Carotid Artery Stiffness as an Early Marker of Vascular Lesions in Children and Adolescents With Cardiovascular Risk Factors

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Introduction and objectives. The availability of a noninvasive marker of vascular lesions that enables their detection in the preclinical phase would be of great benefit for cardiovascular disease prevention. The aim of this study was to investigate the usefulness of a range of indices of arterial wall stiffness in the common carotid artery, as derived using high-resolution Doppler ultrasonography, for identifying vascular damage in children with risk factors.

Methods. The study involved 99 children (age, 8-16 years) divided into two groups: 65 had cardiovascular risk factors (45 obesity, 20 dyslipidemia) and 34 were controls. Family histories of cardiovascular risk factors and anthropometric and biochemical measurements were recorded. Functional parameters of arterial stiffness (ie, arterial compliance, elastic modulus, beta stiffness index, pulse wave velocity, and augmentation index) and the intima-media thickness were also measured.

Results. Some functional vascular parameters were higher in obese children than controls: there were significant differences in beta stiffness index (P<0.02), elastic modulus (P<0.001), and pulse wave velocity (P<0.01). There was a significant difference in arterial compliance between dyslipidemics and controls (P<0.05). No significant difference in intima-media thickness was found between the groups. In obese children, there were positive correlations between body mass index, systolic pressure and triglyceride levels and vascular parameters (ie, elastic modulus and pulse wave velocity); in dyslipidemic children, triglyceride levels same were correlated with these same parameters.

Conclusions. Ultrasonographic measurement of arterial stiffness is a sensitive technique that can detect vascular damage in children with cardiovascular risk factors earlier than intima-media thickness measurement.


Medida de la rigidez de la arteria carótida como marcador precoz de lesión vascular en niños y adolescentes con factores de riesgo cardiovascular

Introducción y objetivos. Para la prevención de la enfermedad cardiovascular resulta de gran interés disponer de un marcador incruento de lesión vascular que permita su detección en fase preclínica. Nuestro objetivo fue analizar diversos índices de rigidez arterial de la carótida común mediante ultrasonografía Doppler de alta resolución para definir su utilidad como detector de daño vascular en niños con factores de riesgo.

Métodos. Se estudió a 99 niños (edades, 8-16 años) divididos en dos grupos: 65 niños con factores de riesgo cardiovascular (45 obesos, 20 dislipémicos) y 34 controles. Se recogieron antecedentes familiares de riesgo cardiovascular y variables antropométricas y bioquímicas. Se midieron parámetros funcionales de rigidez arterial (compliance arterial, módulo elástico, índice beta de rigidez, velocidad de la onda de pulso e índice de aumento) y el grosor íntima-media.

Resultados. Los niños obesos presentaron parámetros vasculares funcionales elevados respecto a los controles, que fueron significativos para el índice beta (p < 0,02), el módulo elástico (p < 0,001) y la velocidad de onda del pulso (p < 0,01). En los dislipémicos constatamos diferencias significativas en la compliance arterial respecto a los controles (p < 0,05). No hubo diferencias significati-
INTRODUCCIÓN

A pesar de los esfuerzos preventivos, la aterosclerosis continúa siendo responsable de la morbimortalidad y mortalidad en el mundo occidental, alcanzando proporciones epidémicas en sociedades tecnológicamente avanzadas. Este proceso bien documentado comienza muy temprano en la infancia con un aumento progresivo de las arterias, a pesar de que no se manifieste clínicamente hasta la edad adulta.

La medida ultrasonográfica de la rigidez arterial es un procedimiento sensible que puede resultar más preciso cuando el grosor íntima-media para detectar daño vascular en niños con factores de riesgo cardiovascular.


ABBREVIATIONS

BMI: body mass index
CVRF: cardiovascular risk factor
DBP: diastolic blood pressure
Dd: diastolic (or minimum) arterial diameter
ds: systolic (or maximum) arterial diameter
SBP: systolic blood pressure
METHODS

Patients

A case-control study was conducted (June 2008-December 2009) in schoolchildren and adolescents aged 8-16 years with CVRF (obesity and dyslipidemia) referred by a primary care pediatrician to the Pediatric Gastroenterology and Nutrition Unit of Hospital Clínico Universitario de Valencia. Patients were recruited from 10 health care centers (Number Area 5 of the Department of Health in Valencia) whose referral hospital is the Hospital Clínico Universitario de Valencia, Spain. Selection was at random and patients presenting dysmorphic syndromes and/or with endocrine disorders were excluded. A control group of children with no CVRF, matched by age and sex was also included in the study (referred from the same health centers for studies of functional murmurs or preoperative study prior to minor surgery). Children were included in the study after the informed consent of a guardian was given. The study protocol was approved by the hospital’s Committee on Ethical Practice.

Clinical Evaluation

For each patient the following data were recorded: a) family history of cardiovascular risk (overweight, obesity, metabolic syndrome, type 2 diabetes mellitus, primary arterial hypertension, inherited, and acquired dyslipidemia, cardiovascular incidents in men <55 years and women <65 years old); b) personal history and growth curve; c) nutritional assessment: anthropometric measurements (weight and height) were collected using a standardized technique; to compare children of different ages and sex, the z-score was calculated for each measurement with reference to WHO standards 2007; to define obesity, the body mass index (BMI) Z-score was calculated and subjects were considered obese when it was ≥2SD, equivalent to a BMI of 30 kg/m² at the age of 19; d) arterial pressure, always measured by the same researcher using the same measuring apparatus (Dinamap® oscillometric method); high values were checked with the auscultatory method and mercury sphygmomanometer; measurements were taken from the right arm with the child sitting comfortably; the sleeve was placed at the level of the heart, adjusting the size to fit the diameter of the arm; 3 measurements were taken for each child and the average of the 3 were calculated; the values for each child were compared with reference tables according to sex, age, and height (The Fourth Report on the Diagnosis, Evaluation, and Treatment of High Blood Pressure in Children and Adolescents). 30

Biochemical Analyses of Plasma

Total cholesterol was determined along with fractions and triglycerides. Dyslipidemia was diagnosed when total cholesterol and LDL-C values were above the 95th percentile for age and sex, with/without HDL-C <35 mg/dL, and triglycerides were above the 95th percentile. 31

High-Resolution Ultrasound Measurement of Arterial Stiffness and Intima-Media Thickness

An Aloka alfa-10 ultrasound apparatus was employed using EchoTracking®. Measurements were taken of the right common carotid artery, 1 cm below the carotid bulb, with the patient in supine position, head turned 45 degrees to the left and an ambient temperature of 22-25ºC, as described previously. 32 The common carotid artery was chosen because it is a central artery, branching directly from the aorta. The patient remained in the supine position for at least 5 minutes before starting the examination. The transducer was positioned so that the carotid artery could be observed longitudinally and was tilted to maximize the echoes from the interface between the medium and adventitia. When this line of division was captured clearly, the 2 system tracers were placed on the diametrically opposed interface points. Once in place, these tracers were oscillated with the arterial wall, thus recording the temporal distance between the two. Patient monitoring by ECG continued throughout the examination so that the system could detect the onset of the pulse wave. Using this technique we recorded (Figure 1): a) graphic chart of the pulse wave; b) maximum or systolic arterial diameter (Ds) corresponding to the peak of each pulse and representing the point at which the blood vessel is subjected to highest pressure (systolic blood pressure [SBP]); c) minimum or diastolic arterial diameter (Dd) corresponding to the lowest point, when the pressure to which the blood vessel is subjected is minimum (diastolic blood pressure [DBP]). All measurements were performed by the same researcher.

After the 3 measurements of blood pressure had been taken, the average of both SBP and DBP were calculated and recorded in the EchoTacking program, assuming the arterial pressure at the level of the brachial artery was the same as that of the carotid. Once all data were recorded, the program implemented a series of equations to calculate the 5 parameters related to arterial wall elasticity:
ship of BMI, arterial pressure, LDL-C, HDL-C, and triglycerides with arterial stiffness parameters and IMT was studied using the Pearson correlation coefficient. For all studies, \( P < .05 \) was considered statistically significant.

RESULTS

The sample included a total of 99 children aged between 8 and 16 years (62 boys and 37 girls, 63\% and 37\%, respectively), distributed into 2 groups: a) study group of 65 children with CVRF of which 45 were obese and 20 had familial dyslipidemia; and b) controls: 34 children without disease who did not have any CVRFs, matched to the study group by age and sex.

Table 1 summarizes anthropometric results, blood pressure and lipid values of the 3 groups, and the differences between them. In obese children, 5 patients had LDL-C levels over 130 mg/dL, and 3 had HDL-C levels below 35 mg/dL, with HDL-C levels significantly lower than in the controls and the dyslipidemic patients (\( P < .001 \)) and triglyceride levels significantly higher than in the controls (\( P < .01 \)). Four patients showed arterial hypertension; the rest had normal values but on average these values were significantly higher than in the controls.

In the dyslipidemic group, all patients were normotensive, with values similar to those of the control group. Obviously, since it was the selection criterion, all had high total cholesterol and LDL-C, with significant differences compared to the other groups. However, HDL-C levels were similar to

Statistical Methods

Sample size was calculated using the Ene 3.0 program (Statistical Department, University of Barcelona, Spain and Department of Biometry, GlaxoSmithKline\textsuperscript{®}). The results of an initial pilot study on PWV and also on Ep were taken as a reference. For both, sample size should be between 23-25 subjects per group.

Statistical processing was performed with SPSS version 15. The results were expressed as mean and standard deviation. Continuous variables were compared using the Student \( t \) test. The relation

Arterial compliance (AC): \( \pi \left( \frac{D_s^2 - D_d^2}{4} \right) / \left( 4SBP-DBP \right) \). Units: mm\(^2\)/kPa

Pressure-strain elasticity modulus (Ep): \( SBP-DBP \frac{D_d}{(D-s-D_d)} \). Units: kPa

Stiffness index (\( \beta \)): \( ln \left( \frac{SBP-DBP}{D_s-D_d} \right) \). Units: mm/s

One-point PWV (PWV): Distance/\( \Delta t \). Units: m/s

Augmentation index (AI): \( D2 - D1 \times 100/(D_s-D_d) \). Units: as a percentage

The IMT was measured at the same point specified for functional examination, determining the thickness in the carotid far wall, according to the criteria established by Mannheim.\textsuperscript{33} The images were recorded digitally and measured again manually. The mean of 3 measurements was obtained, as well as the maximum value; both were analyzed.

Figure 1. Results windows. Top: pulse wave and electrocardiogram. Bottom right: calculated parameters of arterial stiffness in one of the patients. \( \beta \) indicates stiffness parameter; Ep, pressure-strain elasticity modulus; AC, arterial compliance; AI, augmentation index; PWV, one-point pulse wave velocity; D_max, maximum arterial diameter; D_min, minimum arterial diameter; P_max, systolic blood pressure; P_min, diastolic blood pressure; HR, heart rate
those observed in controls. None of them had HDL-C levels below 35 mg/dL.

With respect to measurements of arterial stiffness and IMT (Table 2), on comparing the CVRF groups with the controls we found that all parameters increased in CVRF patients compared to the controls; however, these differences were only statistically significant for the β, Ep, PWV and arterial diameters in obese children. Statistically significant differences in AC also were observed, but only in dyslipemic patients compared to controls. On comparing the 2 CVRF groups, significant differences were found only in AI and arterial diameters.

The IMT measurements were taken in 42 cases (18 obese children, 9 dyslipemic children, and 15 controls). Analysis of results showed no significant differences between groups.

The results of the correlation analysis between anthropometric and biochemical parameters compared with vascular parameters are summarized in Table 3: in controls DBP was correlated with PWV and HDL-C with vascular parameters (β, AC, and PWV); in obese children there was correlation of BMI, SBP and triglyceride levels with vascular parameters (β, Ep, AC, and PWV); in the dyslipidemic subjects, a correlation was only found between triglyceride levels and β, Ep and PWV. IMT was not correlated with any biochemical parameters apart from triglyceride levels in the dyslipidemic group.

**TABLE 1. Anthropometric Results, Arterial Pressure and Lipidic Values of Study Groups**

<table>
<thead>
<tr>
<th>Control (n=34)</th>
<th>CVRF Group (n=65)</th>
<th>Obesity (n=45)</th>
<th>Dyslipidemia (n=20)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sex (male/female)</td>
<td>21/13</td>
<td>32/13</td>
<td>9/11</td>
</tr>
<tr>
<td>Age, mean (SD), y</td>
<td>11.6 (1.9)</td>
<td>12.4 (2.2)</td>
<td>11.1 (2.2)</td>
</tr>
<tr>
<td>Z-score BMI</td>
<td>0.03 (0.7)</td>
<td>2.6 (0.5)</td>
<td>0.44 (1.1)</td>
</tr>
<tr>
<td>SBP, mmHg</td>
<td>101 (10)</td>
<td>115 (12)</td>
<td>104 (12)</td>
</tr>
<tr>
<td>DBP, mmHg</td>
<td>56 (7)</td>
<td>61 (8)</td>
<td>57 (7)</td>
</tr>
<tr>
<td>Total-C, mg/dL</td>
<td>165 (25)</td>
<td>161 (35)</td>
<td>263 (62)</td>
</tr>
<tr>
<td>HDL-C, mg/dL</td>
<td>62 (11)</td>
<td>47 (8)</td>
<td>60 (18)</td>
</tr>
<tr>
<td>LDL-C, mg/dL</td>
<td>88 (21)</td>
<td>97 (29)</td>
<td>188 (52)</td>
</tr>
<tr>
<td>Triglycerides, mg/dL</td>
<td>69 (22)</td>
<td>101 (49)</td>
<td>80 (33)</td>
</tr>
</tbody>
</table>

Abbreviations: BMI, body mass index; CVRF, cardiovascular risk factors; DBP, diastolic blood pressure; HDL-C, high-density lipoproteins cholesterol; LDL-C, low-density lipoproteins cholesterol; SBP, systolic blood pressure.

P<.001 (controls vs CVRF).

P<.001 (controls vs dyslipidemia).

P<.05 (obesity vs dyslipidemia).

P<.001 (controls vs CVRF).

**TABLE 2. Results of the Vascular Parameters of Study Groups**

<table>
<thead>
<tr>
<th>Control (n=34)</th>
<th>CVRF Group (n=65)</th>
<th>Obesity (n=45)</th>
<th>Dyslipidemia (n=20)</th>
</tr>
</thead>
<tbody>
<tr>
<td>β</td>
<td>3.67 (0.84)</td>
<td>4.21 (0.96)</td>
<td>4.08 (1.56)</td>
</tr>
<tr>
<td>Ep, kPa</td>
<td>38.86 (7.79)</td>
<td>48.33 (12.33)</td>
<td>42.26 (14.78)</td>
</tr>
<tr>
<td>AC, mm²/kPa</td>
<td>1.41 (0.32)</td>
<td>1.36 (0.33)</td>
<td>1.2 (0.44)</td>
</tr>
<tr>
<td>Al, %</td>
<td>7.73 (18.67)</td>
<td>2.84 (12.34)</td>
<td>12.68 (15.89)</td>
</tr>
<tr>
<td>PWV, m/s</td>
<td>3.7 (0.34)</td>
<td>4.02 (0.44)</td>
<td>3.72 (0.96)</td>
</tr>
<tr>
<td>Ds</td>
<td>6.48 (0.61)</td>
<td>6.99 (0.5)</td>
<td>6.08 (1.49)</td>
</tr>
<tr>
<td>Dd</td>
<td>5.62 (0.56)</td>
<td>6.08 (0.43)</td>
<td>5.41 (1.33)</td>
</tr>
<tr>
<td>IMT-mean</td>
<td>0.32 (0.05)</td>
<td>0.36 (0.04)</td>
<td>0.33 (0.13)</td>
</tr>
<tr>
<td>IMT-maximum</td>
<td>0.36 (0.05)</td>
<td>0.40 (0.06)</td>
<td>0.33 (0.15)</td>
</tr>
</tbody>
</table>

Abbreviations: β, stiffness parameter; AC, arterial compliance; Al, augmentation index; Dd, diastolic arterial diameter; Ep, pressure-strain elasticity modulus; Ds, systolic arterial diameter; IMT-maximum, maximum intima-media thickness; PWV, one-point pulse wave velocity.

P<.002 (CVRF vs controls).

P<.001 (CVRF vs controls).

P<.01 (CVRF vs controls).

P<.01 (CVRF vs controls).

P<.01 (CVRF vs controls).

Forty-two cases (15 controls, 18 obese, and 9 dyslipidemic)
TABLE 3. Correlation Between Body Mass Index, Blood Pressure and Lipid Values and Vascular Parameters

<table>
<thead>
<tr>
<th></th>
<th>Controls (n=34)</th>
<th>Obese Children (n=45)</th>
<th>Dyslipidemic Children (n=20)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Ep</td>
<td>PWV</td>
<td>IMT-Mean*</td>
</tr>
<tr>
<td>BMI</td>
<td>0.295 (NS)</td>
<td>0.292 (NS)</td>
<td>0.613 (0.02)</td>
</tr>
<tr>
<td>SBP</td>
<td>0.7 (&lt;0.001)</td>
<td>0.655 (&lt;0.001)</td>
<td>0.028 (NS)</td>
</tr>
<tr>
<td>DBP</td>
<td>0.089 (NS)</td>
<td>0.257 (NS)</td>
<td>0.023 (NS)</td>
</tr>
<tr>
<td>LDL-C</td>
<td>0.023 (NS)</td>
<td>0.023 (NS)</td>
<td>0.023 (NS)</td>
</tr>
<tr>
<td>HDL-C</td>
<td>0.367 (NS)</td>
<td>0.43 (0.028)</td>
<td>0.219 (NS)</td>
</tr>
<tr>
<td>TG</td>
<td>0.043 (NS)</td>
<td>0.129 (NS)</td>
<td>0.332 (NS)</td>
</tr>
</tbody>
</table>

Abbreviations: BMI, body mass index; DBP, diastolic blood pressure; Ep, pressure-strain elasticity modulus; HDL-C, high-density lipoprotein cholesterol; IMT-mean, mean intima-media thickness; LDL-C, low-density lipoproteins cholesterol; PWV, one-point pulse wave velocity; SBP, systolic blood pressure; TG, triglycerides.

*42 cases (15 controls, 18 obese, and 9 dyslipidemic).

Other correlations: controls, beta stiffness index versus HDL-C (P=.010); AC versus HDL-C (P=.035); obese children, beta stiffness index versus BMI (P=.006), SBP (P=.002) and TG (P=.007); AC versus SBP (P=.001); dyslipidemic children, beta stiffness index versus TG (P=.031).

DISCUSSION

The preclinical phase of atherosclerosis begins in childhood, when very early alterations that progress slowly have been described1–5 and which do not usually lead to ischemic complications in adulthood. Early detection of the extent of arterial degeneration caused by atherosclerosis using noninvasive techniques, based on ultrasound, has provided an important stimulus to develop early detection programs and to assess the effects of intervention in the pediatric population.

So far most research conducted in children and adolescents has studied the IMT to determine the vascular damage in CVRF patients, mainly in obese children.21,22,34 However, results show there is already structural damage to the arterial wall and thus indicate that measurements of arterial stiffness parameters reflect dysfunction earlier on, which is, therefore, potentially reversible.16,35 Numerous studies have shown that PWV in children and adolescents is significantly correlated with CVRF and increases gradually with age in both sexes. In young adults, this parameter has been established as a good measure of arterial stiffness, proving to be an important predictor of cardiovascular events.32,36

Other parameters to assess arterial stiffness, such as arterial compliance and distensibility, evaluate the role of the artery studied as a hollow structure. The Ep parameter can be regarded as a measure of the intrinsic rigidity of the arterial wall itself and is inversely related to arterial elasticity.32 Two other parameters used in adults have been the beta stiffness index, which assesses the elastic properties of the arterial wall independent of distending pressure and AI as an indirect index to evaluate aortic elasticity.37

In this study, the age range (8 to 16 years) was chosen expecting major changes would take place in carotid elastic properties as compared to the control group. The average age was similar in all groups, even slightly lower in the dyslipidemic group of children; therefore, the differences detected in functional vascular parameters cannot be attributed to age.

On comparing arterial stiffness parameters of healthy children with those of CVRF children, they were higher in the latter, but statistically significant differences were only found in beta stiffness index β, Ep, PWV and arterial diameters in obese children and for AC in dyslipemic patients. Differences in AI and arterial diameters were found between the 2 vascular risk groups. Similar results have been recorded by other authors in both obese and dyslipidemic patients.16,35 However, on assessing whether structural changes were reflected in the IMT, no differences were found in either study group when functional anomalies had already shown up. Bearing in mind the limitations due to sample size, these results seem relevant and support the potential use of these functional indices as early markers of vascular injury in individuals at risk. Studies with larger sample sizes and longer follow-up periods can validate these findings.

The results obtained show the onset of changes in arterial elasticity take place sooner in obese children than in those with dyslipemia. We also found that blood pressure values were significantly higher than other groups, although they did not reach hypertensive levels. This same group also revealed a pattern associating dyslipidemia with significantly lower HDL-C values and higher triglyceride levels, which would suggest they are at an early stage of metabolic syndrome. Moreover, BMI, SBP and triglycerides were significantly correlated with arterial stiffness indices, which would directly indicate adiposity as a trigger of vascular injury. These findings support the need to implement intervention programs for obese school children and adolescents in order to delay the process before adulthood. Similarly, postoperative
follow-up of vascular anomalies would be of great interest to evaluate the possible reversibility of functional alterations and/or IMT.

With regard to the relationship between dyslipidemia and cardiovascular disease, there is no doubt that low levels of LDL-C play a role in the prevention and treatment of coronary disease. The effect of dyslipidemia on arterial elastic properties has been extensively studied in adults, where healthy individuals have shown an inverse relationship between LDL levels and aortic compliance. Elasticity of the common carotid artery was also assessed in normotensive hypercholesterolemic patients (with or without varying degrees of coronary disease), in hypertensive normocholesterolemic patients (without coronary disease) and healthy controls. Results indicated reduced carotid compliance mainly occurred in normotensive hypercholesterolemic patients, suggesting that reduction in compliance is an indicator of severe atherosclerosis.

In children with heterozygous familial hypercholesterolemia, an increase in stiffness of the carotid artery has been observed. This group of patients was found to have LDL-C levels that were positively correlated with PWV. However, no correlation was found between either Ep or IMT and total cholesterol or LDL-C levels. Our patients with dyslipidemia showed significantly higher values of LDL-C than the obese patients and controls, with HDL-C values in all 3 groups over 35 mg/dL. In this group there were statistically significant differences in AC compared with controls; however, we did not observe any correlation between LDL-C levels and vascular damage as reported by other authors, possibly because the sample size of this group was smaller than the others.

Significant changes in functional vascular parameters in the 2 CVRF groups would seem to indicate that the arterial wall components are affected very early on, which probably leads to the onset of IMT found in these patients. Nevertheless, on analyzing this structural parameter in our study groups, we found no significant differences in IMT compared to controls, suggesting that this structural parameter is affected later on. These results lead us to the conclusion that functional vascular parameters are the ultrasonographic indicators which change the earliest in children with CVRF.

CONCLUSIONS

The ultrasonographic measurement of arterial stiffness is a sensitive procedure which can serve as an earlier marker than IMT for detecting vascular changes in children with CVRF. Systematic application of these techniques may have important preventive implications because it would provide a marker of atherosclerosis at the preclinical phase of the disease. Detection of functional alterations in children at risk would enable them to benefit from early therapeutic measures, thereby preventing or delaying the development of atherosclerosis when they reach adulthood.

REFERENCES