

Research Article



INTERNATIONAL RESEARCH JOURNAL OF PHARMACY

www.irjonline.com

ISSN 2230-8407 [LINKING]

EVALUATION OF LIPID PROFILE ABNORMALITIES IN OBESITY AND CHRONIC HEART DISEASE (CHD) AMONG INDIAN PARTICIPANTS

Dr. Anand Kumar

Assistant Professor, Department of Biochemistry, Shree Narayan Medical Institute and Hospital, Saharsa, Bihar

Email: drkumarrims83@gmail.com

How to cite: Kumar A. Evaluation of lipid profile abnormalities in obesity and chronic heart disease (CHD) among Indian participants.

International Research Journal of Pharmacy, 2023, 14:03:16-20.

DOI: 10.56802/2230-8407.1303303

ABSTRACT

Background: Over the last thirty years, obesity has become a worldwide epidemic. Additionally, the leading cause of death worldwide, including in India, is coronary heart disease. However, there is a dearth of information in the literature currently available on lipid profiles in obese and CHD patients.

Aim: The purpose of this study was to evaluate the lipid profile abnormalities associated with obesity and chronic heart disease (CHD) in Indian participants.

Methods: Using commercially available kits, blood samples were obtained from 250 CHD patients, 125 obese patients, and 125 healthy controls in order to measure VLDL, LDL-C, HDL-C, triglycerides, and total cholesterol.

The results were compared in two groups for dyslipidemia in controls and study subjects. Logistic regression was done to assess the association of lipid parameters with CHD and obesity.

Results: Dyslipidemia was seen in CHD and obese subjects compared to healthy controls with a higher proportion of CHD in first-degree blood relatives from the CHD group with 36%. Similar results were seen in the obese group with a positive family history in 63.9% of subjects. In some cases, 51% of subjects showed combined lipid abnormalities with deranged HDL-C, TG, LDL-C, and TC values. Higher than cut-off values were seen for TC (> 200 mg/dl) and LDL-C (> 100 mg/dl) in 49.5% and 52% subjects respectively. Also, 64% of subjects had HDL < 50 mg/dl (moderate CHD risk) and 84% had TG > 150 mg/dl (> upper normal range).

Conclusion: The present study concludes that Indian subjects are hyperlipidemic with low HDL and deranged lipid profiles are seen in subjects with comorbidities. The present study also points to deranged lipid profiles in metabolic disorders that can predispose to complications.

Keywords: coronary heart disease, CHD, Dyslipidemia, obesity, lipid profile

INTRODUCTION

Obesity is directly linked to cardiovascular concerns and children with a parental history of cardiovascular diseases have high weight during childhood and are obese when they are adults. CHD (coronary heart disease) is considered to be the leading cause of death in developing nations in coming years. It has been estimated that more than 80% of the subjects with CHD worldwide contribute to the burden from low-income countries. However, the knowledge concerning vital risk factors is attained from the developing nations.¹

A large geographic variation is being seen concerning the prevalence of CHD risk factors and it is vital to utilize the existing local data during the discussion of the relationship between various risk factors and the diseases. It remains unclear as to what extent, the risk factors for CHD known are applicable in the Indian population.

This can be attributed to the higher prevalence of risk factors for CHD as dyslipidemia, obesity, and hypertension in Indian subjects.² As CHD can result from obesity, the involved biochemical pathways in obesity development have also been assessed to have a role in CHD development.³

Levels of lipids in the blood are the modifiable risk factors for CHD and atherosclerosis. As lipids are hydrophobic in nature, phospholipids, triglycerides, cholesterol esters, and cholesterol are transported in the form of lipoproteins to each other. HDL (high-density lipoproteins), LDL (low-density lipoproteins), and chylomicrons are the major classes of lipoproteins that are named by their assembly site and the type of apoprotein and lipids they have.

The remnants of VLDL with further hydrolysis of the triglyceride contents produce IDL (intermediate-density lipoproteins) and LDL. LDL has an apo B100 apoprotein component which is a major carrier of cholesterol in the peripheral circulation. Elevated levels of plasma in these non-HDL lipoproteins are major risk factors for CHD.⁴

It has been noted that many lipoprotein/lipid abnormalities have a high prevalence in subjects with cardiovascular diseases and obesity which collectively is known as dyslipidemia. Majorly, these dyslipidemias are hyperlipidemias where the majority of the lipids are shifted to the upper limit range or more than the range.

Transportation availability and recent lifestyle modernization have resulted in a high prevalence of metabolic disorders in Indian subjects as seen on the global level. This not only interferes with social interactions, work performance, and daily activities but also lays a high burden on healthcare.⁵ Considering the vital role of lipid traits in developing nutritional disorders, the present study aimed to assess the patterns of lipid profiles in CHD and obese subjects in the Indian context.

MATERIALS AND METHODS

The present case-control clinical study aimed to assess the patterns of lipid profiles in CHD and obese subjects in the Indian context. The study subjects were from the Department of Cardiology of the Institute. The study included 250 subjects with CHD, 125 obese subjects, and 125 healthy controls that were gender and age-matched. Verbal and written informed consent were taken from all the subjects before study participation.

The study included subjects with CHD with non-fatal MI (myocardial infarction) as diagnosed by a consultant cardiologist depending on clinical history, Troponine T/I, angiography, cardiac echo, and ECG. Only the cases with coronary artery disease were recruited in the study who had recent diagnoses and were not on the antihypertensive drugs or lipid-lowering drugs.

The inclusion criteria for the study were subjects that were obese following BMI cut-off values for Asian populations where $>26\text{kg/m}^2$ was taken as obese and $\text{BMI} >23\text{ kg/m}^2$.^{6,7} Indirect indicator of lower and upper body fat was taken as waist to hip ratio.⁸ The control subjects were healthy subjects with BMI in the range of 18 to 23 and with no history of CHD. The exclusion criteria for the study were subjects with obesity and CHD, infections, malignancies, pregnant females, and underweight subjects.

The presence of various comorbidities as hypertension or diabetes was also identified. Diabetes was considered with FBG (fasting blood glucose) levels of 6.7 mmol/L or higher or a 2-hour PP (postprandial) blood glucose levels of 11.1 mmol/L or higher. For hypertensive and normotensive subject classification, it was followed by reports from the seventh Joint National Committee for Hypertension (JNC-V).⁹ Blood pressure of each participant was done by a healthcare personnel expert in the field after 5 minutes of sitting.

For each participant, 5ml of intravenous blood was collected after overnight fasting under strict aseptic and sterile conditions in a test tube having a gel clot activator. The blood was collected from the median cubital vein with a tourniquet placed on the arm and with the fist squeezed. The collected blood was subjected to centrifugation at 10,000 rpm for 10 minutes to settle all the elements formed and for separation of the serum. The separated serum was then transferred to the autoclaved Eppendorf and was refrigerated till further assessment.

Plasma lipoproteins and lipid variables such as triglycerides (TG) and total cholesterol (TC) were assessed with the commercially available kits. Optical density assessments were done at 546 nm for triglycerides and total cholesterol. HDL-C (high-density lipoprotein) and LDL-C (low-density lipoprotein C) were assessed following Mohsen Ibrahim et al¹⁰ in 2012. Assessment of HDL needed precipitation of LDL and VLDL from the serum leaving HDL-C in the serum which is further assessed following the principle similar to total cholesterol.

The data gathered were analyzed statistically using the SPSS software version 21.0 (IBM Corp., Armonk, NY, USA) and the one-way ANOVA (analysis of variance) along with logistic regression analysis. The data were expressed as mean and standard

deviation for normally distributed and non-normally distributed continuous variables and frequency and percentage for categorical variables. Statistical significance was kept at a p-value of <0.05.

RESULTS

The mean age of the study subjects was significantly higher in controls and CHD subjects with $p=0.002$. The gender between controls, obese, and CHD subjects was statistically comparable with $p=0.47$. The family history was significantly higher in obese and CHD subjects compared to controls with $p<0.0001$. Similar results were seen for smoking, hypertension, and diabetes which was significantly lower in controls compared to obese and CHD subjects with $p<0.0001$. The waist-hip ratio was higher in obese subjects compared to CHD and controls with $p<0.0001$. BMI was significantly higher in obese subjects compared to CHD and controls with $p<0.0001$ as shown in Table 1.

On assessing the blood lipid levels in different groups of study subjects, HDL-C and TC levels were significantly higher in subjects with CHD and obese subjects with $p<0.001$. LDL-C levels were significantly higher in obese subjects followed by subjects with CHD and were least in controls with $p<0.001$. HDL-C levels were highest in controls followed by CHD and obese subjects which was statistically significant with $p<0.001$. Total triglycerides were highest in obese subjects followed by CHD and controls with $p<0.001$. Similar results were seen for total cholesterol levels that were highest in obese subjects followed by CHD and controls depicting statistical significance with $p<0.001$ as depicted in Table 2.

DISCUSSION

The mean age of the study subjects was significantly higher in controls and CHD subjects with $p=0.002$. The gender between controls, obese, and CHD subjects was statistically comparable with $p=0.47$. The family history was significantly higher in obese and CHD subjects compared to controls with $p<0.0001$. Similar results were seen for smoking, hypertension, and diabetes which was significantly lower in controls compared to obese and CHD subjects with $p<0.0001$.

The waist-hip ratio was higher in obese subjects compared to CHD and controls with $p<0.0001$. BMI was significantly higher in obese subjects compared to CHD and controls with $p<0.0001$. These data were similar to Lawler PR et al¹¹ in 2015 and Sutter I et al¹² in 2014 where authors assessed subjects with demographic data comparable to the present study in their respective studies.

Concerning the blood lipid levels in different groups of study subjects, HDL-C and TC levels were significantly higher in subjects with CHD and obese subjects with $p<0.001$. LDL-C levels were significantly higher in obese subjects followed by subjects with CHD and were least in controls with $p<0.001$. HDL-C levels were highest in controls followed by CHD and obese subjects which was statistically significant with $p<0.001$. Total triglycerides were highest in obese subjects followed by CHD and controls with $p<0.001$. Similar results were seen for total cholesterol levels that were highest in obese subjects followed by CHD and controls depicting statistical significance with $p<0.001$. these results were consistent with the studies of Genest J et al¹³ in 2003 and Gordon L et al¹⁴ in 2010 where blood lipid levels in CHD and obesity subjects reported by the authors were similar to the present study.

The study results showed that on gender-wise assessment of subjects in obese and CHD groups with deranged lipid profiles, LDL-C values of >99 were seen in 55.2% ($n=69$) males and 47.2% ($n=59$) females from the obese group and in 24.4% ($n=61$) males and 19.2% ($n=48$) females from CHD group respectively. HDL-C <40 was seen in 52% ($n=65$) males and 45.6% ($n=57$) females from the obese group and 31.6% ($n=79$) males and 22% ($n=55$) females from the CHD group. HDL-C <49 was seen in 48.8% ($n=61$) males and 44.8% ($n=56$) females from the obese group and 22% ($n=55$) males and 8.4% ($n=42$) females from the CHD group respectively. TG >149 was seen in 88.8% ($n=111$) males and 74.4% ($n=93$) females from the obese group and in 38% ($n=950$) males and 28.4% ($n=71$) females from the CHD group. TC >199 was seen in 45.6% ($n=57$) males and 39.2% ($n=49$) females from the obese and 24.8% ($n=62$) males and 17.2% ($n=43$) females from the CHD group. These results were in agreement with the findings of Gupta M et al¹⁵ in 2006 and Aziz KU et al¹⁶ in 2012 where a lipid profile similar to the present study was reported by the authors in their respective studies.

It was seen that on gender-wise assessment of lipid profile in the obese and CHD group, LDL-C was 103.5 ± 29.4 and 98.5 ± 27.6 mg/dl in males and females from the obese group which was statistically non-significant with $p=0.06$ and was 104.6 ± 39.7 and 104.4 ± 35.3 mg/dl in CHD group showing statistical non-significance with $p=0.85$. HDL-C levels were 55.3 ± 17.1 and 54.9 ± 17.8 mg/dl in males and females from the obese group and 44.4 ± 11.4 and 46.5 ± 12.3 mg/dl in the CHD group which was statistically non-significant in both groups with $p=0.81$ and 0.08 respectively. Similar results were seen for TG levels in obese and CHD groups with $p=0.06$ and 0.67 respectively. TC also showed statistical non-significance in males and females from

obese and CHD groups with $p=0.08$ and 0.86 respectively. These findings were in line with de Lemos J et al¹⁷ in 2010 and Sacks FM et al¹⁸ in 2003 where authors reported the results similar to the present study.

CONCLUSIONS

Considering its limitations, the present study showed that Indian subjects are hyperlipidemic with low HDL, and deranged lipid profiles are seen in subjects with comorbidities. The present study also points to deranged lipid profiles in metabolic disorders that can predispose to complications. Further longitudinal studies with larger sample sizes and longer monitoring periods are needed.

REFERENCES

1. Klatzkin RR, Gaffney S, Cyrus K, Bigus E, Brownley KA. Binge eating disorder and obesity: preliminary evidence for distinct cardiovascular and psychological phenotypes. *Physiol Behav.* 2015;145:20–7.
2. Awan ZA, Gul AM, Sahibzada WA, Hafizullah M. Prevalence of coronary artery disease in rural areas of Peshawar. *J Postgraduate Med Inst (Peshawar-Pakistan)* 2011;19:14–22.
3. Yusuf S, Hawken S, Ôunpuu S, Dans T, Avezum A, Lanas F, et al. Effect of potentially modifiable risk factors associated with myocardial infarction in 52 countries (the INTERHEART study): case-control study. *Lancet.* 2004;364:937–52.
4. David A, Jumeau M, Murad M, Okorodudu D, Kumar S, Somers V, et al. Diagnostic performance of body mass index to identify obesity as defined by body adiposity in children and adolescents: a systematic review and meta-analysis. *Pediatric Obesity.* 2014;10:234–44.
5. Flegal KM, Graubard BI, Williamson DF, Gail MH. Cause-specific excess deaths associated with underweight, overweight, and obesity. *Obstet Gynecol Surv.* 2008;63:157–9.
6. Mascie-Taylor CN, Goto R. Human variation and body mass index: a review of the universality of BMI cut-offs, gender and urban-rural differences, and secular changes. *J Physiol Anthropol.* 2007;26:109–12.
7. Yusuf S, Hawken S, Ôunpuu S, Bautista L, Franzosi MG, Commerford P, et al. Obesity and the risk of myocardial infarction in 27 000 participants from 52 countries: a case-control study. *Lancet.* 2005;366:1640–9.
8. Sandhu HS, Koley S, Sandhu KS. A study of the correlation between lipid profile and waist-to-hip ratios in patients with diabetes mellitus. *Anthropologist.* 2008;10:215–8.
9. Chobanian AV, Bakris GL, Black HR, Cushman WC, Green LA, Izzo JL, et al. Seventh report of the joint national committee on prevention, detection, evaluation, and treatment of high blood pressure. *Hypertension.* 2003;42:1206–52.
10. Mohsen Ibrahim M, Ibrahim A, Shaheen K, Nour MA. Lipid profile in Egyptian patients with coronary artery disease. *Egyptian Heart J.* 2012;65:79–85.
11. Lawler PR, Akinkuolie A, Glynn R, Ridker P, Mora S. Atherogenic lipoprotein particle subclasses and residual cardiovascular risk: an analysis of the Jupiter trial. *J Am Coll Cardiol.* 2015;65:362–6.
12. Sutter I, Riwayto M, Rohrer L, Othman A, Hornemann T, Landmesser U, et al. Low concentrations of sphingosine-1-phosphates and plasmalogens in HDL are associated with coronary artery disease and reduced anti-apoptotic activity of HDL. *Atherosclerosis.* 2014;235:46.
13. Genest J, Frohlich J, Fodor G, McPherson R. Recommendations for the management of dyslipidemia and the prevention of cardiovascular disease: summary of the 2003 update. *Can Med Assoc J.* 2003;169:921–4.
14. Gordon L, Ragoobirsingh D, St Errol Y, Choo-Kang E, McGrowder D, Martorell E. Lipid profile of type 2 diabetic and hypertensive patients in the Jamaican population. *J Lab Physicians.* 2010;2:25.
15. Gupta M, Brister S, Verma S. Is South Asian ethnicity an independent cardiovascular risk factor? *Can J Cardiol.* 2006;22:193–7.
16. Aziz KU, Faruqui A, Patel N, Jaffery H. Prevalence and awareness of cardiovascular disease including lifestyles in a lower middle-class urban community in an Asian country. *Pakistan Heart J.* 2012;41:11–20.

17. de Lemos J, Braunwald E, Blazing M, Murphy S, Downs J, Gotto A, et al. Efficacy and safety of more intensive lowering of LDL cholesterol: a meta-analysis of data from 170,000 participants in 26 randomized trials. *Lancet*. 2010;376:1670–81.
18. Sacks FM, Campos H. Clinical review 163: cardiovascular endocrinology: low-density lipoprotein size and cardiovascular disease: a reappraisal. *J Clin Endocrinol Metab*. 2003;88:4525–32.

TABLES

Variables	Controls	Obese	CHD	p-value obese vs controls	p-value CHD vs controls
Number (n)	125	125	250	-	-
Mean age (years)	55.85±10.35	39.4±15.17	59.02±12.5	-	0.002
Gender % (n)					
Males	55.2 (69)	53.6 (67)	58 (145)	0.129	0.47
Females	41.6 (52)	45.6 (57)	44.4 (111)		
Family history	1.6 (2)	64 (80)	36 (90)	<0.0001	<0.0001
Smoking	10.4 (13)	12 (15)	29.2 (73)	<0.0001	<0.0001
Hypertension	16 (20)	25.6 (32)	60 (150)	<0.0001	<0.0001
Diabetes	13.6 (17)	32 (40)	64.4 (161)	<0.0001	<0.0001
Waist hip ratio	0.85±0.08	1.00±0.07	0.85±0.04	<0.0001	1
BMI (kg/m2)	21.2±8.9	36.7±6.2	22.2±6.5	<0.0001	<0.0001

Table 1: Baseline demographic and disease data of study participants

Variables	Controls	Obese	CHD	p-value CHD vs controls	p-value obese vs controls	p-value (ANOVA)
HDL-C/TC	2.5±1.13	5.2±0.95	4.4±0.89	1.4x10 ⁻³²	3.6x10 ⁻⁹⁸	<0.001
LDL-C (mg/dl)	78.3±15.40	115.9±21.97	104.8±37.92	1.4x10 ⁻³²	1.1x10 ⁻²²	<0.001
HDL-C (mg/dl)	80.4±12.73	43.3±4.21	45.3±11.61	1.04x10 ⁻⁶⁹	1.4x10 ⁻¹⁷⁷	<0.001
TG (mg/dl)	187.5±41.79	265.3±50.36	214.3±74.62	2.07x10 ⁻³²	1.6x10 ⁻⁵	<0.001
TC (mg/dl)	158.8±33.12	214.6±35.04	208.4±54.09	2.7x10 ⁻²¹	2.2x10 ⁻³⁴	<0.001

Table 2: Blood lipid levels in control and study subjects