Dyslipidemia prevalence, awareness, treatment and control in Mexico: results of the Ensanut 2012

César Hernández-Alcaraz, MS,⁽¹⁾ Carlos A Aguilar-Salinas, MD, DSc,⁽²⁾ Kenny Mendoza-Herrera, MS,⁽¹⁾ Andrea Pedroza-Tobías, MS,^(1,3) Salvador Villalpando, PhD,⁽¹⁾ Teresa Shamah-Levy, DrPH,⁽⁴⁾ Juan Rivera-Dommarco, PhD,⁽⁵⁾ Mauricio Hernández-Ávila, PhD,⁽⁶⁾ Simón Barquera, MD, PhD.⁽¹⁾

Hernández-Alcaraz C, Aguilar-Salinas CA, Mendoza-Herrera K, Pedroza-Tobías A, Villalpando S, Shamah-Levy T, Rivera-Dommarco J, Hernández-Ávila M, Barquera S. Dyslipidemia prevalence, awareness, treatment and control in Mexico: results of the Ensanut 2012. Salud Publica Mex. 2020;62:137-146. https://doi.org/10.21149/10520

Abstract

Objective. To describe in a national sample 1) the prevalence, awareness, treatment and control of dyslipidemias 2) the prevalence of dyslipidemias through previous national surveys. Materials and methods. We analyzed data of the National Health and Nutrition Survey 2012, a representative cross-sectional study. Serum samples of 9 566 adults ≥20 years old with fasting ≥8 hours were analyzed for lipid fractions. Age-adjusted prevalences were calculated, by sociodemographic variables. Prevalence of awareness, treatment and control was estimated. A description of the dyslipidemia prevalence reported in previous surveys is reported. Results. Hypoalphalipoproteinemia and elevated LDL-C are the most prevalent dyslipidemias in Mexican adults. One in four adults had hypercholesterolemia at the moment of the interview without previous diagnosis. Awareness, treatment and control of dyslipidemia were 12.6, 3.7 and 3.1%, respectively. Conclusions. Dyslipidemias are the most prevalent risk factor for cardiovascular diseases in Mexico. Public policies to increase awareness, access to therapy and sustained control are urgently needed.

Keywords: dyslipidemias; prevention and control; nutrition surveys; Mexico

Hernández-Alcaraz C, Aguilar-Salinas CA, Mendoza-Herrera K, Pedroza-Tobías A, Villalpando S, Shamah-Levy T, Rivera-Dommarco J, Hernández-Ávila M, Barquera S. Prevalencia de dislipidemias, diagnóstico previo, tratamiento y control: resultados de la Ensanut 2012. Salud Publica Mex. 2020;62:137-146. https://doi.org/10.21149/10520

Resumen

Objetivo. Describir en una muestra nacional I) la prevalencia de dislipidemias, su diagnóstico previo, tratamiento y control, y 2) la prevalencia de dislipidemias en las encuestas previas. Material y métodos. Se analizaron datos de la Encuesta Nacional de Salud y Nutrición 2012 con representatividad nacional. Se análizaron fracciones de lípidos séricos de 9 566 adultos ≥20 años con ayuno ≥8 horas. Se estimaron las prevalencias de dislipidemias, diagnóstico previo (DP), tratamiento y control, ajustadas por edad. Se presenta un análisis comparativo de las prevalencias de dislipidemias reportadas previamente. **Resultados.** Las dislipidemias más prevalentes en adultos mexicanos fueron hipoalfalipoproteinemia y LDL-C elevado. Uno de cada cuatro adultos tenía hipercolesterolemia al momento de la entrevista, sin DP. El DP, tratamiento y control de dislipidemias fue de 12.6, 3.7 y 3.1%, respectivamente. Conclusiones. Las dislipidemias son el factor de riesgo para enfermedades cardiovasculares más prevalente en adultos mexicanos. Se necesitan políticas públicas para incrementar el diagnóstico, acceso a terapia y control.

Palabras clave: dislipidemias; prevención y control; encuestas nutricionales; México

- (I) Centro de Investigación en Salud y Nutrición, Instituto Nacional de Salud Pública. Cuernavaca, Morelos, Mexico.
- (2) Departamento de Endocrinología y Metabolismo, Instituto Nacional de Ciencias Médicas y Nutrición Salvador Zubirán Mexico City, Mexico.
- 3) Institute for Global Health Sciences, University of California. San Francisco, United States.
- (4) Centro de Investigación en Evaluación y Encuestas, Instituto Nacional de Salud Pública. Cuernavaca, Morelos, Mexico.
- (5) Instituto Nacional de Salud Pública. Cuernavaca, Morelos, Mexico.
- (6) Dirección de Prestaciones Económicas y Sociales, Instituto Mexicano del Seguro Social. Mexico City, Mexico.

Received on: April 30, 2019 • Accepted on: August 30, 2019
Corresponding author: Simón Barquera. Instituto Nacional de Salud Pública.
Av. Universidad 655, col. Santa María Ahuacatitlán. 62100 Cuernavaca, Morelos, Mexico.
email: sbarquera@insp.mx

ortality rates of cardiovascular diseases (CVD) Mave been increasing since 1990 in every region of the world, except for countries with high sociodemographic index.1 Particularly in Mexico, CVD has been the main cause of mortality from 1990 to 2012, and have moved from position four to two as a cause for disability-adjusted life years (DALYs) in the same time period. Several risk calculators estimate the risk of CVD based on well-known risk factors such as age, gender, smoking status, high blood pressure as well as high density lipoprotein cholesterol (HDL-C) and total cholesterol.²⁻⁵ Furthermore, dyslipidemias are modifiable risk factors for the development and control of CVD.6 By 1994, high levels of low density lipoproteins (LDL) cholesterol were considered the second most common risk factor for CVD mortality in Mexico and worldwide, only after arterial hypertension. Also, high cholesterol is one of the main risk factors for mortality in the Mexican population.6 Lipid disorders are among the main contributors to the development of atherosclerosis.⁷

Since 1988, several National surveys in Mexico have been able to document the prevalence of dyslip-idemias accounting for sociodemographic variables.⁸⁻¹¹ According to the 2006 National Health and Nutrition Survey (Ensanut, by its acronym in Spanish), 43.6% of adults older than 20 years had high levels of cholesterol, 31.5% of triglycerides and 60.5% low levels of HDL-C.¹¹ Prevalence of awareness has been reported previously by body mass index categories and diabetes.¹¹ However, prevalence of treated and controlled population has not been reported previously.

The present study aims to describe the prevalence of major dyslipidemias in Mexican adults by sociodemographic characteristics, as well as the status of awareness, treatment and control in our population study with Ensanut 2012 data. Furthermore, a comparative of the prevalence through several national surveys since 1994 is presented.

Materials and methods

Study design

This is an analysis of the data collected in the Ensanut 2012. This survey is representative at national, state, urban and rural level. The objective of the Ensanut is to quantify the frequency, distribution and trends of health and nutrition conditions and its determinants. The collection of information from Ensanut 2012 was carried out between October 2011 and May 2012, and information was obtained from 50 528 households, and 96 031 individuals of different age groups. Data on risk factors and sociodemographic characteristics were

collected through questionnaires. Also, anthropometric measurements, blood pressure, diet, and physical activity information were collected.

Regarding the information obtained in adults 20 years of age and older, previously trained and standardized personnel made anthropometric and clinical measurements that included: weight, height, waist and hip circumference, blood pressure measurements, as well as validated questionnaires to obtain information on risk factors, personal history of chronic diseases, and affiliation to health systems, among others. Detailed information on the methodology of the survey has been previously published.¹²

Definition and selection of population

Blood samples were obtained from a random subsample of about 33% calculated through probabilistic methods. The present analysis included 9 566 adults \geq 20 years old in Ensanut 2012, with fasting \geq 8 hours and with complete information on the concentration of triglycerides, total cholesterol and HDL-C. In addition, the concentrations of non-HDL cholesterol (nonHDL-C) (n=9 565) and LDL-C (n=9 006) were estimated.

Lipid assessment

A blood sample was collected in one tube with ethylenediaminetetra-acetic acid (EDTA) and another sample without any substance to obtain a tube of whole blood and another tube of serum. To obtain serum, samples were centrifuged in-situ. Subsequently, they were stored in cryotubes and transported in nitrogen tanks to the INSP (Mexico's *Instituto Nacional de Salud Pública*) laboratory, where they were stored at -70°C until analysis.

A minimum volume of 250 microliters of serum was used to determine triglycerides, total cholesterol and HDL-C concentrations. For the determination of triglycerides, the enzymatic method of glycerol phosphate oxidase was used, for cholesterol the enzymatic colorimetric method of quinonimine was used and HDL-C was determined by the enzymatic method via peroxidase that additionally uses accelerated and selective washing with a surfactant. For these determinations, the Abbott Architect CI8200 equipment was used. Non HDL-C concentrations were obtained from subtracting total cholesterol minus HDL-C concentrations. While concentrations of LDL-C were obtained with the Friedewald formula only in participants with triglycerides levels below 400 mg/dL.¹³

National Cholesterol Education Program recommendations were used as the criteria to define the presence of dyslipidemias. These thresholds were applied for the analyses of the previous Mexican health surveys; the definitions have not been modified in the current guidelines. Hypercholesterolemia, hypertriglyceridemia, and hypoalphalipoproteinemia were defined as total cholesterol levels equal or greater than 200 mg/dl, triglycerides levels equal or greater than 150 mg/dl and HDL-C levels less than 40 mg/dl, respectively. In addition, the cut points for elevated non HDL-C and LDL-C were 130 mg/dL and 100 mg/dL, respectively.

Dyslipidemia phenotypes were classified according to combinations of total cholesterol, triglyceride and HDL-C thresholds.

Statistical analysis

The mean values of the lipid traits (mg/dL), the prevalence of dyslipidemias and their phenotypes were obtained in the total sample, by sex, and age group, with their respective confidence intervals. Means were compared with Student's t-test. The prevalence of dyslipidemias by education level, socioeconomic status (SES), and affiliation to health service, area, and region in the country were estimated.

Prevalence of awareness, treated and controlled adults were calculated in the study population and by gender. Awareness was considered as the self-report of previous diagnosis of hypercholesterolemia. In adults that reported previous diagnosis, treatment for hypercholesterolemia was inquired. This variable included statins (pravastatin, simvastatin, atorvastatin, rosuvastatin or fluvastatin), recommendation to reduce the intake of fat or cholesterol in foods, and to increase physical activity in the last 12 months. Specifically, statin treatment variable was analyzed separately from dietary and physical activity recommendations. Uncontrolled adults were considered as a report of previous diagnosis of hypercholesterolemia or treated with statins, with hypercholesterolemia at the moment of the interview.

Analyses were adjusted for age and design of complex surveys. In order to maintain the representative nature of the sample at the national level and by strata, an expansion factor was included, using the SVY module for complex samples, with the statistical software Stata version 14.

Ethical considerations

During the data collection of the Ensanut 2012, the study subjects were informed of the objective and management of the information as well as the confidentiality of the data. Those who agreed to participate were asked to sign an informed consent form, in which they agreed to the interview and to the blood sample collection.

These forms and the draft were approved by the Ethics Committee of the National Institute of Public Health (*Instituto Nacional de Salud Pública*, INSP). The authors of this manuscript did not have access to the personal data of the study participants.

Results

The overall mean of total cholesterol, HDL-C, LDL-C, nonHDL-C and triglycerides was 180.7 mg/dL (95%CI 178.8-182.6), 39.8 mg/dL (95%CI 39.3-40.2), 106.6 mg/dL (95%CI 105.0-108.2), 141.3 (95%CI 139.5-143.1) and 179.2 (95%CI 174.2-184.3), respectively (table I). After adjusting for age, men had higher mean levels of triglycerides and nonHDL-C, and lower mean levels of HDL-C (p<0.05). Total cholesterol and LDL-C concentrations were significantly higher only in men in the age group of 20-29 years compared to women in the same age group. In contrast, women aged 60-79 had higher mean levels of total cholesterol and women aged 50-79 had higher LDL-C mean levels, compared to men in the same age groups (table I).

Table II shows the prevalence of five lipid abnormalities. The most prevalent dyslipidemia among Mexican adults was hypoalphalipoproteinemia (55.2%, 95% CI 53.2-57.3), followed by high LDL-C (56.1%, 95%CI 53.9-58.2), hypertriglyceridemia (47.4%, 95%CI 45.3-49.6) and hypercholesterolemia (30.6%, 95%CI 28.7-32.6). Regardless of the age group, the prevalence of hypoalphalipoproteinemia was significantly higher in men, with values between 55.5 and 74.0%. Likewise, the prevalence of hypertriglyceridemia was higher in men aged 20-59 years and in the study population (p<0.05). On the other hand, the prevalence of hypercholesterolemia was significantly lower in women aged 20-29 years old and higher in women aged 60 years or older (p<0.05). The highest prevalence of elevated LDL-C was observed in men and women between 60 and 79 years old, with a higher prevalence in women. A higher prevalence of elevated nonHDL-C was found in men and, specifically, men aged 20-39 years old. The highest prevalence of elevated nonHDL-C was observed in women between 70-79 years old (52.5%, 95%CI 42.8-62.1)

Mixed hyperlipidemia is associated to an atherogenic risk profile (table III). This phenotype was observed in one in every five Mexican adults (women 19.9%, 95%CI 17.8-22.0 vs. men 24.6%, 95%CI 22.1-27.1, p<0.05), with a higher prevalence in the 60-69 years old group. (31.0%, 95%CI 25.6-37.0). Similarly, it was observed that more than one third of the adults had hypertriglyceridemia and hypoalphalipoproteinemia (34.5%, 95%CI 32.5-36.6%), and the prevalence was significantly higher in men compared to women. The highest prevalence of this

Table I

LIPID CONCENTRATIONS OF MEXICAN ADULTS, STRATIFIED BY AGE AND SEX. MEXICO, ENSANUT 2012

Age group (years)*	Both sexes (n= 9 566) Mean (95%CI)	Men (n= 3 716) Mean (95%Cl)	Women (n= 5 850) Mean (95%Cl)	
Cholesterol (mg/dl)				
20-29	164.3 (160.6-168.0)	168.0 (161.3-174.6)	160.4 (156.9-163.8) [‡]	
30-39	178.1 (174.9-181.4)	180.9 (174.5-187.3)	176.1 (172.2-179.9)	
40-49	188.8 (183.9-193.8)	190.1 (181.9-198.4)	187.6 (182.0-193.3)	
50-59	191.4 (185.6-197.1)	188.1 (180.6-195.6)	194.3 (186.4-202.2)	
60-69	197.8 (192.8-202.8)	189.6 (182.2-196.9)	205.0 (198.6-211.5) [‡]	
70-79	195.2 (189.1-201.3)	181.3 (172.9-189.7)	203.9 (196.4-211.4)‡	
≥ 80	183.8 (172.1-195.6)	178.8 (160.1-197.4)	190.4 (182.4-198.4)	
Total ≥ 20	180.7 (178.8-182.6)	180.9 (177.9-184.0)	180.5 (178.1-182.9)	
HDL-cholesterol (mg/dl)				
20-29	40.1 (39.3-40.9)	38.3 (37.0-39.6)	42.0 (41.0-43.0) [‡]	
30-39	39.2 (38.3-40.1)	36.3 (35.1-37.5)	41.4 (40.1-42.6) [‡]	
40-49	38.7 (37.8-39.6)	36.5 (35.4-37.7)	40.7 (39.5-41.9) [‡]	
50-59	38.6 (37.5-39.7)	35.2 (33.9-36.5)	41.6 (40.3-42.9) [‡]	
60-69	41.6 (40.6-42.7)	40.1 (38.8-41.3)	43.0 (41.5-44.5) [‡]	
70-79	44.4 (41.7-47.1)	41.0 (38.4-43.5)	46.6 (42.9-50.3) [‡]	
≥ 80	40.7 (38.1-43.3)	37.9 (34.4-41.3)	44.3 (41.8-46.9) [‡]	
Total ≥ 20	39.8 (39.3-40.2)	37.3 (36.7-37.9)	41.9 (41.3-42.6) [‡]	
Triglycerides (mg/dl)				
20-29	149.5 (140.0-159.1)	167.3 (150.0-184.6)	130.5 (122.6-138.5) [‡]	
30-39	176.0 (167.4-184.6)	205.4 (189.5-221.3)	154.2 (144.6-163.8) [‡]	
40-49	207.3 (195.3-219.4)	220.2 (200.4-239.9)	195.6 (181.7-209.5)‡	
50-59	206.6 (190.9-222.3)	243.3 (214.3-272.3)	173.8 (162.3-185.2) [‡]	
60-69	181.8 (170.9-192.6)	178.8 (163.5-194.1)	184.4 (169.0-199.7)	
70-79	171.0 (157.2-184.9)	164.3 (142.6-185.9)	175.3 (156.6-193.9)	
≥ 80	187.3 (151.6-222.9)	203.4 (146.6-260.1)	166.3 (153.6-178.9)	
Total ≥ 20	179.2 (174.2-184.3)	199.0 (189.9-208.1)	161.7 (156.8-166.6) [‡]	
Non-HDL-cholesterol (mg/dl)	(n = 9 565)	(n = 3 715)	(n = 5 850)	
20-29	125.3 (122.2-128.5)	132.0 (126.6-137.4)	118.3 (115.0-121.7)‡	
30-39	138.9 (135.7-142.1)	144.6 (138.3-150.8)	134.7 (130.8-138.6) [‡]	
40-49	150.1 (145.2-155.1)	153.6 (145.6-161.6)	147.0 (140.8-153.1)	
50-59	152.8 (147.4-158.2)	152.9 (145.4-160.3)	152.7 (145.4-160.0)	
60-69	156.2 (151.3-161.1)	149.5 (142.6-156.4)	162.0 (155.4-168.6) [‡]	
70-79	150.8 (145.6-156.0)	140.3 (131.5-149.2)	157.3 (151.2-163.5) [‡]	
≥ 80	143.1 (133.1-153.1)	140.9 (124.4-157.3)	146.1 (138.6-153.5)	
Total ≥ 20	141.3 (139.5-143.1)	144.3 (141.5-147.1)	138.6 (136.2-141.0) [‡]	
LDL-cholesterol (mg/dl)	(n = 9 006)	(n = 3 390)	(n = 5 616)	
20-29	96.5 (94.2-98.8)	99.4 (95.7-103.2)	93.5 (90.7-96.4)‡	
30-39	104.6 (101.8-107.4)	104.6 (99.2-110.0)	104.6 (101.4-107.7)	
40-49	110.3 (105.9-114.7)	110.5 (103.4-117.6)	110.1 (104.7-115.5)	
50-59	113.7 (108.4-118.9)	107.1 (100.2-113.9)	119.0 (112.2-125.7)‡	
60-69	120.4 (116.6-124.1)	113.5 (108.5-118.4)	126.3 (121.3-131.4)‡	

(continues...)

(continuation)

70-79	117.7 (112.5-122.9)	107.6 (99.6-115.6)	124.2 (118.3-130.1) [‡]	
≥ 80	107.6 (91.3-123.9)	102.8 (75.3-130.2)	113.6 (107.2-120.0)	
Total ≥ 20	106.6 (105.0-108.2)	105.5 (102.9-108.1)	107.5 (105.4-109.5)	

Ensanut: Encuesta Nacional de Salud y Nutrición

LDL: low density lipoprotein HDL: high density lipoprotein

Table II Prevalence of Lipid disorders by age and sex groups in Mexican adults. Mexico, Ensanut 2012

Age group (years)*	Both sexes (n= 9 566)	Men (n= 3 716)	Women (n= 5 850)	
Cholesterol ≥ 200 mg/dl				
19.2 (16.1-22.6)		23.4 (18.2-29.4)	14.7 (11.7-18.3) [‡]	
30-39 27.9 (24.2-31.8)		30.5 (24.5-37.2)	25.9 (21.4-31.1)	
40-49	37.4 (33.4-41.6)	40.0 (34.0-46.2)	35.0 (29.9-40.5)	
50-59	36.8 (31.5-42.4)	36.9 (29.6-44.8)	36.7 (29.7-44.4)	
60-69	43.1 (36.8-49.6)	33.0 (24.4-42.9)	51.9 (43.1-60.5) [‡]	
70-79	45.4 (37.4-53.6)	28.6 (20.2-38.7)	55.9 (45.6-65.7) [‡]	
≥ 80	32.6 (22.2-45.0)	20.3 (10.5-35.7)	48.5 (37.5-59.7) [‡]	
Total ≥ 20	30.6 (28.7-32.6)	31.4 (28.5-34.2)	29.9 (27.5-32.4)	
HDL-cholesterol < 40 mg/dl				
20-29	52.3 (47.9-56.6)	60.6 (53.7-67.1)	43.3 (38.6-48.2) [‡]	
30-39	58.5 (54.3-62.6)	68.0 (60.6-74.6) 51.5 (4		
40-49	58.9 (54.6-63.1)	68.7 (63.0-73.8)	50.0 (44.2-55.8) [‡]	
50-59	58.7 (53.2-63.9)	74.0 (67.1-79.9)	44.9 (37.9-52.1) [‡]	
60-69	47.3 (41.2-53.4)	56.2 (48.5-63.7)	39.4 (31.4-48.0) [‡]	
70-79	41.9 (33.7-50.5)	55.5 (42.7-67.5)	33.4 (24.5-43.6) [‡]	
≥ 80	51.0 (36.9-65.0)	61.6 (39.3-79.9)	37.2 (28.1-47.4) [‡]	
Total ≥ 20	20 55.2 (53.2-57.3)		46.4 (43.7-49.1) [‡]	
Triglycerides ≥ 150 mg/dl				
20-29 32.3 (28.7-36.1)		38.7 (32.4-45.4) 25.5 (2		
30-39	45.4 (41.2-49.7)	55.9 (48.3-63.2)	37.7 (32.9-42.7) [‡]	
40-49 58.4 (53.9-62.8)		63.7 (57.0-69.8)	53.7 (47.9-59.3) [‡]	
50-59 60.1 (54.7-65.3)		68.5 (60.7-75.4)	52.6 (45.5-59.6) [‡]	
60-69	53.4 (46.1-60.6)	53.3 (42.1-64.2) 53.5 (44.2-6		
70-79	79 51.5 (43.2-59.6)		53.3 (42.3-64.0)	
≥ 80	51.5 (35.9-66.8)	58.3 (35.5-78.0)	42.7 (28.3-58.5)	
Total ≥ 20	47.4 (45.3-49.6)	54.1 (50.9-57.3)	41.5 (38.9-44.1) [‡]	
Non-HDL-cholesterol ≥ 130 mg/dl	(n = 9 565)	(n = 3 715) (n = 5 8		
20-29	40.4 (36.6-44.1)	46.3 (39.7-52.9) 34.1 (29.7-38		
30-39	57.8 (53.9-61.7)	63.5 (56.7-70.4)	53.5 (48.3-58.8) [‡]	

(continues...)

Lipid concentrations presented as means (95%CI). * Adults were in a \geq 8h fasting state. Estimates were adjusted for age and the complex survey design.

[‡] p<0.05 compared to men.

(continuation)

40-49	63.7 (59.6-67.9)	66.8 (60.3-73.2)	60.9 (55.1-66.8)	
50-59	65.6 (60.5-70.6)	67.1 (58.6-75.6)	64.2 (58.0-70.4)	
60-69	69.6 (63.3-75.9)	67.8 (60.9-74.6)	71.2 (61.2-81.2)	
70-79	68.9 (62.7-75.1)	57.1 (45.1-69.1)	76.3 (68.9-83.7) [‡]	
≥ 80	60.7 (44.7-76.7)	55.6 (29.5-82.0)	67.0 (58.5-75.6)	
Total ≥ 20	56.8 (54.9-58.7)	59.9 (56.8-63.0)	54.0 (51.4-56.6) [‡]	
LDL-cholesterol ≥ 100 mg/dl	(n = 9 006)	(n = 3 390)	(n = 5616)	
20-29	41.7 (37.7-45.8)	45.4 (38.6-52.3)	38.1 (33.4-42.9)	
30-39	56.8 (52.3-61.1)	58.2 (50.5-65.5)	55.8 (50.3-61.1)	
40-49	61.3 (56.7-65.6)	64.1 (57.0-70.6)	58.8 (52.5-64.8)	
50-59	61.7 (55.6-67.4)	55.6 (45.4-65.4)	66.5 (60.3-72.2)	
60-69	73.1 (68.2-77.5)	66.4 (59.1-73.0)	78.9 (72.5-84.1) [‡]	
70-79	68.0 (61.2-74.1) 58.1 (45.8-69.4)		74.3 (65.7-81.3) [‡]	
≥ 80	61.1 (43.7-76.0)	53.9 (28.1-77.7)	70.03 (60.5-78.1)	
Total ≥ 20	56.1 (53.9-58.2)	55.8 (52.1-59.5)	56.3 (53.6-58.9)	

Data are presented as percentages (95%CI).

Ensanut: Encuesta Nacional de Salud y Nutrición

Table III

Prevalence of Lipid disorders by Phenotypes in Mexican adults. Mexico, Ensanut 2012

	High TG/Low HDL-C	NormalTG/ Low HDL-C	High TG/High Chol	High TG/Normal Chol	Normal TG/High Chol	Normal TG/Normal Chol
Total (n= 9 566)	34.5 (32.5-36.6)	20.7 (18.9-22.5)	22.1 (20.5-23.7)	25.3 (23.3-27.3)	8.5 (7.3-9.7)	44.1 (42.0-46.1)
Men (n= 3 716)	43.1 (39.8-46.4)	22.0 (19.3-24.8)	24.6 (22.1-27.1)	29.5 (26.3-32.7)	6.8 (5.0-8.5)	39.1 (35.9-42.3)
Women (n= 5 850)	26.9 (24.5-29.3)*	19.5 (17.3-21.7)	19.9 (17.8-22.0)*	21.6 (19.5-23.8)*	10.0 (8.4-11.7)*	48.5 (45.9-51.1)*
Age, years‡						
20-29	24.1 (20.8-27.6)	28.2 (24.3-32.4)	13.7 (11.1-16.9)	18.6 (15.8-21.7)	5.5 (3.9-7.6)	62.2 (58.2-66.1)
30-39	35.2 (31.3-39.2)	23.4 (19.8-27.4)	19.2 (16.6-22.1)	26.3 (22.7-30.2)	8.7 (5.9-12.6)	45.9 (41.7-50.1)
40-49	43.3 (38.8-48.0)	15.6 (12.7-19.0)	29.1 (25.3-33.1)	29.4 (25.1-34.1)	8.3 (6.7-10.4)	33.3 (29.2-37.6)
50-59	43.7 (37.9-49.7)	14.9 (12.0-18.4)	28.5 (23.7-33.9)	31.6 (25.8-38.0)	8.2 (6.0-11.2)	31.6 (27.1-36.6)
60-69	32.3 (26.8-38.5)	14.9 (10.6-20.6)	31.0 (25.6-37.0)	22.5 (18.0-27.6)	12.1 (8.8-16.4)	34.5 (27.1-42.7)
70-79	29.9 (22.2-38.9)	12.0 (8.2-17.3)	27.0 (21.1-33.7)	24.5 (17.1-33.9)	18.4 (11.6-27.9)	30.1 (24.2-36.8)
≥ 80	36.8 (21.4-55.4)	14.2 (8.5-23.0)	19.7 (13.4-28.0)	31.8 (16.5-52.4)	12.9 (6.1-25.0)	35.6 (24.5-48.6)

Data are presented as percentages (95%CI).

^{*} Adults were in a \geq 8h fasting state. Estimates were adjusted for age and the complex survey design.

[‡] p<0.05 compared to men.

^{*} p<0.05 compared to men.

[‡] Adults were in a ≥8h fasting state. Estimates were adjusted for age and the complex survey design. Ensanut: Encuesta Nacional de Salud y Nutrición

phenotype was observed in the 50-59 years age group (43.7%, 95%CI 37.9-49.7). Less than half of the Mexican adults had normal levels of triglycerides and total cholesterol (44.1%, 95%CI 42.0-46.1).

Higher prevalence of hypercholesterolemia, elevated LDL-C and nonHDL-C were observed in adults with high school level compared to adults with studies less than elementary. Hypertriglyceridemia and hypoal-phalipoproteinemia lowest prevalences were observed in high SES compared to low SES and medium SES categories, respectively. The prevalence of hypercholesterolemia, elevated LDL-C and nonHDL-C was lower in Mexico City and the metropolitan area, compared to the northern region of the country. On the contrary, higher prevalence was observed in the same region compared to the northern region of the country. Hypertriglyceridemia prevalence was higher in southern and center regions compared to northern region.¹⁵

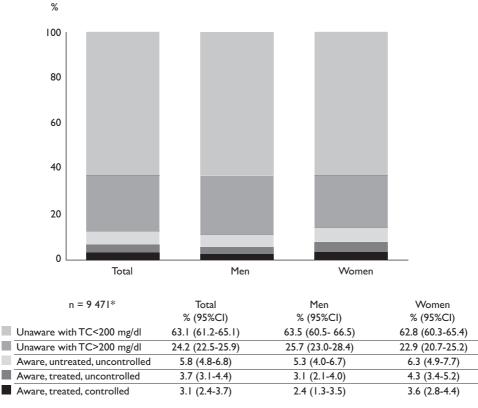
Awareness, treatment and control prevalence are presented in figure 1. Almost one in four Mexican adults had hypercholesterolemia at the moment of the survey and did not reported previous diagnosis (24.2%, 95%CI

22.5-25.9). From those aware of having hypercholesterolemia, less than half did not have treatment. From the treated adults, less than half were controlled at the moment of the survey. There were no statistically significant differences in awareness, treatment and control prevalence between men and women. The prevalences of awareness, treatment and control of dyslipidemia were 12.6, 3.7 and 3.1%, respectively.

Based on four national surveys (ENN 1994, ENSA 2000, Ensanut 2006 and 2012), no trend was observed in the prevalence of hypercholesterolemia, hypertriglyceridemia and hypoalphalipoproteinemia in men and women.¹⁵

Discussion

Previous national surveys have reported dyslipidemias as one of the most prevalent cardiovascular disease risk factor in Mexican adults. Our main findings were that hypoalphalipoproteinemia and elevated LDL-C were the most prevalent dyslipidemias among Mexican adults. There were differences in the prevalence of all



^{* 95} adults did not have information of previous diagnosis Ensanut MC: Encuesta Nacional de Salud y Nutrición de Medio Camino

FIGURE 1. AGE-ADJUSTED PREVALENCE OF AWARENESS, TREATMENT AND CONTROL OF HYPERCHOLESTEROLEMIA IN MEXICAN ADULTS. ENSANUT 2012

dyslipidemias by gender, age group, and sociodemographic characteristics. Also, almost a quarter of the population was identified with hypercholesterolemia in the survey without previous diagnosis. A low proportion of awareness is reported, with a very low treatment and control rate. No observable trend along surveys was observed.

One in two Mexican adults has a LDL-C concentration that is considered above the optimal and it is well established that such levels confer an increased risk for cardiovascular disease. According to the Global Burden of Disease project, LDL-C is ranked as the second metabolic risk factor associated to CVD mortality and DALYs. Solid evidence from diverse sources associate elevated LDL-C with a higher risk of heart disease. Along with this, clinical trials have demonstrated that therapy focused on reducing LDL level, reduces the risk of coronary heart disease (CHD). The most recent version of current guidelines considers LDL-C as the primary target of therapy, along with a risk assessment that includes other risk factors. Solid exists a see second

However, no consensus has been made regarding the use of LDL-C in primary care as a biomarker of atherogenic lipoproteins. LDL-C measurements and calculations have limitations associated, mainly, with triglyceride levels and fasting samples. The recent recommendation for lipid assessment considered non-HDL-C over LDL-C as the best biomarker of atherogenic lipoproteins. The benefits of non-HDL-C are that it only requires total cholesterol and HDL-C for its calculation, and it does not rely on triglyceride levels. NonHDL-C can be used in settings of non-fasting and it does not incur additional expense to the patient or health care system. The prevalence of high nonHDL-C and high LDL-C in our population study are quite similar (56.8% vs. 56.1%), and combined prevalence of high nonHDL-C/LDL-C accounts for almost half of the adults with altered nonHDL-C or LDL-C (47.5%, data not presented in tables). The recommendation for clinical laboratories is to proactively report calculated nonHDL-C with LDL-C on all lipid profiles, and physicians' conceptual understanding of this combination needs to be addressed. 17

Additionally, more than half of the Mexican adults had hypoalphalipoproteinemia. Although in some observational studies, HDL-C levels were considered as an independent risk factor for CHD, the combination of low HDL-C and high triglyceride levels is considered an atherogenic profile. Up to a third of the study population had this profile, with a higher proportion of men compared to women. Currently, no drug therapy is focused to increase HDL-C levels; treatment of hypoalphalipoproteinemia is based on the correction of hypertriglyceridemia, excess body weight, physical

inactivity and cigarette smoking. Also, genetic factors and diets with carbohydrate intake >60% of the total energy intake are major determinants of the low concentrations of HDL-cholesterol in Mexicans. ¹⁶ According to national reference data for 2012, obesity and overweight are present in 73% of the adults, 12% of the population are current smokers, 30% of the adults are physically inactive and have diets with excessive intake of added sugars, saturated fats and insufficient intakes of fiber and other micronutrients. ^{12,18} In this scenario, the approach to reduce the prevalence of hypoalphalipoproteinemia must consider a therapy that includes lifestyle changes to reduce the risk of CHD.

Differences among age groups and by gender were observed, although no patterns were identified. Education level, SES and region of the country were environmental factors associated with differences in dyslipidemias. Hypercholesterolemia was mainly found in adults with high school level or living in northern Mexico. Hypertriglyceridemia was less prevalent in high SES levels and more prevalent in the southern region of the country. Hypoalphalipoproteinemia prevalence was lower in high SES and in the Mexico City area. The highest prevalence of elevated LDL-C was found in adults with high school or more, or living in the northern region.

Almost 25% of the Mexican adults had hypercholesterolemia at the moment of the interview and did not report a previous diagnosis of this condition. The prevalence of awareness is 12.6% and, although more than half are treated, less than half are controlled. US and China data from national surveys report higher and lower prevalences of hypercholesterolemia, respectively. But the prevalences of awareness in Mexico is a third of the awareness reported in China and 15% of that of the US.¹⁹ Despite treatment, almost half of the adults have cholesterol levels above 200 mg/dl. The positive effects of statins may not be enough in countries like Mexico, where quality of nutrition is low, and there is still a high prevalence of physical inactivity and sedentarism. 18,20,21 Along with the low adherence to statin treatment, low adherence to recommendations of diet and physical activity may contribute to our results.²²⁻²⁴ Previous surveys have not reported this information and acknowledging the situation of undetected, untreated and uncontrolled population may impact the current guidelines and screening processes in primary care medical services. Public policies to increase awareness, access to therapy and sustained control are urgently needed.

Some methodological aspects need to be considered in the interpretation of these results. The design and sampling methods of the study allow having representative results at national and regional level. The selected cut point for high LDL-C considers <100mg/dL as the optimal level for LDL-C.¹⁴ Future survey analyses may address the comparison of different LDL-C cut points along with different cardiovascular risk profiles to identify the proportion of the population that need to be treated under these considerations. Direct comparisons between surveys are limited by the differences in the analytical methods in each survey. Awareness and treatment variables had self-reported information. Although it is not the most precise method to assess adherence therapy, similar instruments to collect information have been used in surveys in other countries.

Our data confirm that hypertriglyceridemia and elevated LDL-C are highly prevalent in Mexican adults and some environmental characteristics are associated with higher prevalence of dyslipidemias. Screening processes are deficient and preventive strategies should focus on reducing the prevalence of unaware cases of dyslipidemias.

Acknowledgments

Authors express their gratitude to Luz E. Guillen Pineda, Carmen Moreno Villatoro, Guadalupe Lopez and Adriana Cruz Lopez for the technical assistance in the analyses of the samples.

Funding

This research is a secondary analysis of the data obtained from the National Health and Nutrition Survey, implemented with funding of the Ministry of Health in Mexico. Sanofi provided additional funding to analyze serums in a subsample, under registration number DIREGL07856 (MHA, SB). Partially, the Fogarty International Center of the National Institutes of Health (http://www.fic.nih.gov/Pages/Default.aspx) supported the work with funding under Award Number R03TW009061 (SB, AP); and the Robert Wood Johnson Foundation with a grant under Award Number 74155 (CHA, KM). The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

Declaration of conflict of interests. The authors declare that they have no conflict of interests.

References

I. Institute for Health Metrics and Evaluation. GBD Compare [Internet]. Seattle: IHME, 2015 [cited November 21, 2018]. Available from: http://vizhub.healthdata.org/gbd-compare

- 2. Ho KKL, Pinsky JL, Kannel WB, Levy D.The epidemiology of heart failure:The Framingham Study. J Am Coll Cardiol. 1993;22(4):A6-13. https://doi.org/10.1016/0735-1097(93)90455-A
- 3. D'Agostino RB, Vasan RS, Pencina MJ, Wolf PA, Cobain M, Massaro JM, et al. General cardiovascular risk profile for use in primary care. Circulation. 2008;117(6):743-53. Available from: https://www.ahajournals.org/doi/10.1161/CIRCULATIONAHA.107.699579
- 4. Cook NR, Paynter NP, Eaton CB, Manson JE, Martin LW, Robinson JG, et al. Comparison of the framingham and reynolds risk scores for global cardiovascular risk prediction in the multiethnic women's health initiative. Circulation. 2012;125(14):1748-56.
- Sahle BW, Owen AJ, Chin KL, Reid CM. Risk prediction models for incident heart failure: a systematic review of methodology and model performance. J Card Fail. 2017;23(9):680-7. https://doi.org/10.1016/j. cardfail.2017.03.005
- 6. Lozano R, Gómez-Dantés H, Pelcastre B, Ruelas M, Montañez J, Campuzano J, et al. Carga de la enfermedad en México, 1990-2010. Nuevos resultados y desafíos [Internet]. Cuernavaca: Secretaría de Salud, Instituto Nacional de Salud Pública, 2014. Available from: https://www.insp.mx/images/stories/Produccion/pdf/150122_cargaEnfermedad.pdf
- 7. Aikawa M, Libby P.The vulnerable atherosclerotic plaque: pathogenesis and therapeutic approach. Cardiovasc Pathol. 2004;13(3):125-38. https://doi.org/10.1016/S1054-8807(04)00004-3
- 8. Posadas-Romero C, Tapia-Conyer R, Lerman-Garber I, Zamora-González J, Cardoso-Saldaña G, Salvatierra-Izaba B, et al. Cholesterol levels and prevalence of hypercholesterolemia in a Mexican adult population. Atherosclerosis. 1995;118(2):275-84.
- 9. Aguilar-Salinas CA, Olaiz G, Valles V, Torres JM, Gómez Pérez FJ, Rull JA, et al. High prevalence of low HDL cholesterol concentrations and mixed hyperlipidemia in a Mexican nationwide survey. J Lipid Res [Internet]. 2001;42(8):1298-307. Available from: http://www.ncbi.nlm.nih.gov/pubmed/J 1483632
- 10. Barquera S, Flores M, Olaiz-Fernández G, Monterrubio E, Villalpando S, González C, et *al.* Dyslipidemias and obesity in Mexico. Salud Publica Mex. 2007;49(suppl 3):S338-47.
- I I. Aguilar-Salinas CA, Gómez-Pérez FJ, Rull J, Villalpando S, Barquera S, Rojas R. Prevalence of dyslipidemias in the Mexican National Health and Nutrition Survey 2006. Salud Publica Mex [internet]. 2010;52 (suppl I):S44-53. Available from: http://www.ncbi.nlm.nih.gov/pubmed/20585729 I 2. Gutiérrez JP, Rivera-Dommarco JÁ, Shamah-Levy T, Villalpando-Hernández S, Franco A, Cuevas-Nasu L, et al. Encuesta Nacional de Salud y Nutrición 2012. Resultados nacionales. Cuernavaca: Instituto Nacional de Salud Pública, 2013.
- Friedewald WT, Levy RI, Fredrickson DS. Estimation of the concentration of low-density lipoprotein cholesterol in plasma, without use of the preparative ultracentrifuge. Clin Chem [Internet]. 1972;18(6):499-502.
 Available from: http://www.ncbi.nlm.nih.gov/pubmed/4337382
- 14. Stone NJ, Robinson JG, Lichtenstein AH, Bairey-Merz CN, Blum CB, Eckel RH, et al. 2013 ACC/AHA Guideline on the treatment of blood cholesterol to reduce atherosclerotic cardiovascular risk in adults. J Am Coll Cardiol [Internet]. 2014;63(25):2889-934. Available from: https://linkinghub.elsevier.com/retrieve/pii/S0735109713060282
- 15. Hernandez-Alcaraz C, Aguilar Salinas CA, Mendoza-Herrera K, Pedroza-Tobías A, Villalpando S, Shamah-Levy T, et al. Supplementary material. Dyslipidemia prevalence and trends. DRAFT VERS. Harvard Dataverse; 2019. p. 1-2. https://doi.org/10.7910/DVN/D3BA8V
- 16. Goodman DS, Hulley SB, Clrak LT, Davis CE, Fuster V, LaRosa JC, et al. Report of the National Cholesterol Education Program Expert Panel on Detection, Evaluation, and Treatment of High Blood Cholesterol in Adults. Arch Intern Med [Internet]. 1988;148(1):36-69. Available from: http://archinte.jamanetwork.com/article.aspx?doi=10.1001/archinte.1988.00380010040006

17. Langlois MR, Chapman MJ, Cobbaert C, Mora S, Remaley AT, Ros E, et al. Quantifying atherogenic lipoproteins: Current and future challenges in the era of personalized medicine and very low concentrations of Idl cholesterol. a consensus statement from EAS and EFLM. Clin Chem. 2018;64(7):1006-33.

18. Rivera JA, Pedraza LS, Aburto TC, Batis C, Sánchez-Pimienta TG, González de Cosío T, et al. Overview of the dietary intakes of the mexican population: results from the National Health and Nutrition Survey 2012. J Nutr [Internet]. 2016;146(9):1851S-1855S. Available from: https://academic.oup.com/jn/article/146/9/1851S-1855S/4584848

mic.oup.com/jn/article/146/9/1851S-1855S/4584848

19. Lu Y, Wang P, Zhou T, Lu J, Spatz ES, Nasir K, et al. Comparison of prevalence, awareness, treatment, and control of cardiovascular risk factors in China and the United States. J Am Heart Assoc [Internet]. 2018;7(3):1-16. Available from: http://www.ncbi.nlm.nih.gov/pubmed/29374046

20. Medina C, Janssen I, Campos I, Barquera S. Physical inactivity prevalence and trends among Mexican adults: results from the National Health and Nutrition Survey (ENSANUT) 2006 and 2012. BMC Public Health

[Internet]. 2013;13(1):1063. Available from: http://bmcpublichealth.biomed-central.com/articles/10.1186/1471-2458-13-1063

21. Medina C, Tolentino-Mayo L, López-Ridaura R, Barquera S. Evidence of increasing sedentarism in Mexico City during the last decade: Sitting time prevalence, trends, and associations with obesity and diabetes. PLoS One. 2017;12(12):e0188518. https://doi.org/10.1371/journal.pone.0188518 22. Berra K. Lipid-lowering therapy today. J Cardiovasc Nurs [Internet]. 2008;23(5):414-21. Available from: http://content.wkhealth.com/linkback/openurl?sid=WKPTLP:landingpage&an=00005082-200809000-00007 23. Schwiesow SJ, Nappi JM, Ragucci KR. Assessment of compliance with lipid guidelines in an Academic Medical Center. Ann Pharmacother [Internet]. 2006;40(1):27-31. Available from: http://journals.sagepub.com/doi/10.1345/aph.1G303

24. Zafrir B, Cohen S. Primary prevention in high-risk dyslipidemic patients without an established cardiovascular disease: Undertreatment and rationale for lipid-lowering therapy. Eur J Intern Med [Internet]. 2006;17(7):495-9. https://doi.org/10.1016/j.ejim.2006.03.005