EDITORIAL

Predictors of Carotid Atherosclerosis in Young Adults: Insights From the Bogalusa Heart Study

Ching Lung Cheung, PhD; Bernard M. Y. Cheung D, PhD

therosclerotic cardiovascular disease (ASCVD) develops early in life, but becomes clinically manlifest later on in adulthood. The Bogalusa Heart Study, which was initiated in 1972 by the pediatric cardiologist, Dr Gerald Berenson, and is still ongoing, has shown that ASCVD begins in childhood. In this issue of the Journal of the American Heart Association (JAHA), Razavi et al¹ reported on the predictors of absence of carotid plague among 508 young adult participants of the Bogalusa Heart Study, who underwent ultrasound imaging at baseline and a decade later. As expected, young age and a total cholesterol/high-density lipoprotein cholesterol ratio <3.5 were associated with absence of plaque. Besides a favorable traditional risk factor profile and high glomerular filtration rate, they observed that low serum phosphate and calcium-phosphate product and dietary sodium <2300 mg/d were associated with nondevelopment of plaque. This is a refreshing and important message, because it suggests that lifestyle changes early in life may protect against ASCVD many years later. However, because of the small sample size and the large number of variables collected, the findings would need to be confirmed in another cohort.

See Article by Razavi et al.

Dietary sodium intake is linked to the risk of developing hypertension² and so the present study adds to the evidence supporting the current advice to cut sodium intake to <2300 mg/d. Nevertheless, a dietary sodium <2300 mg/d is substantially lower than the average sodium intake in the United States and other developed countries. Such a low intake is hard to sustain even in research studies. As a low sodium intake may activate the renin-angiotensin system, Dr Michael Alderman and others have questioned whether this may be more harmful than beneficial.³ Moreover, a low sodium intake can be a marker of healthy diet or healthy lifestyle, because achieving a low sodium intake in Bogalusa would probably require fewer fast food meals and more fresh food ingredients.

Chronic kidney disease is well known to be associated with ASCVD, which is the leading cause of death in this patient group. Glomerular filtration rate and albuminuria are predictors of cardiovascular outcomes.⁴ Although the addition of glomerular filtration rate improves the prediction of cardiovascular events beyond traditional risk factors, it has not been included in popular cardiovascular risk scores, such as the Pooled Cohort Equation or the Systematic Coronary Risk Evaluation. Therefore, the study by Razavi et al confirms glomerular filtration rate as a predictor of carotid atherosclerosis independent of traditional risk factors. In young adults, a faster decline in the glomerular filtration rate may reflect the renal damage caused by chronically elevated blood pressure and glucose, especially when the young person is untreated.

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Correspondence to: Bernard M. Y. Cheung, PhD, Department of Medicine, Queen Mary Hospital, 102 Pokfulam Rd, Hong Kong. E-mail: mycheung@hku.hk The opinions expressed in this article are not necessarily those of the editors or of the American Heart Association. For Disclosures, see page 3.

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In chronic kidney disease, serum calcium, phosphate, and the calcium-phosphate product are elevated and are thought to contribute to arterial calcifications and cardiovascular events.⁵ Thus, phosphate binders are used clinically in patients with advanced chronic kidney disease, although it is still unclear whether non-calcium-based phosphate binders are safer than calcium-based phosphate binders in terms of cardiovascular outcome. Nevertheless, serum phosphate is also associated with vascular dysfunction and ASCVD, even in people without chronic kidney disease.⁶ Calcium-phosphate homeostasis is tightly regulated by the levels of calcium, phosphate, parathyroid hormone, vitamin D, as well as newcomers, such as klotho and fibroblast growth factor 23, via the interplay between parathyroid gland, bone, kidney, and gut. The relationship between vitamin D and ASCVD has aroused much interest in the past decade. Circulating vitamin D levels were associated with a reduced risk of cardiovascular events in cohort studies^{7,8}; but clinical trials of vitamin D supplementation showed beneficial effects on cardiovascular risk factors only,9 but not on the incidence of CVD,^{10,11} even with a high dose of vitamin D.¹² Crucially, the VITAL (Vitamin D and Omega-3 Trial) involving 25 871 participants followed up for 5 years found no benefit in vitamin D supplementation compared with placebo.¹³

Because other mineral markers are closely correlated with vitamin D levels, the association observed in those epidemiological studies could have been contributed by other minerals, such as calcium and phosphate. The role of calcium in ASCVD is intriguing. Two meta-analyses by Ian Reid's group showed that calcium supplementation increased the risk of myocardial infarction and cardiovascular diseases.^{14,15} A Mendelian randomization study supported a possible casual role for serum calcium,¹⁶ although a subsequent meta-analysis did not support a relationship between calcium supplementation and ASCVD.¹⁷ In the study by Razavi et al, serum phosphate and calcium-phosphate product, but not serum calcium, were shown to be predictors of carotid atherosclerosis. Most of the previous studies on calcium-phosphate and ASCVD risk were conducted in an older population and people with chronic kidney disease. The findings in the study by Razavi et al are of interest because these electrolyte measurements preceded clinical ASCVD. Interestingly, despite the difference in serum phosphate and calcium-phosphate product, the baseline renal function was comparable between those with and without incident carotid atherosclerosis. This implies that young people with elevated phosphate and calcium-phosphate product might be more prone to carotid atherosclerosis development. Further studies, using Mendelian randomization for instance, would be needed to establish causality.

The study by Razavi et al does not negate the importance of traditional risk factors. Although it showed that the 5 components of the metabolic syndrome could not explain nearly three quarters of the premature carotid atherosclerosis, age, smoking, blood pressure, low-density lipoprotein, triglyceride, and waist circumference were all significantly associated with carotid atherosclerosis before multivariable adjustment. This brings home the point that ASCVD risk factors can be different in different age groups. Logically, the risk factors in the young are less likely to be diseases, such as hypertension and diabetes mellitus, but more likely to be related to genetics and lifestyle factors, including diet. Studies of ASCVD risk factors in children and young adults may reveal more novel risk factors and offer new approaches to stopping ASCVD before it develops.

ARTICLE INFORMATION

Affiliations

Department of Pharmacology and Pharmacy (C.L.C.) and Department of Medicine (B.M.C.), Li Ka Shing Faculty of Medicine, University of Hong Kong, Hong Kong, China.

Disclosures

None.

REFERENCES

- Razavi AC, Bazzano LA, He J, Krousel-Wood M, Chen J, Fernandez C, Whelton SP, Kelly TN. Early contributors to healthy arterial aging versus premature atherosclerosis in young adults: the Bogalusa Heart Study. J Am Heart Assoc. 2021;10:e020774. DOI: 10.1161/ JAHA.121.020774.
- Whelton PK, Appel LJ, Sacco RL, Anderson CAM, Antman EM, Campbell N, Dunbar SB, Frohlich ED, Hall JE, Jessup M, et al. Sodium, blood pressure, and cardiovascular disease: further evidence supporting the American Heart Association sodium reduction recommendations. *Circulation*. 2012;126:2880–2889. DOI: 10.1161/CIR.0b013e3182 79acbf.
- Graudal N, Jürgens G, Baslund B, Alderman MH. Compared with usual sodium intake, low-and excessive-sodium diets are associated with increased mortality: a meta-analysis. *Am J Hypertens*. 2014;27:1129– 1137. DOI: 10.1093/ajh/hpu028.
- Matsushita K, Coresh J, Sang Y, Chalmers J, Fox C, Guallar E, Jafar T, Jassal SK, Landman GWD, Muntner P, et al; CKD Prognosis Consortium. Estimated glomerular filtration rate and albuminuria for prediction of cardiovascular outcomes: a collaborative meta-analysis of individual participant data. *Lancet Diabetes Endocrinol.* 2015;3:514– 525. DOI: 10.1016/S2213-8587(15)00040-6.
- Mathew S, Tustison KS, Sugatani T, Chaudhary LR, Rifas L, Hruska KA. The mechanism of phosphorus as a cardiovascular risk factor in CKD. J Am Soc Nephrol. 2008;19:1092–1105. DOI: 10.1681/ASN.2007070760.
- Ginsberg C, Houben A, Malhotra R, Berendschot T, Dagnelie PC, Kooman JP, Webers CA, Stehouwer CDA, Ix JH. Serum phosphate and microvascular function in a population-based cohort. *Clin J Am Soc Nephrol.* 2019;14:1626–1633. DOI: 10.2215/CJN.02610319.
- Grandi NC, Breitling LP, Brenner H. Vitamin D and cardiovascular disease: systematic review and meta-analysis of prospective studies. *Prev Med.* 2010;51:228–233. DOI: 10.1016/j.ypmed.2010.06.013.
- Leung RY, Han Y, Sing CW, Cheung BM, Wong IC, Tan KC, Kung AW, Cheung CL. Serum 25-hydroxyvitamin D and the risk of stroke in Hong Kong Chinese. *Thromb Haemost*. 2017;117:158–163. DOI: 10.1160/ TH16-07-0551.

- Mirhosseini N, Rainsbury J, Kimball SM. Vitamin D supplementation, serum 25(OH)D concentrations and cardiovascular disease risk factors: a systematic review and meta-analysis. *Front Cardiovasc Med.* 2018;5:87. DOI: 10.3389/fcvm.2018.00087.
- Swart KMA, Lips P, Brouwer IA, Jorde R, Heymans MW, Grimnes G, Grübler MR, Gaksch M, Tomaschitz A, Pilz S, et al. Effects of vitamin D supplementation on markers for cardiovascular disease and type 2 diabetes: an individual participant data meta-analysis of randomized controlled trials. *Am J Clin Nutr.* 2018;107:1043–1053. DOI: 10.1093/ ajcn/ngy078.
- Barbarawi M, Kheiri B, Zayed Y, Barbarawi O, Dhillon H, Swaid B, Yelangi A, Sundus S, Bachuwa G, Alkotob ML, et al. Vitamin D supplementation and cardiovascular disease risks in more than 83000 individuals in 21 randomized clinical trials: a meta-analysis. *JAMA Cardiol.* 2019;4:765–776. DOI: 10.1001/jamacardio.2019.1870.
- Scragg R, Stewart AW, Waayer D, Lawes CMM, Toop L, Sluyter J, Murphy J, Khaw KT, Camargo CA Jr. Effect of monthly high-dose vitamin D supplementation on cardiovascular disease in the vitamin D assessment study: a randomized clinical trial. *JAMA Cardiol.* 2017;2:608–616. DOI: 10.1001/jamacardio.2017.0175.

- Manson JE, Cook NR, Lee I-M, Christen W, Bassuk SS, Mora S, Gibson H, Gordon D, Copeland T, D'Agostino D, et al. Vitamin D supplements and prevention of cancer and cardiovascular disease. *N Engl J Med.* 2019;380:33–44. DOI: 10.1056/NEJMoa1809944.
- Bolland MJ, Grey A, Avenell A, Gamble GD, Reid IR. Calcium supplements with or without vitamin D and risk of cardiovascular events: reanalysis of the Women's Health Initiative limited access dataset and meta-analysis. *BMJ*. 2011;342:d2040. DOI: 10.1136/bmj.d2040.
- Bolland MJ, Avenell A, Baron JA, Grey A, MacLennan GS, Gamble GD, Reid IR. Effect of calcium supplements on risk of myocardial infarction and cardiovascular events: meta-analysis. *BMJ*. 2010;341:c3691. DOI: 10.1136/bmj.c3691.
- Larsson SC, Burgess S, Michaelsson K. Association of genetic variants related to serum calcium levels with coronary artery disease and myocardial infarction. *JAMA*. 2017;318:371–380. DOI: 10.1001/ jama.2017.8981.
- Chung M, Tang AM, Fu Z, Wang DD, Newberry SJ. Calcium intake and cardiovascular disease risk: an updated systematic review and meta-analysis. *Ann Intern Med.* 2016;165:856–866. DOI: 10.7326/ M16-1165.