The –514 C/T polymorphism in the hepatic lipase gene promoter is associated with insulin sensitivity in a healthy young population

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Abstract

Impaired insulin action has been associated with diabetes, dyslipidemia and atherosclerotic vascular disease. The expression of insulin resistance results from the interaction of environmental and genetic factors. Human hepatic lipase (HL) is a lipolytic enzyme that plays a role in the metabolism of several lipoproteins, while insulin up-regulates the activity of HL via insulin-responsive elements in the HL promoter. We have examined the influence of –514 C/T polymorphism in the hepatic lipase gene promoter on insulin sensitivity in 59 healthy young subjects (30 males and 29 females). The volunteers were subjected to three dietary periods, each lasting four weeks. During the first period all subjects consumed a saturated fat (SFA)-enriched diet with 38% as fat (20% SFA, 12% monounsaturated fatty acids (MUFA) and 6% polyunsaturated fatty acids (PUFA)). In the second and third dietary periods, a randomized crossover design was used, consisting of a low fat, high carbohydrate diet (CHO diet) (<10% SFA, 12% MUFA and 6% PUFA) and a high-MUFA, or Mediterranean diet, with <10% SFA, 22% MUFA and 6% PUFA. We determined the *in vivo* insulin resistance using the insulin suppression test with somatostatin. Steady-state plasma glucose (SSPG) concentrations (a measure of insulin sensitivity) were significantly higher in men carriers of the –514T allele after the consumption of the SFA diet than after the CHO diet and the Mediterranean diet. This effect was not observed in women. Moreover, there were no significant differences in insulin sensitivity after the three diets in men and women with the CC genotype. In summary, our results show an improvement in insulin sensitivity in men with the –514T allele of the HL promoter polymorphism, when MUFA and carbohydrates are consumed instead of SFA fat.

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Introduction

Impaired insulin action has been associated with diabetes, obesity, dyslipidemia, hypertension and also with atherosclerotic vascular disease. Insulin resistance is determined by the interaction between genetic and environmental factors. In general, high intake of dietary fat has been associated with obesity and its comorbid conditions, including heart disease and diabetes. All these factors are related to the global process of ‘Westernization’ of lifestyles and dietary habits, especially the high intake in calories from saturated fat. Insulin resistance usually precedes the diagnosis of type 2 diabetes mellitus by decades and, in most cases, the clinical expression of the disease could be prevented by dietary and lifestyle modification (Mayer-Davis *et al.* 1997). On the other hand, the genetic base of type 2 diabetes mellitus is very heterogeneous and has been related to various genetic mutations that codify proteins connected with glucose and insulin metabolism, such as the insulin receptor (Kusari *et al.*

1991), insulin receptor substrate-1 (Almind *et al.* 1993), the Rad protein (Reynet & Kahn 1993), the glycogen synthase (Bjorbaek *et al.* 1994) and the β -3-adrenergic receptor (Walston *et al.* 1995), to mention the most well-known. However, the genetic background of insulin resistance and type 2 diabetes mellitus is more complex and it can also involve other genes seemingly unrelated with carbohydrate metabolism.

Hepatic lipase (HL) is an enzyme anchored to the vascular endothelium in the liver as well as to the surface of hepatocytes, where it catalyzes the hydrolysis of various lipids in lipoprotein particles. Several studies have reported that the –514T allele of the HL gene promoter appears to be associated with decreased HL activity, increased high density lipoprotein cholesterol (HDL-C) (Guerra *et al.* 1997, Zambon *et al.* 1998), and increased low density lipoprotein (LDL) buoyancy (Zambon *et al.* 1998). However, at present, the effects of this common hepatic lipase variant –514C/T on lipid and lipoprotein levels are not well known. Moreover, as HL is involved in the clearance of triglyceride-rich

lipoproteins, the primary effects of the variant genes influence the metabolism of triglyceride-rich lipoproteins which may, secondarily, affect glucose homeostasis. Also, insulin has been assumed to upregulate the activity of the HL promoter. This has been proposed to explain the associations between hyperinsulinemia and high HL activity (Romano *et al.* 1997). Recently, in a Finnish population, an association of the HL variant with insulin resistance was observed in patients with familial combined hyperlipidemia (Pihlajamäki *et al.* 2000).

Whether the HL promoter variant contributes to impaired insulin sensitivity in the normal population has not been investigated. In the present study we have explored the possible influence of the -514 C/T variant in the promoter of the HL gene on insulin sensitivity in a healthy young population. In addition, the interaction between this polymorphism and diet on insulin sensitivity was evaluated.

Materials and methods

Subjects and diets

Fifty-nine healthy normolipemic subjects, 39 homozygous for the most common allele (C/C) and 20 carriers of the T allele (CT/TT), attending the University of Cordoba volunteered to participate in the study. They ranged in age from 22.6 ± 1.4 years. None of them had diabetes, or liver, renal or thyroid disease. All underwent a comprehensive medical history, physical examination, and clinical chemistry analysis before enrollment. None of the subjects was taking medication or vitamins known to affect plasma lipids. Dietary information, including alcohol consumption, was collected over seven consecutive days. Individual energy requirements were calculated by taking into consideration each subject's weight and physical activity. Subjects were encouraged to maintain their regular physical activity and life-style and were asked to record in a diary any event that could affect the outcome of the study, such as stress, change in smoking habits, and alcohol consumption or foods not included in the experimental design as described in our previous studies (Fuentes *et al.* 2001, Pérez-Jiménez *et al.* 2001, Pérez-Martínez *et al.* 2001).

The study design included an initial 28-day period during which all the subjects consumed a saturated fat (SFA)-enriched diet with 15% of energy as protein, 47% as carbohydrate, and 38% as fat (20% SFA, 12% monounsaturated fatty acids (MUFA) and 6% polyunsaturated fatty acids (PUFA)). All participants were then randomized in a crossover design and exposed to two new dietary periods: a low fat, high carbohydrate diet (CHO diet), and a high-MUFA diet, with a typical 'Mediterranean diet' enriched with olive oil. The two groups of subjects were assigned one of the two dietary

regimes for 28 days each. Group 1 (30 subjects) was placed on a Mediterranean diet followed by a CHO diet. For group 2 (29 subjects) the order was reversed. Assignment of volunteers to the sequence of diets was carried out at random. The CHO diet (National Cholesterol Education Program 1994) contained 15% of energy as protein, 57% as CHO, and 28% as fat (<10% SFA, 12% MUFA, and 6% PUFA). The Mediterranean diet contained 15% of energy as protein, 47% as CHO, and 38% as fat (<10% SFA, 22% MUFA, and 6% PUFA). Olive oil provided 75% of total MUFA consumed during this last dietary period. Dietary cholesterol was maintained constant in our experimental design and the mean cholesterol intake was 115 mg/1000 kcal during the three periods. The Human Investigation Review Committee approved this study at the Reina Sofia University Hospital. Informed consent was obtained from all participants.

The composition of the experimental diets was calculated using the United States Department of Agriculture (USDA) (Human Nutrition Information Service of Department of Agriculture 1987) food tables, or the Spanish food composition tables for local foodstuffs. Fourteen menus, prepared with regular solid foods, were rotated during the experimental period. We used virgin olive oil for cooking and salad dressing during the Mediterranean diet, and palm oil and butter for the high-SFA diet. During the CHO diet period, biscuits, bread and jam replaced some olive oil or palm oil. Lunch and dinner were consumed in the hospital dining room, whereas breakfast and an afternoon coffee break were eaten in the medical school cafeteria. A dietician supervised all meals. Duplicate samples from each menu were collected, homogenized, and stored at -80°C . Protein, fat and CHO content of the diet were analyzed using standard methods (Association of Official Analytical Chemists, Arlington 1990). Evaluation of dietary compliance was also performed by examining the food diaries and by analyzing the fatty acid content of the cholesterol ester fraction in LDL (Ruiz-Gutierrez *et al.* 1993).

Blood sampling and biochemical determinations

Venous blood for insulin, glucose, lipid and lipoprotein analysis was collected in EDTA-containing tubes from the subjects after a 12-h overnight fast at the end of each dietary period. Each analysis was performed three times. Total cholesterol (TC) and triglycerides (TG) were assayed by enzymatic procedures (Allain *et al.* 1974, Bucolo & David *et al.* 1973). HDL-C was measured by analyzing the supernatant obtained after precipitation of a plasma aliquot with dextran sulfate- Mg^{2+} (Warnick *et al.* 1982). The LDL-cholesterol (LDL-C) level was calculated from total cholesterol, triglyceride, and HDL-C values using the Friedewald formula (Friedewald *et al.* 1972). Unesterified free fatty acid (FFA)

levels were determined by an enzymatic colorimetric assay (Boehringer Mannheim) as described by Shimizu *et al.* (1979). To reduce inter-assay variation, plasma for biochemical determinations was stored at -80°C and analyzed in duplicate at the end of the study.

Glucose suppression test

At the end of each dietary period all subjects underwent a modified insulin suppression test (Harano *et al.* 1977, Laws *et al.* 1994). The technique used in the present study to quantify insulin sensitivity was the insulin suppression test, a simple and cost-effective test for the measurement of insulin resistance, which has been used increasingly often in recent years. The study began at 0800 h, after a 12-h fast. A continuous infusion of somatostatin (214 nmol/h), insulin (180 pmol/m²/min), and glucose (13.2 mmol/m²/min) was administered in the same vein. Somatostatin was used to inhibit endogenous insulin secretion. Blood was sampled every 30 min for the first 2.5 hours, by which time steady-state plasma glucose (SSPG) and steady-state plasma insulin (SSPI) levels were achieved. Blood was then sampled at 10-min intervals for the last 30 min (at 150, 160, 170 and 180 min) for measurement of plasma glucose and insulin concentrations. These four values determined the SSPG and SSPI concentrations. Since SSPI levels were similar in all subjects, SSPG levels provided a measure of the ability of insulin to promote disposal of infused glucose. Subjects with high SSPG are relatively more insulin-resistant than those with lower SSPG.

Genotyping of hepatic lipase gene polymorphism

DNA was extracted from 10 ml EDTA-containing blood. Amplification of a 299-bp region of the hepatic lipase gene was carried out by polymerase chain reaction (PCR) with 250 ng genomic DNA and 0.2 μmol of each oligonucleotide primer (P1; 5'-AAGAAGTGTGTTTA CTCTAAGGATCA-3', and P2, 5'-GGTGGCTTCCA CGTGGCTGCCTAAG-3') in 50 μl . DNA was denatured at 95°C for 5 min followed by 30 cycles of denaturation at 95°C for 1 min, annealing at 58°C for 1.5 min, and extension at 72°C for 2 min. The PCR product (10 μl) was digested with 5 units of restriction enzyme NlaIII (BRL, Baltimore, MA, USA) in a total volume of 35 μl . Digested DNA was separated by electrophoresis in an 8% non-denaturing polyacrylamide gel at 150 V for 2 h. Bands were visualized after silver staining. Samples containing the T allele were amplified a second time to verify the genotype.

Statistical analysis

Statistical analyses were carried out using the SPSS statistical package (SPSS, Chicago, IL, USA). ANOVA

for repeated measures was used to analyze the differences in plasma lipid, glucose, SSPG levels and basal glucose and insulin-stimulated glucose uptake between dietary phases. When statistically significant effects were demonstrated, Tukey's post-hoc test was used to identify between-group differences. The general linear models for repeated measures procedure was used to test gene and diet interactions. A value of $P < 0.05$ was considered significant.

Results

Dietary composition was analyzed in duplicate meals portions, and the results are shown in Table 1. The results were in good agreement with values obtained from the food composition tables. Analysis of the cholesterol ester fraction of plasma LDL showed good adherence to the different diets. During consumption of the SFA diet, there was a significant increase in palmitic acid (16:0) compared with that recorded during the high CHO and Mediterranean dietary periods (27.2 ± 1.4 vs 18.9 ± 3.9 , $P < 0.004$ and 27.2 ± 1.4 vs 15.1 ± 0.4 , $P < 0.004$ respectively). We also observed a significant increase in oleic acid (18:1) when subjects switched from the CHO diet to the Mediterranean diet (38.5 ± 9.0 vs 49.7 ± 4.7 , $P < 0.05$).

The baseline characteristics of the subjects according to the -514C/T polymorphism are shown in Table 2. Carriers of the -514T allele ($n=20$) had greater total cholesterol (TC) and LDL-C plasma levels compared with homozygotes for the C allele ($n=39$). There were no significant differences between subjects with the CC genotype and those with the CT/TT genotype for any of the lipid parameters after the three dietary periods (Table 3).

Steady state plasma glucose (SSPG) concentrations (a measure of insulin sensitivity) were significantly higher in men carriers of the -514T allele after the consumption of the saturated diet than after the CHO diet ($P < 0.05$) and the Mediterranean diet ($P < 0.05$) (Fig. 1A), but this was not so in women (Fig. 1B). Moreover, there were no significant differences in insulin sensitivity after the three diets in subjects with the CC genotype, either in men or women (Fig. 1A,B). Analysis of plasma FFAs revealed that homozygotes for the C allele showed lower levels of FFAs following the CHO and the Mediterranean diets, as compared with the saturated diet (0.47 ± 0.08 vs 0.58 ± 0.09 , $P < 0.02$ and 0.41 ± 0.06 vs 0.58 ± 0.09 , $P < 0.001$ respectively). The same effect was observed in carriers of the -514T allele, but a significant difference was noted only between the SFA diet and the CHO diet (0.61 ± 0.09 vs 0.54 ± 0.01 , $P = 0.01$). Furthermore, total cholesterol, LDL-C, and apolipoprotein B plasma levels were correlated positively with SSPG ($r = 0.22$, $P < 0.001$, $r = 0.30$, $P < 0.001$, and $r = 0.35$, $P < 0.001$ respectively).

Table 1 Mean daily intake during each experimental diet period (mean±S.D.). SFA diet, saturated fat enriched diet; CHO-diet, low fat, high carbohydrate diet

	High SFA diet	CHO diet	Mediterranean diet
Protein (% of energy intake)			
Calculated	15	15	15
Analyzed	18.1±2.5	17.6±1.5	17.5±2.0
Fat (% of energy intake)			
Saturated			
Calculated	20	10	10
Analyzed	22.6±4.1	9.2±3.5	9.2±4.2
Monounsaturated			
Calculated	12	12	22
Analyzed	10.1±2.9	13.5±1.2	24.4±2.2
Polyunsaturated			
Calculated	6	6	6
Analyzed	5±1.5	5.2±2.0	4.8±1.1
Carbohydrates (% of energy intake)			
Calculated	47	57	47
Analyzed	44.2±8.3	54.5±8.6	44.1±7.8
Cholesterol (mg/day)			
Calculated	115	115	115
Analyzed	112±39	113±48	117±42
Energy (MJ)			
Calculated	10.2	10.2	10.2
Analyzed	10.8±1.1	10.6±1.0	10.8±1.5
Dietary fiber (g/day)			
Calculated	30	30	30
Analyzed	25.9±7.0	26.1±6.1	24.9±8.2

Additionally, there was a significant inverse correlation between mean glucose levels of the SSPG and HDL-C plasma levels ($r = -0.20$, $P < 0.05$).

Discussion

Several exogenous factors such as diet play an important role in the peripheral effect of insulin, although it is now thought that there is genetically determined individual

variability in the development of insulin resistance (Moller *et al.* 1996). The novel finding of the present study is that the -514C/T polymorphism in the HL promoter is associated with a decrease in insulin sensitivity in normal young males after consuming an SFA diet. Thus the replacement of this diet by a CHO diet or a Mediterranean diet improved insulin sensitivity in male -514T carriers.

The saturated fatty acids reduce insulin sensitivity in type 2 diabetes patients (Moller *et al.* 1996, Purnell & Brunzell 1997) and in healthy subjects as shown in a recent study carried out by our group (Pérez-Jiménez *et al.* 2001). Furthermore, it has also been suggested that hypertriglyceridemia and an SFA diet might favor insulin resistance (Steiner *et al.* 1991, Storlien *et al.* 1993), increasing FFAs, which may inhibit glucose utilization by peripheral cells, thus reducing insulin sensitivity (Bjorntorp 1994). In addition, there is a stronger interindividual variability in the response to dietary fat, as we show in our present study. This suggests that differences in gene products involved in candidate metabolic pathways produce phenotypic differences in response to dietary changes. Although it is well known that nutrients are involved in modulating the metabolism of lipoproteins, these aspects have been poorly

Table 2 Baseline characteristics of the subjects according to the -514C/T polymorphism. Value are means±S.D.

	CC (n=39)	CT/TT (n=20)	P*
Age	22.3±1.30	23.0±1.50	0.21
Body mass index	20.9±2.70	22.2±3.28	0.21
Total cholesterol	4.02±0.65	4.47±0.56	0.01
Triglycerides	0.69±0.31	0.71±0.22	0.83
LDL-C	2.33±0.58	2.72±0.70	0.03
HDL-C	1.36±0.31	1.42±0.43	0.60
Apo B	1.74±0.65	1.66±0.48	0.66
Apo AI	1.38±0.49	1.26±0.45	0.41

*P values calculated by ANOVA.
Apo, apolipoprotein.

Table 3 Levels of plasma lipids, lipoprotein and other metabolic parameters according to -514C/T polymorphism after each of the dietary periods. SFA diet, saturated fat enriched diet; CHO-diet, low fat, high carbohydrate diet. Values are means±s.d.

	Diet	Total cholesterol	LDL-C	ApoB	Triglycerides	HDL-C	ApoA-I
Genotype CC (n=39)	SFA	4.10±0.61	2.51±0.51	71±12	0.57±0.26	1.27±0.23	148±18
	CHO	3.59±0.60	2.11±0.53	64±12	0.71±0.30	1.13±0.24	138±16
	Mediterranean	3.65±0.64	2.13±0.53	64±10	0.71±0.28	1.18±0.25	142±18
CC/TT (n=20)	SFA	4.43±0.61	2.75±0.71	75±14	0.69±0.24	1.30±0.30	153±13
	CHO	3.76±0.64	2.22±0.65	65±15	0.73±0.21	1.19±0.25	164±13
	Mediterranean	3.93±0.73	2.31±0.69	67±17	0.74±0.28	1.26±0.30	149±16
<i>P</i> -value*	Genotype	0.129	0.216	0.454	0.740	0.445	0.203
	Diet	0.000	0.000	0.000	0.268	0.000	0.000
	Interaction	0.260	0.136	0.238	0.977	0.344	0.621

**P* values calculated by ANOVA.

Apo, apolipoprotein.

investigated as regards HL. Previous studies in rats have shown that HL activity is inhibited by diets rich in saturated fats (Summerfield *et al.* 1984).

Our study clearly shows that gender interacts with genotype and diet to determine dietary modifications in insulin resistance. Likewise, in the present study, while the female carriers of the T allele displayed a tendency towards lower insulin resistance following the three dietary periods, no significant differences were observed when compared with the females homozygous for the C allele. The lack of effect of the T allele on insulin sensitivity in women is similar to the phenomenon observed in plasma HDL-C levels in response to changes in diet (Gómez *et al.* 2004). The association of the -514C/T polymorphism with HDL-C levels in a specific gender may be due to differences in exogenous administration or endogenous levels of sex hormones, which may differentially modulate lipoprotein metabolism between males and females. Moreover, because HL activity is regulated by sex-steroid hormones and because HL activity is higher in men than in women, divergent findings could be caused by gender difference, as we observed in our study. The gender difference in HL activity has led some to hypothesize that HL activity is a major determinant of the more atherogenic lipoprotein profile in men compared with women.

Insulin has been assumed to upregulate the activity of HL via insulin-responsive elements in the HL promoter. This has been proposed to explain the associations between hyperinsulinemia and high HL activity (Jansen *et al.* 1997, Romano *et al.* 1997). Jansen *et al.* (1997) observed a positive correlation between plasma levels of insulin and HL activity in non-carriers of the -514T allele, whereas no such relation was noted in carriers of the mutation. The same effect was recorded by Pihlajamäki *et al.* (2000) who observed an association between insulin resistance and the -250 G/A polymorphism in the HL promoter region. Therefore, variants in

HL promoter activity may abolish the ability of insulin to stimulate HL activity. However, whether the promoter polymorphisms of the HL gene could regulate other actions of insulin is not known. Theoretically, changes in serum FFA levels could regulate the expression of peroxisome proliferator-activator receptors and, therefore, insulin sensitivity. Alternatively, the effect of HL on insulin sensitivity could be partly mediated via changes in the amount or distribution of body lipid storage. A third possibility is that changes in HL activity primarily alter serum lipids and secondarily lead to changes in intramyocellular lipid storage; therefore, the HL promoter variant could affect insulin sensitivity in skeletal muscle. Finally, HL may have other currently unknown functions that could affect the ability of insulin to stimulate glucose uptake. In addition, it is noteworthy that the -514 site is at the center of a CAC*GGG sequence, almost analogous to the CACGTG motif characteristic of an E-box onto which the upstream stimulatory factors (USF) 1/2 can bind. The latter are transcription factors involved in the regulation of glucose and lipid metabolism in the liver. For instance, USFs are part of the insulin responsive complexes that interact with the fatty acid synthase gene (Wang & Sul 1997). It is tempting to speculate that the -514C/T substitution would disrupt the E-box analogous sequence and impair the stimulatory regulation exerted by insulin. Interestingly, it has recently been reported that USF proteins can bind to the -514 region, and that the affinity is reduced fourfold by the -514C/T substitution (Botma *et al.* 2001).

In summary, after ingestion of a saturated fat-enriched diet, male carriers of the -514T allele in the promoter region of the HL gene show decreased insulin sensitivity which improves on consuming a carbohydrate-rich diet and a Mediterranean diet. Our results suggest that male carriers of the -514T allele are at greater risk of developing the insulin resistance

Figure 1 Steady state plasma glucose concentrations in relation to the -514C/T HL promoter polymorphism after different diets in (A) men and (B) women. The saturated fat enriched diet (SFA) is represented by solid bars; the low-fat, high-carbohydrate diet (CHO diet) is represented by shaded bars and the monounsaturated fat-enriched Mediterranean diet is represented by open bars. Multivariate ANOVA for repeated measures are shown: P1, diet effect; P2, genotype effect; P3, genotype by diet interaction. Bars with different letters are significantly different ($P < 0.05$).

syndrome and type 2 diabetes mellitus when they ingest a diet rich in saturated fat. Detection of this group of subjects may be one way to inhibit the development of type 2 diabetes.

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