Relationship between plasma and lymphocyte linoleate in multiple sclerosis

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SYNOPSIS The linoleate level in total lipids was measured in lymphocytes from control subjects and patients with multiple sclerosis. A small but significant decrease was found in cases of multiple sclerosis. The percentage composition of lymphocyte fatty acids was determined in rats fed diets with various linoleate contents. There was a correlation between lymphocyte linoleate and plasma linoleate in both humans and rats.

Previous work from this laboratory has shown that in patients with multiple sclerosis there is a lowering of the linoleate levels in platelets and erythrocytes (Gul *et al.*, 1970) in addition to a lowering of the linoleate level in the plasma lipids (Belin *et al.*, 1971), the latter result confirming earlier work (Baker *et al.*, 1964; Tichy *et al.*, 1969) and according with more recent reports (Love *et al.*, 1974; Kalofoutis and Jullien, 1974; Crawford and Hassam, 1975; Paty *et al.*, 1975), although some authors have failed to find lower linoleate in multiple sclerosis patients (Cumings *et al.*, 1965; Karlsson *et al.*, 1971; Wolfgram *et al.*, 1975). We have now extended the studies on linoleate levels to lymphocytes and to the free (non-esterified) fatty acids of the plasma. This study is relevant to reports that free fatty acids, especially when they are polyunsaturated, can inhibit certain *in vitro* activities of lymphocytes (Mertin *et al.*, 1974; Offner and Clausen, 1974) and *in vivo* can prolong skin graft rejection in rodents (summarised in Hughes *et al.*, 1975). Mertin *et al.* (1973) suggested that *in vivo* these unsaturated acids may play an immunoregulatory role, which may form the basis of the beneficial effect of dietary sunflower seed oil in the treatment of multiple sclerosis (Millar *et al.*, 1973).

Further support for the concept that a relationship may exist between fatty acid status and immunological activity arises from work with animals. It has been shown (Clausen and Møller, 1967; Selivonchick and Johnston, 1975) that animals fed a diet deficient in essential fatty acids (EFA) are more susceptible to the development of experimental allergic encephalomyelitis than those on normal diets and that the addition of linoleate to the diet exerted a pronounced protective effect.

In view of these various findings we thought that it would be valuable to examine the levels of linoleate in the lymphocytes and non-esterified fatty acids of patients with multiple sclerosis and to establish the levels of linoleate in the lymphocytes of rats fed diets in which the linoleate was varied over a wide range.

METHOD

PREPARATION OF HUMAN LYMPHOCYTES Fifty millilitre blood samples were drawn from patients with multiple sclerosis who had fasted overnight, using heparin as anticoagulant. So far as could be ascertained the patients were not supplementing their diet with polyunsaturated fatty acid. No account was taken in the study of the state of the disease. Lymphocytes were prepared by the method of Bøyum (1968). To remove platelets the lymphocytes were washed three times in 0.9% saline, being centrifuged at 100 g for 10 minutes between washes.

PREPARATION OF FATTY ACID FRACTIONS Lipids were extracted from the lymphocyte pellet after the final saline wash by resuspending the pellet in 0.5 ml saline and then using the extraction procedure of Folch *et al.* (1957). After taking the chloroform layer to dryness the lipid fraction was saponified and the fatty acid composition determined, as described by Belin *et al.* (1971). To obtain the free fatty acids 5-ml samples of plasma were extracted by the method of Hagenfeldt (1966). The methods previously described (Belin *et al.*, 1971) were used for the methylation of the free fatty acids and also for the estimation of the total fatty acids.

EXPERIMENTS WITH RATS The animals were of the Wistar strain. The mothers were fed a fat-free diet (Nutritional Biochemicals) from the time of mating. At weaning the rats were divided into three groups and fed the fat-free diet supplemented with oil as follows: group A, 10% hydrogenated coconut oil; group B, 9% hydrogenated coconut oil plus 1% sunflower seed oil; group C, 10% sunflower seed oil. Control rats were fed the normal cubed diet *ad libitum*.

The animals were killed when aged 34–36 weeks. Lymphocytes were prepared from the spleen by gentle teasing with forceps in RPMI Medium 1640, then layered on to Lymphoprep (Nyegaard & Co., Oslo), and centrifuged at 400 *g* for 10 minutes. The cells at the interface were collected and washed three times with RPMI medium 1640 (Flow Laboratories), the first centrifugation being at 180 *g* for 10 minutes and the subsequent two (to remove platelets) at 100 *g* for 10 minutes. The pellet was treated in the same way as described above for human lymphocytes.

Rat blood was drawn by heart puncture under light ether anaesthesia and treated for fatty acid determination in the same way as described for human blood.

RESULTS

OBSERVATIONS IN PATIENTS The level of total plasma linoleate when compared with 14 normal control subjects was again found to be significantly lower in this sample of 19 patients (Table 1), who were randomly selected apart from the need to take those who were not using oil supplements in the diet. A decrease in linoleate was also found in the free fatty acids. Again, a small but significant decrease was found in the lymphocyte fraction. As might be expected, a very close correlation was found

TABLE 1
MEAN ($\pm$ SD) PERCENTAGE OF LINOLEATE IN SERUM TOTAL LIPIDS, SERUM FREE FATTY ACIDS, AND LYMPHOCYTES IN CONTROL SUBJECTS AND PATIENTS WITH MULTIPLE SCLEROSIS

	Controls	MS patients	P
Total lipids	35.0 $\pm$ 1.0 (14)	28.8 $\pm$ 0.77 (19)	< 0.001
Non-esterified fatty acids	17.0 $\pm$ 1.1 (14)	12.1 $\pm$ 0.50 (18)	< 0.001
Lymphocytes	10.6 $\pm$ 0.64 (12)	8.3 $\pm$ 0.33 (17)	< 0.005

between the linoleate levels in the free and total fatty acids (Fig. 1).

When a scatter diagram was prepared of the lymphocyte linoleate levels against the serum free fatty acid linoleate levels it was found that if results from the multiple sclerosis patients were plotted alone a poor correlation resulted because of the narrow range over which the values extended. The same applied if the control results were plotted alone. When control and patient results were plotted together, however, good correlation resulted, which was further strengthened when results from a few patients and controls who had taken oil were added (Fig. 2). It thus appears that there is a correlation

FIG. 1 Percentage linoleate in serum total lipids and in serum free fatty acids in humans.

FIG. 2 Percentage linoleate in serum free fatty acids and in lymphocyte total lipids in humans. ○ Controls. □ Controls on oil. ● MS. ■ MS on oil.

($r=0.62$) between linoleate levels in the serum free fatty acids and in the lymphocytes which can be demonstrated provided values extend over a wide enough range. A good correlation also exists between linoleate levels in the blood total fatty acids and in the lymphocytes ($r=0.66$).

EXPERIMENTS WITH RATS As support for the work with human lymphocytes and as the basis for a projected study of the relationship between reduced linoleate levels and sensitivity to the encephalitogenic protein, comparable studies of the lymphocyte linoleate level were made in rats on appropriate dietary regimes. The percentage

compositions of the lymphocyte fatty acids as a result of the different dietary treatments (EFA deficient or supplemented with 1% or 10% sunflower seed oil) are shown in Table 2. The linoleate level can be varied from about 2–20% in animals fed the diets for many months. We are currently investigating the extent to which these changes can be brought about over shorter periods.

Over this range of values a close correlation ($r=0.93$) is again found between linoleate levels in lymphocytes and blood (Fig. 3). It is perhaps not surprising that if the animal is severely deprived of linoleate the acid will be lacking in both lymphocytes and blood. However, even using the results from animals fed sufficient linoleate in the diet—that is, omitting the results from those animals on the EFA deficient diet—a highly significant correlation is found ($r=0.75$).

DISCUSSION

These and previous results show that lowered linoleate levels are found in the plasma fatty acids, both esterified and free, and in three of the major blood cell types in patients with multiple sclerosis.

The finding, reported for the first time, of a decrease of the linoleate level in the free fatty acids may well be of importance because these are the more likely to influence the level of linoleate in the membrane, since it seems from work with cells in culture that they derive their membrane fatty acids primarily from free fatty acids, although other types of lipids make a

TABLE 2

MEAN ($\pm$ SD) WEIGHT PERCENTAGE COMPOSITION OF LYMPHOCYTE FATTY ACIDS IN RATS ON DIFFERENT DIETS

Diet	No. of rats	Fatty acid						
		Palmitic	Palmitoleic	Stearic	Oleic	Eicosatrienoic	Linoleic	Arachidonic
Normal	5	30.5 ± 0.31	3.9 ± 0.24	17.6 ± 0.36	18.2 ± 0.67	—	12.3 ± 0.44	17.5 ± 0.67
Diet A (deficient)	3	27.3 ± 0.71	6.8 ± 0.24	16.2 ± 1.23	28.4 ± 0.72	13.0 ± 0.11	2.2 ± 0.4	6.0 ± 0.69
Diet B (1% sunflower seed oil)	5	28.0 ± 0.51	5.0 ± 0.32	18.2 ± 0.62	17.3 ± 0.89	—	13.3 ± 0.43	18.2 ± 1.75
Diet C (10% sunflower seed oil)	5	27.2 ± 0.70	3.3 ± 0.38	16.9 ± 0.90	14.9 ± 0.62	—	18.2 ± 1.0	19.5 ± 0.74

FIG. 3 Percentage linoleate in serum free fatty acids and lymphocyte total lipids in rats on diets of varying linoleate content. □ Normal diet. ● Deficient diet. ■ Deficient diet supplemented with 1% sunflower seed oil. ▲ Deficient diet supplemented with 10% sunflower seed oil.

small contribution (Bailey *et al.*, 1973). Furthermore, the observed changes in linoleate in the free fatty acids are more relevant to the *in vitro* conditions used by Mertin *et al.* (1974) in their study of the effect of polyunsaturated fatty acids on lymphocyte function.

The question remains to be resolved whether the low levels in the cells arise because of the low plasma levels, since, as we have shown here and previously (Gul *et al.*, 1970), there is a correlation between levels in plasma and each of these cell types, or whether there is some other cause of the low linoleate levels in cells.

The results further show that lymphocyte linoleate levels are susceptible to change by alteration in diet, and in particular that the feeding of sunflower seed oil to patients will

increase the level of linoleate in their lymphocytes. To what extent this is the basis of the beneficial effects reported for the oil is another question to be resolved in the future. It will also be of interest to investigate the effect of these changes on a number of lymphocyte function tests with a view to assessing the extent to which it may be possible to modify by dietary supplementation with linoleate lymphocyte reactivity in patients with multiple sclerosis.

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