

COMUNICACIÓN BREVE

Early atherosclerotic lesions and post-mortem serum cholesterol level in a group of Colombian children

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Introduction: Atherosclerosis is an asymptomatic chronic disease, which begins at early age and is difficult to detect during this stage. Prospective studies suggest a causal relationship between total serum cholesterol levels during childhood and early adolescence and the development of cardiovascular diseases in adulthood.

Objective: The aim of this study was to evaluate the prevalence of early atherosclerotic lesions in the aorta artery of children and its relationship with post-mortem serum cholesterol levels.

Material and methods: Post-mortem samples of blood and aorta were taken from 43 subjects aged less than 17 years old. Histopathological analysis (intimal thickening and inflammatory infiltrates) of the thoracic aorta and measurement of total serum cholesterol were performed.

Results: The analysis showed thickening of the intima and lymphocyte infiltrates in 93% of children, and macrophage infiltrates in 79.1% of cases. A relationship between the highest terciles of total serum cholesterol levels and the presence of multiple lesions in the aorta wall was found ($P < 0.05$).

Conclusion: This group of children had a high prevalence of early inflammatory atherosclerotic lesions positively related with serum cholesterol levels. To our knowledge this study represents the first report of a relationship between post-mortem total serum cholesterol levels and pathological findings of macrophages and lymphocytes infiltrates in the aorta wall.

Key words: Atherosclerosis, child, adolescent, cholesterol, post-mortem changes, aorta.

doi: <http://dx.doi.org/10.7705/biomedica.v33i3.1443>

Lesiones ateroscleróticas tempranas y nivel de colesterol post mórtem en un grupo de niños colombianos

Introducción. La aterosclerosis es una enfermedad crónica asintomática que se inicia a edad temprana y es de difícil detección en esta etapa. Los estudios prospectivos sugieren una relación causal entre el nivel de colesterol sérico total en la niñez y la adolescencia, y el desarrollo de enfermedades cardiovasculares en la adultez.

Objetivo. El objetivo de este estudio fue evaluar, post mórtem, la prevalencia de lesiones ateroscleróticas tempranas en la arteria aorta de niños y su relación con los niveles de colesterol.

Materiales y métodos. Se tomaron muestras de sangre y de aorta de 43 sujetos con edades menores de 17 años. Se hizo el análisis histopatológico (engrosamiento de la íntima e infiltrado inflamatorio) de la aorta torácica y la medición de colesterol en suero.

Resultados. El análisis mostró engrosamiento de la íntima e infiltrado linfocitario en 93 % de los niños, e infiltrado de macrófagos en 79,1 %. Se encontró relación entre los terciles superiores de colesterol sérico y la presencia de múltiples lesiones en la pared de la aorta ($p < 0,05$).

Conclusión. Este grupo de niños presentó una alta prevalencia de lesiones ateroscleróticas inflamatorias tempranas relacionada positivamente con niveles de colesterol sérico. Para nuestro conocimiento, este estudio representa el primer reporte de la relación entre el colesterol sérico

Contribución de los autores:

José Guillermo Ortega-Ávila, Isabella Echeverri-Jiménez, Lorena Jiménez-Bastidas, Mildrey Mosquera: estandarizaron las técnicas.

Milton Fabián Suárez-Ortegón y Alberto Pradilla: analizaron los datos.

Alberto Pradilla, Luis Eduardo Bravo, Cecilia Aguilar-de Plata: formularon el proyecto.

Todos los autores participaron en la escritura del artículo.

y los hallazgos histopatológicos de infiltrado de macrófagos y linfocitos en la pared de la aorta, post mórtem.

Palabras clave: aterosclerosis, niño, adolescente, colesterol, cambios post mórtem, aorta.

doi: <http://dx.doi.org/10.7705/biomedica.v33i3.1443>

Atherosclerosis is an asymptomatic chronic process, which begins at early age and progresses into adulthood (1,2). Post-mortem studies in children and fetuses have reported the presence of atherosclerotic lesions (3). These findings have been confirmed by the presence of fatty streaks in aortas of children below three years old and in coronary arteries of children between 10 and 14 years of age (1-3). Histological observations have shown that the lesions developed from six months of age and may progress until forming fibrous plaques in coronary arteries (4).

Atherosclerosis is difficult to detect during childhood because the lesions, in case of being present, are not easily observable, and conventional risk factors (e.g., sedentary lifestyle, obesity, smoking, alcoholism), which have a harmful effect on vascular endothelium, are still not evident. However, prospective studies suggest a causal relationship of serum cholesterol levels during childhood and early adolescence with coronary disease during adulthood (5).

Total serum cholesterol is considered a good post-mortem biomarker because of its stability and sensitivity as a cardiovascular risk factor (6). Some autopsy studies have showed that early atherosclerosis is related with lipid fractions like high serum total cholesterol levels, LDL cholesterol, and low HDL cholesterol (7-8). However, reports of these associations in Latin American populations are unknown. Likewise, there are no evaluation reports of the relationship between post-mortem serum cholesterol and lesions of cellular infiltration (macrophages and lymphocytes). Therefore, the aim of this study was to evaluate the post-mortem prevalence of early atherosclerotic lesions (intimal thickening and presence of lymphocytes and macrophages) in the thoracic aorta and its relationship with serum cholesterol in a group of Colombian children.

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Recibido: 12/09/12; aceptado:21/05/13

Materials and methods

Subjects

This study analyzed 43 samples of thoracic aorta and blood of autopsies of children done at the *Instituto Nacional de Medicina Legal y Ciencias Forenses* in the city of Cali. We included autopsies of children less than 17 years old, with accidental or violent cause of death and without severe damage to the cardiac cavity. Parents or relatives gave a written consent. The research protocol was approved by the Human Ethics Committee in the *Facultad de Salud* at *Universidad del Valle* in Cali, Colombia.

Histopathology

The thoracic aortas were removed from a 2 cm proximal point to the ligamentum arteriosum to a 2 cm distal point to the iliac bifurcation. The samples were incubated in 10% formalin for 24 hours. Then, they were dehydrated and embedded in paraffin. Tissue sections (6 µm) were cut and stained with hematoxylin-eosin (9) and were examined by a certified pathologist. Microscopic evaluation considered intimal thickening, as well as the presence of inflammatory cells (macrophages, lymphocytes) using a visual analogue scale (10). The presence of macrophages and lymphocytes was defined as having a moderate and abundant amount of these cells, while a low amount or nothing was considered as absence.

Post-mortem serum cholesterol

The blood samples were obtained by cardiac puncture with a 10 ml syringe and transferred into a tube without any anticoagulant. The sample was centrifuged for 10 minutes at 4,000 rpm; the serum was extracted and stored at -20°C. The serum cholesterol was measured by the Abell-Kendall method recommended by the Center for Disease Control (CDC, Atlanta GA), modified with previous standardization (11). For processing, 0.5 ml of serum were mixed with 5 ml of KOH at 33% ethanol and incubated at 50°C for 1 hour. Afterwards, 10 ml of n-hexane were added and centrifuged for 5 minutes at 2,500 rpm. The n-hexane layer was desiccated at 62°C and the precipitate was suspended again in 6 ml of the Liebermann-Burchard stable reactive (12). The absorbance was read at 620 nm (UV-1700

SHIMADZU). As a standard, we used cholesterol (Sigma, HPLC grade), and for quality control, we used clinical biochemistry control serums (Biosystems Spain).

Statistical analysis

Data analysis was performed using SPSS software (version 11.0). Post-mortem serum cholesterol was divided in terciles. The proportions difference was estimated by χ^2 and Fisher's exact tests, and the relationship between age and post-mortem serum cholesterol by Spearman correlation. A p value less than 0.05 was considered significant, a p value between 0.05 and 0.1 was considered as marginal significance.

Results

Characteristics of the subjects are shown in table 1. Almost the totality of the group was male and 84.4% were adolescents. The mean total serum cholesterol of the subjects (133 mg/dl) was in the normality range. Prevalence of early atherosclerotic lesions, i.e. intimal thickening and presence of abundant lymphocytes and macrophages was found in almost all of the subjects. In 75% of the

subjects, intimal thickening, lymphocytes and macrophages were found concomitantly.

We found a higher prevalence of atherosclerotic lesions in the second and third tercile of cholesterol (figure 1). By specific lesion, the presence of macrophages was higher in terciles 3 (156-257 mg/dl) and 2 (109-156 mg/dl) in relation to

Table 1. Population characteristics

	n (%)
Sex	
Male	39 (90.7)
Female	4 (9.3)
Age group	
≤5 years old	1 (2.3)
>5 and <12 years old	4 (9.3)
≥12 years old	38 (84.4)
Early atherosclerotic lesions	
Intimal thickening	40 (93.0)
Presence of lymphocytes	40 (93.0)
Presence of macrophages	34 (79.1)
All three early atherosclerotic lesions	32 (74.4)
Serum total cholesterol (mg/dL)¹	134±45

¹Mean ± standard deviation

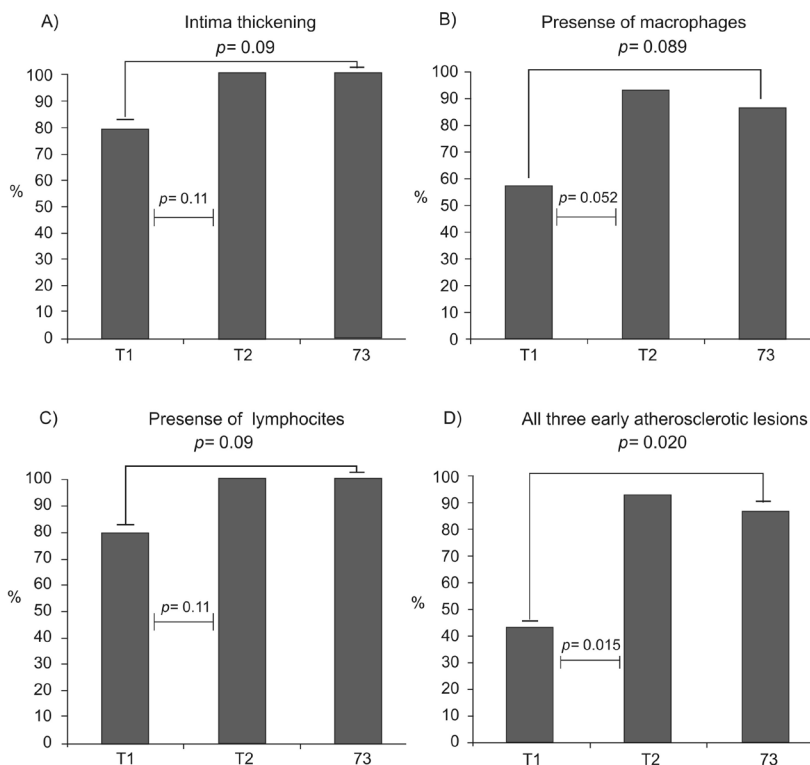


Figure 1. Prevalence of early atherosclerotic lesions by terciles (T) of serum cholesterol. (A) Intimal thickening, B) presence of macrophages, C) presence of lymphocytes, and D) all three early atherosclerotic lesions. Differences between T1 (53 -108 mg/dl), T2 (109- 156 mg/dl) and T3(156- 257mg/dl) are shown.

cholesterol tercile 1 (53-108 mg/dl) with marginal significances ($p=0.089$ and $p=0.052$, respectively) (figure 1B). The infiltrate lymphocytes and intimal thickening were significantly higher in tercile 3 in relation to tercile 1 ($p=0.089$ (marginal) and $p=0.09$, respectively) (figures 1A and 1C). While the prevalence of all atherosclerotic lesions evaluated was also significantly higher in the second and third terciles of serum cholesterol vs first tercile (figure 1D), differences in the prevalence of lesions between the second and the third terciles of serum cholesterol were not observed.

Age showed a weak correlation with postmortem serum cholesterol levels with marginal significance ($\rho=0.278$, $p=0.071$), but it was not associated with any early atherosclerotic lesions (data not shown).

Discussion

In this study, we found an unexpected high prevalence of early atherosclerotic post-mortem lesions in a group of Colombian children, and this prevalence was associated to a higher serum cholesterol level. Post-mortem studies have revealed that the atherosclerosis process starts from childhood (2-6), where the earliest vascular finding is the fatty streak, as a result of the accumulation of cholesterol inside the macrophages and other lipids in the intimal-medial of the artery. This lesion could progress to a chronic inflammatory state because the continuous infiltration and accumulation of macrophages and lymphocytes, accompanied by the proliferation of smooth muscle cells, lead to intimal-medial thickening.

We found correlations with marginal significance between elevated levels of post-mortem serum cholesterol and the degree of infiltration in the aorta vascular wall by macrophages and lymphocytes, which are compatible with a chronic inflammatory state, characteristic of the atherosclerotic process. This finding agrees with other studies that reported a relationship between serum cholesterol, endothelial dysfunction, and intimal-medial thickness (13,14).

Our study found a mean post-mortem serum cholesterol of 133 mg/dl, lower than that reported in a population of Colombian children of the same age range (168 mg/dl) (15). A possible explanation for this difference is the fact that most of the subjects we studied (85%) were in the age of puberty development, a stage that has been associated to diminished serum cholesterol (16). Likewise, most of the subjects (76.7%) had levels of cholesterol <170 mg/dl, the cut-off point recommended by

the United States National Cholesterol Education Program (NCEP) for pediatric populations (17,18).

Regarding lesions in the aorta, we observed an association between high tercile serum cholesterol with the presence of all histopathological alterations evaluated. The Bogalusa Heart study reported an association between total cholesterol and the presence of fatty streaks in the aorta and the coronary arteries (19). These findings show a possible early relationship between cholesterol levels and the chronic inflammatory process in the artery wall, even when the cholesterol levels are in the normal range.

To our knowledge this study represents the first report of a relationship between post-mortem serum cholesterol and pathologic findings of macrophages and lymphocytes infiltration in aorta wall, and the first evaluation precedent of an association between post-mortem serum cholesterol and early atherosclerotic lesions in Latin American children. Although our sample size was small, this limitation can be lessened by the high prevalence of incipient atherosclerotic lesions found in the children group.

The findings of this study suggest the need to implement preventive measures in pediatric populations to reduce the morbimortality related with inflammatory atherosclerotic process and also the importance to measure cholesterol in childhood as a possible predictor of early atherosclerosis and provide rationale for recent recommendations for pediatric lipid screening (20,21) and, furthermore, the modification of unhealthy life styles in early ages in order to reduce the serum cholesterol levels(22).

Acknowledgments

This work was supported by Colciencias (grant number 1106-04-16540).

Conflict of interest

The authors declare no conflict of interest

References

1. McMahan CA, Gidding SS, Malcom GT, Tracy RE, Strong JP, McGill HC Jr.; Pathobiological Determinants of Atherosclerosis in Youth Research Group. Pathobiological determinants of atherosclerosis in youth risk scores are associated with early and advanced atherosclerosis. *Pediatrics*. 2006;118:1447-55. <http://dx.doi.org/10.1016/j.atherosclerosis.2008.07.002>
2. Pathobiological Determinants of Atherosclerosis in Youth (PDAY) Research Group. Natural history of aortic and coronary atherosclerotic lesions in youth. Findings from the

- PDAY study. *Arterioscler Thromb Vasc Biol.* 1993;13:1291-8. <http://dx.doi.org/10.1161/01.ATV.13.9.1291>
3. **Charakida M, Tousoulis D, Stefanadis C.** Early atherosclerosis in childhood: Diagnostic approaches and therapeutic strategies. *Int J Cardiol.* 2006;109:152-9. <http://dx.doi.org/10.1016/j.ijcard.2005.06.010>
 4. **Newman WP, Wattigney W, Berenson GS.** Autopsy studies in United States children and adolescents relationship of risk factors to atherosclerotic lesions. *Ann N Y Acad Sci.* 1991;623:16-25. <http://dx.doi.org/10.1111/j.1749-6632.1991.tb43715.x>
 5. **Berenson GS, Srinivasan S.** Cholesterol as a risk factor for early atherosclerosis: The Bogalusa Heart Study. *Prog Pediatr Cardiol.* 2003;17:113-22. [http://dx.doi.org/10.1016/S1058-9813\(03\)00048-1](http://dx.doi.org/10.1016/S1058-9813(03)00048-1)
 6. **Uemura K, Shintani-Ishida K, Saka K, Nakajima M, Ikegaya H, Kikuchi Y, et al.** Biochemical blood markers and sampling sites in forensic autopsy. *J Forensic Leg Med.* 2008;15:312-7. <http://dx.doi.org/10.1016/j.jflm.2007.12.003>
 7. **Srinivasan SR, Myers L, Berenson GS.** Distribution and correlates of non-high-density lipoprotein cholesterol in children: The Bogalusa Heart Study. *Pediatrics.* 2002;110:29-34. <http://dx.doi.org/10.1542/peds.110.3.e29>
 8. **Ford ES, Li C, Zhao G, Mokdad AH.** Concentrations of low-density lipoprotein cholesterol and total cholesterol among children and adolescents in the United States. *Circulation.* 2009;119:1108-15. <http://dx.doi.org/10.1161/CIRCULATIONAHA.108.816769>
 9. **Fernández-Britto JE, Falcon L, Campos R, Contreras D, Guski H.** Cardiac sudden death: A morphometric study applying an atherometric system. *Path Res Pract.* 1987;182:489-90.
 10. **Holman RL, McGill HC, Strong JP, Geer JC.** Technics for studying the atherosclerotic lesions. *Lab Invest.* 1958;7:42-7.
 11. **Abell LL, Levy BB, Brodie BB, Kendall FE.** A simplified method for the estimation of total cholesterol in serum and demonstration of its specificity. *J Biol Chem.* 1952;195:357-66.
 12. **Kim E, Goldberg M.** Serum cholesterol assay using a stable Liebermann-Burchard reagent. *Clin Chem.* 1969;15:1171-9.
 13. **Sorensen KE, Celermajer DS, Georgakopoulos D, Hatcher G, Betteridge DJ, Deanfield JE.** Impairment of endothelium-dependent dilation is an early event in children with familial hypercholesterolemia and is related to the lipoprotein level. *J Clin Invest.* 1994;93:50-5. <http://dx.doi.org/10.1172/JCI116983>
 14. **Järvisalo MJ, Rönnemaa T, Volanen I, Kaitosaari T, Kallio K, Hartiala JJ, et al.** Brachial artery dilatation responses in healthy children and adolescents. *Am J Physiol Heart Circ Physiol.* 2002;282:H87-92.
 15. **Gracia B, de Plata C, Méndez F, Cruz M, Leiva J, Conde L, et al.** Evaluation of early manifestations of chronic non transmitted diseases risk in school population in Cali, Colombia. *Arch Latinoam Nutr.* 2005;55:267-78.
 16. **Berenson GS, Srinivasan SR, Cresanta JL, Foster TA, Webber LS.** Dynamic changes of serum lipoproteins in children during adolescence and pubertal maturation. *Am J Epidemiol.* 1981;113:157-70.
 17. **NCEP Expert Panel on Blood Cholesterol Levels in Children and Adolescents.** National Cholesterol Education Program: Report of the expert panel on blood cholesterol levels in children and adolescents. *Pediatrics.* 1992;89:525-84.
 18. **American Academy of Pediatrics.** Committee on Nutrition. Cholesterol in childhood. *Pediatrics.* 1998;101:141-7.
 19. **Freedman DS, Dietz WH, Srinivasan SR, Berenson GS.** The relation of overweight to cardiovascular risk factors among children and adolescents: The Bogalusa Heart Study. *Pediatrics.* 1999;103:1175-82.
 20. **Magnussen CG, Venn A, Thomson R, Juonala M, Srinivasan SR, Viikari JS, et al.** The association of pediatric low- and high-density lipoprotein cholesterol dyslipidemia classifications and change in dyslipidemia status with carotid intima-media thickness in adulthood evidence from the cardiovascular risk in Young Finns study, the Bogalusa Heart study, and the CDAH (Childhood Determinants of Adult Health) study. *J Am Coll Cardiol.* 2009;53:860-9. <http://dx.doi.org/10.1016/j.jacc.2008.09.061>
 21. **Juonala M, Viikari JS, Rönnemaa T, Marniemi J, Jula A, Loo BM, et al.** Associations of dyslipidemias from childhood to adulthood with carotid intima-media thickness, elasticity, and brachial flow-mediated dilatation in adulthood: The Cardiovascular Risk in Young Finns Study. *Arterioscler Thromb Vasc Biol.* 2008;28:1012-7. <http://dx.doi.org/10.1161/ATVBAHA.108.163329>
 22. **Ferranti SD.** Childhood cholesterol disorders: The iceberg base or nondisease? *Med Clin North Am.* 2012;96:141-54. <http://dx.doi.org/10.1016/j.mcna.2012.01.011>

