Differential Effect of B-Vitamin Therapy by Antiplatelet Use on Risk of Recurrent Vascular Events After Stroke

Baback Arshi, MD, MSc; Bruce Ovbiagele, MD, MSc, MAS; Daniela Markovic, MSc; Gustavo Saposnik, MD, MSc; Amytis Towfighi, MD

Background and Purpose—Although several randomized controlled trials failed to show a benefit of B vitamin therapy on composite outcomes of cardiovascular death, myocardial infarction, and stroke among individuals with elevated homocysteine, recent post hoc analyses have suggested that several factors may interact with the effects of vitamin treatment. One post hoc analysis revealed an interaction between B vitamin therapy and antiplatelet use; however, those results have not been replicated in other studies or populations.

Methods—We conducted a post hoc analysis of the Vitamin Intervention for Stroke Prevention (VISP) trial, a randomized controlled trial evaluating treatment with high- versus low-dose B vitamin therapy for secondary prevention of vascular events among stroke survivors with elevated homocysteine. Cox regression models were used to assess primary (recurrent stroke) and secondary (stroke, myocardial infarction, or vascular death) outcomes among individuals on high- versus low-dose vitamin therapy, categorized by antiplatelet use, after adjusting for covariates.

Results—Among 3680 participants, 52% took antiplatelet medications. When compared with low-dose therapy, high-dose vitamin therapy was associated with higher stroke risk among individuals on antiplatelets (hazard ratio, 1.43; 95% confidence interval, 1.02–2.01), but trended toward lower risk among those not on antiplatelets (hazard ratio, 0.86; 95% confidence interval, 0.62–1.19).

Conclusions—High-dose B vitamin therapy may be associated with a higher risk of recurrent stroke among stroke survivors taking antiplatelets, but does not have a significant effect on recurrent stroke risk in those who are not on antiplatelets. Future randomized controlled trials may consider evaluating the effect of homocysteine lowering among stroke survivors with elevated homocysteine who are not on antiplatelet therapy. (Stroke. 2015;46:870-873. DOI: 10.1161/STROKEAHA.114.006927.)

Key Words: antiplatelet agents ■ homocysteine ■ myocardial infarction ■ stroke ■ vitamin B complex
given to lower total homocysteine levels would reduce the incidence of recurrent cerebral infarction in patients with a nondisabling ischemic stroke versus low-dose vitamin therapy. The study design has been described in detail previously. Briefly, demographic, clinical, and laboratory data were collected at baseline, with subsequent clinical and laboratory information obtained at follow-up visits 6, 12, 18, and 24 months. Patients were enrolled after undergoing written informed consents. The trial was approved by the ethics committee at each participating center. The primary outcome was recurrent stroke and secondary outcome was stroke, MI, or death from a vascular cause.

**Statistical Analysis**

Multivariate cox regression models were used before and after adjusting for covariates including age, sex, previous stroke, diabetes mellitus, coronary artery disease, mean systolic blood pressure during the trial, congestive heart failure, alcohol use, carotid endarterectomy, smoking, body mass index, and stroke severity. Secondary analyses were performed with Cox regression models to assess whether there were any interactions between vitamin doses versus each of the other factors in predicting adverse vascular events among patients not using antiplatelet medications.

**Results**

Among 3680 participants in the VISP trial, 1907 (52%) individuals were taking antiplatelet medications. The group on antiplatelet medications was equally matched with the group not on antiplatelet medications with respect to stroke severity, demographic and vascular risk factors (Table 1). Subjects taking antiplatelets, however, were less likely to smoke, more likely to take antilipid medications, and more likely to have a history of stroke before the index stroke.

For the group taking antiplatelet medications, high-dose vitamin replacement therapy was associated with a higher risk of stroke when compared with low-dose vitamin therapy (hazard ratio [HR], 1.43; 95% confidence interval, 1.02–2.01; Table 2; Figure). On the contrary, among subjects not taking antiplatelet medications, there was a trend toward lower risk of stroke among those who received high-when compared with those who received low-dose vitamins (HR, 0.86; 95% confidence interval, 0.62–1.19; Table 2; Figure). High-dose vitamin therapy did not affect the odds of the secondary outcome of stroke, MI, or vascular death regardless of antiplatelet use.

The highest cumulative incidence of recurrent stroke occurred in subjects not taking antiplatelet medications on low-dose vitamin therapy (Figure). Conversely, the lowest cumulative incidence of recurrent stroke occurred in subjects taking antiplatelet medications and low-dose vitamin therapy. The risk of recurrent stroke was similar among (1) those taking antiplatelet medications and high-dose vitamin therapy and (2) those not taking antiplatelet medications but on high-dose vitamin therapy (Figure).

**Discussion**

In this retrospective analysis of the VISP trial, we found that high-dose B vitamin therapy was associated with a higher rate of recurrent stroke among individuals on antiplatelet therapy. Individuals not on antiplatelet therapy, however, tended to have a lower risk of recurrent stroke on high-dose B vitamin therapy.

The findings in this study corroborate the post hoc analysis of the VITATOPS trial. In VITATOPS, they found that for participants taking antiplatelets at baseline, B vitamins did not affect the primary outcome of stroke, MI, or death from vascular cause (HR, 0.94; 95% confidence interval, 0.83–1.07). However, for the participants not taking antiplatelet drugs at baseline, B vitamins did have a significant effect on the primary outcome (HR, 0.76; 95% confidence interval, 0.60–0.96).

These findings have 2 potential ramifications. First, if patients with elevated homocysteine have a contraindication to antiplatelet therapy, they may potentially benefit from taking high-dose B vitamin replacement therapy. Second, high-dose B vitamin therapy may need to be used with caution in those on antiplatelet medications given the potentially deleterious effects.

The cause for the interaction between B vitamins and antiplatelet medications remains unclear, but may be related to the roles of these compounds at the endothelium. At the endothelium, homocysteine inhibits nitric oxide, regulates prostanooids, activates endothelin-1, activating angiotensin II receptors, and causing oxidative stress. By lowering homocysteine, B vitamins may reduce endothelial damage. Antiplatelets inhibit platelet aggregation at the endothelium. Additional studies and scientific models are needed to elucidate the interaction between antiplatelets and homocysteine.

### Table 1. Descriptive Summary Statistics of Patients in VISP Trial (n=3680)

<table>
<thead>
<tr>
<th>Value</th>
<th>No Antiplatelet Use</th>
<th>Concurrent Antiplatelet Use</th>
</tr>
</thead>
<tbody>
<tr>
<td>Avg age, y</td>
<td>65.7±0.4 66.1±0.4</td>
<td>66.7±0.3 66.6±0.3</td>
</tr>
<tr>
<td>Sex, men, %</td>
<td>61.7 59.8</td>
<td>63.8 64.6</td>
</tr>
<tr>
<td>Race, %</td>
<td>White 75.4 77.9</td>
<td>Black 17.3 16.6</td>
</tr>
<tr>
<td></td>
<td>Other 7.3</td>
<td>Hypertension, % 72.9 73.7</td>
</tr>
<tr>
<td></td>
<td>Diabetes mellitus, % 28.6 26</td>
<td>Smoking, % 18.3 19.1</td>
</tr>
<tr>
<td></td>
<td>Antilipid use, % 35.3 36</td>
<td>Previous stroke, % 17.9 15.9</td>
</tr>
<tr>
<td>NIHSS*</td>
<td>0</td>
<td>Avg systolic BP, mmHg 140.7 142.5</td>
</tr>
<tr>
<td></td>
<td>1–4</td>
<td>Avg LDL† 127.2 124.9</td>
</tr>
<tr>
<td></td>
<td>≥5</td>
<td>Avg total cholesterol† 207.9 203.2</td>
</tr>
<tr>
<td>BP indicates blood pressure; LDL, low-density lipoprotein; NIHSS, National Institutes of Health Stroke Scale; and VISP, Vitamin Intervention for Stroke Prevention.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>NIHSS as a marker for stroke severity.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>†Cholesterol measured in mg/dL.</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

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This study has limitations. First, because this was a post hoc analysis of a clinical trial, it was not adequately powered for this subgroup analysis; therefore, any interactions seen could be because of chance. Second, the trial was performed in North America and Scotland during a time where folate fortification occurred. This lowered the baseline serum homocysteine and altered potential benefits of supplementing with B vitamins. A recent meta-analysis suggested that there is more likely to be risk reduction in trials lowering homocysteine when the baseline homocysteine is decreased by >20%.15 Third, because stroke subtypes were not predefined in this trial, we were not able to determine whether the group on antiplatelet medications was equally matched with the group not on antiplatelet medications with respect to stroke subtype. There remains a possibility that the interaction between vitamin replacement therapy and antiplatelet use occurred as a result of unbalanced stroke subtypes. Fourth, we were not able to differentiate between type and dose of antiplatelet used for all individuals during the trial. Fifth, as mentioned, the elevated stroke risk among individuals taking antiplatelets combined with high-dose B vitamins has not been noted previously and may be a spurious finding, given the wide confidence interval. Finally, the results of this study are not generalizable to all patients with stroke because the trial included individuals with only nondisabling, noncardioembolic stroke. Nevertheless, the strengths of the study include its rigorous trial design and detailed information on covariates.

Given the negative findings in 3 large RCTs, subgroup analyses have suggested that certain subgroups of individuals, such as younger individuals or those living in areas without folate fortification, may benefit from homocysteine lowering after stroke. The findings from this study, combined with the post hoc analysis from VITATOPS, suggest that individuals with contraindications to antiplatelet therapy may be another subgroup worthy of investigation in prospective RCTs of homocysteine lowering.

Table 2. Association Between Vitamin Therapy and Primary and Secondary Outcomes by Antiplatelet Use

<table>
<thead>
<tr>
<th>Outcome</th>
<th>Group</th>
<th>Vitamin Dose</th>
<th>Unadjusted Model</th>
<th>P Value</th>
<th>Adjusted Model*</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Stroke†</td>
<td>Not on antiplatelet</td>
<td>Low</td>
<td>Ref</td>
<td>…</td>
<td>Ref</td>
<td>…</td>
</tr>
<tr>
<td></td>
<td></td>
<td>High</td>
<td>0.81 (0.59–1.12)</td>
<td>0.206</td>
<td>0.86 (0.62–1.19)</td>
<td>0.365</td>
</tr>
<tr>
<td></td>
<td>On antiplatelet</td>
<td>Low</td>
<td>Ref</td>
<td>…</td>
<td>Ref</td>
<td>…</td>
</tr>
<tr>
<td></td>
<td></td>
<td>High</td>
<td>1.35 (0.98–1.88)</td>
<td>0.070</td>
<td>1.43 (1.02–2.01)</td>
<td>0.038</td>
</tr>
<tr>
<td>Stroke, MI, or vascular death‡</td>
<td>Not on antiplatelet</td>
<td>Low</td>
<td>Ref</td>
<td>…</td>
<td>Ref</td>
<td>…</td>
</tr>
<tr>
<td></td>
<td></td>
<td>High</td>
<td>0.85 (0.66–1.09)</td>
<td>0.204</td>
<td>0.90 (0.70–1.17)</td>
<td>0.449</td>
</tr>
<tr>
<td></td>
<td>On antiplatelet</td>
<td>Low</td>
<td>Ref</td>
<td>…</td>
<td>Ref</td>
<td>…</td>
</tr>
<tr>
<td></td>
<td></td>
<td>High</td>
<td>1.16 (0.90–1.50)</td>
<td>0.264</td>
<td>1.18 (0.90–1.54)</td>
<td>0.237</td>
</tr>
</tbody>
</table>

CI indicates confidence interval; HR, hazard ratio; and MI, myocardial infarction.

*Adjusted for age, sex, previous stroke, diabetes mellitus, coronary artery disease, mean systolic blood pressure, congestive heart failure, alcohol use, carotid endarterectomy, smoking, and stroke severity.

†Antiplatelet×vitamin dose interaction: P=0.038.

‡Antiplatelet×vitamin dose interaction: P=0.157.
Disclosures
None.

References
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