

1954, v66, 776.

10. Coates, M. E., Davies, M. K., Dawson, R., Harrison, G. F., Holdsworth, E. S., Kon, S. K., and Porter, J. W. G., *Biochem. J.*, 1956, v64, 682.

11. Bernhauer, K., Blumberg, K., and Petrides, P., *Arzneim. Forsch.*, 1955, v5, 442.

12. Brown, F. B., Cain, J. C., Gant, D. E., Parker,

L. F. J., and Smith, E. L., *Biochem. J.*, 1955, v59, 82.

13. Briggs, G. M., and Fox, M. R. Spivey, *PROC. SOC. EXP. BIOL. AND MED.*, 1955, v89, 318.

14. Holdsworth, E. S., and Coates, M. E., *Nature*, 1956, v177, 701.

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Effects of Unsaturated Fat on Serum Lipids in Idiopathic Hyperlipemia. (23266)

MARK ALLEN EVERETT, WALTER D. BLOCK, F. A. J. KINGERY, AND ARTHUR C. CURTIS

Department of Dermatology and Syphilology, Medical School, and Institute of Industrial Health, University of Michigan, Ann Arbor.

It has been known for some time that dietary fat influences serum lipid levels. Beveridge *et al.*(1) found that diets high in animal fat produced elevations in serum lipid levels, while diets high in fats of vegetable origin did not. Ahrens and his associates(2) also produced lower serum lipid levels with vegetable oils, especially corn oil. The serum lipid lowering effect of vegetable fat has been attributed by Kinsell(3) and more recently by Bronte-Stewart(4) to its higher content of unsaturated fatty acids. Bronte-Stewart(4, 5), in particular, has shown that the lipid lowering effect of administered fats in normal individuals may be directly proportional to the unsaturated fatty acid content of the diet. It was reported by Lever(6), and confirmed in this laboratory(7), that the intravenous administration of a cottonseed oil preparation (Lipomul I-V)* on some occasions led to a drop in serum lipids in patients with idiopathic hyperlipemia. Administration of the oral product (Lipomul Oral)* failed to lower serum lipids. Since the oral preparation is primarily coconut oil, containing almost no unsaturated fatty acids, this may explain the lack of success with the oral product. According to Deuel(8), the iodine number of coconut oil is 8-40, while that of cottonseed oil is 105-115.

Methods. Two patients with idiopathic hyperlipemia were placed on diets carefully

controlled with respect to fat content. One patient, E.B., was a male 29 years old, the other, D.L., a male 31 years old. Neither patient was hospitalized during the course of the experiments. The following diets were used in this study: Diet I, uncontrolled with respect to fat content; and Diet II, which contained 120 g fat consisting of 50 g of highly purified soybean oil,[†] 30 g of soy phospholipid concentrate,[‡] and 40 g animal fat. Diet III, administered to only one patient (D.L.) had no added soy phospholipid concentrate. This diet, which was isocaloric with Diet II, contained 50 g of soybean oil,[†] 40 g of animal fat, and 10 g of vegetable fat. The caloric content of all diets was 2400-2600. Serum lipid partitions were done on these patients on fasting blood samples taken

[†] Supplied by the Procter and Gamble Co., Cincinnati, O. This soybean oil had the following composition:

Iodine No.	131.9
Total fatty acids	95.9 %
Free " "	.0
OH number	1.6
Unsaponifiable	.5
Tocopherol	.12
Trans fatty acids	.2

Spectrographic analysis yielded the following values:

Saturated fatty acids	16.6 %
Oleic acid	21.2
Linoleic acid	50.4
Linolenic acid	7.7

[‡] Purchased from Associated Concentrates, Woodside, L. I., N. Y.

* Supplied by the Upjohn Co., Kalamazoo, Mich.

TABLE I. Effect of Unsaturated Fats on Serum Lipids. Patient E. B.

	Diet I*	Diet II*				Diet I*	
	10/15	(Fed 10/16-12/22)				(Fed 12/22-3/1)	
		11/3	12/3	12/21		1/31	3/21
Total lipid†	2137	1314	1219	1012		1454	1705
Total cholesterol†	560	471	302	274		448	478
% cholesterol esters	77	81	76	80		77	80
Phospholipids†	413	418	366	263		369	463
Triglycerides†	888	184	395	325		403	505

* See text for description of diets.

† Values expressed as mg %.

at intervals before, during, and after the administration of the special diets. Total lipids were determined by the method of Bragdon (9), total cholesterol and cholesterol esters by the Schoenheimer-Sperry method (10), and phospholipids by the Fiske-Subbarrow method (11).

Results. The serum lipid values resulting from the present study are listed in Tables I and II. It can be seen from Table I that the first patient, E.B., showed a marked drop in serum values for total lipid, total cholesterol, phospholipid, and triglycerides during the period of administration of Diet II. When the patient resumed eating Diet I, these values all showed marked increases. Although the absolute values did change, there was no significant change in percentage of cholesterol present in esterified forms.

Table II shows the results obtained on the second patient, D.L. The same qualitative picture is repeated. The extraordinarily high total lipid values obtained on this patient under Diet I showed marked decreases during the administration of Diet II. Resumption of Diet I produced very great elevations in all serum lipid values studied. This patient was then placed on Diet III, which contained high levels of unsaturated fatty acids, but without added phospholipid concentrate.

During the period of Diet III, the patient again demonstrated marked decreases in all the serum lipid fractions studied. It should be noted that this patient also demonstrated no significant change in the percentage of cholesterol present in esterified forms. Both patients had opaque, milky sera, which cleared during the periods on Diets II and III. Both patients had tuberous xanthomata of the extensor joint surfaces. One patient, D. L., had generalized papular or eruptive xanthomas and lipemia retinalis. The papular xanthoma and lipemia retinalis disappeared, and the tuberous xanthoma diminished in this patient during the administration of Diet II. Patient E.B. showed some disappearance of the tuberous xanthomata.

Discussion. This dietary study would seem to support the concept that the lipid lowering effect of administered fats may be dependent on the unsaturated fatty acid content of these fats. The recent findings of Bronte-Stewart (4,5) suggest that unsaturated dietary fat may prevent elevations of serum lipid in the presence of moderately large amounts of dietary animal fat. We believe that the study of hyperlipemic patients provides a very convenient means of evaluating this hypothesis, because of the very high initial serum lipid values found in idiopathic

TABLE II. Effect of Unsaturated Fats on Serum Lipids. Patient D. L.

	Diet I*	Diet II*		Diet I*	Diet III*
	10/15	(Fed 10/16-12/21)		(Fed 12/21-2/1)	(Fed 2/1-3/1)
		11/21	12/20	1/31	3/1
Total lipid†	20316	1985	3627	23327	3724
Total cholesterol†		454	489	2344	582
% cholesterol esters		75	72	71	71
Phospholipids†		429	728	2104	877
Triglycerides†		871	2171	17741	1984

* See text for description of diets.

† Values expressed as mg %

hyperlipemia.

Summary. Two patients with idiopathic hyperlipemia were placed on 2400-2600 calorie diets containing 50 g of highly purified soybean oil, with or without the addition of a phospholipid concentrate. Marked decreases in serum lipids occurred during the period of administration of both diets. The serum lipid decreases were independent of phospholipid concentration in the diet, and believed attributable to the presence of unsaturated fatty acids in the dietary fat. Clinical signs of the disease regressed or disappeared coincident with the decrease in serum lipids.

1. Beveridge, J. M. R., Connell, W. F., Mayer, G. A., Firstbrook, J. B., and DeWolfe, M. S., *J. Nutrition*, 1955, v56, 311.

2. Ahrens, E. H., Jr., Tsaltas, T. T., Hirsch, J.,

Insull, W., Jr., *J. Clin. Invest.*, 1955, v34, 918.

3. Kinsell, L. W., Michaels, G. D., Cochrane, G. C., Partridge, J. W., Jahn, J. P., and Balch, H. E., *Diabetes*, 1954, v3, 113.

4. Bronte-Stewart, B., Antonis, A., Eales, L., and Brock, J. F., *Lancet*, 1956, v270, 521.

5. Bronte-Stewart, B., Moodie, A. D., Antonis, A., Eales, L., and Brock, J. F., *African Med. J.*, 1955, v29, 1151.

6. Lever, W. F., and Waddell, W. R., *J. Invest. Derm.*, 1955, v25, 233.

7. Everett, M. A., Block, W. D., and Curtis, A. C., unpublished data.

8. Deuel, H. F., Jr., *The Lipids*, Interscience Publishers, N. Y., 1951, v1, p197.

9. Bragdon, J. H., *J. Biol. Chem.*, 1951, v190, 513.

10. Schoenheimer, R., and Sperry, W. M., *ibid.*, 1954, v106, 745.

11. Fiske, C. H., and Subbarow, Y., *ibid.*, 1925, v66, 375.

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Effect of 5-Hydroxytryptophane Upon Electroencephalogram in Hepatic Coma.* (23267)

S. P. BESSMAN, J. K. MERLIS AND F. BORGES

Department of Pediatrics, University of Maryland School of Medicine.

Liver failure is usually associated with serious disturbances of consciousness. These episodes are accompanied by electroencephalographic changes of a diffuse sort with an increase in amount and amplitude of slower frequencies and a diminution of faster frequencies. Some investigators have reported what they considered to be characteristic neurologic signs(1) and electroencephalographic patterns(2-3) in liver disease, but these have not been observed uniformly(4). Observations of increased ammonia content of blood in liver failure(5-6), caused by inability of the liver to remove or "detoxify" ammonia from the portal blood because of shunt or deficiency of the urea synthesizing system have led to an hypothesis for the mechanism of ammonia-genic coma(7). The liver, however, does far more than detoxify substances. It has been pointed out recently that some metabolites

essential to brain are probably synthesized in the liver and that liver failure may lead to serious deficiency of these substances(8). Among these may be the precursor material for serotonin, which appears to play a significant role in brain metabolism(9-11). Parenteral injection of moderate amounts of serotonin causes no change in the brain content of this intermediate, but administration of small amounts of a precursor substance, 5 hydroxytryptophane, (5HTP), does cause a large increase of serotonin in brain(12-13) together with symptoms similar to those seen after a block in serotonin metabolism in brain (14) caused by lysergic acid diethylamide. Although the synthesis of 5 HTP in liver preparations has not been demonstrated, other hydroxylations of a similar nature have occurred in that organ. Indirect evidence for failure of supply of the serotonin precursor comes from observations that urinary excretion of 5 hydroxyindole acetic acid, (5HIAA),

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