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## The Use of Medium-Chain Triglyceride in Children with Intestinal Malabsorption

by

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Medium-chain triglycerides (MCT) are fats in which the chain length is six to twelve carbon atoms. This is in contrast to the more usual dietary sources of fats, long-chain triglycerides (LCT), which contain more than 14 carbon atoms in their atomic chain. Most naturally occurring triglycerides are LCT and are found in common dietary constituents such as animal fats and dairy products. However coconut oil contains more than 10% MCT and is used as the raw material for the commercial production of MCT oils which are obtained by fractionation.

The differences in chemical structure between LCT and MCT are important because they are related to major differences in the characteristics of absorption and metabolism of these two classes of lipids (Figure 1).

Unlike LCT, significant absorption of MCT occurs in the absence of bile

salts and pancreatic lipase. Intestinal uptake and transport of MCT are rapid, even when very little intraluminal lipase is present (Isselbacher, 1966). MCT cross the epithelial cells of the gut rapidly, the later stages of absorption are different from LCT as most of the MCT travel directly to the portal blood stream without going through the steps of intracellular re-esterification and formation of chylomicrons (Bloom et al., 1951). After reaching the liver, MCT are rapidly oxidised to 2-carbon fragments which are available for different metabolic processes (Scheig, 1968).

These physiological differences between LCT and MCT have led to the use of MCT in various disorders where malabsorption of LCT occurs but where MCT can be absorbed because they bypass several of the

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steps necessary for the absorption of long chain fats. These include pancreatic disease, chronic liver disease, disorders of the biliary tract and diseases of the small intestinal epithelium and lymphatic vessels. Over recent years commercial preparations of MCT have become widely available, particularly in Europe, North America and Australia, and the usefulness of these preparations in various clinical situations has been well established (Holt, 1967; Senior, 1969). The place of MCT in paediatrics has also been reviewed recently (Gracey, Burke and Anderson, 1967). The comments which follow are a brief summary of the therapeutic place of MCT in children with malabsorption. This will be followed by some remarks about the potential value of MCT in countries like Indonesia where the commonest cause of chronic diarrhoea and malabsorption in children is malnutrition and chronic and repeated gastrointestinal infections and infestations and where MCT has not been used widely.

#### *Pancreatic disease*

In Westernized countries, such as Australia, the commonest cause of exocrine pancreatic insufficiency in children is cystic fibrosis of the pancreas in which there is usually severe steatorrhoea due to lack of pancreatic enzyme secretion. MCT are a useful additional source of calories for these children but do not appear to improve long-term weight gain or

eventual prognosis. In developing or pre-industrialized countries, malnutrition is probably the commonest cause of exocrine pancreatic insufficiency. As can be seen from Figure 2, an MCT diet leads to improvement of faecal fat excretion in children with pancreatic insufficiency and on theoretical grounds could be useful in various forms of malabsorption due to pancreatic disease, whatever the underlying cause.

#### *Liver disease*

Burke and Danks (1966) showed that steatorrhoea is common in infants and children with liver and biliary disease such as neonatal hepatitis, biliary atresia and cirrhosis and that MCT are effective in reducing the amount of fat lost in the stools in these patients (Figure 3).

#### *Small bowel diseases*

MCT are able to be absorbed rapidly by the small intestine despite the presence of diffuse mucosal disease and loss of surface area, as in coeliac disease. In some severely ill patients MCT may help induce weight gain whilst awaiting response to a gluten-free diet which is the specific treatment for this disorder. It can also be used as a source of calories in infants and children with intestinal malabsorption of uncertain aetiology where the main aim is to improve weight gain and the patient's clinical state before complete investigation can be undertaken safely.

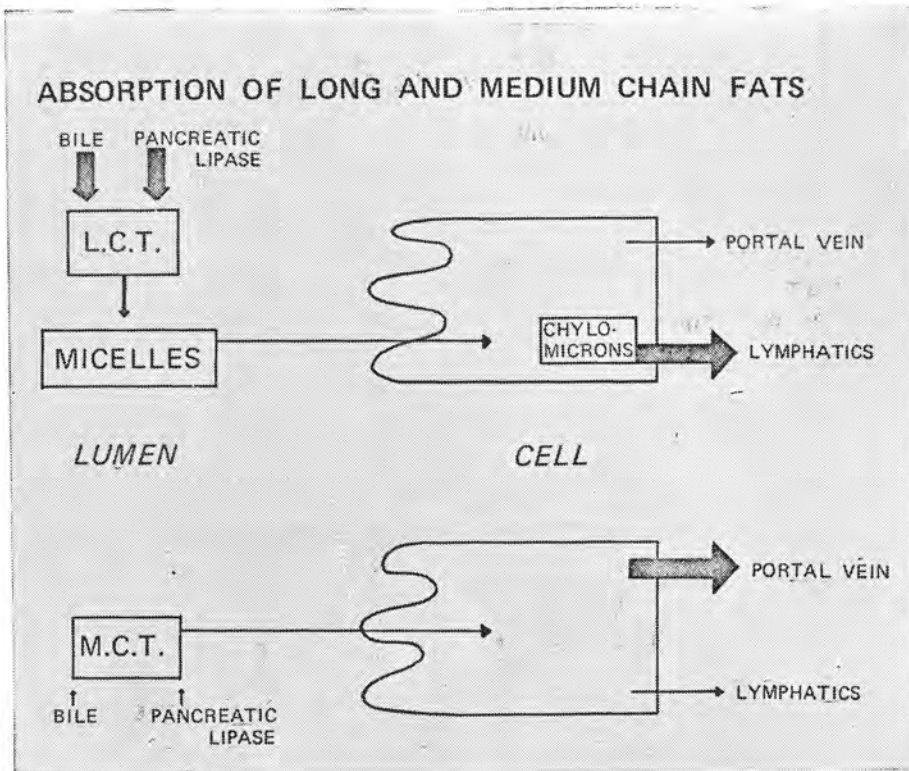


FIG. 1 : Differences between absorption of long-chain triglycerides (LCT) and medium-chain triglycerides (MCT) as explained in the text. Reproduced from the Australian Paediatric Journal with kind permission of the Editor and the authors, Burke and Anderson (1967).

CYSTIC FIBROSIS  
EFFECT ON MCT ON FAT EXCRETION

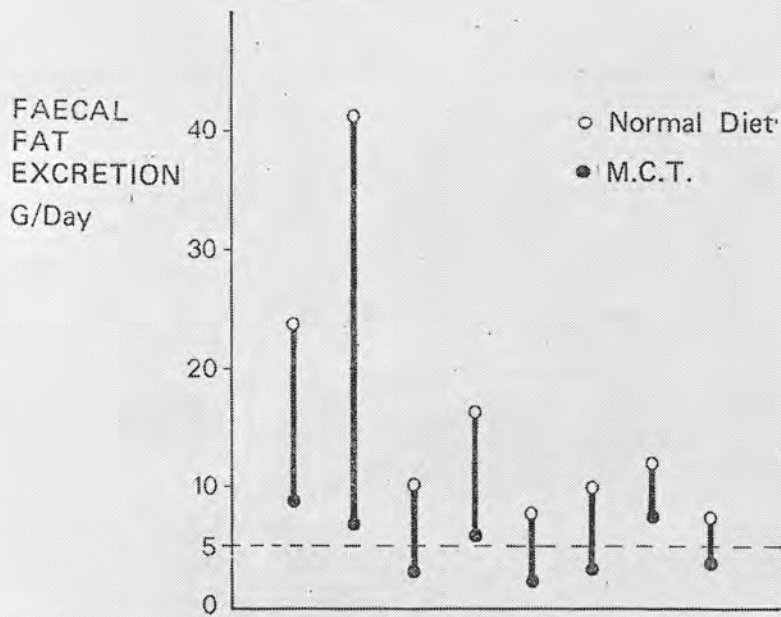


FIG. 2 : Decrease in stool fat excretion in children with cystic fibrosis of the pancreas when on an MCT-containing diet. Reproduced with kind permission of the Editor of the Australian Paediatric Journal and the authors, Burke and Anderson, 1967.

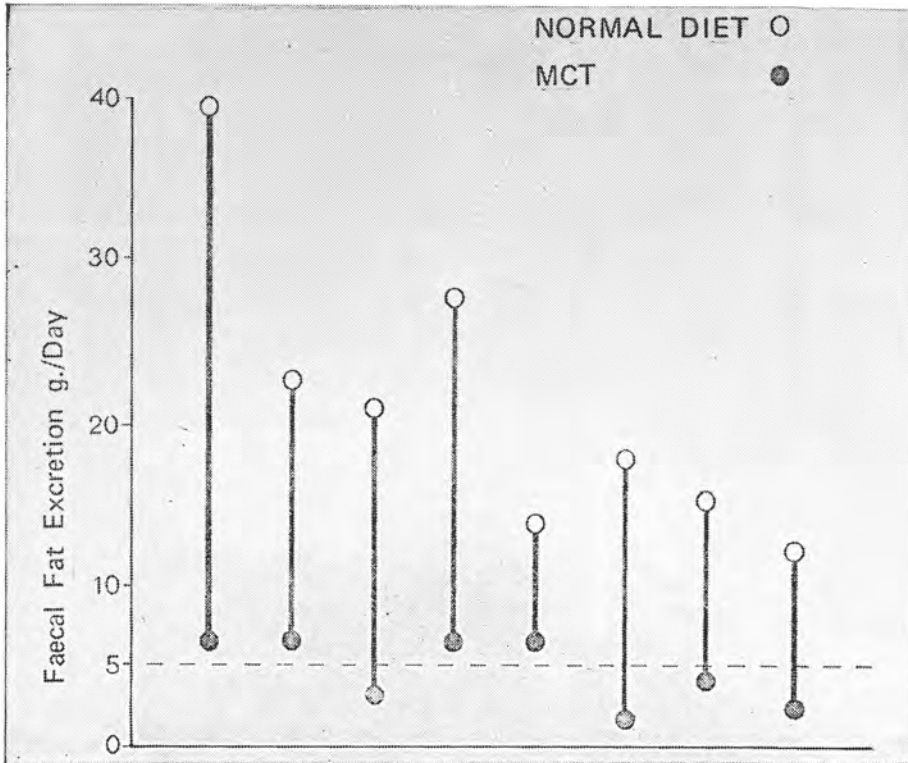


FIG. 3 : Reduction of stool fat excretion in four patients with biliary atresia (left) and four patients with neonatal hepatitis when put on an MCT-containing diet. Reproduced with kind permission of the Editor the Australian Paediatric and the authors, Burke and Anderson, 1967.

MCT have also been useful in the dietary management of young patients following extensive intestinal resection which can lead to malnutrition, impaired growth and deficiency states (Burke and Anderson, 1967). When using MCT preparations in patients following gastrointestinal surgery, particular attention must be paid to the sugar content of the formula as these patients may have sugar intolerance which will cause severe diarrhoea if it is not recognized and treated by removal of the affected sugar from the patient's diet—in most cases lactose will be the sugar not tolerated and a lactose-free, MCT-containing formula will need to be used. Such preparations are available commercially.

#### *Lymphatic diseases*

As mentioned above, MCT are not transported through the intestinal lymphatics but travel to the portal blood stream. An MCT diet may, therefore, be useful in lymphatic disorders such as congenital lymphangiectasia or acquired (e.g. trauma or tuberculosis) of the intraabdominal lymphatic vessels. (Tamir et al., 1968; Gracey and Gooch, 1968). In such situations MCT will be absorbed into the portal blood and transported to the liver whereas LCT would either leak out of the damaged lymphatic vessel or would not be absorbed because of blockage of the lymphatic drainage system.

#### *MCT in children with malnutrition*

Diarrhoea is a characteristic feature of malnourished children; its pathogenesis is complex and multifactorial.

Included amongst these factors are:

- (i) histological abnormalities of the small intestinal mucosa,
- (ii) secondary disaccharidase deficiency caused by the histological abnormalities,
- (iii) impaired absorption of nutrients,
- (iv) repeated infections and infestations in the gut,
- (v) exocrine pancreatic insufficiency,
- (vi) liver dysfunction, and
- (vii) microbial contamination of the upper gut.

In a recent study (Gracey et al., in press) steatorrhoea was found commonly in young Indonesian children with protein-calorie malnutrition, using the short term lipiodol absorption test as a screening procedure for malabsorption of fat. Although it is not possible to say which factors are most important in causing malabsorption in these children, it is reasonable to suspect that a combination of small bowel disease and impairment of hepatic and pancreatic function are involved. On theoretical grounds, it could be anticipated that MCT would be absorbed much better than LCT in such children and that MCT would be a useful source of dietary calories in the early

stages of nutritional rehabilitation in these patients. It must be remembered, of course, that these children have a high incidence of sugar intolerance as well as malabsorption of fat and this must be taken into account when planning their dietary management; because lactase deficiency is so common in these children feeding formulae containing a significant quantity of lactose ought to be avoided. Recently, a lactose-free, MCT-containing preparation has been used with success in malnourished children in South America (Graham et al., 1973). Sutedjo (1974) has very recently advocated the use of MCT in malnourished children in Indonesia. Admittedly, the limiting factor

about this suggestion is the high cost of MCT preparations and their relative unavailability. Despite this, it seems desirable that carefully planned clinical trials should be conducted to test the therapeutic value of MCT in this part of the world. Furthermore, as pointed out by Sutedjo (1974) attempts should be made to provide predominantly MCT-containing formulae from vegetable oils, such as coconut oil, which are readily available in Indonesia and should, therefore, be much less expensive. It is hoped that the technical expertise of the pharmaceutical industry will accept this important challenge to their inventiveness.

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ful pregnancy and delivery. All other children were considered to be normal throughout their neonatal period and infancy except for this patient who had a chronic and severe rhinitis in infancy. On admission the patient denied a past history of renal disease, edema, hematuria, hematuria, diabetes, diabetes, toxin exposure, apathetic eruption or thrombosis.

Three months prior to admission she had developed a rapid succession of enlargement of her abdomen, oliguria, and the passage of cloudy and reddish urine. She was then admitted to a small hospital in Tjuring Rejo. Because there was no improvement after one month therapy and hospitalization she was referred to Surabaya.

On admission she weighed 12.6 kg. The blood pressure was 140/110.

Among the diseases considered to be the possible etiologic causes of nephrotic syndrome are renal vein thrombosis and congenital syphilis (Hill et al., 1972; Straus and Well, 1971). Renal involvement in syphilis has been recognized for at least 150 years. Scattered reports of congenital syphilis presenting as nephrotic syndrome or nephritis in children have appeared in the medical literature (Bhobade et al., 1971; Papanicolaou et al., 1961).

Underneath is a report of a case with nephrotic syndrome that might be due to congenital syphilis.

#### Case Report

A 9-year-old Indonesian girl was admitted to the Dr. Soetomo Hospital, Surabaya, on September 15, 1973. The patient, the eldest of five children was the product of an uneven

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