weaned piglets* (A, n 68; B, n 61; C, n 60; D, n 55 piglets) (Expt II) (Dierick et al. 2002b)

<table>
<thead>
<tr>
<th>Post-weaning period (d)</th>
<th>0–7</th>
<th>7–14</th>
<th>14–21</th>
<th>0–21</th>
<th>0–21 (relative to A)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Feed intake (g/d)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>A (soyabean oil)†</td>
<td>156</td>
<td>365</td>
<td>472</td>
<td>331</td>
<td>100</td>
</tr>
<tr>
<td>B (MCTAG2 oil)</td>
<td>191</td>
<td>376</td>
<td>536</td>
<td>368</td>
<td>111</td>
</tr>
<tr>
<td>C (MCTAG2 oil + lipase‡)</td>
<td>180</td>
<td>391</td>
<td>533</td>
<td>361</td>
<td>110</td>
</tr>
<tr>
<td>D (soyabean oil + acid mix§)</td>
<td>189</td>
<td>355</td>
<td>469</td>
<td>338</td>
<td>102</td>
</tr>
<tr>
<td>Growth rate (g/d)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>A (soyabean oil)</td>
<td>127a</td>
<td>127a</td>
<td>300a</td>
<td>185ab</td>
<td>100</td>
</tr>
<tr>
<td>B (MCTAG2 oil)</td>
<td>164b</td>
<td>160b</td>
<td>301a</td>
<td>206a</td>
<td>112</td>
</tr>
<tr>
<td>C (MCTAG2 oil + lipase‡)</td>
<td>165b</td>
<td>161b</td>
<td>297a</td>
<td>207a</td>
<td>111</td>
</tr>
<tr>
<td>D (soyabean oil + acid mix§)</td>
<td>141ab</td>
<td>129ab</td>
<td>280b</td>
<td>181b</td>
<td>98</td>
</tr>
<tr>
<td>FCR (kg feed/kg live-weight gain)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>A (soyabean oil)</td>
<td>1·23</td>
<td>2·88</td>
<td>1·57</td>
<td>1·79</td>
<td>100</td>
</tr>
<tr>
<td>B (MCTAG2 oil)</td>
<td>1·16</td>
<td>2·35</td>
<td>1·78</td>
<td>1·77</td>
<td>99</td>
</tr>
<tr>
<td>C (MCTAG2 oil + lipase‡)</td>
<td>1·09</td>
<td>2·43</td>
<td>1·79</td>
<td>1·74</td>
<td>97</td>
</tr>
<tr>
<td>D (soyabean oil + acid mix§)</td>
<td>1·34</td>
<td>2·89</td>
<td>1·68</td>
<td>1·87</td>
<td>104</td>
</tr>
</tbody>
</table>

**MCTAG, medium-chain fatty acid-containing triacylglycerols.**

* Mean values within a row with unlike superscript letters were significantly different (P < 0·05).
† 2·5 % oil in diets A–D. For details of MCTAG2, see Table 3.
‡ Lipase ‘L5’ was of microbial origin (6563 U/g; Kemin Europa, Herentals, Belgium). Concentration was 500 parts per million in diets.
§ 1·5 % mixture containing 25 % citric acid, 75 % fumaric acid, 50 % calcium formate.
The concentration of lipase ‘L5’ was of microbial origin (6563 U/g; Kemin Europa, Herentals, Belgium); concentration was 500 parts per million in diet.

The enzymically released antimicrobial MCFA (1.7 g/kg fresh gastric contents) decreased the number of coliforms in the proximal small intestine, _Streptococci_ in the whole small intestine and _Lactobacilli_ in the stomach and the proximal and distal small intestine. No effects were noted on the total anaerobic microbial load. Most probably, better results could have been obtained by using other _Cuphea_ varieties (Graham, 1989), for example, _C. painteriana_, _C. painteri_, _C. cyanea_, containing more than 50 % C8:0 and 25 % C10:0 in the fat fraction and/or by using a higher exogenous lipase activity for a better and faster release of the fat more proximally in the foregut. Indeed, the endogenous lipase activity in the ground seeds tested here was very low in comparison with that found in other pig-feed raw materials (cereals, legumes) (Dierick & Decuyper, 2002; Dierick et al. 2003). In comparison with the control diet containing no extra lipase, feeding _Cuphea_ + lipase further resulted in a significantly greater villus height (proximal small intestine) and a lesser crypt depth (proximal and distal small intestine), a greater villus:crypt ratio (proximal and distal small intestine) and a lower number of intra-epithelial lymphocytes (IEL) in the villous epithelium, which is indicative for a more healthy and well-differentiated intestinal mucosa (Table 8). IEL are T-lymphocytes, mainly of the CD8+ phenotype, of which the functions have not been entirely elucidated. It is accepted that they play a central role as a first line of defence against foreign luminal antigens (pathogens, proteins), in the induction of apoptosis of epithelial cells and in the conservation of the mucosal integrity (Vega-Lopez et al. 2001). This is in line with the observations of Czernichow et al. (1996) showing that enterally infused MCTAG enhanced mucosal mass and favoured epithelial cell renewal in the proximal intestine in rats. The findings of a more slender villus structure in piglets when antibacterial growth promoters were added to the feed are possibly related to the same mechanism (Van Leeuwen et al. 2001). Potential interference arising from the mucilage-containing hairs and/or secondary plant metabolites (Salatino et al. 2000; Puupponen-Pimiä et al. 2002) of the _Cuphea_ seeds with the gut flora and/or the intestinal surfaces cannot be excluded but further research is required to elucidate this point.

---

**Table 7. Effect of feeding whole _Cuphea_ seeds and lipase on the zootechnical performances of early weaned piglets (fifteen piglets per treatment) (Expt III) (Dierick et al. 2003)**

<table>
<thead>
<tr>
<th></th>
<th>Diet</th>
<th>Days post-weaning</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>0–7</td>
</tr>
<tr>
<td>Feed intake (g/d) (per pen)</td>
<td>Control</td>
<td>212</td>
</tr>
<tr>
<td></td>
<td><em>Cuphea</em> + lipase*</td>
<td>194</td>
</tr>
<tr>
<td>Growth (g/d) (individual)</td>
<td>Control</td>
<td>135</td>
</tr>
<tr>
<td></td>
<td><em>Cuphea</em> + lipase*</td>
<td>135</td>
</tr>
<tr>
<td></td>
<td><em>P</em> value</td>
<td>1.00</td>
</tr>
<tr>
<td>FCR (kg feed/kg LWG) (per pen)</td>
<td>Control</td>
<td>1.57</td>
</tr>
<tr>
<td></td>
<td><em>Cuphea</em> + lipase*</td>
<td>1.43</td>
</tr>
</tbody>
</table>

FCR, feed conversion ratio; LWG, live-weight gain.

* The concentration of _Cuphea_ seeds was 50 g/kg diet. Lipase ‘L5’ was of microbial origin (6563 U/g; Kemin Europa, Herentals, Belgium); concentration was 500 parts per million in diet.
Feeding swine with antibacterial growth promoters has been documented to increase weight gain by 3–9 % and improve feed efficiency by 2–7 % (Thomke & Elwinger, 1998a). The exact mechanisms by which this occurs are complex and not completely understood (Vishek, 1978). The authors of the present study believe that the benefits of growth-promoting antibiotics result from a substantial decrease in bacterial load, whether commensal or pathogenic (Vervaeke et al. 1979) and the consequent direct or indirect alterations in epithelial functions (for example, increased villus height; decreased crypt depth) in the small intestine. This leads to an enhanced uptake and use of nutrients and an enhancement of the activity of the immune system (for example, decreased number of IEL) (Dierick et al. 1979) and the consequent direct or indirect effects on the immune system (for example, increased villus height; decreased crypt depth) in the small intestine. It can be concluded that MCFA can be a valuable alternative to nutritional antibiotics because the same underlying factors affecting growth promotion are affected.

Metabolic and dietary constraints for the use of intact MCTAG in comparison with commonly used fats.

Intact medium-chain fatty acid-containing triacylglycerols for pregnant sows. No differences were found in the productive efficiency of sows and growth of the sucking pigs between feeding MCTAG or long-chain fatty acid-containing TAG (LCTAG) in late gestation (Averette Gatlin et al. 2002). However, Stahly (1983) and, more recently, Newcomb et al. (1991), Azain (1993) and Jean & Chiang (1999) found an increased survival of neonatal piglets by supplementing the sow diet with MCTAG in comparison with soyabean oil. They explained the effects by citing increased blood glucose levels, enhanced glycogen stores and maturity of the piglets at birth. The 4-fold increase of the MCTAG in the sows’ milk, which normally contains only traces of MCFA (Table 1), could also have been beneficial for the piglets in the pre-weaning period. However, the transfer of MCFA into milk fat by feeding sows with diets containing as much as 10 % MCTAG remains rather low (2 % of total milk fat) (Newcomb et al. 1991; Azain, 1993).

Intact medium-chain fatty acid-containing triacylglycerols for neonatal and pre-weaning piglets. In his excellent reviews, Odle (1997, 1999) concluded that MCTAG should have the desired characteristics to supplement the low energy reserves in neonatal piglets and to increase their survival. However, the narcotic effect of high doses of MCTAG might cause piglets to be less vigorous and thus increase their mortality, especially with newborn piglets weighing less than 1 kg at birth.

Intact medium-chain fatty acid-containing triacylglycerols for post-weaning piglets. Also for post-weaning piglets, in which a growth lag of more than 14 d, combined with a rise in mortality, is not uncommon, MCFA-containing fat sources may be beneficial, but again the results are rather inconsistent. With weaned piglets of 10 kg during a 21 d experimental period, Allee et al. (1972) did not find differences in feed intake, gain or gain:feed ratio with rations containing 10 % MCTAG compared with tallow, lard, coconut oil or maize oil. This is in accordance with the results of Newport et al. (1979) with diets containing 15 % MCTAG and pigs of 2 or 28 d old. Neither did Mahan

Table 8. Effect of feeding whole *Cuphea* seeds and lipase on the mean villus height (VH), crypt depth (CD) (μm) and number of intra-epithelial villous lymphocytes (IEL) in the proximal small intestine of weaned piglets (five piglets per treatment) (Expt III) (Dierick et al. 2003)

<table>
<thead>
<tr>
<th>Diet</th>
<th>Proximal†</th>
<th>Distal†</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>VH</td>
<td>CD</td>
</tr>
<tr>
<td>Control</td>
<td>381·9</td>
<td>244·2</td>
</tr>
<tr>
<td><em>Cuphea</em> + lipase§</td>
<td>365·7</td>
<td>201·0</td>
</tr>
<tr>
<td>Significance or P value</td>
<td>0·14</td>
<td>***</td>
</tr>
</tbody>
</table>

The mean value for the *Cuphea* + lipase diet was significantly different to that for the Control diet: *P* < 0·05, **P** < 0·01, ***P** < 0·001.
† Sampled sites were 3 m distal from the pylorus and 3 m proximal of the caecum.
‡ Number of IEL per 100 enterocytes.
§ The concentration of *Cuphea* seeds was 50 g/kg diet. Lipase ‘LS’ was of microbial origin (6563 U/g; Kemin Europa, Herentals, Belgium); concentration was 500 parts per million in diet.
(1991) find differences in performance comparing soyabean oil with coconut oil in the diet of early-weaned pigs. Furthermore, Fakler et al. (1992) did not observe differences in performance when weaned piglets were fed a diet containing 8 % MCTAG or 10 % soyabean oil. Recently, Léon et al. (1998) stated that the provision of MCTAG did not improve the energy status of the bottle-fed newborn piglet in comparison with LCTAG. On the other hand, Cera et al. (1989a) and Jin et al. (1998) found that coconut oil was superior for weaned piglets compared with soyabean oil, maize oil or tallow. These results agreed well with those of De Rodas & Maxwell (1990) who obtained a significantly increased average daily gain, feed intake and feed efficiency during the first week post weaning (weaning at 21 to 28 d) by feeding 4 % lard combined with 6 % MCTAG oils, compared with 10 % lard or butterfat. Finally, Dove (1993), feeding 5 % soyabean oil, MCTAG or animal fat, obtained the highest growth rate with the MCTAG source. There are discrepancies between the effects of MCTAG on pig performance noted in most of the cited literature and the results obtained in the authors’ research. These discrepancies may be related to the absence or very low levels of endogenous gastric or plant lipases, or to the fact that no exogenous lipases were added to the diets, resulting in levels of MCFA in the stomach and duodenum that were too low to influence the gut flora.

**Intact medium-chain fatty acid-containing triacylglycerols for growing pigs and pre-ruminant calves.** In growing pigs (30–90 kg), Glaps (1970) did not observe any difference in performance when feeding 2 ml MCTAG/d per kg compared with LCTAG. This was also the conclusion of Takada et al. (1992), feeding 8 % MCTAG in comparison with 8 % LCTAG. However, substituting part of the tallow in the milk replacer by tricaprin or tricaprylin, combined with coconut oil, for preruminant calves resulted in a 30–40 % increase in growth rate and energy efficiency (Aurossseau et al. 1984).

Feeding non-esterified MCFA instead of feeding intact (i.e. non-hydrolysed) MCTAG to weaned piglets had no effect (Cera et al. 1989b), but with growing pigs Rys et al. (1969/70) fed 5 % pure non-esterified MCFA (C5–C12) and obtained an increase in the growth rate of 6 %, in comparison with an isoenergetic control diet. The lack of a positive effect of non-esterified MCFA in young pigs may be related to the fatty acid level and profile of the fat sources used and to negative effects on feed intake, as mentioned earlier.

**Effects on product quality**

The concept of using TAG containing proper amounts of MCFA combined with exogenous lipolytic enzymes as an alternative to nutritional antibiotics has initially been targeted at piglet and early grower nutrition. However, applying higher doses of MCTAG, alone or in combination with lipases, with the objective to increase the energetic value of the pig feed, may selectively increase the firmness of the carcass fat by MCFA chain elongation (Takada et al. 1992). In order to avoid increases in the degree of saturation of the carcass fat in slaughter pigs and because the major changes in fatty acid composition due to diet influences will occur within 4–5 weeks (Wiseman & Agunbiade, 1998), high levels of MCTAG (4 % or more) are not recommended in the finishing phase of pigs.

Finally, some reports indicate that high levels of MCFA in the diet may reduce the deposition of fat and decrease protein catabolism in mammals, poultry and fish (Aurossseau et al. 1984; Crozier et al. 1987; Chiang et al. 1990b; Mabayo et al. 1993; Resjø et al. 2000). This may be related to the lower gross energy and net energy content (~20 %) of MCTAG (gross energy of 34·9 and net energy of 28·6 kJ/g) compared with LCTAG (Ingle et al. 1999).

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**Effects on the gastrointestinal mucosa and physiology**

Bacteria differ from eukaryotic cells in that they have rigid cell walls. Sheu et al. (1975) reported growth inhibition and morphological alterations in mammalian cell cultures (HeLa, human fibroblasts and mouse neuroblastoma cells) by millimolar concentrations of C6–C10 fatty acids, most probably due to alterations in cell attachment structures or processes. Nevertheless, Odle et al. (1991), studying the metabolism of even- and odd-C MCFA as Na salts (1 mmol/l) in isolated piglet hepatocytes, did not mention any alterations. However several authors warned of epithelial cell damage and disorders in phospholipid bilayers and membranes by MCFA and related compounds (Wargovich et al. 1984; Van Hoogdalem et al. 1989; Bergner & Sommer, 1994; Shima et al. 1998, 1999; Kimura et al. 2001). Kanai & Kondo (1979) concluded that, because MCFA and their acylglycerols are anionic surface-active compounds, in vitro cytotoxicity and membrane disorders and perturbations in tissue culture cells can occur. However, it is generally believed that the same events would not occur in living bodies in which various neutralizing agents such as serum, chyme and mucins are abundant. A higher susceptibility of isolated cells in vitro compared with tissue-associated cells in a natural environment was also observed with volatile fatty acids (Wächtershäuser & Stein, 2000).

Today, based on clinical observations in human subjects, it is clear that MCTAG and MCFA may have several positive effects on gut physiology. First, they improve intestinal morphology and function, through their positive effects on crypt cell activation and reactive villous hyperplasia (malnutrition, ageing) (Galluser et al. 1993; Jenkins & Thompson, 1993; Czernichow et al. 1996; Iba et al. 1998). Second, they have positive effects on epithelial cell membrane-bound enzyme activities (Takase & Goda, 1990). Third, they enhance absorption as measured in Caco-2 cell monolayers (Lindmark et al. 1995, 1998). Fourth, they are an excellent fuel as a source of acetate for small-intestinal enterocytes (Greenberger et al. 1965; Guillot et al. 1993) and a more suitable energy source than LCFA for epithelial cells in the treatment of Crohn’s disease (Andoh et al. 2000). As already mentioned, Odle (1997, 1999) did not indicate any deleterious effect of MCFA when studying their metabolism in isolated hepatocytes of neonatal piglets. Finally, Traul et al. (2000) reviewed the toxicological prop-
properties of MCTAG for pigs. There was no evidence that dietary administration of MCTAG adversely affected the reproductive performance of sows or resulted in maternal or fetal toxicity and teratogenic effects in doses up to 4 g/kg live weight per d in the diet, in accordance with the results of Hendrich et al. (1993) obtained with three generations of mice. For all these reasons, negative effects of MCFA on host epithelial cells are unlikely. This is in line with our results (Dierick et al. 2003) where an increased villus:crypt ratio and a lower number of IEL, indicative of a more healthy and better functional status of the mucosa, was observed when feeding a *Cuphea* + lipase-containing diet to weaned piglets. It can be argued that the combined use of TAG and lipases reduces the potential for tissue irritation and toxicity that could be produced by the ingestion of large amounts of ionized NEFA together with large quantities of damaging cations such as Na$^+$ (Bergner & Sommer, 1994; Wächtershäuser & Stein, 2000).

Although MCTAG and MCFA are “generally regarded as safe” (GRAS) and of benefit for oral and enteral use in human nutrition by the Food and Drug Administration in the USA, their effects on mucosal integrity need further *in vivo* research, especially with large doses or sudden incorporation in the diet.

**Gut flora, pathogens and resistance**

The antibacterial mechanism(s) of organic acids and SCFA are not fully understood and activity may vary depending on the growth phase of the organism and on environmental characteristics. An autolytic enzyme (autolysin) seems to be involved in the bacterial death and cellular lysis induced by MCFA (Tsuchido et al. 1985). The fate of MCFA, once inside the microbial cell, is not clear. According to Fay & Farias (1975), MCFA are not metabolised by *E. coli*. In contrast, Cherrington et al. (1991) claimed that Gram-negative bacteria are capable of metabolising MCFA. According to these authors, MCFA penetrate the membrane via porins and once inside the cell they should be degraded via the β-oxidation cycle, of which the necessary enzymes are induced by the acid.

At present there is some evidence that (potentially) pathogenic bacteria may be inactivated by MCFA or their monoacylglycerols (Kabara et al. 1972; Kabara, 1984; Isaacs et al. 1990, 1992; Boddie & Nickerson, 1992; Wang & Johnson, 1992; Guthery, 1993; Oh & Marshall, 1993; Kindlerer et al. 1996; Petrone et al. 1998; Petschow et al. 1998; Sprong et al. 2002).

An emerging potential problem is that organic acids and SCFA have been observed to enhance the survivability of acid-sensitive food-borne pathogens (*Salmonella, E. coli, Listeria, etc.*) exposed to low pH (gastric contents) by the induction of an acid-tolerance response, linked to an increased virulence (Ricke, 2003). According to Petschow et al. (1998), it is important to note that MCFA do not induce a notable resistance and, in any case, the induction of resistance must be very low in comparison with antibiotics. The exact mode of action of MCFA in inhibiting the growth and colonization of pathogens (*E. coli, Salmonella, Serpulina, Clostridium, etc.*) needs to be further elucidated, however.

**Role of preduodenal lipases in the release of medium-chain fatty acids in the stomach and proximal gut**

In most mammals there is a more or less pronounced preduodenal lipase activity, originating from lingual or gastric secretions. These lipases are active in a broad pH range with a preference for MCFA in milk fat (Table 2). Endogenous gastric lipase (optimal pH range 5–9; Höller, 1970) located in the cardiac region of the pig stomach, is thought to play a significant role in the hydrolysis of fats in the stomach but its quantitative contribution to overall fat digestion still remains to be elucidated. The enzyme seems to be resistant to acid and pepsin and its action is independent of bile acids or cofactors. The degree of fat hydrolysis (17 % NEFA in total fat, Table 4; 25–35 % NEFA in total fat, Table 5) noted in the stomach of piglets after feeding diets without added lipases is in accordance with the scarce literature data (Newport & Howarth, 1985; Chiang et al. 1990a). Jensen et al. (1997) followed the development of lipases in pigs and noted a pronounced decrease in pancreatic and an increase in stomach lipase activity (only 0.2 % of the pancreatic activity) in newly weaned piglets. Höller (1970) estimated the gastric lipase activity to be only 5 % of the pancreatic lipase activity, while Newport & Howarth (1985) reported that the total lipase activity in stomach tissue was only about 3 % of that found in the pancreas. Recently Li et al. (2001), investigating the development of lipase in nursing piglets, reported that total gastric lipase activity was fully developed on day 21, but reached only about 5 % of the pancreatic lipase activity at that time. However, the fact that fatty digesta are retained for a longer time in the stomach than in the small intestine could explain why gastric lipase, despite its low activity, may actually hydrolyse a considerable part of the fat in the stomach. Furthermore, Hunt & Knox (1986) found that fatty acids are more effective than the corresponding TAG in delaying gastric emptying and that increasing the chain length up to C14:0 (primarily MCFA) led to a progressively slower emptying. Gastric lipase also remains active in the duodenum, acting in synergy with pancreatic lipase (Edwards-Webb & Thompson, 1977).

However, it appears that this endogenous lipase activity in the stomach of piglets is too low (with a degree of hydrolysis of 15–30 %; Tables 4 and 5) to generate enough MCFA (0.025 m) to control the gut flora. This could be related to the lower (4–5) than optimal pH (5–9) for pig gastric lipase activity. As no contaminating bacteria were detected in any of the prepared feeds, lipolytic activity from microbial origin also seemed unlikely. Therefore, the addition of exogenous lipolytic activity, through lipase supplementation of the diet, with an optimal pH in the acid range (4–5), seems to be advantageous for releasing sufficient amounts of MCFA for an effective antimicrobial activity.

**Role of endogenous lipases, originating from raw materials, in the release of medium-chain fatty acids in the stomach and proximal gut**

Besides preduodenal lipases, a second source of endogenous lipases, originating from raw materials or prepared...
feeds, may interfere with the application of the concept. Information on endogenous lipolysis in raw and processed materials during storage is very scarce (O’Connor et al. 1992). Plant seeds store TAG in intracellular organelles called oil bodies or oleosomes, which are oil droplets covered by a coat of phospholipids and proteins serving as high-energy C reserves (Beisson et al. 2001). Lipolysis and subsequent rancidity caused by oxidation is usually not a problem in intact whole grains or seeds, stored at normal temperature (< 20°C) and moisture levels (< 12 %). During germination and in sprouted, cracked, broken or ground seeds, however, lipase activity may increase considerably. Subsequently, it is evident that the NEFA content of milled feedstuffs, raw materials and compound feeds and the subsequent lipolysis during the storage of compound feeds are important parameters, which will influence the subsequent nutritional and economic value of the fat and the feed.

From the authors’ own results, NEFA levels of more than 50 % in the lipid fraction were found in milled raw materials and in compound feeds, after a few weeks of storage. Normally endogenous lipolysis remains low (5–15 % NEFA in fat) in stored heat-treated cereals, in milk (products), fish products and Cuphea seeds (Dierick & Decuypere, 2002; Dierick et al. 2003). It is recommended that lipolysis should be prevented both in feedstuffs and compound feeds during storage. Indeed, most post-weaning diets are supplemented with fats and oils to increase palatability and energy intake. For the application of the proposed concept, preference should be given to heat-treated raw materials or mixed feeds (for example, pelleting, expansion, extrusion), showing no or low levels of endogenous lipase activity. Also an appropriate choice of the exogenous lipase used can greatly influence the subsequent lipolysis in the compound feed and is essential in order to prevent any aggravation of the inevitable endogenous lipolysis in the feed. The release of small amounts of MCFA during the storage of feed is not disadvantageous per se, however, because it can prevent the growth of deleterious microbial contaminants in the feed before ingestion. High amounts of non-esterified MCFA, however, may produce an adverse odour, as already mentioned.

The question may arise as to whether the lipase activity results, entirely or in part, from surface-associated microorganisms such as Penicillium, Pseudomonas and Candida, rather than from the feedstuff itself, as suggested by Pettersson et al. (1999). However, the total microbial count on the feeds and feedstuffs used in all our experiments (Dierick et al. 2002a,b) was about 1000 colony forming units/g. A simple calculation indicates that such populations are much too small to contribute significantly to lipolytic activity, which is in line with the results of Petersen (1999).

Our results clearly demonstrate that the activity of all those endogenous lipases (predouodenal, plant raw materials, in-feed microbes) in normal circumstances (fresh feed; piglets in post-weaning period) is too low for releasing sufficient amounts of MCFA for the antimicrobial management in the stomach and foregut of piglets.

Mode of application

The antibacterial activity of the classic non-esterified organic acids can be reduced by a decrease in feed intake, because some of them (for example, propionic acid) have a bad taste. This is very important in early-weaned pigs where feed intake is already seriously impaired. Another cause for reduced efficacy is the direct absorption in the stomach and upper small intestine (Clark et al. 1969; Dierick et al. 2002b). Moreover, the use of some of these acids (acetic acid, formic acid) is limited by problems of handling, strong odour and corrosion during feed processing and during its use on the farm. The use of specific preparations that gradually release the active acids (for example, micro-encapsulated or protected acids) at the desired site of action represents a strategy to overcome this problem (Cerchiarri, 2000). Taking all these arguments into consideration, a gradual enzymic release of MCFA, eventually together with MCFA-containing monoacylglycerols from TAG, in situ in the foregut, seems to be preferable to the supplementation of the feed with fixed doses of more common organic acids or MCFA or their salts. This mode of application avoids taste aversion, disturbance of the acid–base balance or possible mucosal damage, which can also decrease performance (Ostrowski et al. 1972).

Concluding remarks and needs for further research

During the last decade a lot of research has been directed to the use of MCTAG in human and animal nutrition, especially for piglets. Almost no research has been focused, however, on their use as a potential source of antimicrobial compounds, when liberated in situ in the stomach and foregut by appropriate lipases.

The present review illustrates similarly strong activity of MCFA and the classical antimicrobial growth promoters and therapeutics on the gut flora (total flora, Gram-positive flora, Gram-negative flora, potential pathogens) especially in the stomach and the foregut, as well as on the gut function of piglets. This means that these naturally occurring antimicrobial agents, which have little or no toxicity, can be an effective alternative to in-feed antibiotics.

A progressive enzymic release of MCFA, eventually combined with MCFA-containing monoacylglycerols from TAG, in situ in the foregut, seems to be preferable to the supplementation of the feed with non-esterified classical organic acids or non-esterified MCFA or their salts, avoiding taste aversion and disturbance of the acid–base balance in the animal.

However, when applying this concept, the choice of feedstuffs and lipases should be done very carefully and needs further exploration.

Because the recovery rates of bacteria by classical culturing methods, as compared with direct microscopic counts, have been reported to range from 30 to 60 %, more research is needed based on molecular techniques. There are three techniques used extensively in microbial ecology, based on the variability in the 16S rRNA-gene (rDNA) or on the use of specific primers and probes based on 16S rRNA: fluorescence in situ hybridization, quantitative polymerase chain reaction and denaturing gradient gel electrophoresis or tem-
perature gradient gel electrophoresis (Snel et al. 2002). These techniques most probably will provide more precise identification and enumeration of microbial populations, independent of cultivating on plates. They will enable a reassessment of the microbial ecology of the pig gastrointestinal tract (Simpson et al. 1999; Van Den Bossche et al. 2001; Leser et al. 2002; Akkermans et al. 2003) and the microbial populations as altered by growth-promoting antibacterials (Gaskins et al. 2002), including MCFA. As there are also limitations to these molecular techniques, a complete picture of the diversity and the role of the complex microbial ecosystem in the pig gut will need a combination of both classical and molecular techniques (Knarreborg et al. 2002).

The role and potential of MCFA in inhibiting the growth and colonization of an autochthonous microflora, including food-borne pathogens (E. coli, Salmonella, Clostridium, etc) needs further investigation, especially with regard to the possible induction of acid tolerance, mechanisms of resistance and linked virulence.

More attention should also be given to meat quality, especially when applying high doses of MCTAG in growing and finishing pig diets.

Alternatives to the common sources of MCTAG oils, which have the very serious constraint of being rather highly priced, should be explored.

Finally, as the concept is not limited to a specific medium, further development in warm-blooded production and companion animals and in cold-blooded animals as well as in plants for treatment of microbial infections awaits further research.

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Alternative to in-feed antibiotics in piglets


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