

The Effect of Estrogen on the Lipoprotein Lipase Activity of Rat Adipose Tissue

MARGIT HAMOSH and PAUL HAMOSH

From the Department of Anatomy and Department of Physiology & Biophysics, Georgetown University School of Medicine, Washington, D. C. 20007

ABSTRACT The effect of 17β -estradiol or progesterone administration on adipose tissue lipoprotein lipase activity was studied in male and ovariectomized female rats. Lipoprotein lipase activity was measured in acetone-ether-extracted preparations of adipose tissue with doubly labeled (^{14}C -fatty acid, ^3H -glyceryl) chylomicron triglyceride as substrate. Administration of 17β -estradiol to male rats lowered adipose tissue lipoprotein lipase activity from 8.22 ± 1.8 U/g (1 U = 1 μmol triglyceride hydrolyzed per h) to 4.96 ± 0.5 U/g in the treated group. Ovariectomy increased adipose tissue lipoprotein lipase activity from 10.4 ± 1.8 U/g in controls to 22.7 ± 4.3 U/g. 17β -Estradiol administration to ovariectomized rats caused a marked fall in adipose tissue lipoprotein lipase activity: 17β -estradiol (2.5 $\mu\text{g}/\text{day}$) lowered the enzyme activity to 9.00 ± 1.2 U/g, whereas 25 $\mu\text{g}/\text{day}$ further decreased lipoprotein lipase activity to 3.2 ± 0.6 U/g. Blood triglyceride levels increased from 0.8 ± 0.05 $\mu\text{mol}/\text{ml}$ in ovariectomized rats to 1.4 ± 0.09 $\mu\text{mol}/\text{ml}$ in 25 $\mu\text{g}/\text{day}$ 17β -estradiol-treated rats. Progesterone administration did not affect adipose tissue lipoprotein lipase activity in either male or ovariectomized rats. Heart and lung lipoprotein lipase activity was unaffected by hormone treatment. We suggest that the rise in blood triglyceride concentrations, which accompanies high plasma estrogen levels, could be due to the marked inhibition of adipose tissue lipoprotein lipase activity.

INTRODUCTION

There is now considerable evidence that plasma triglycerides are hydrolyzed to free fatty acids during their removal from blood. This process is catalyzed and regulated by lipoprotein lipase, an enzyme active in or near the capillary wall (1).

This work was presented in part at the 1975 meeting of the Southern Section of the American Federation for Clinical Research, New Orleans, La.

Received for publication 16 November 1974 and in revised form 13 February 1975.

Lipemia of pregnancy, which is essentially a rise in plasma triglyceride concentration, is related to changes in the level of lipoprotein lipase activity in adipose tissue and mammary gland in the rat (2, 3). A few days before parturition, during the peak of hyperlipemia adipose tissue lipoprotein lipase is almost completely inhibited. Disappearance of the lipemia just before parturition is directly related to a marked rise in mammary gland lipoprotein lipase activity. In addition to pregnancy, increased plasma triglyceride levels have been reported in women on mixed contraceptives (4, 5); the lipemia seems to be caused by the estrogen component only (5).

There are marked changes in blood estrogen concentration during the estrous cycle in the rat (6). A comparison of lipoprotein lipase activity in adipose tissue of male with that of female rats showed that in males the activity level was almost constant, whereas wide fluctuations were observed in females.¹ This observation raised the possibility that female sex hormones might affect the activity of adipose tissue lipoprotein lipase.

METHODS

Animals. Adult male and female Sprague-Dawley rats delivered by cesarean section were used in this study. The rats were purchased from Charles River Breeding Laboratories, Inc., Wilmington, Mass. The animals were caged individually and had free access to food (Purina Laboratory Chow, Ralston Purina Co., St. Louis, Mo.) and water.

Hormone preparations. 17β -Estradiol (Depo-estradiol cypionate) and progesterone (Depo-Provera [medroxyprogesterone acetate suspension U.S.P.]) were purchased from the Upjohn Co., Kalamazoo, Mich. Male rats were given the hormones by intraperitoneal injection once weekly for 8 wk: 17β -estradiol 50 μg , or progesterone 5 mg, in 0.1 ml saline. Hormone treatment of ovariectomized rats was started 2 wk after surgery. The rats were given 17β -estradiol or progesterone in 0.1 ml corn oil by subcutaneous

¹Hamosh, M., P. Hamosh, and R. O. Scow. Unpublished observations.

TABLE I
Effect of 17 β -Estradiol* and Progesterone† on Adipose Tissue Lipoprotein Lipase Activity of Male Rats

| Treatment‡ | No. of animals | Lipoprotein lipase activity¶ U/g |
|-----------------------|----------------|-------------------------------------|
| None | 12 | 8.22±0.88 |
| 17 β -Estradiol | 12 | 4.96±0.52** |
| Progesterone | 12 | 8.36±0.67 |

Results are mean±SE.

* 17 β -Estradiol was injected i.p. 50 μ g in 0.1 ml once weekly.

† Progesterone was injected i.p. 5.0 mg in 0.1 ml once weekly.

‡ Hormones were administered for a period of 8 wk.

|| Weight gain during the 8-wk period of treatment was 335±10.7 g in control, 320±24 g in progesterone-treated, and 290±7.8 g in 17 β -estradiol-treated rats.

¶ 1 unit of lipoprotein lipase = 1 μ mol triglyceride hydrolyzed per h. The activity rate for each tissue is the result of four separate determinations.

** Values significantly different from control ($P < 0.05$).

injection daily for 7 days. One group of animals received 2.5 μ g 17 β -estradiol, a second group 25 μ g, a third group was given 5 mg progesterone; controls received 0.1 ml corn oil daily. The dosage of 17 β -estradiol was chosen to be in the range administered by Chen and Meites (7) and lower than that given by Cairns and Constantinides (8).

Lipoprotein lipase assay. The animals were anesthetized with sodium pentobarbital (12.5 mg/rat) and exsanguinated through the heart. Ovariectomized rats were examined at autopsy to insure the complete removal of the ovaries. The uteri were dissected and weighed in order to check the effect of hormone treatment. Dried defatted powders of adipose tissue, heart, and lung were prepared and assayed

for lipoprotein lipase activity as described previously (3, 9). Lipoprotein lipase activity was measured by the amount of rat chylomicron triglyceride hydrolyzed to fatty acids and glycerol with 1 U of activity = 1 μ mol of triglyceride hydrolyzed per h (3, 9).

RESULTS

Effect of 17 β -estradiol and progesterone on lipoprotein lipase activity in adipose tissue of male rats (Table I). There was a marked decrease in lipoprotein lipase activity in epididymal adipose tissue of male rats treated with 17 β -estradiol for a period of 8 wk. Progesterone treatment had no effect on adipose tissue lipoprotein lipase activity. The weight gain in the three groups of rats was not markedly different. Lipoprotein lipase activity in heart and lung of male rats was unaffected by estrogen or progesterone treatment.

Effect of 17 β -estradiol and progesterone on lipoprotein lipase activity in adipose tissue of ovariectomized rats (Table II). The level of lipoprotein lipase activity in adipose tissue rose after ovariectomy from 10.40±1.8 U/g in control female unoperated rats to 22.72±4.3 U/g 3 wk after the operation. Plasma triglyceride concentration was in the normal range in both control and ovariectomized rats. Administration of 2.5 μ g 17 β -estradiol daily for a period of 7 days to ovariectomized rats decreased lipoprotein lipase activity in adipose tissue to levels similar to those in control unoperated animals. Increasing the dose of 17 β -estradiol to 25 μ g daily caused a further decrease in the level of adipose tissue lipoprotein lipase activity and a concomitant rise in plasma triglyceride concentration. Animals in the five different groups had a similar weight gain during the

TABLE II
Effect of 17 β -Estradiol and Progesterone on Lipoprotein Lipase Activity in Adipose Tissue of Ovariectomized Rats

| Animals | Treatment* | No. of animals‡ | Lipoprotein lipase activity§ U/g | Plasma triglyceride μ mol/ml |
|----------------|-----------------------------------|-----------------|-------------------------------------|-------------------------------------|
| Control | None | 6 | 10.40±1.8 | 0.85±0.05 |
| Ovariectomized | None | 9 | 22.72±4.3 | 0.80±0.05 |
| | 17 β -Estradiol, 25 μ g | 5 | 9.00±1.17¶ | — |
| | 17 β -Estradiol, 25 μ g | 5 | 3.20±0.60 ¶ | 1.40±0.09 ¶ |
| | Progesterone, 5 mg | 8 | 22.28±1.71 | — |

Results are mean±SE.

* 17 β -Estradiol or progesterone were dispersed in corn oil and injected daily (0.1 ml subcutaneous) for a period of 7 days.

‡ Weight gain during the 7 days of treatment was 29±5 g in control, 32±4 g in ovariectomized, 24±3 and 21±2 g in 17 β -estradiol-treated rats (2.5 and 25 μ g/day, respectively), and 21±3.2 g in progesterone-treated rats.

§ The activity rate for each tissue is the result of four separate determinations.

|| Values significantly different from control ($P = 0.01$).

¶ Values significantly different from ovariectomized ($P = 0.01$).

hormone treatment period. Lipoprotein lipase activity in heart and lung was not affected by 17 β -estradiol administration.

Heart lipoprotein lipase was 41.3 \pm 5.3 U/g in ovariectomized rats, 33.3 \pm 3.0 U/g in 2.5 μ g and 43.9 \pm 3.3 U/g in 25 μ g 17 β -estradiol-treated animals, and 49.2 \pm 1.4 U/g in progesterone-treated rats. Lipoprotein lipase in the lung was 13.0 \pm 0.88 in control unoperated rats, 11.6 \pm 1.6, 12.00 \pm 2.54, 10.50 \pm 0.60, and 12.2 \pm 2.0 U/g in ovariectomized untreated, 2.5 μ g and 25 μ g 17 β -estradiol- and progesterone-treated rats, respectively. Blood triglyceride levels were 0.85 \pm 0.05 μ mol/ml in control unoperated rats, 0.80 \pm 0.05 μ mol/ml in ovariectomized rats and 1.4 \pm 0.09 μ mol/ml in ovariectomized rats treated with 25 μ g/day 17 β -estradiol. There was a sevenfold increase in the wet weight of the uterus after treatment of ovariectomized rats with either 2.5 or 25 μ g 17 β -estradiol daily for a period of 7 days; the uterus wet weight increased less than twofold after progesterone treatment.

DISCUSSION

The effect of sex hormones on lipoprotein lipase activity has been limited to the study of the enzyme present in postheparin plasma (4, 5, 10, 11). These studies have been carried out mainly in the human (4, 5, 10, 11). Postheparin lipolytic activity is a mixture of enzymes containing lipoprotein lipase of extrahepatic origin and triglyceride lipase of hepatic origin (12). The two lipolytic activities, hepatic and extrahepatic, have different substrate specificity, triglyceride emulsions and lipoprotein-triglyceride, respectively, and respond differently to enzyme inhibition (12, 13). It has recently been shown that postheparin lipolytic activity of extrahepatic origin (12) is highest in postmenopausal women (13). In rats ovariectomy increases and estrogen administration decreases postheparin lipolytic activity (8). There is no information, however, on the effect of estrogen and progesterone on tissue levels of lipoprotein lipase in either man or other mammals. The present study shows that estrogen administration caused a marked decrease in the level of lipoprotein lipase activity in adipose tissue of male or ovariectomized rats (Tables I and II). There was a concomitant increase in plasma triglyceride concentration (Table II). Administration of progesterone did not affect the enzyme activity in adipose tissue of either male or ovariectomized rats.

Studies in vivo and in vitro in the rat have shown that lipoprotein lipase activity in adipose tissue is markedly decreased during fasting (14), lactation (2, 3), and in diabetes (14). The decrease in lipoprotein lipase activity after estrogen treatment was not caused by inadequate food intake since the rats in the different experimental groups (control, ovariectomized, and

hormone treated) had a similar weight gain (Tables I and II).

We have recently shown that the very low activity of lipoprotein lipase in adipose tissue of lactating rats is the result of inhibition of enzyme synthesis by prolactin (15). We do not know at present whether the low adipose tissue lipoprotein lipase activity in estrogen-treated rats is caused directly by this hormone or whether it is mediated through the secretion of prolactin.

Estrogen administration to ovariectomized rats has been shown to increase both anterior pituitary and serum prolactin concentrations (7). The high levels of adipose tissue lipoprotein lipase activity in some of the control females and in pregnant rats during the first 2 wk of gestation (3) could be due to the low levels of both estrogen (6) and prolactin (16) during metestrus and diestrus and during the first 2 wk of gestation. Estrogen or progesterone treatment did not affect lipoprotein lipase activity in heart and lung.

We do not know at present what is the quantitative contribution of hepatic and extrahepatic tissues to the clearing of blood triglyceride. However, the presence of very low levels of lipase of extrahepatic origin and normal levels of hepatic lipase in postheparin plasma of patients with hyperchylomicronemia (13) indicates that the extrahepatic tissues play a major role in the clearing of blood triglyceride. This assumption agrees well with the data of Bragdon and Gordon, who showed that in the rat, 32% of chylomicron-triglyceride fatty acids were present in adipose tissue 10 min after intravenous injection of ¹⁴C-fatty acid-labeled chylomicrons (17). The decrease in adipose tissue lipoprotein lipase activity without the concomitant increase in enzyme activity in other tissues could lead to a rise in plasma triglyceride levels as a result of impaired clearing of blood triglyceride.

ACKNOWLEDGMENTS

The authors are grateful to Mr. Michael Blunda for excellent assistance, and Ms. Mary Jordan for her secretarial assistance.

This work was supported by grants from the Council for Tobacco Research—U. S. A., Inc., and the Washington Heart Association.

REFERENCES

1. Scow, R. O., M. Hamosh, E. J. Blanchette-Mackie, and A. J. Evans. 1972. Uptake of blood triglyceride by various tissues. *Lipids*. 7: 497-505.
2. Otway, S., and D. S. Robinson. 1968. The significance of changes in tissue clearing-factor lipase activity in relation to the lipaemia of pregnancy. *Biochem. J.* 106: 677-682.
3. Hamosh, M., T. R. Clary, S. S. Chernick, and R. O. Scow. 1970. Lipoprotein lipase activity of adipose and mammary tissue and plasma triglyceride in pregnant

- and lactating rats. *Biochim. Biophys. Acta.* **210**: 473-482.
4. Glueck, C. J., D. Scheel, J. Fishback, and P. Steiner. 1972. Progestagens, anabolic-androgenic compounds, estrogens: effects on triglycerides and postheparin lipolytic enzymes. *Lipids.* **7**: 110-113.
 5. Hazzard, W. R., M. J. Spiger, J. P. Bagdade, and E. L. Bierman. 1969. Studies on the mechanism of increased plasma triglyceride levels induced by oral contraceptives. *N. Engl. J. Med.* **280**: 471-474.
 6. Yoshinega, K., R. A. Hawkins, and J. F. Stocker. 1969. Estrogen secretion by the rat ovary *in vivo* during the estrous cycle and pregnancy. *Endocrinology.* **85**: 103-112.
 7. Chen, C. L., and J. Meites. 1969. Effects of estrogen and progesterone on pituitary and serum prolactin levels as measured by radioimmunoassay. *Fed. Proc.* **28**: 505. (Abstr.)
 8. Cairns, A., and P. Constantinides. 1955. Endocrine effects on heparin-induced lipemia-clearing activity (LCA) of rat plasma. *Can. J. Biochem. Physiol.* **33**: 530-538.
 9. Hamosh, M., and P. Hamosh. 1975. Lipoprotein lipase in rat lung. The effect of fasting. *Biochim. Biophys. Acta.* **380**: 132-140.
 10. Hazzard, W. R., D. T. Notter, M. J. Spiger, and E. L. Bierman. 1972. Oral contraceptives and triglyceride transport: acquired heparin resistance as the mechanism for impaired post-heparin lipolytic activity. *J. Clin. Endocrinol. Metab.* **35**: 425-437.
 11. Fabian, E., A. Stork, J. Kobilova, and J. Sponarova. 1967. The activity of the lipoprotein lipase and estrogens. *Enzymol. Biol. Clin.* **8**: 451-455.
 12. LaRosa, J. C., R. I. Levy, H. G. Windmueller, and D. S. Fredrickson. 1972. Comparison of the triglyceride lipase of liver, adipose tissue and postheparin plasma. *J. Lipid Res.* **13**: 356-363.
 13. Krauss, R. M., R. I. Levy, and D. S. Fredrickson. 1974. Selective measurement of two lipase activities in postheparin plasma from normal subjects and patients with hyperlipoproteinemia. *J. Clin. Invest.* **54**: 1107-1124.
 14. Robinson, D. S. 1970. The function of plasma triglyceride in fatty acid transport. *Comprehen. Biochem.* **18**: 51-116.
 15. Zinder, O., M. Hamosh, T. R. Clary Fleck, and R. O. Scow. 1974. Effect of prolactin on lipoprotein lipase in mammary gland and adipose tissue of rats. *Am. J. Physiol.* **226**: 744-748.
 16. Amenomori, Y., C. L. Chen, and J. Meites. 1970. Serum prolactin levels in rats during different reproduction states. *Endocrinology.* **86**: 506-510.
 17. Bragdon, J. H., and R. S. Gordon, Jr. 1958. Tissue distribution of C¹⁴ after the intravenous injection of labeled chylomicrons and unesterified fatty acids in the rat. *J. Clin. Invest.* **37**: 574-578.