Triglycerides and high-density lipoprotein cholesterol are associated with insulinemia in adolescents

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Association of lipids and insulin concentration in adolescents

ARTÍCULO ORIGINAL

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Triglycerides and high-density lipoprotein cholesterol are associated with insulinemia in adolescents

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Abstract

Objective. The aim of this study was to evaluate the association between lipids and insulin concentration in adolescents. Material and Methods. A cross-sectional study of 350 adolescents aged 14-19 years old from a public high school in Guadalajara, in the state of Jalisco, Mexico, was conducted. Fasting insulin concentration was determined using microparticle enzyme immunoassay; total cholesterol and triglycerides were detected by standard enzymatic procedures; and low- and high-density lipoproteins were found using standard precipitation methods. Statistical analysis included linear multivariate regression. Results. Serum triglycerides were associated positively with insulin fasting (β = 0.003, p = 0.0001) and high-density lipoprotein cholesterol was negatively associated with insulin fasting in male adolescents 18-19 years old (β = -0.03, p = 0.012). Conclusions. The relationships between triglycerides and insulin and between high-density lipoprotein cholesterol and insulin are already present in adolescence.

Key words: triglycerides, high-density lipoprotein cholesterol, lipids, insulin, obesity, adolescents, Mexico

Resumen

Objetivo. Evaluar la relación entre lípidos e insulina en adolescentes. Material y métodos. Estudio transversal en 350 adolescentes de 14 a 19 años de preparatoria pública en Guadalajara, Jal., México. La insulina se midió por inmunoensayo enzimático de micropartícula, el colesterol total y triglicéridos por pruebas enzimáticas estándares, y las lipoproteínas de baja y alta densidad por métodos de precipitación. Se utilizó análisis de regresión lineal multivariada. Resultados. Los triglicéridos se asociaron en forma positiva con insulina (β = 0.003, p = 0.0001) y la lipoproteína de alta densidad se asoció en forma negativa con insulina en varones de 18-19 años de edad (β = -0.03, p = 0.012). Conclusiones. En los adolescentes, los triglicéridos y las lipoproteínas de alta densidad están asociados con la insulinemia.

Palabras claves: triglicéridos, lipoproteínas de alta densidad; lípidos; insulina; obesidad; adolescentes, México
Type 2 diabetes and cardiovascular disease are the leading causes of mortality in Mexico. Both appear to begin early in life. Considering the recent findings of high incidence of type 2 diabetes in the Mexican population, the problem is likely to reach more serious proportions. Moreover, it has been shown that type 2 diabetes incidence in adolescents has increased in recent years and a higher prevalence of metabolic syndrome has been shown among Mexican-American adolescents. Hyperinsulinemia is the strongest predictor for type 2 diabetes and is associated with lipid-profile abnormalities. Obesity is related with hyperinsulinemia and altered lipid and lipoprotein concentrations in adults, especially in those with abdominal fat distribution. Insulin, lipid, and lipoprotein concentrations might be altered by environmental as well as genetic factors. In adults, triglycerides, low-density lipoprotein cholesterol (LDL), and very-low-density lipoprotein cholesterol (VLDL) are positively associated with insulin resistance, while high-density lipoprotein cholesterol (HDL) is negatively associated with insulin resistance.

Hyperinsulinemia, obesity, and lipid and lipoprotein metabolism abnormalities may be present during adolescence and may persist through adulthood, at which time they contribute to the establishment of metabolic syndrome and eventually influence the development of diabetes along with cardiovascular risks later in life. The higher concentrations of insulin, lipids, and lipoproteins observed during early adolescence as compared with late adolescence are explained mainly by pubertal growth spurt related insulin secretion increases. The association among insulin, lipids, and lipoproteins during this period of life is not completely understood. There is controversy concerning this matter, as some researchers have found that triglycerides are positively associated with insulin, whereas others have also found associations with LDL, HDL, and VLDL. Total cholesterol has not been associated with insulin. Furthermore, the association between insulin, abdominal fat, blood lipids, and lipoproteins in adolescents is not as clear as it is in adults.

The aim of this study was to determine the relationship among lipids, lipoprotein profile, and fasting insulin concentration, as well as the relationships between lipids, lipoproteins, insulin, and abdominal obesity in adolescents, as evaluated by waist circumference (WC) and waist-to-hip-ratio (WHR).

### Material and Methods

**Subjects.** The relationship among lipids, lipoprotein profile, insulin, and obesity in adolescents was investigated using a cross-sectional study. Recruitment and data collection were carried out from October 1998 to March 1999. Methods and procedures pertinent to this study have been reported previously. Briefly, 350 adolescents (113 boys and 237 girls) were randomly recruited from 10th to 12th-grade classes at a state-funded Mexican high school in the Jalisco state capital of Guadalajara. Within the sample, 45.4% of adolescents were classified as high socioeconomic status, 33.4% as middle, and 21.1% as low. Adolescents who worked comprised 27.0% of the sample. Both adolescents and parents gave written informed consent in accordance with the Institutional Review Board of the Instituto Mexicano del Seguro Social (Mexican Institute of Social Security) in Guadalajara.

**Main outcome measurements.** Fasting blood samples were obtained for serum lipid, lipoprotein, and insulin determinations after 12 h. Serum samples were stored frozen at -25°C until analyzed. Serum insulin was determined based on microparticle enzyme immunoassay using anti-insulin (mouse-monoclonal)-coated microplates. Regarding lipids and lipoproteins, total serum cholesterol and triglycerides were determined by standard enzymatic procedures, while LDL and HDL were determined using standard precipitation methods with polyvinyl sulphate and phosphotungstic acid, respectively. A BMI analyzer (Hitachi 705, Indianapolis, IN, USA) was used for all assays. VLDL was estimated according to the following equation: $VLDL = \frac{Total\ cholesterol - (HDL + LDL)}{2}$. Intra- and interassay variation coefficients were 4.0 and 6.2% for insulin and <2.7 and <3.7% for all lipids, respectively.

A systematic physical examination was performed on all study participants, and body weight, height, and waist and hip circumference measurements were assessed according to standardized protocols. Overweight was defined as body mass index (BMI, kg/m²) for age >85th percentile. Unfavorable abdominal fat distribution was defined as a WHR ≥0.85 for females and ≥0.94 for males.

In addition, a validated 7-day physical-activity recall questionnaire was administered by a trained dietitian. Activities were classified according to metabolic equivalents (METS) (ratio of working metabolic rate/resting metabolic rate). Energy expenditure (EE) was estimated as follows: \[ EE = \sum [weight \times \text{activity intensity}] \text{METs} \times \text{duration (days)} \]
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(1) To assess energy and lipid intake, a validated semi-quantitative food-frequency questionnaire was administered containing 124 items.30 Average daily energy and lipid intake were derived using the System of Nutritional Habits and Nutrients Intake (SNUT) evaluation program.31 Socioeconomic status, family history of type 2 diabetes, and smoking status were assessed using a standardized questionnaire.

Statistical analysis. Descriptive analysis was performed to estimate the study population’s clinical and anthropometric characteristics; results are expressed as mean ± standard deviation (X ± SD) or as percentages. Differences between means were obtained with the Student t test for continuous variables and proportions were compared using a χ² test. Partial correlation coefficients were computed to determine any correlation among serum lipids, lipoproteins, and insulin, adjusted for age and sex. Adjusted blood lipids and lipoprotein means were obtained across anthropometric variables (BMI, WC, and WHR). Confounders included age, sex, BMI, WC, EE, caloric intake, and smoking status. Linear trend was tested with multivariate analysis. Adjusted means of lipid and lipoprotein concentrations were obtained across anthropometric variables (BMI, WC, and WHR). Confounders included age, sex, BMI, WC, EE, caloric intake, and smoking status. Under partial correlation analysis adjusting for age and sex, fasting insulin concentration correlated positively with lipids and lipoproteins, except VLDL, and correlated negatively with HDL. BMI and WC correlated positively with insulin, lipids, and lipoproteins, and negatively with HDL. BMI showed strongest correlations for nearly all parameters, WHR also correlated positively with all these parameters except total cholesterol and LDL, and WC showed stronger correlations with these parameters in comparison with WHR (Table I).

Linear trend results of lipids and lipoproteins across anthropometric parameters demonstrated that total cholesterol, triglycerides, and LDL increased from the lowest to the highest BMI quartile (147.7-168.5 mg/dl, p= 0.02, 91.2-118.4 mg/dl, p= 0.008, and 85.2-120.9 mg/dl, p=0.07). Table I

Table I

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Insulin (pmol/l)</th>
<th>Cholesterol (mg/dl)</th>
<th>Triglycerides (mg/dl)</th>
<th>LDL (mg/dl)</th>
<th>VLDL (mg/dl)</th>
<th>HDL (mg/dl)</th>
<th>HL (mg/dl)</th>
<th>HDL/CHOL (mg/dl/mg/dl)</th>
<th>LDL/CHOL (mg/dl/mg/dl)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Insulin (pmol/l)</td>
<td>0.207*</td>
<td>0.357*</td>
<td>0.259*</td>
<td>0.099*</td>
<td>-0.23*</td>
<td>0.343*</td>
<td>0.363*</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Body mass index (kg/m²)</td>
<td>0.610*</td>
<td>0.187*</td>
<td>0.377*</td>
<td>0.250*</td>
<td>0.099</td>
<td>-0.23*</td>
<td>0.343*</td>
<td>0.363*</td>
<td></td>
</tr>
<tr>
<td>Waist circumference (cm)</td>
<td>0.587*</td>
<td>0.152*</td>
<td>0.383*</td>
<td>0.216*</td>
<td>0.145*</td>
<td>-0.295*</td>
<td>0.340*</td>
<td>0.381*</td>
<td></td>
</tr>
<tr>
<td>Waist-to-hip ratio (cm/cm)</td>
<td>0.386*</td>
<td>0.029</td>
<td>0.281*</td>
<td>0.099</td>
<td>0.112*</td>
<td>-0.281*</td>
<td>0.247*</td>
<td>0.288*</td>
<td></td>
</tr>
</tbody>
</table>

* p< 0.0001
† p< 0.05
‡ p< 0.01

LDL (low-density lipoprotein cholesterol); VLDL (very low-density lipoprotein cholesterol); HDL (high-density lipoprotein cholesterol); LDL/HDL (low-density lipoprotein cholesterol/high-density lipoprotein cholesterol ratio; CHOL/HDL (total cholesterol/high-density lipoprotein cholesterol ratio)

Results

The study population consisted of 350 adolescents, 68% female, with a mean age of 16 ± 1.19 years (range, 14-19 years old). Nineteen percent of all adolescents were overweight, 3% had unfavorable abdominal fat distribution, 22% were current smokers, and 8.3% had a family history of type 2 diabetes. Mean fasting serum insulin concentration was 54.8 ± 29.4 pmol/l; triglycerides, 100.3 ± 35.4 mg/dl; total cholesterol (CHOL), 157.2 ± 27.0 mg/dl; LDL, 95.2 ± 26.3 mg/dl; VLDL, 18.5 ± 7.3 mg/dl; and HDL, 43.5 ± 8.9 mg/dl. Lipid and lipoprotein profile distribution by sex and socioeconomic status in this population has been reported elsewhere.32

Under partial correlation analysis adjusting for age and sex, fasting insulin concentration correlated positively with lipids and lipoproteins, except VLDL, and correlated negatively with HDL. BMI and WC correlated positively with insulin, lipids, and lipoproteins, and negatively with HDL. BMI showed strongest correlations for nearly all parameters, WHR also correlated positively with all these parameters except total cholesterol and LDL, and WC showed stronger correlations with these parameters in comparison with WHR (Table I).
Triglycerides increased from the lowest to the highest WC quartile (97.6-114.5 mg/dl, p = 0.046), while HDL decreased from the lowest to the highest WHR quartile (47.1-39.9 mg/dl, p < 0.001).

Table II shows median values of fasting lipid and lipoprotein concentrations according to insulin quartiles after adjustment for known confounders. After conducting multivariate regression analysis as well as a trend test run for each lipid and lipoprotein, only triglycerides were found to increase across insulin quartiles from lowest to highest (93.1 mg/dl-110.4 mg/dl, p = 0.043).

Total cholesterol, triglycerides, LDL, VLDL, and HDL, and total CHOL/HDL ratio were associated with insulin after adjustment for age and sex. However, the relationship among insulin and total cholesterol, LDL, VLDL, and HDL was not statistically significant when BMI was included in the model. Moreover, when WC was included triglycerides and CHOL/LDL ratio remained independently associated with insulin concentration. Finally, when EE was included in the model only triglycerides were associated with insulin concentration (Table III).

When no significance was found in the model adjusted by age, sex, BMI, WC, and EE, additional multiple linear analyses were performed by age group (14-15, 16-17, and 18-19 years) and sex (male and female). These analyses demonstrated a significant relationship between insulin and HDL in males 18-19 years old when BMI (β = -0.03, p = 0.012), WC (β = -0.03, p = 0.009), and EE (β = -0.03, p = 0.02) were included. The variance inflation factor was 3.8 and no problems were detected for multicollinearity. These associations were not statistically significant in younger adolescents.

Table IV shows the relationship between serum triglycerides and insulin, adjusted for socioeconomic and environmental factors. Triglycerides were associated with fasting insulin concentration (β = 0.003,
Table IV  
**Relationship between serum triglycerides and insulin in adolescents in Guadalajara, Jalisco, Mexico, 1998-1999 (n=348)**

<table>
<thead>
<tr>
<th>Predictors</th>
<th>β</th>
<th>95% CI</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Triglycerides (mg/dl)</td>
<td>0.00267</td>
<td>0.0012, 0.0041</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Body mass index (kg/m²)</td>
<td>0.07582</td>
<td>0.0401, 0.1116</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Waist circumference (cm)</td>
<td>0.01586</td>
<td>0.0007, 0.0310</td>
<td>0.040</td>
</tr>
<tr>
<td>Energy expenditure (kcal/day)</td>
<td>-0.00038</td>
<td>-0.0005, -0.0002</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Energy intake (kcal/day)</td>
<td>0.00001</td>
<td>-0.0001, 0.0001</td>
<td>0.800</td>
</tr>
</tbody>
</table>

*Results of multivariate linear regression [R²= 0.41, F= 14.23] Adjusted for age (years), sex (female/male), family history of diabetes (yes/no), cigarette smoking categories (never, 1-20, 21-100, 101-250, >250 cigarettes per year), saturated fat (g/day), monosaturated fat (g/day), and polyunsaturated fat (g/day). 95% CI

Discussion

The findings of this study suggest that insulin resistance syndrome factors are already present in adolescence and, as demonstrated in adult studies, triglyceridemia, HDL, and obesity are related with insulinemia independently of other relevant factors. The results show that hyperinsulinemia correlates negatively with HDL and positively with total cholesterol, triglycerides, LDL, and the CHOL/HDL ratio, after age and sex adjustments. A further adjustment including BMI, WC, EE and smoking status was carried out. With the exception of triglycerides and HDL in 18-19-year-old male adolescents, associations among insulin, lipids, and lipoproteins disappeared. These results are similar to those reported by Bonora et al.,15 except that those authors found no association with HDL. Other studies have found that insulin resistance is also related with LDL and VLDL,16,17 but these associations may have been confounded because BMI, WC, and EE were not controlled. Moreover, adolescents in those studies were younger than adolescent participants in the study presented here, which also may have contributed to the differences. Although only triglycerides and HDL in males aged 18-19 years were significantly associated with insulin in this study, it is important to point out the tendency toward higher fasting insulin with ad-verse lipid or lipoprotein concentration, as observed in Table II. Although some metabolic alterations persist from adolescence to young adulthood,33 the possible biological implications for late adulthood of tendencies observed in the adolescents representative of this study are yet to be established.

During adolescence, insulin, lipid, and lipoprotein concentrations change according to sex and puberty.14,33-34 Although puberty was not evaluated in this study, it is assumed that adolescents are mainly at Tanner stages 4 and 5, because in other series it has been found that 56% of 14- to 19-year-old adolescents from Guadalajara were at Tanner stage 4 and 42% at Tanner stage 5 (unpublished data). Total cholesterol and LDL were not associated with insulin probably because of total cholesterol decreases during Tanner stages 4 and 5,34,35 possibly due to the effect of steroid hormone increases. For example, in this study, total cholesterol values fell from 165.5 in 14-15 years old to 157.6 mg/dl in 18-19 years old females. The same pattern occurred with LDL which decreased from 102.4-92.4 mg/dl in 14-15 years old to 92.4 mg/dl in 18-19 years old.32

One interesting finding in this study comprised the negative association between HDL and insulin, particularly in 18-19-year-old males compared with younger males or females. These results differ from the results of a study of 18-year-old healthy young men in Italy in which HDL was not associated negatively with insulin. However, HDL concentration was higher in adolescents in that study than in those in the study presented here (53.7 vs. 36.3 mg/dl).15 This is probably due to the Italian adolescents’ diet, which was rich in monounsaturated fat. The present finding may suggest that atherogenesis development could be triggered at this young age, which warrants further investigation.

High abdominal fat may increase free fatty acid flux to the liver, thereby increasing its oxidation, modifying glucose disposal to extra-hepatic tissues, and increasing insulin resistance.36 This high free fatty acid flow to hepatocytes may modify the number of insulin receptors, alter receptor function, and modify insulin internalization and degradation. These effects may account for altered hepatic insulin extraction and sensitivity.37 It was expected that BMI, WC, and WHR would be independent predictors of serum insulin, as had been published,7,19 nevertheless, as demonstrated elsewhere15 this study found that only BMI and WC were independently associated. This study controlled for potential confounders, which might explain these discrepancies. Furthermore, lack of an association might be explained by the early development of abdominal fat deposits at this stage of life, or because WHR does not really capture visceral adiposity as it...
does in adults, in whom it was shown to be an important risk factor for insulin and lipid concentrations.

The relationship between serum insulin concentration and lipid and lipoprotein profile is better understood in adults than in adolescents. Perhaps there is a critical time-frame that could enable identification of an evolving dysfunctional metabolism. Improved understanding of several factors, including the genetics and physiology of puberty, would explain associations between lipids and insulin in adolescents. Due to the cross-sectional design of this study, no causal relationship between triglycerides and insulin or between HDL and insulin can be established. However, the data in this study support the hypothesis that high triglyceride concentration, low HDL concentration, obesity, and abdominal fat may be independently associated with hyperinsulinemia. Future cohort studies and interventions in adolescents would be useful to establish causality between triglycerides and insulin, and HDL and insulin. Meanwhile, the results of this study suggest that it is important to initiate primary prevention programs to reduce, in adolescence, these risk factors for chronic diseases, such as type 2 diabetes mellitus.

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