Plasma lipids and their interrelationship in Turkish adults

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Abstract

Study objective—The aim was to describe the plasma total cholesterol and triglyceride profiles in a random sample of Turkish adults and analyse the effects of certain coronary risk factors on these levels.

Design—This was a cross sectional population based survey.

Setting—59 communities scattered in all seven geographical regions of Turkey were surveyed in the summer of 1990.

Subjects—A random sample of 3689 men and women 20 years of age and over was studied.

Measurements and main results—Plasma total cholesterol, triglycerides, glucose (using Reflotron and with partial validation in reference laboratory), weight, height, and blood pressure were measured, and information on smoking, physical activity, and family income obtained. Hypercholesterolaemia (≥6.5 mmol/litre, 250 mg/dl) prevailed in 8.5%, and hypertriglyceridaemia (≥2.25 mmol/litre, 200 mg/dl) in 16.6% among men and women aged 40–59 years of age. Age adjusted total cholesterol values were 4.8 mmol/litre (185 mg/dl) in men and 5 mmol/litre (192 mg/dl) in women. A steep rise appeared in mean cholesterol levels between the ages of 20–29 and 40–49 years, in a ratio greater than the available data from some other populations indicated. Mean total cholesterol values increased substantially in both genders with diminishing grades of physical activity, rising serum triglyceride levels, in urban (opposed to rural) residents, in men with increasing income levels, and in the younger adults with rising body mass index.

Conclusions—Turkish adults have comparatively low levels of total cholesterol and medium to moderately high levels of triglycerides. Lifestyle factors affect these levels in Turks as in other populations.

Methods

The survey on the prevalence of cardiac disease and risk factors in adults in Turkey includes 3689 men and women 20 years of age and over residing in 59 different communities scattered over all the seven geographical regions of Turkey. The criteria for selecting the urban and rural communities, participating subjects, the surveying teams and the steering committee, methods of data collection, and the data obtained in the questionnaire have been presented in a separate report. Briefly, a random sample of the Turkish adult population was surveyed with the purpose of determining the prevalence of heart diseases and the risk factors for coronary artery disease. The sample was representatively stratified for sex, age, and geographical region as well as for the rural-urban distribution.

In this study a community was defined as rural when it had a population less than 8000 and urban when its population exceeded this figure. Communities were selected in the sample so that the same proportion of the rural population (43%) and of cities with a population over 500 000 (27%) was present as in the whole of Turkey (43.7% and 23.4% respectively). Towns with intermediate populations were also represented proportionately in the sample.

In the selection of participants from the various communities, the number of subjects in each age group was predetermined. When each surveying team reached the sample community, they first obtained information from the local authority about the socioeconomic distribution of the living quarters, and then rang randomly preselected doors in the evening and gave appointments for an
examination the next morning. Roughly 60–90 persons were invited for examination to obtain a mean of 62 persons per community to be surveyed. The ratio of responders exceeded 85%. Males and females were partly selected from the same, and partly from different households. The sample did not include participants using lipid lowering drugs. In over 65% of participants (1151 men and 1257 women) blood was sampled in the postabsorptive state 12–15 h after the last meal for total cholesterol, triglycerides, and glucose. In the remaining persons cholesterol alone was determined 1–5 h after the ingestion of a breakfast.

Three teams were formed, each consisting of two physicians in their fourth year of specialisation in internal medicine and a laboratory technician receiving postgraduate training in medical biology. A schedule was designed to examine 24–30 persons daily for 5–6 days a week to allow the completion of the survey within 8–11 weeks. The survey started on July 13th and ended late in September, 1990. The task of each physician was to examine the cardiovascular system and record the ECG. The technician obtained blood samples by finger prick using disposable lancets and determined the plasma concentrations of cholesterol, triglycerides, and glucose using the Reflotron apparatus. Logistics were provided by the Ministry of Health of the Turkish Republic, each team having a large ambulance with a driver at its disposal.

The Reflotron operators were given a course over two full days in Istanbul by Boehringer representatives to train them to operate the instrument and to become familiar with the use of pipettes and reagent sticks. By the end of this time it was felt they demonstrated competence in carrying out these operations. Subsequently all team members were given a training course over a period of two days by the supervising staff to acquaint them with the technique of random sampling, the approach to adopt in the communities, and other pertinent items.

Internal quality control was made with control sticks obtained from the manufacturer. These were utilised at the beginning and commonly also at the end of each day of surveying. Any sampled value above 7.76 mmol/litre (300 mg/dl) or below 2.59 mmol/litre (100 mg/dl) for cholesterol was remeasured at the same sitting, and both were recorded. External quality control was made with reference to the biochemistry laboratory of the Admiral Bristol Hospital, Istanbul, certified for accuracy by the Center for Disease Control, Houston Branch, USA. Venous blood samples were drawn by each team once or twice a week into tubes containing 1 mg/dl of EDTA as anticoagulant and were placed into dry tubes after centrifugation, to be forwarded the same day via aeroplane to the reference laboratory, which determined the cholesterol concentration by the enzymatic method using Boehringer (Mannheim) kits. A total of 212 samples was sent. These samples were obtained on 17 different days from participants residing in 13 cities which were served by an airline. A random selection of one in four blood samples was selected on that day for validation.

The Reflotron apparatus indicated triglyceride levels below 0.79 mmol/litre (70 mg/dl) as such, without specifying accurately. Subjects with such values comprised 23.8%, of those in whom blood was sampled for triglycerides. By plotting a histogram of the log normal distribution it was estimated that values shown as less than 0.79 mmol/litre corresponded to a mean of 0.62 mmol/litre. This presumed figure was used in calculating mean values in the respective age groups in men and women.

Physical examination consisted of weighing and measuring the surveyed person without shoes and heavy outer garments; palpation of the character, site, and size of cardiac impulses; auscultating the heart sounds and identifying murmurs; and measuring the blood pressure twice in the sitting position in the right arm. Body mass index was calculated by the computer as weight divided by height squared (kg/m²) to characterise the relative weight of the participants. All survey data were checked by the supervising staff before being included in the database, so as to ensure data quality and completeness.

The effect of physical activity upon the blood concentration of cholesterol was studied in this survey in four categories: grade I (minimal): white collar worker, sewing-knitting, walking <1 km daily; grade 2 (slight): repair worker, housework, walking 1–2 km daily; grade 3 (moderate): mason, carpenter, truck driver, cleaning floors and windows, walking 4 km daily; grade 4 (heavy): heavy labour, farming, regular sports activity. The correlation coefficient between two studied variables was determined and when found sufficiently high the regression equation between them was computed according to the method of Pearson-Bravais.

### Results

#### VALIDATION STUDY

In 212 plasma samples in which the total cholesterol concentration was measured by both the Reflotron and the reference laboratory, the correlation coefficient between individual values was 0.90. The mean concentration determined in the reference laboratory was 4.36 mmol/litre (168.6 mg/dl) resulting in a bias of +0.063 mmol/litre (+1.42 mg/dl). This bias, representing a deviation of +1.14%, is acceptable.³

#### MEAN PLASMA CHOLESTEROL

Mean plasma total cholesterol values of 3687 participants, stratified into sex, age group, and urban or rural residence, are presented in table I.

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### Table I Mean plasma total cholesterol and triglyceride concentrations in Turkish adults in various age groups (in mmol/litre)

<table>
<thead>
<tr>
<th>Age groups (years)</th>
<th>Men Cholesterol</th>
<th>Men Triglyceride</th>
<th>Women Cholesterol</th>
<th>Women Triglyceride</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>n</td>
<td>Mean (SD)</td>
<td>n</td>
<td>Mean (SD)</td>
</tr>
<tr>
<td>20–29</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>30–39</td>
<td>424</td>
<td>4.51 (0.98)</td>
<td>263</td>
<td>1.69 (1.12)</td>
</tr>
<tr>
<td>40–49</td>
<td>296</td>
<td>4.86 (1.07)</td>
<td>198</td>
<td>1.98 (1.14)</td>
</tr>
<tr>
<td>50–59</td>
<td>258</td>
<td>4.89 (1.14)</td>
<td>172</td>
<td>1.62 (0.99)</td>
</tr>
<tr>
<td>60–69</td>
<td>206</td>
<td>4.75 (1.06)</td>
<td>146</td>
<td>1.46 (0.89)</td>
</tr>
<tr>
<td>70 and over</td>
<td>82</td>
<td>4.58 (1.01)</td>
<td>56</td>
<td>1.30 (0.56)</td>
</tr>
</tbody>
</table>

Total urban       | 1085 | 4.47 (1.09) | 646 | 1.56 (1.04) | 1034 | 4.63 (1.09) | 706 | 1.34 (0.78) |

Total rural       | 802 | 4.37 (1.05) | 505 | 1.47 (0.90) | 786 | 4.53 (0.99) | 551 | 1.29 (0.83) |
and graphically depicted in fig 1. Low average cholesterol concentrations prevailed in Turkish men and women in the age group 20–29 years (3.82 and 3.97 mmol/litre, respectively, or 148 and 153 mg/dl). These values rise rapidly in the age group 30–39 years to 4.5 mmol/litre (174 mg/dl) in men and 4.43 mmol/litre (171 mg/dl) in women, and further to 4.86 mmol/litre (188 mg/dl) in both sexes in the age bracket 40–49 years. Within two age groups 20 years apart this represents a 27% rise in Turkish males and 23% in females. As in surveys of other communities males in this survey reached a plateau in the 40–49 year age period, with a mean value of 4.9 mmol/litre (189 mg/dl) in the 50–59 year age group and a slight decline to 4.76 mmol/litre (184 mg/dl) in the 60–69 year age group, and further to 4.58 mmol/litre (177 mg/dl) at age 70 and over. In women, mean total cholesterol reached a plateau of 5.27 mmol/litre a decade later, in the 50–59 year age group, and this persisted in the subsequent decade before declining to 5 mmol/litre (194 mg/dl) at age 70 and over. From age 40 years onwards, total cholesterol values were on average 0.35 mmol/litre (13.5 mg/dl) higher in urban males than in their rural counterparts (p < 0.001), a difference of 7%. Urban women of the same age also had higher total cholesterol than rural women, but the difference was less, namely 0.22 mmol/litre (8.4 mg/dl), equivalent to 4.4% (p < 0.01).

Female participants over 40 years of age had higher mean total cholesterol concentrations than males; the difference was more pronounced among rural residents at 0.38 mmol/litre (14.4 mg/dl) (p < 0.001) than among the urban population (0.27 mmol/litre, 9.6 mg/dl, p < 0.01).

**Prevalence of Hypercholesterolaemia**

In the general adult sample population of 3687 subjects, the following total cholesterol percentile values were observed, in mmol/litre (mg/dl): 10th percentile: 3.16 (122); 25th percentile: 3.73 (144); median: 4.45 (172); 90th percentile: 5.92 (229); 95th percentile: 6.44 (249). The following cut off values were noted: 4.7% over 6.45 mmol/litre (250 mg/dl), 0.7% over 7.76 mmol/litre (300 mg/dl). The prevalence of hypercholesterolaemia (defined as 6.5 mmol/litre and over, or over 250 mg/dl) in various age groups in each sex is presented in fig 2.

The interrelations between plasma cholesterol concentration and certain variables such as body mass index, physical activity, triglycerides, and income level were investigated. With the purpose of eliminating age as a joint determinant of serum cholesterol, the relation between the latter and body mass index was sought in various age groups. This relationship was significant in men solely in the age groups 20–29 years (γ = [63 + 3.48x] ± 33.3; r = 0.32) and 30–39 years (γ = [107 + 2.69x] ± 36.6; r = 0.26), and in women in the age group 20–29 years (γ = [107 + 1.95x] ± 34.5; r = 0.24). Hence, as indicated by the coefficients, in Turkish men aged 20–39 years each unit of body mass index raised the mean cholesterol level by slightly over 0.08 mmol/litre (3 mg/dl) and in women by 0.05 mmol/litre (2 mg/dl) (fig 3).

Age standardisation was performed using fixed weights of five age groups in each category of physical activity in urban and rural population with separate but minimally different weighting for men and women. The mean total cholesterol values in each grade of physical activity among adults of age 20–69 years are represented in fig 4. As activity grade increased from 1 to 4, cholesterol concentrations declined in men from 4.76 to 4.19 mmol/litre, diminishing at each activity grade by a mean of 0.57 mmol/litre. Similarly, plasma cholesterol decreased in women from 4.74 mmol/litre to 4.3 mmol/litre, by an average of 0.15 mmol/litre for each increase in physical activity grade. Figure 4 clearly indicates that
lower cholesterol values occur with increasing grades of physical activity in both the rural and the urban population, and in women as well as in men. Due to the relatively small size of the sample of rural women in activity grade 1 and of urban women in grade 4, the related separate urban-rural data may have limited reliability. However, data in the other three activity grades in women and all those in men possess a sufficiently broad base and are consistent in showing a definite trend of decreasing cholesterol levels with rising physical activity. Over three activity grades, a reduction in the mean cholesterol value of 12% was observed in urban and rural men. An effect of similar magnitude prevailed in rural women, but it was less in urban women, at only 7%.

In examining the effect of physical activity on plasma cholesterol, another approach was to plot the regression line between age and cholesterol in the various grades of physical activity. Though at grade 1 there was no significant correlation, the correlation coefficients for higher grades were of sufficient magnitude (0.24 to 0.40 in men; 0.43 to 0.52 in women) to define the relationship between age and cholesterol (table I). Worthy of note is that between grades 2 and 4 of activity, mean plasma cholesterol values in men aged 25 to 50 years by 0.17 to 0.61 mmol/litre, and in women of similar age by 0.20 to 0.41 mmol/litre. It appears that the role of physical activity in moderating serum cholesterol greatly diminishes after age 50 years.

The income level also appeared to influence the mean plasma cholesterol concentrations, in men more clearly than in women. When the regression equation was determined between average serum cholesterol value and a particular family income bracket for certain ages, it was observed that for a man aged 40 years (or 55 years) the mean cholesterol concentration in the highest income bracket exceeded by 0.55 mmol/litre (or 0.79 mmol/litre) that in the lowest bracket. The magnitude of the effect of income level on plasma cholesterol in women was about two fifths of that in men at age 40 years, and about one third at age 55 years (fig 5). When the influence of urban-rural living was examined, it was found that the positive relation between family income and mean cholesterol did not exist in rural women, but persisted in urban women as well as in urban and rural men. Between the lowest and highest income brackets in a 40 (or 55) year old man, the mean cholesterol gradient amounted to 0.48 (or 0.60) mmol/litre in both the urban and the rural sectors. (The subset of rural men with the highest income bracket was not taken into account, since no valid regression equation was obtained due to small sample size).

Plasma cholesterol data for men and women aged 40–59 years (with identical weighting of the two age groups) in various geographical regions of Turkey are presented in table III. When assessed for men and women combined, the Marmara region ranks top, the Mediterranean and south-east Anatolia regions rank lowest, while the remaining four regions rank in the middle with a mean cholesterol value varying closely around 5 mmol/litre.

TRIGLYCERIDE CONCENTRATIONS

Plasma triglyceride concentrations showed a log normal distribution. Mean triglyceride values in the Turkish adult sample are presented by gender and age groups in table 1, and certain percentile values are shown graphically in fig 6. The latter data indicate that median values in men attain a plateau between the ages 35–55 years of around 1.55 mmol/litre (137 mg/dl) before declining, whereas in women a progressive rise is observed, from 0.8 mmol/litre in the age group 20–29 years to a peak of 1.58 mmol/litre in age group 60–69.

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**Figure 4** Mean plasma cholesterol after adjustment for age, by grades of physical activity in urban and rural men and women aged 20 to 69 years.

**Figure 5** Relation between mean plasma cholesterol values and family income levels in Turkish men and women.
years. Mean values standardised for age 30–69 years were 1.68 mmol/litre in men and 1.42 mmol/litre in women. Hypertriglyceridaemia, defined as 2.26 mmol/litre (200 mg/dl) and over, prevailed in 9.8% of all adult women and 14.8% of men. In the decades spanning the ages 40–69 years, 14.7% of women and 19.3% of men were found to be hypertriglyceridaemic.

A positive relationship existed between body mass index and mean plasma triglyceride concentration in adults aged 20 to 49 years, but not in those aged 50 years and over (fig 7). In men, a rise of 0.08–0.09 mmol/litre (7–8 mg/dl) was noted for each unit increment in body mass index, while the corresponding rise in women was barely half that (0.04 mmol/litre, or 3.5 mg/dl).

Data available on plasma concentrations of both total cholesterol and triglyceride in 2408 individuals were computed to provide regression equations between the cholesterol values and those of triglyceride and age. The following equations were obtained (units = mg/dl):

**In men:**
\[ \text{cholesterol} = 114.6 + (0.201 \times \text{triglyceride}) + (0.77 \times \text{age}) \]
\[ r = 0.50, p < 0.001 \]

**In women:**
\[ \text{cholesterol} = 112.3 + (0.191 \times \text{triglyceride}) + (1.08 \times \text{age}) \]
\[ r = 0.55, p < 0.001 \]

The equation implies a change of 1 mg/dl in cholesterol for each change of about 5 mg/dl in plasma triglyceride in our sample population.

**Discussion**

Two indices of an acceptable accuracy of cholesterol measurement exist in this study: (1) Though a conventional biochemical method was used in the reference laboratory in contrast to a “dry chemistry” method by the Reflotron apparatus in the survey, the bias between the mean of the two sets of measurements was small (+1.4%), and the correlation between individual measurements was high \( r = 0.90 \). (2) The “interior consistency” of the data is apparent: the rise of cholesterol values during early adulthood, the delayed attainment of the plateau in women compared to men, the fall past the age of 70 years, the relationship with triglycerides and body mass index, etc.

It is important to bear in mind that the median age of the population surveyed was only 37 years, so that the overall mean of cholesterol values has limited significance. When age adjustment is performed according to World Health Organization criteria, Turkish men had a mean value of 4.8 mmol/litre (185 mg/dl), and women 5.0 mmol/litre (192 mg/dl). These cholesterol concentrations are far lower than are found in northern European countries and lie in the lower range of the Mediterranean populations. For an overall comparison one may recall that, excluding Beijing, China, the mean age adjusted blood cholesterol values in the 30 communities included in the MONICA project ranged between 5.3 and 6.4 mmol/litre in men and about 0.1 mmol/litre higher in women. The areas of Bremen, Belfast, and Charleroi assumed a median position among these communities, with mean values of 6.0 to 6.1 mmol/litre. Thus Turkish men have age adjusted average total cholesterol values of 1.1 mmol/litre lower, and women 0.9–1.0 mmol/litre lower, than in the mentioned cities of the developed world.

When comparison is made with the data of the Lipid Research Clinics5,6 a striking parallel was seen during adult life between the cholesterol curve of white Americans and that of Turkish men between the ages of 20 and 59 years, the latter being on average 0.6 mmol/litre lower than the former (and somewhat more from the age of 60 on). The difference between Turkish and American women in the age range 20 to 59 years was about 0.4 mmol/litre (15 mg/dl) in favour of the former.

Although the present study is merely of a cross sectional nature, with limited longitudinal implications, it is worth pointing out the steep rise in mean cholesterol levels in Turkish subjects observed between the age groups 20–29 years and 40–49 years. Young male and female Turks appear to enter adulthood with very low cholesterol levels, only 0.4 mmol/litre higher than those of the Chinese,7 yet their mean cholesterol concentration rises by 1 mmol/litre (39 mg/dl), equivalent to 25%, in this period of two decades. This is an appreciably steeper rate of rise than in available studies for other populations. As calculated by us, data from US adults5,6 show an 18% rise, almost identical to the findings from
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Italy\(^8\) (17.4\%). Data from Japan\(^9\) show a rise of 12\% and from China\(^7\) a rise of 11\%. In the study of 12 092 British men and women reported by Mann et al.,\(^10\) mean plasma total cholesterol values in both sexes were 5.15 mmol/litre at age 25–29 years and increased to 6 mmol/litre at age 45–49 years. The “rise” within the age groups two decades apart was 16.5\%. The increase in total cholesterol concentration in this earlier period of adulthood is known to stem primarily from an augmentation in low density lipoprotein (LDL) cholesterol,\(^5\)\(^11\) and this is in turn believed to reflect a progressive decrease in LDL catabolism with age which, particularly in women, could reflect hormonal changes.\(^11\) However, the observation that a significant rise in cholesterol concentrations in early adulthood is lacking in some African sample populations\(^12\) implies that underlying environmental factors also play a role in this phenomenon after the age of about 25 years.

This survey revealed significantly higher mean cholesterol values in the urban population than in the rural population, by on average 4\% in women and 8\% in men. Between the urban and rural sample population in Augsburg, Germany, the difference was far less, namely 0.1 mmol/litre both in men and women.\(^4\) It is worth noting that no significant difference existed in our survey between urban and rural participants with regard to the prevalence of hypertension in either sex, or smoking in men.

The prevalence of hypercholesterolaemia (defined as a level greater than 6.5 mmol/litre or 250 mg/dl) in Turkish men of 40–59 years of age was 8\% and was thus comparable to that of Yugoslavia and Japan, but moderately lower than that of Italy (13\%) and Greece (14\%), found in the Seven Country Study.\(^13\)

A recent study estimated the average consumption of “visible” fats (butter, fats, and oils) in Turkey as 37 g/caput/d. Reliable data on consumption of “invisible” fats (cheese, other dairy products, and meat) are not available, and data on the regional distribution of consumed “visible” fats have limited reliability. Nonetheless, it is generally considered that the people of Marmara and Black Sea regions consume most fat, and the intake of those inhabiting the Mediterranean and southeast Anatolia regions is the least. Our data on the mean serum cholesterol concentrations by geographical region indicate a parallel trend.

Combined assessment both of work and leisure activity by questionnaire appeared to affect mean serum cholesterol values substantially, though it failed to affect the percentage of cigarette smokers and, in women, the body mass index; it marginally influenced the latter in men and the blood pressure in both sexes. Our age adjusted findings indicated that, independent of urban–rural differences and over a wide age range, as physical activity increased from grade 1 to 4, a decline in mean plasma cholesterol by 12\% in men and by 10\% in women occurred. A similar effect of leisure activity was observed by Hickey and coworkers\(^14\) in a study of coronary risk factors related to physical activity in 15 171 Irish men: the magnitude of reduction in serum cholesterol in men younger than 41 years (0.06 mmol/litre) was comparable to that in the present survey. However, in the MRFIT trial,\(^15\) in which 12 138 men were classified by questionnaire into three groups according to their leisure activity, no significant difference of mean serum total cholesterol was observed in the three activity levels.

Increasing net family income from the lowest to the highest of five brackets was associated with a successive and substantial rise by 13.2\% in average plasma cholesterol values in middle aged men and by a smaller rise in women. This appeared to be independent of urban life in men, but was related to urban life in women. Various factors, including climate, higher dietary saturated fatty acids, diminished physical activity, and psychosocial stress, are likely determinants for this observation. In 14 677 men aged 40–49 years included in the Oslo study, it was noted that serum cholesterol and triglyceride concentrations decreased with increasing socioeconomic status.\(^16\) The decrease in serum cholesterol was 4.7\% and in serum triglyceride, 19.5\%. This study, combined with our present survey, support the view that improved economic status from a low base is accompanied by a rise in blood cholesterol values up to a level when further improvement in socioeconomic status, resulting in more widespread awareness of health issues and adoption of low fat diets, leads to a decline in the lipid values.

The triglyceride profile in Turkish adults seems not to be as favourable as that of cholesterol. The mean serum triglyceride concentration of Turkish men up to the age of 55 years exceed by almost 0.1 mmol/litre that of their American\(^5\)\(^6\) and Italian\(^8\) counterparts, though the values are slightly lower than those in British\(^10\) and German\(^17\) men. Triglyceride levels in Turkish women exceed by about 0.1 mmol/litre those found in American women and in women in western European nations. The relatively high triglyceride values are probably related to the higher relative body weight of Turkish women; thus 47% of Turkish women aged 40–59 years had a body mass index in excess of 29 kg/m\(^2\), while 13–31\% of European women of the same age had a body mass index exceeding 30 kg/m\(^2\) in the WHO ERICA project.\(^18\)

As regards the relation between the levels of plasma triglyceride and cholesterol, it is reassuring to note in our sample population that for each rise of 5 mg/dl triglyceride an increase in cholesterol of 1 mg/dl occurred in both men and women, which confirmed the well known Friedewald formula\(^19\) concerning the very low density lipoprotein (VLDL) required for the transport of triglyceride in plasma and VLDL-cholesterol.

In conclusion, the study of a representative sample of Turkish adults, comprising women as well as men over a full age range of adult life, allowed us to characterise the cholesterol profile of the inhabitants of a developing Mediterranean country and to analyse the relationship with total cholesterol of variables such as sex, age, urban residence, physical activity, body mass index, family income, and plasma triglyceride level.

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