

# Relationship between Lipoprotein Lipase Activity and Plasma Sex Steroid Levels in Obese Women

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## Abstract

In obese women ( $n = 16$ ) at their usual weight, fasting adipose tissue lipoprotein lipase (LPL) activity, obtained by elution with serum and heparin at 4° and 37°C, was inversely correlated to plasma estradiol levels ( $r = -0.724$ ;  $P = 0.002$ ) and ( $r = -0.641$ ;  $P = 0.010$ ), respectively. Furthermore, fasting postheparin plasma LPL activity during a heparin infusion, showed an even stronger inverse correlation to plasma estradiol when measured at 60 min ( $r = -0.815$ ;  $P < 0.001$ ). None of the above parameters was correlated to the body mass index.

Postprandial LPL activity in postheparin plasma, measured 10 min after a heparin injection, showed a strong positive correlation with plasma free testosterone ( $r = 0.780$ ;  $P = 0.001$ ). Neither of these parameters was correlated with the body mass index. The origin of this LPL activity is presently unknown but could conceivably represent a pool of LPL from skeletal muscle.

Since it has been shown convincingly that estrogen decreases adipose tissue LPL activity in the rat, the present studies strongly suggest that estradiol is a major negative regulator of fasting adipose tissue LPL activity in women.

## Introduction

Although men and women are clearly different regarding circulating plasma lipoprotein concentrations (1) and body fat distribution (2) the underlying mechanisms are incompletely understood. In both these aspects of lipid metabolism, lipoprotein lipase (LPL)<sup>1</sup> probably has a critical role. The enzyme hydrolyzes the triglyceride transported in very low density lipoproteins (VLDL) and chylomicrons thereby making their triglyceride fatty acids available for uptake in various tissues including adipose tissue (3). Thus the enzyme participates in the metabolism of major lipoprotein species as well as in the regulation of fat cell size.

The concentration of most of the hormones reported to either stimulate (4–9) or inhibit (5, 10–12) rat adipose tissue LPL activity, are the same in both sexes. Sex steroids on the other hand account for many differences between the sexes,

making it likely that these hormones also account for sex differences in LPL regulation. Convincing evidence for their effect on LPL has so far mainly been obtained in the rat. Thus, estradiol reduces adipose tissue LPL activity in both intact and gonadectomized animals (13–20), while progesterin, only affecting female adipose tissue (16), counteracts the estrogen effects on LPL (16–17). Androgen inhibition of adipose tissue LPL appears to result from estradiol formed by aromatization rather than from a direct effect (21).

In man, some studies have failed to demonstrate effects of oral estrogen on either adipose tissue (22) or postheparin plasma LPL (22–25), while others have reported that oral estrogen inhibits postheparin plasma LPL in postmenopausal women (26, 27). Oral or percutaneous progesterin alone (24, 28, 29), or in combination with estrogen (30), may enhance LPL activity in adipose tissue (28, 29) and postheparin plasma (24), respectively, while no effect on postheparin plasma LPL has been observed with low dose oral contraceptive (31) and anabolic steroid (31, 32).

Only a few human studies have explored the relationship between endogenous sex steroid levels in plasma and LPL activity. Thus no relationship has been demonstrated between either estradiol and postheparin plasma LPL (33) or estrone and adipose tissue LPL activity (28). Correlations between testosterone levels and LPL have been reported to be negative for adipose tissue (33), and either positive (28, 34) or absent (33), for postheparin plasma.

In summary, available data regarding sex steroid effects on adipose tissue and postheparin plasma LPL activity in man are contradictory and have failed to reproduce the strong and consistent effects observed in the rat. However, endogenous sex steroid levels in plasma were measured only in a few of the human studies.

The present study was undertaken in an attempt to resolve some of the discrepancies and to explore the role of endogenous sex steroids in the regulation of adipose tissue and postheparin plasma LPL in women. Precise and specific assays for LPL were employed. In addition, postheparin plasma lipase activities were determined after a standard bolus heparin injection and during prolonged heparin infusion, to take into account the possibility that there are differences in the kinetics of LPL release from different tissues. The results strongly suggest that circulating estradiol is an important negative regulator of adipose tissue LPL in fasting women. Furthermore, postheparin plasma in the postprandial state contains an LPL fraction of unknown origin, that is strongly correlated to plasma testosterone.

## Methods

**Human subjects.** Obese females ( $n = 16$ ) within 10% of their usual weight were admitted to the University of Washington Clinical Re-

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Received for publication 2 October 1987 and in revised form 21 March 1988.

1. Abbreviations used in this paper: LPL, lipoprotein lipase (EC 3.1.1.3).

Table I. Anthropometric Measures and Laboratory Screening

Subject	Age	Height	Weight	Body mass index	Glucose	Thyroxine	TSH	Triglycerides	Cholesterol
	yr	cm	kg	kg/m <sup>2</sup>	mg/dl	μg/dl	μU/ml	mg/dl	mg/dl
1	25	165	106.2	39.2	102	6.5	3.6	112	180
2	56	157	78.1	31.7	101	5.7	6.0	96	218
3*	22	164	73.8	27.4	81	5.8	5.2	73	320
4	43	162	103.0	39.2	94	5.8	3.2	71	169
5	41	158	91.6	36.7	107	7.0	4.2	87	231
6	39	173	98.2	32.8	109	6.5	4.6	83	226
7	46	163	80.5	30.3	71	5.4	2.6	146	184
8	34	169	87.3	30.6	85	6.8	3.5	127	193
9	27	173	76.5	25.5	90	5.6	4.0	110	197
10 <sup>‡</sup>	57	163	84.4	32.0	89	7.7	12.5	106	176
11	24	180	115.6	35.7	82	5.8	3.7	118	183
12 <sup>§</sup>	37	163	130.2	49.0	81	12.5	—	127	204
13	61	168	96.6	34.2	89	8.0	9.0	119	246
14	22	164	100.9	37.5	74	6.8	—	61	149
15 <sup>§</sup>	66	155	72.6	30.2	92	9.0	4.3	41	197
16*	33	172	117.9	40.1	85	7.1	4.0	195	229

\* Hyperlipidemia: subject 3 has hypercholesterolemia and subject 16 both hypertriglyceridemia and hypercholesterolemia (44). <sup>‡</sup> Borderline hypothyroidism. <sup>§</sup> Treated hypothyroidism: subjects 12 and 15 were maintained on 0.2 and 0.15 mg/d, respectively, of L-thyroxine.

search Center after informed consent had been obtained as approved by the Human Subjects Review Committee at the University of Washington. The subjects were fed a formula diet consisting of 15% (kcal/total kcal) protein, 40% fat and 45% carbohydrate, which was split in five isocaloric feedings between 8 a.m. and 8 p.m. The formula diet was supplemented with multivitamins, vitamin C and ferrous sulfate. Caloric intake for weight maintenance was calculated from an empirical formula (unpublished observation).<sup>2</sup> Subjects were weighed daily in nightgown with empty bladder before breakfast. Over a 2-wk study period the weight change in the 16 subjects ranged from -68 to +49 g/d (mean±SD, -6.5±38.3). The subjects, except two on thyroid (Table I) and one on low-dose estrogen replacement (Table IV), were taken off their usual medications during the study. Other data from these subjects have been reported previously (35).

**Clinical chemistry.** Fasting morning blood samples were obtained after the initial weight adjustment period had been completed. An automated chemical profile and assays for thyroxine, T<sub>3</sub>-resin uptake, and thyrotropin were done in the clinical laboratory at the University of Washington Hospital. Radioimmunoassays for total serum estrone (36) and testosterone (37) as well as determinations of free testosterone by equilibrium dialysis (38) were performed by Nichols Institute, San Juan Capistrano, CA. Total serum estradiol was measured by a kit (unpublished observations) from Pantex Laboratories, Santa Monica, CA. In menstruating subjects, these samples were obtained on the fourteenth day from the start of the last menstrual period. Plasma lipid determinations were performed at the Northwest Lipid Research Clinic following a standard protocol (39).

**Lipoprotein lipase.** Fasting LPL activity in gluteal adipose tissue needle biopsies from below the postero-lateral iliac crest, and postheparin plasma was determined as described previously (35, 40). Before assay, the enzyme was recovered from the adipose tissue by elution with serum and heparin at 4° and 37°C, and by extraction with detergent. The fractions obtained by elution supposedly represent mainly extracellular enzyme and are therefore a better index of the physiologi-

cally active enzyme than detergent-extracted enzyme, which measures the total LPL activity in the tissue (35). Adipose tissue LPL activity was related to either wet tissue weight or cell number after determination of average fat cell size (41).

Postheparin plasma LPL activity was calculated as the fraction of total lipolytic activity that was inhibited by an antiserum to LPL whereas the noninhibited fraction was taken as hepatic triglyceride lipase activity (42). Fasting activity was measured 10 min after the injection of heparin (60 U/kg) as well as at several time points during a high-dose 6-h heparin infusion (35, 43). Postprandial postheparin plasma activity was measured by injecting heparin (60 U/kg) 120 min after receiving the first feeding of the day and plasma was sampled 10 min after the injection.

**Data analysis.** An AT&T Personal Computer 6300 equipped with the SPSS/PC+ statistical software (SPSS Inc., Chicago, IL) and Sigma-Plot (version 3.0) scientific graph software (Jandel Scientific, Sausalito, CA) was used for statistical computations and production of graphs.

## Results

**Characterization of study population.** Anthropometric measures and results of laboratory screening are shown in Table I. None of the subjects had diabetes, two had hyperlipidemia, two had treated hypothyroidism and one had borderline hypothyroidism (Table I). In addition, four subjects were postmenopausal and two had oligomenorrhea (Table IV). The results of LPL determinations in adipose tissue and postheparin plasma as well as plasma sex steroid levels are displayed in Tables II-IV, respectively.

**Relationship between LPL activity and plasma estrogen levels.** The correlation matrix displayed in Table V demonstrates that plasma estradiol but not estrone levels showed significant negative correlations with adipose tissue as well as fasting postheparin plasma LPL activity. Only adipose tissue LPL activity eluted at either 4°C or 37°C and expressed per unit weight, showed a significant correlation (Figs. 1 A and 2). The LPL activity in postheparin plasma sampled at 10, 30, and

2. Equation describing the relationship between caloric intake (I; kcal/kg per d) required for weight maintenance on a metabolic ward and body mass index (BMI; kg/m<sup>2</sup>):  $I = 52.2 - 0.752 \times \text{BMI}$ .

Table II. Adipose Tissue LPL Activity and Fat Cell Size

Subject	Activity per weight			Activity per cell			Cell size
	Elution at 4°C	Elution at 37°C	Detergent extraction	Elution at 4°C	Elution at 37°C	Detergent extraction	
	<i>nmol · min<sup>-1</sup> · g<sup>-1</sup></i>			<i>nmol · min<sup>-1</sup> · 10<sup>-6</sup> cells</i>			
1	42.2	48.2	169	30.0	34.3	120	0.710
2	39.1	87.2	228	25.2	56.1	146	0.644
3	43.2	124.0	273	27.0	77.1	171	0.624
4	33.4	76.9	190	21.3	49.1	121	0.639
5	27.4	43.8	211	17.0	27.2	130	0.619
6	29.0	78.1	177	20.6	55.5	126	0.711
7	61.4	112.0	316	40.5	74.0	209	0.660
8	11.0	31.8	157	7.1	20.5	101	0.645
9	22.1	65.3	223	12.0	35.6	121	0.544
10	50.2	107.0	305	32.7	70.0	199	0.651
11	27.1	52.1	145	16.7	32.1	89	0.616
12	37.1	92.0	226	18.4	45.5	112	0.496
13	59.4	106.0	267	34.5	61.8	155	0.581
14	33.5	77.2	216	18.8	43.3	121	0.561
15	68.4	154.0	298	46.8	106.0	204	0.685
16	22.5	66.6	223	20.9	61.7	207	0.927

All values were obtained after a 12-h overnight fast.

60 min after the start of a heparin infusion showed significant negative correlations at all three time points (Table V). The strongest and most significant was the correlation between fasting 60 min postheparin plasma LPL activity and plasma estradiol (Fig. 3 A). Since the relationships in Figs. 1 A and 3 A

appear to be nonlinear, the data were also plotted with a logarithmic scale on the abscissa (Fig. 1 B and 3 B). This maneuver improved the correlations from  $r = -0.724$  ( $P = 0.002$ ;  $n = 15$ ) and  $r = -0.815$  ( $P = 0.0002$ ;  $n = 15$ ) to  $r = -0.781$  ( $P = 0.0006$ ) and  $r = -0.860$  ( $P = 0.0000$ ), respectively.

Table III. Postheparin Plasma LPL Activity

Subject	Fasting			Fed	Difference fasting-fed
	10 min*	30 min	60 min	10 min*	10 min
	<i>nmol · min<sup>-1</sup> · ml<sup>-1</sup></i>				
1	180	239	212	169	11
2	287	456	509	—	—
3	213	262	256	147	66
4	179	137	148	154	25
5	177	171	99	104	73
6	218	285	266	156	62
7	244	332	332	177	67
8	102	147	146	124	-22
9	213	262	256	147	66
10	218	296	315	—	—
11	259	170	175	194	65
12	239	259	324	163	76
13	267	433	516	133	134
14	280	357	354	244	36
15	360	630	609	184	176
16	234	304	244	132	102

\* The means±SD of paired cases ( $n = 14$ ) were  $226.1 \pm 59.9$  and  $159.1 \pm 34.5$ , respectively, and significantly different by the paired  $t$  test ( $t = 5.09$ ;  $P = 0.000$ , two-tailed probability).

Table IV. Plasma Levels of Sex Steroids

Subject	Estradiol	Estrone	Testosterone	Free testosterone
	<i>pg/ml</i>		<i>ng/dl</i>	<i>pg/ml</i>
	1	—	—	27
2*	58	2.7	10	0.9
3	141	88.0	21	3.4
4	227	91.0	11	2.3
5	207	35.0	15	2.6
6‡	141	55.0	36	4.4
7§	65	56.0	20	2.9
8	197	76.0	18	3.0
9	90	20.0	21	4.3
10*	55	75.0	24	3.1
11	136	107.0	28	4.3
12‡	110	80.0	22	3.7
13*	25	16.0	10	1.4
14	90	115.0	43	7.6
15*	45	—	32	3.6
16	235	148.0	10	1.8

\* Postmenopause.

‡ Oligomenorrhea ( $\leq 3$  menstrual periods per year).

§ Status posthysterectomy (ovaries intact).

|| Maintained on low dose estrogen replacement (Premarin, 0.3 mg/d) throughout the study.

Table V. Correlations between LPL Activity and Plasma Sex Steroid Levels

Enzyme source	Estradiol	Estrone	Testosterone	Free testosterone
<b>Adipose tissue</b>				
Per gram				
Elution at 4°C	-0.7239*	-0.2903	0.0766	-0.1331
Elution at 37°C	-0.6414*	-0.1265	0.1144	-0.1434
Detergent extraction	-0.6179	-0.2141	-0.0923	-0.2558
Per 10 <sup>6</sup> cells				
Elution at 4°C	-0.6135	-0.1814	0.0623	-0.1725
Elution at 37°C	-0.4818	0.0380	0.0729	-0.2166
Detergent extraction	-0.2935	0.0868	-0.1685	-0.3600
<b>Postheparin plasma</b>				
Fasting				
10 min	-0.6643*	-0.0667	0.2780	0.0508
30 min	-0.7198*	-0.3642	0.1590	-0.1021
60 min	-0.8146 <sup>‡</sup>	-0.4191	0.0890	-0.1572
Fed				
10 min	-0.4520	0.4005	0.7775*	0.7804 <sup>‡</sup>
Difference fasting-fed				
10 min	-0.5298	-0.2134	-0.0581	-0.3263

Pearson correlation coefficients calculated after pairwise exclusion of missing variables.

\*  $P \leq 0.01$ ; <sup>‡</sup>  $P \leq 0.001$  (two-tailed significance).

There were no significant correlations between degree of obesity, measured as the body mass index, and either of the above parameters of LPL activity or estradiol levels (data not shown).

**Relationship between LPL activity and plasma androgen levels.** A strong positive correlation was demonstrated between postprandial LPL activity in 10 min postheparin plasma and total as well as free plasma testosterone (Table V; Fig. 4). The postprandial LPL activity in 10 min postheparin plasma was significantly lower than the fasting activity (Table III) and the difference was strongly correlated to fasting adipose tissue LPL activity eluted at 37°C and expressed per unit weight ( $r = 0.713$ ;  $P = 0.004$ ;  $n = 14$ ). Thus, feeding may have decreased the contribution from the adipose tissue to the postheparin plasma activity and thereby increased the proportion of en-

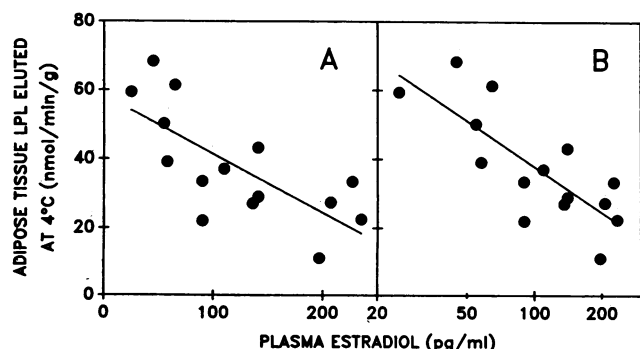


Figure 1. Linear (A) and semilogarithmic (B) regression of fasting adipose tissue LPL activity eluted at 4°C, on plasma estradiol concentration. The linear ( $r = -0.724$ ;  $P = 0.002$ ) and semilogarithmic ( $r = -0.781$ ;  $P = 0.0006$ ) correlation coefficients were calculated after pairwise exclusion of a missing variable ( $n = 15$ ).

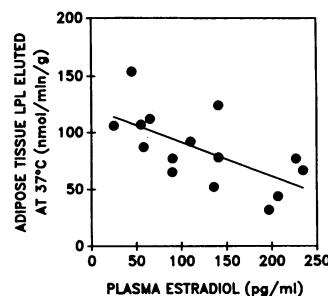


Figure 2. Linear regression of fasting adipose tissue LPL activity eluted at 37°C, on plasma estradiol concentration. The correlation coefficient ( $r = -0.641$ ;  $P = 0.010$ ) was calculated after pairwise exclusion of a missing variable ( $n = 15$ ).

zyme from other sites allowing a previously masked correlation to emerge.

There were no significant correlations between the body mass index and either the postprandial 10 min postheparin plasma enzyme or total and free testosterone levels (data not shown).

## Discussion

The strong negative statistical correlations in women between plasma estradiol levels and LPL activity in both adipose tissue and postheparin plasma, have not been reported previously. These findings raise the question as to where the postheparin plasma enzyme originates. Since the bulk of the total body LPL probably resides in adipose tissue and skeletal muscle, each tissue contributing approximately equal amounts of enzyme (45), it is likely that both tissues are major contributors to postheparin plasma LPL activity. Previously, using the same subjects as in the present study, fasting postheparin plasma LPL activity at various times during a heparin infusion was shown to correlate positively with adipose tissue LPL activity eluted at 4° and 37°C, expressed per unit weight. Values obtained at 60 min after starting the infusion showed the best correlations (35). It would therefore seem plausible that the strong negative correlations between postheparin plasma LPL and estradiol (Table V) at least partly reflect inhibition of adipose tissue LPL by estrogen. Obviously, LPL from sources other than adipose tissue may also be inhibited by estrogens and contribute to the postheparin plasma lipase activity. For instance, some animal experiments have resulted in decreased

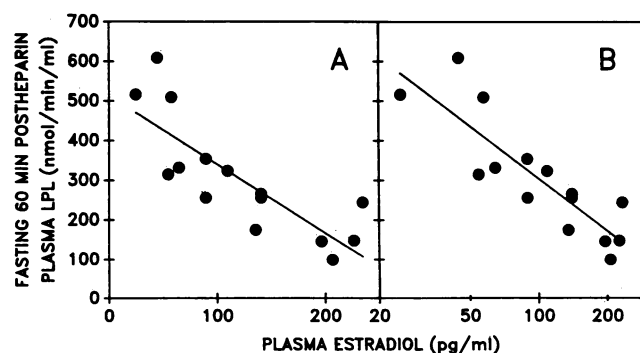


Figure 3. Linear (A) and semilogarithmic (B) regression of fasting postheparin plasma LPL activity, sampled at 60 min into a continuous heparin infusion, on plasma estradiol concentration. The linear ( $r = -0.815$ ;  $P = 0.0002$ ) and semilogarithmic ( $r = -0.860$ ;  $P = 0.0000$ ) correlation coefficients were calculated after pairwise exclusion of a missing variable ( $n = 15$ ).

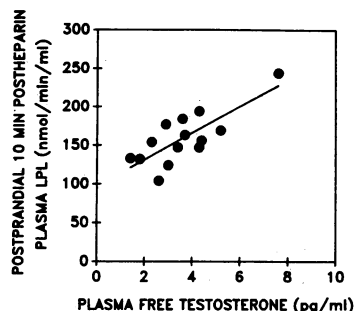


Figure 4. Linear regression of postprandial plasma LPL activity, sampled 10 min after the injection of heparin, on plasma free testosterone concentration. The correlation coefficient ( $r = 0.780$ ;  $P = 0.001$ ) was calculated after pairwise exclusion of two missing variables ( $n = 14$ ).

skeletal muscle LPL activity by estrogen (18). Other experiments, however, have demonstrated the opposite effect (15). The noticeable correlation between postheparin plasma LPL and estrogen therefore remains to be fully explained.

Adipose tissue LPL activity from the gluteal site, also employed in the present study, is similar to that from the femoral site (46). In menstruating women, the femoral site has higher LPL activity than tissue from the abdominal site and only LPL from the former site is responsive to the hormonal changes of pregnancy and lactation (47). After the menopause, however, abdominal and femoral adipose tissue have similar LPL levels (30). The inverse correlation between gluteal adipose tissue LPL and estradiol suggests that this tissue site also is similar to femoral adipose tissue with regard to hormone responsiveness.

It should be noted that only the adipose tissue LPL recovered by elution, showed significant correlations with estradiol, while detergent-extracted enzyme did not. Since the former parameters presumably represent extracellular enzyme better than the latter (35), it might be argued that a putative direct effect of estradiol on the tissue could involve inhibition of enzyme secretion, stimulation of uptake and degradation, or removal of extracellular binding sites.

The population studied was heterogeneous in several respects. Although seven of the sixteen women did not have regular periods (Table IV), it was expected that these subjects would not pose a sampling problem since their estrogen levels could be expected to fluctuate less than in menstruating females. The one subject (No. 7) who had undergone hysterectomy would most likely be postmenopausal given her age (Table I) and estrogen level (Table IV). One postmenopausal subject (No. 10) was treated with 0.3 mg/d of conjugated equine estrogens (Premarin), a dose that is approximately one-third or less of full replacement. The main component of this preparation is estrone sulfate (48), which should act by conversion to estradiol. It is not known, however, to what extent the equine steroids contribute to unmeasurable estrogenic activity.

Heterogeneity of the study population was particularly apparent for age and degree of obesity (Table I). Nevertheless, the observed correlations should still be valid unless age and obesity were confounding variables. Since the subjects with the lowest estradiol levels were older than those with higher levels (Tables I and IV), it may be argued that age was a confounding variable. However, the correlations between age and adipose tissue LPL eluted at 4°C ( $r = 0.643$ ;  $P = 0.007$ ;  $n = 16$ ), at 37°C ( $r = 0.567$ ;  $P = 0.022$ ;  $n = 16$ ), and 60 min postheparin plasma LPL ( $r = 0.670$ ;  $P = 0.005$ ;  $n = 16$ ) were all weaker than the corresponding correlations with estradiol (Figs. 1 A, 2, and 3 A, respectively). Moreover, age was not significantly

correlated to estradiol ( $r = -0.502$ ;  $P = 0.057$ ;  $n = 15$ ). Finally, previous animal data have provided convincing evidence for inhibition of adipose tissue LPL by estradiol (see Introduction). The present data, interpreted in the latter context, therefore suggest that estradiol also inhibits adipose tissue LPL in man. It cannot be excluded, however, that age may have a minor effect on LPL, independent of estradiol.

Why then, have several previous human studies with two exceptions (26, 27) not found this striking correlation? There are at least two factors that alone or in combination conceivably could explain this discrepancy.

First, the likelihood of detecting an inhibition by exogenous estrogen would be the greatest in the postmenopausal female who has low endogenous estrogen levels and high lipase activity. Indeed, studies in which such effects have been reported were carried out in postmenopausal subjects (26–27). On the other hand, the administration of pure estrogen or birth control pills to menstruating women with moderate to high endogenous estrogen levels is likely not to alter total estrogenic activity in plasma due to suppression of ovarian production. Furthermore, the inhibitory effect of estradiol on adipose tissue LPL eluted at 4°C and 60 min postheparin plasma LPL, appears to reach a plateau at estradiol concentrations in the high normal range (Figs. 1 A, and 3 A) where all the cases have regular menstrual periods (Table IV). As expected, effects of exogenous estrogen on LPL have not been observed in menstruating women (23, 31).

Second, the sampling time is important in order to obtain a postheparin plasma fraction that shows a strong correlation with adipose tissue LPL activity. While most earlier studies have used postheparin plasma samples obtained at 10 to 15 min after a heparin injection, it was recently demonstrated that samples obtained 60 min after the start of a continuous heparin infusion shows the best correlation with the adipose tissue enzyme (35).

The absence of any relationship between degree of obesity and LPL activity expressed per unit weight of adipose tissue (see Results) has been reported previously in men (49). Although all the subjects were studied at their usual weight and in the eucaloric state, the difference between the highest and lowest LPL activity in adipose tissue measured by the different parameters varied from two- to severalfold (Table II). It may be surmised that these differences could reflect the different propensities for rate of weight gain should the caloric intake be increased above the level for weight maintenance.

The strong positive correlation between plasma testosterone and LPL activity in postprandial 10 min postheparin plasma (Fig. 4) was an unexpected finding. Since this enzyme fraction probably represents tissues other than the adipose tissue (see Results), skeletal muscle could conceivably be a major source. The fact that this tissue is a target organ for testosterone (50) as well as a documented source of LPL (51) lends further support for this notion. It has also been observed previously that postheparin plasma LPL activity released early (15 min) after a heparin injection is positively correlated to LPL activity in skeletal muscle (52). Contrary to the recent report by Lithell et al. (53), no correlation between any measure of adipose LPL activity and plasma testosterone levels were observed in the present study (Table V). It is possible, that the negative correlation noted by these authors in fact represents inhibition of adipose tissue LPL by estradiol formed by aroma-

tization in that tissue of testosterone present in supranormal concentrations (21, 54).

Since steroid hormones do not display major short-term fluctuations in response to feeding, it is unlikely that testosterone alone was responsible for the decrease in LPL activity that took place in the 10-min postheparin plasma fraction over 2 h. If testosterone is involved in the regulation of the postprandial 10-min postheparin plasma enzyme, the hormone would be more likely to have a long-term effect permissive for short-term regulation. Conceivably, testosterone could control any of the events involved in the regulation of a secretory protein, e.g., synthesis, secretion, and degradation as well as the number of extracellular binding sites for the enzyme.

The notion that human LPL may be regulated by endogenous estrogens as well as androgens requires a comment about hepatic triglyceride lipase. In a number of studies, the latter enzyme measured in postheparin plasma has been shown to be inhibited by exogenous estrogens (22–23, 27, 55) and stimulated by exogenous androgens and progestins (32, 56). The few studies addressing the effects of endogenous sex steroids on postheparin plasma hepatic triglyceride lipase found suppression of the enzyme during the luteal phase of the menstrual cycle (57) and during pregnancy (58). A negative correlation between estradiol levels and enzyme activity could be demonstrated only in the latter study (58).

In this study, there was no significant correlation between fasting 10 min postheparin plasma hepatic triglyceride lipase activity and either estradiol ( $r = 0.264$ ;  $P = 0.341$ ;  $n = 15$ ) estrone ( $r = 0.264$ ;  $P = 0.362$ ;  $n = 14$ ) or testosterone levels ( $r = -0.229$ ;  $P = 0.394$ ;  $n = 16$ ). Furthermore, there was no significant correlation between hepatic triglyceride lipase activity and any of the parameters of LPL activity in adipose tissue or postheparin plasma presented in this communication (data not shown), making it highly unlikely that any significant part of the hepatic triglyceride lipase was measured as LPL or vice versa. Thus, available data suggest that the hepatic triglyceride lipase is very sensitive to regulation by exogenous sex steroids, perhaps because the oral route of administration leads to transiently supranormal hormone concentrations in the hepatic circulation, whereas the role of endogenous sex steroids is still unclear. By contrast, the reverse situation seems to apply for LPL.

In summary, this study corroborates previous experiments in animals and strongly suggests that estradiol is a major negative regulator of fasting adipose tissue LPL in women, independent of degree of obesity. Together with the finding of a postheparin plasma LPL fraction that is positively correlated to plasma testosterone and probably unrelated to adipose tissue LPL, the present work also hints that both male and female sex steroids may participate in the regulation of LPL in more tissues and to a larger extent than previously thought.

## Acknowledgments

The excellent technical assistance of Martha Kimura and Steven Hashimoto is gratefully acknowledged.

Support for this work was provided by the American Diabetes Association, the National Institutes of Health (DK-02456 and HL-30086), the General Clinical Research Center (RR 37) at the University of Washington, the Biomedical Research Support Grant (2 S07 RR0 5428) at the University of Utah, and the Veterans Administration.

*Note added in proof.* A recent report appearing in press since this paper was submitted has demonstrated that the rise in endogenous testosterone and estradiol levels following the injection of human chorionic gonadotropin (hCG) to prepubertal or early pubertal boys is accompanied by decreased fasting LPL activity in postheparin plasma (59).

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