Doppler study of cerebral arteries in hypercholesterolemia

Mehdi Farhoudi1
Kaveh Mehrvar2
Naser Aslanabadi3
Kamyar Ghabili1
Nazila Rasi Baghmishe4
Farzad Ilkhchoei4

1Neuroscience Research Center, 2Razi Hospital, 3Department of Cardiology, 4Faculty of Medicine, Tabriz University of Medical Sciences, Tabriz, Iran

Background: Hypercholesterolemia is one of the major modifiable risk factors for atherosclerosis of the coronary and carotid arteries. Although transcranial Doppler (TCD) studies of the cerebral arteries are indicative of decreased cerebral blood flow velocities in patients with hypercholesterolemia, the number of these studies has been limited. The aim of this study was to assess the hemodynamic status of the cerebral arteries in patients with hypercholesterolemia using TCD.

Methods: In a case-control study, 60 individuals, including 30 hypercholesterolemic cases (low-density lipoprotein [LDL] > 160 mg/dL) and 30 healthy controls were enrolled. Other arterial risk factors, including diabetes mellitus, hypertension, smoking, and obesity (body mass index > 30), were evaluated and matched as well. TCD was used to assess the hemodynamics of the intracranial arteries as well as the internal carotid arteries. The mean blood flow velocity, pulsatility index, and resistance index were recorded in all the arteries.

Results: The mean blood flow velocity, pulsatility index, and resistance index of the intracranial arteries and internal carotid arteries were not significantly different between the two groups (P > 0.05). However, those with higher levels of LDL (>180 mg/dL) showed significantly lower mean blood flow velocity and resistance index of the internal carotid arteries than the healthy controls. In addition, individuals with high-density lipoprotein (HDL) <35 mg/dL had significantly lower mean blood flow velocity in the internal carotid arteries.

Conclusion: Hypercholesterolemia (LDL > 160 mg/dL) does not seem to have a detrimental effect on the hemodynamic status of the intracranial arteries and internal carotid arteries. However, in cases of higher LDL (>180 mg/dL) and lower HDL, decreased TCD parameters in the internal carotid arteries, as a surrogate of the peripheral arteries, are prominent.

Keywords: hypercholesterolemia, transcranial Doppler, cerebral arteries

Introduction
Cerebrovascular disease and subsequent stroke is one of the most important neurological diseases. Several modifiable (eg, hypertension, hyperlipidemia, obesity, smoking, and diabetes) and nonmodifiable (eg, gender, age, and race) risk factors for stroke have been established.1 Hypercholesterolemia is one of the major modifiable risk factors for atherosclerosis of the coronary and carotid arteries. However, its role in the pathogenesis of stroke is unknown.2–4 Although early epidemiological studies failed to find an association between cholesterol levels and overall stroke risk,5,6 later investigations revealed a relationship between dyslipidemia and risk of ischemic stroke.7–9

Assessments of human and animal arterial vasculature revealed an impaired endothelial function in hypercholesterolemia.10–12 In addition, transcranial Doppler
Farhoudi et al

(TCD) studies of the major cerebral arteries have shown decreased cerebral blood flow velocities in patients with familial hypercholesterolemia. Nonetheless, studies of the hemodynamics of the cerebral vasculature in hypercholesterolemic patients are limited. The present study aimed at assessing the hemodynamic status of the cerebral arteries in patients with hypercholesterolemia using TCD.

Methods

In this case-control study, 60 individuals comprising 30 hypercholesterolemic cases (low-density lipoprotein [LDL] > 160 mg/dL) and 30 healthy controls were enrolled in a university-affiliated hospital in Tabriz, Iran. The study was approved by the local ethics committee and written informed consent was obtained for each subject. Both groups were age- and gender-matched. Other arterial risk factors including diabetes mellitus, hypertension, smoking, and obesity (body mass index > 30) were evaluated and matched as well. Individuals who had given up smoking for more than five years were considered to be nonsmokers. TCD was used to assess the hemodynamics of the cerebral arteries (DWL®, Multi-Dop®, Germany). The standard method of using a 2 mHz probe for the intracranial arteries and a 4 mHz probe for the cervical carotids was applied in the present study. The Doppler signals were detected through the temporal window for the middle cerebral artery, anterior cerebral artery and two segments of the posterior cranial artery (PCA1, PCA2), through the suboccipital window for the vertebral arteries and basilar artery, and through the cervical window for the internal carotid arteries. Mean blood flow velocity, pulsatility index, and resistance index were recorded in all the arteries.

Data were presented as means ± standard deviations. Statistical analysis was performed with SPSS for Windows version 13.0 (Chicago, IL) and by using the Chi-square test, Fisher’s Exact test, and independent-samples t-test wherever appropriate. P < 0.05 was considered statistically significant.

Results

Sixty individuals were included in the study, ie, 30 in the hypercholesterolemic group and 30 in the healthy control group. There were no differences in gender, age, and vascular risk factors, including hypertension, diabetes mellitus, smoking, and obesity between the groups (Table 1, P > 0.05). Moreover, the mean blood flow velocity, pulsatility index, and resistance index of the intracranial arteries and internal carotid arteries were not significantly different between the two groups (Table 2, P > 0.05).

To find out whether higher levels of LDL alter the hemodynamics of the cerebral arteries, individuals with LDL levels of less and more than 180 mg/dL were compared (Table 3). The mean blood flow velocity and resistance index of the internal carotid arteries in those with LDL > 180 mg/dL was significantly lower than that in individuals with LDL < 180 mg/dL (P < 0.05, Table 3). In addition, individuals with high-density lipoprotein (HDL) < 35 mg/dL were compared regarding TCD indices in both groups. The analysis revealed a significant difference only in mean blood flow velocity of the internal carotid arteries.

Discussion

The hemodynamics of the cerebral vasculature in both healthy individuals and patients have been of great research interest. Impaired cerebral vasomotor reactivity to L-arginine and/or CO2 has been reported in patients with recent stroke, severe internal carotid artery stenosis, lacunar infarction, and high blood pressure. Nonetheless, studies of the hemodynamics of the cerebral vasculature in hypercholesterolemic patients are limited. The current literature mainly focuses on the effect of lipid-lowering methods on hemodynamic alterations in the cranial arteries. Sander et al22 and Sterzer et al23 reported an improvement in cerebral vasoreactivity after statin administration. However, the results of PROSPER (Prospective Study of Pravastatin in the Elderly at Risk) failed to confirm this finding. On the other hand, Pfefferkorn et al25 revealed that cholesterol reduction through

| Patients’ demographic data and clinical characteristics (mean ± standard deviation) |
|---------------------------------|---------------------------------|----------------|
| Hypercholesterolemia group (n = 30) | Control group (n = 30) | P value |
| Gender (male:female) | 14:16 | 14:16 | 1.00 |
| Age (years) | 54.66 (36–70) | 51.60 (30–75) | 0.55 |
| Hypertension, n (%) | 13 (43.3) | 8 (26.6) | 0.27 |
| Diabetes mellitus, n (%) | 4 (13.3) | 5 (16.6) | 1.00 |
| Smoker, n (%) | 11 (36.6) | 4 (13.3) | 0.07 |
| Obesity, n (%) | 8 (26.6) | 5 (16.6) | 0.53 |
Dopplers of cerebral arteries in hypercholesterolemia

Table 2 Transcranial Doppler indices of the intracranial artery (mean ± standard deviation)

| Artery | Index | Hypercholesterolemia group (n = 30) | Control group (n = 30) | P value |
|--------|-------|------------------------------------|------------------------|---------|
| MCA MV | 60.5 ± 18.8 | 56.2 ± 9.2 | 0.98 |
| PI     | 0.82 ± 0.17  | 0.78 ± 0.15 | 0.45 |
| RI     | 0.53 ± 0.08  | 0.51 ± 0.05 | 0.46 |
| ACA MV | 46.4 ± 22.3  | 63.9 ± 10.1 | 0.86 |
| PI     | 0.76 ± 0.35  | 0.83 ± 0.5  | 0.70 |
| RI     | 0.46 ± 0.19  | 0.48 ± 0.14 | 0.45 |
| ICA MV | 22 ± 17.9    | 28.7 ± 9.9  | 0.15 |
| PI     | 0.59 ± 0.45  | 0.83 ± 0.27 | 0.08 |
| RI     | 0.36 ± 0.26  | 0.51 ± 0.15 | 0.06 |
| PCA1 MV| 38.9 ± 12.4  | 37 ± 9.6    | 0.91 |
| PI     | 0.08 ± 0.22  | 0.76 ± 0.2  | 0.43 |
| RI     | 0.51 ± 0.11  | 0.49 ± 0.1  | 0.44 |
| PCA2 MV| 40.1 ± 13    | 38.6 ± 11.3 | 0.51 |
| PI     | 0.79 ± 0.21  | 0.79 ± 0.16 | 0.51 |
| RI     | 0.51 ± 0.11  | 0.53 ± 0.12 | 0.75 |
| VA MV  | 35.2 ± 11.6  | 32.9 ± 8.9  | 0.60 |
| PI     | 0.82 ± 0.17  | 0.76 ± 0.13 | 0.21 |
| RI     | 0.52 ± 0.06  | 0.51 ± 0.08 | 0.32 |
| BA MV  | 41.8 ± 13.2  | 41.2 ± 11.3 | 0.62 |
| PI     | 0.85 ± 0.17  | 0.76 ± 0.22 | 0.14 |
| RI     | 0.54 ± 0.06  | 0.49 ± 0.11 | 0.12 |

Abbreviations: MCA, middle cerebral artery; ACA, anterior cerebral artery; ICA, internal carotid artery; PCA1, first segment of posterior cranial artery; PCA2, second segment of posterior cranial artery; VA, vertebral artery; BA, basilar artery; MV, mean blood flow velocity; PI, pulsatility index; RI, resistive index.

Table 3 Transcranial Doppler indices of intracranial arteries in LDL > 180 mg/dL versus < 180 mg/dL groups (mean ± standard deviation)

| Artery | Index | LDL >180 mg/dL | LDL <180 mg/dL | P value |
|--------|-------|----------------|----------------|---------|
| MCA MV | 56.8 ± 12.56 | 59.1 ± 15.9 | 0.54 |
| PI     | 0.79 ± 0.19  | 0.81 ± 0.15 | 0.50 |
| RI     | 0.52 ± 0.09  | 0.52 ± 0.05 | 0.41 |
| ACA MV | 44.5 ± 23.06 | 60.4 ± 88   | 0.69 |
| PI     | 0.71 ± 0.35  | 0.84 ± 0.46 | 0.23 |
| RI     | 0.44 ± 0.19  | 0.49 ± 0.15 | 0.27 |
| ICA MV | 19.3 ± 16    | 28.4 ± 13.4 | 0.04* |
| PI     | 0.56 ± 0.44  | 0.79 ± 0.34 | 0.055 |
| RI     | 0.34 ± 0.26  | 0.48 ± 0.19 | 0.02* |
| PCA1 MV| 36 ± 13      | 38.9 ± 10   | 0.12 |
| PI     | 0.74 ± 0.24  | 0.8 ± 0.2   | 0.25 |
| RI     | 0.48 ± 0.01  | 0.51 ± 0.1  | 0.26 |
| PCA2 MV| 39.8 ± 15.4  | 39.1 ± 10.4 | 0.96 |
| PI     | 0.74 ± 0.23  | 0.82 ± 0.16 | 0.22 |
| RI     | 0.48 ± 0.12  | 0.54 ± 0.11 | 0.18 |
| VA MV  | 34.7 ± 10    | 33.6 ± 10.6 | 0.66 |
| PI     | 0.78 ± 0.15  | 0.8 ± 0.15  | 0.55 |
| RI     | 0.51 ± 0.05  | 0.52 ± 0.08 | 0.45 |
| BA MV  | 39.7 ± 9.1   | 42.3 ± 13.6 | 0.23 |
| PI     | 0.81 ± 0.16  | 0.81 ± 0.22 | 0.80 |
| RI     | 0.52 ± 0.06  | 0.51 ± 0.1  | 0.80 |

Note: *Statistically significant (P < 0.05).

Abbreviations: MCA, middle cerebral artery; ACA, anterior cerebral artery; ICA, internal carotid artery; PCA1, first segment of posterior cranial artery; PCA2, second segment of posterior cranial artery; VA, vertebral artery; BA, basilar artery; MV, mean blood flow velocity; PI, pulsatility index; RI, resistive index.

heparin-mediated extracorporeal LDL precipitation resulted in an improvement of cerebrovascular CO2 reactivity in patients with coronary heart disease and hyperlipidemia. Interestingly, Rubba et al13 and Iannuzzi et al26 noted an increasing trend in cerebral blood flow velocities after LDL apheresis in patients with familial hypercholesterolemia.

The present study showed that the hemodynamic status of the cerebral arteries, including the internal carotid arteries, was similar in healthy individuals and those with LDL > 160 mg/dL. Nonetheless, in cases of a higher cholesterol level (LDL > 180 mg/dL), decreased mean blood flow velocity and resistance index of the internal carotid artery were noted, indicating the effect of high LDL levels only on the peripheral arteries. Reduced mean blood flow velocity of the internal carotid arteries was also seen with lower levels of HDL, probably indicative of the protective effect of HDL in prophylaxis of arterial damage. Kerenyi et al27 failed to find any significant difference between healthy controls and hyperlipidemic patients in cerebrovascular reactivity or reserve capacity after intravenous administration of acetazolamide. In a study of stroke-free Chinese individuals, hypercholesterolemia was associated with decreased blood flow velocity in the extracranial arteries, eg, the common carotid artery and internal carotid artery. However, this association was an inverse one in the intracranial arteries.28 Nevertheless, a few studies of patients with familial hypercholesterolemia have reported contrasting results in this regard. Rubba et al13 and Iannuzzi et al26 found an abnormally low cerebral blood flow velocity in the middle cerebral artery prior to LDL apheresis in patients with familial hypercholesterolemia.

A later study by Rubba et al28 demonstrated lower diastolic blood flow velocities and a higher pulsatility index in the middle cerebral artery in familial hypercholesterolemic patients compared with control subjects. It seems that higher levels of LDL (about 600 mg/dL) in studies of patients with familial hypercholesterolemia have reported controversial results regarding the hemodynamic status of the cranial arteries.

Although the relationship between hypercholesterolemia and coronary and carotid artery atherosclerosis is established, its role in the pathogenesis of stroke is unknown. An ultrasound study has shown a direct relationship between lipid levels and atherosclerosis of the carotid artery.1,30 Moreover, formation of reactive oxygen
species has been deemed to play a role in the pathogenesis of cerebral vascular dysfunction during hypercholesterolemia. Reactive oxygen species impair the response to acetylcholine, enhance superoxide activity, and lead finally to endothelial dysfunction.31

This study has certain limitations. It was a single-center study with a quite small sample size. A multicenter study including a large number of patients may give more concrete results. We did not exclude patients with a drug history of lipid-lowering agents. Further investigations excluding individuals on lipid-lowering agent therapy, e.g., statins are recommended. Furthermore, other sources of potential bias, including anxiety disorders, were not controlled for. On the other hand, some advantages of the current study might be highlighted. We matched the hypercholesterolemic and control groups with regard to arterial risk factors, including diabetes mellitus, hypertension, smoking, and obesity. We applied two different definitions of hypercholesterolemia (LDL 160 versus 180 mg/dL) to achieve further results.

In conclusion, hypercholesterolemia (LDL > 160 mg/dL) does not seem to have a detrimental effect on the hemodynamic status of the intracranial arteries and internal carotid arteries. However, in cases of higher cholesterol levels (LDL > 180 mg/dL) and lower levels of HDL, decreased TCD parameters for the internal carotid arteries, as a surrogate for the peripheral arteries, are prominent.

Disclosure
The authors report no conflicts of interest in this work.

References
1. Goldstein LB, Adams R, Alberts MJ, et al. Primary prevention of ischemic stroke: A guideline from the American Heart Association/American Stroke Association Stroke Council: Cosponsored by the Atherosclerotic Peripheral Vascular Disease Interdisciplinary Working Group; Cardiovascular Nursing Council; Clinical Cardiology Council; Nutrition, Physical Activity, and Metabolism Council; and the Quality of Care and Outcomes Research Interdisciplinary Working Group: The American Academy of Neurology affirms the value of this guideline. Stroke. 2006;37:1583–1633.
2. Amarenco P, Tonkin AM. Statins for stroke prevention: Disappointment and hope. Circulation. 2004;109:III44–III49.
3. Demchuk AM, Hess DC, Brass LM, et al. Is cholesterol a risk factor for stroke?: Yes. Arch Neurol. 1999;56:1518–1520.
4. Landau WM. Is cholesterol a risk factor for stroke? No. Arch Neurol. 1999;56:1521–1524.
5. Iso H, Jacobs DR Jr, Wentworth D, Neaton JD, Cohen JD. Serum cholesterol levels and six-year mortality from stroke in 350,977 men screened for the Multiple Risk Factor Intervention Trial. N Engl J Med. 1989;320:904–910.
6. Kagan A, Popper JS, Rhoads GG. Factors related to stroke incidence in Hawaiian Japanese men. The Honolulu Heart Study. Stroke. 1980;11:14–21.
7. Zhang X, Patel A, Horibe H, et al. Cholesterol, coronary heart disease, and stroke in the Asia Pacific region. Int J Epidemiol. 2003;32:563–572.
8. Bots ML, Elwood PC, Nikitin Y, et al. Total and HDL cholesterol and risk of stroke. EUROSTROKE: A collaborative study among research centres in Europe. J Epidemiol Community Health. 2002;56 Suppl 1:i19–i24.
9. Horenstein RB, Smith DE, Mosca L. Cholesterol predicts stroke mortality in the Women’s Pooling Project. Stroke. 2002;33:1863–1868.
10. Busse R, Fleming I. Endothelial dysfunction in atherosclerosis. J Vasc Res. 1996;33:181–194.
11. d’Uschio LV, Smith LA, Katosis ZS. Hypercholesterolemia impairs endothelium-dependent relaxations in common carotid arteries of apolipoprotein e-deficient mice. Stroke. 2001;32:2658–2664.
12. d’Uschio LV, Baker TA, Mantilla CB, et al. Mechanism of endothelial dysfunction in apolipoprotein E-deficient mice. Arterioscler Thromb Vasc Biol. 2001;21:1017–1022.
13. Rubba P, Facenda F, Di Somma S, et al. Cerebral blood flow velocity and systemic vascular resistance after acute reduction of low density lipoprotein in familial hypercholesterolemia. Stroke. 1993;24:1154–1161.
14. Farhoudi M, Kermani S, Sadeghi-Bazargani H. Relatively higher norms of blood flow velocity of major intracranial arteries in North-West Iran. BMC Res Notes. 2010;3:174.
15. Ghabili K, Khosroshahi HT, Shahker A, Tubbs RS, Bahluli A, Shoja MM. Can Doppler ultrasonographic indices of the renal artery predict the presence of supernumerary renal arteries? Transplant Proc. 2009;41:2731–2733.
16. Nemati M, Aslanabadi S, Bavel AS, et al. Diagnostic accuracy of Doppler ultrasonography in differentiation between malignant and benign cervical lymphadenopathies in pediatric age group. Pak J Biol Sci. 2010;13:757–760.
17. Ansarin K, Bavel AS, Ghabili K, et al. Are Doppler ultrasonography parameters symmetric between the right and left kidney? Int J Gen Med. 2010;3:371–373.
18. Carod-Artal FJ. Statins and cerebral vasomotor reactivity: Implications for a new therapy? Stroke. 2006;37:2446–2448.
19. Zvan B, Zaleiel M, Pogacnik T, Kliaut T. Testing of cerebral endothelium function with L-arginine after stroke. Int Angiol. 2002;21:256–259.
20. Miceli G, Bosone D, Zappoli F, Marcheselli S, Argerenti A, Nappi G. Vasomotor response to CO2 and L-arginine in patients with severe internal carotid artery stenosis; pre- and post-surgical evaluation with transcranial Doppler. J Neurol Sci. 1999;163:153–158.
21. Pretnar-Oblak J, Zaleiet M, Zvan B, Sabovic M, Pogacnik T. Cerebrovascular reactivity to L-arginine in patients with lacunar infarctions. Cerebrovasc Dis. 2006;21:180–186.
22. Sander K, Hof U, Poppurt H, Conrad B, Sander D. Improved cerebral vasoreactivity after statin administration in healthy adults. J Neuroimaging. 2005;15:266–270.
23. Sterzer P, Meintzschel F, Rösler A, Lantermann H, Steinmetz H, Sitzer M. Pravastatin improves cerebral vasomotor reactivity in patients with subocital small-vessel disease. Stroke. 2001;32:2817–2820.
24. ten Dam VH, Box FM, de Craen AJ, et al. Lack of effect of pravastatin on cerebral blood flow or parenchymal volume loss in elderly at risk for vascular disease. Stroke. 2005;36:1633–1636.
25. Pfefferkorn TK, Knüppel HP, Jaeger BR, Thiery J, Hamann GF. Increased cerebral CO2 reactivity after heparin-mediated extracorporeal LDL precipitation (HELP) in patients with coronary heart disease and hyperlipidemia. Stroke. 1999;30:1802–1806.
26. Iannuzzi A, Bianciardi G, Faccenda F, et al. Correction of erythrocyte density lipoprotein in familial hypercholesterolemia. Stroke. 1997;28:115–121.
28. Zhang P, Huang Y, Li Y, et al. Gender and risk factor dependence of cerebral blood flow velocity in Chinese adults. *Brain Res Bull*. 2006;69:282–287.

29. Rubba P, Mercuri M, Faccenda F, et al. Premature carotid atherosclerosis: Does it occur in both familial hypercholesterolemia and homocysteinuria? Ultrasound assessment of arterial intima-media thickness and blood flow velocity. *Stroke*. 1994;25:943–950.

30. Heiss G, Sharrett AR, Barnes R, Chambless LE, Szklo M, Alzola C. Carotid atherosclerosis measured by B-mode ultrasound in populations: Associations with cardiovascular risk factors in the ARIC study. *Am J Epidemiol*. 1991;134:250–256.

31. Kitayama J, Faraci FM, Lentz SR, et al. Cerebral vascular dysfunction during hypercholesterolemia. *Stroke*. 2007;38:2136–2141.