Hyperhomocysteinemia and vascular access thrombosis in hemodialysis patients: a retrospective study

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Background: Elevated total plasma homocysteine is an independent risk factor for arterial and venous thrombosis in patients with normal renal function. Patients on hemodialysis have a high prevalence of mild to moderate hyperhomocysteinemia. Conflicting retrospective analyses and prospective studies have been reported regarding the association between total homocysteine levels and hemodialysis vascular thrombosis. The purpose of this retrospective study was to investigate the relationship between hyperhomocysteinemia and vascular access thrombosis (VAT) in patients on hemodialysis.

Methods: One hundred and twenty-five patients undergoing dialysis were selected as subjects. The experimental group participants were identified as those having one or more VAT during the previous 13 months and the control group participants had no access thrombosis during the same period. Additional subgroup analysis included the presence of hypertension, diabetes, low-density lipoprotein levels, sex, and use of aspirin.

Results: No statistically significant difference was found in total homocysteine levels between the two groups (P = 0.27). No association was found between VAT and sex (P = 0.09), VAT and hypertension (P = 0.96), VAT and diabetes (P = 0.49), nor VAT and low-density lipoprotein level (P = 0.04). A lower rate of VAT was associated with aspirin intake (P = 0.04).

Conclusion: This study did not demonstrate a relationship between total homocysteine concentrations and risk of VAT in patients with end-stage renal disease on hemodialysis. There were no significant differences in the number of VAT across additional variables of sex and previous morbidity. Aspirin intake was associated with a lower incidence of VAT.

Keywords: hyperhomocysteinemia, vascular access thrombosis, hemodialysis

Introduction
Vascular access failure is a major cause of morbidity and mortality in hemodialysis patients. Little is understood about the risk factors associated with vascular access thrombosis (VAT). Currently, placement of a synthetic graft and central venous catheter rather than a native arteriovenous fistula represents the better defined risk factor for vascular access failure. Other risk factors identified in single studies include location of the graft, time of use of graft after surgical creation, diabetes, age older than 65 years, hypoalbuminemia, elevated lipoprotein-a level, lupus anticoagulant, female sex, elevated low-density lipoprotein (LDL) cholesterol, alterations in genes regulating the coagulation cascade, and stenosis within the graft and draining vein. Hyperhomocysteinemia has been reported as a risk factor for VAT. An elevated total plasma homocysteine level is associated with premature atherosclerosis, adversely affecting the endothelium by damaging cells, inhibiting anticoagulants, enhancing
procoagulants, and impairing the bioavailability of the vasodilator, nitric oxide. Hyperhomocysteinemia is an independent risk factor for arterial and venous thrombosis in patients with normal renal function. While there is an excess prevalence of mild to moderate hyperhomocysteinemia in hemodialysis patients, the association with VAT has not been universally confirmed. Conflicting retrospective analysis and prospective studies have been reported regarding the association between total homocysteine levels and hemodialysis vascular thrombosis.

The purpose of this retrospective study was to determine if a significant association exists between hyperhomocysteinemia, age, sex, previous morbidity (diabetes and hypertension), and hemodialysis access thrombosis.

Materials and methods

One hundred and twenty-five patients undergoing dialysis were selected as subjects from two different dialysis clinics. Patients with a history of antiphospholipid syndrome or on anticoagulation were excluded from the study. Information was obtained from their medical files and included the following for the past 13 months: number of VAT, age, sex, past medical history, duration of hemodialysis, mean LDL level, mean albumin level, mean ferritin level, and aspirin intake. Each patient included in the study was assigned a number to maintain anonymity of the patient and to preserve confidentiality. The plasma homocysteine level was checked once after obtaining informed consent from the patients. The sample was divided into an experimental (VAT, n = 53) group and control (no VAT, n = 72) group. Participants in the experimental group were identified as those having one or more VAT during the previous 13 months (December 2003 to January 2005) and participants in the control group were those with no access thrombosis during the same period. Additional subgroup analysis included: presence of hypertension, diabetes, LDL level, sex, (male versus female), and use of aspirin.

Statistical analysis

The data were analyzed using the chi-square test for categorical variables and t-test for interval data. Statistical significance was set at $P = 0.05$. Logistic regression was used to examine the relationship between the model of dependent variables and the outcome variable of VAT.

Results

No significant difference in total homocysteine levels was found between the VAT group and control group, with mean values being $25.2 \pm 8.38$ and $25.69 \pm 1.66$, respectively ($P = 0.27$). No significant association was found between VAT and hypertension ($P = 0.96$), VAT and diabetes ($P = 0.49$), VAT and LDL level ($P = 0.40$), nor VAT and sex ($P = 0.09$, see Table 1). The only variable that contributed to explaining the model was aspirin intake, with a $P$-value of 0.04 when considered with the effect of the other variables removed. Aspirin intake had a relative risk of 0.74 (0.49–1.13). This indicated a reduction in risk for access thrombosis of 26%.

Discussion

Elevated total homocysteine concentrations are common in patients with end-stage renal disease on hemodialysis. The etiology of high homocysteine levels is not well understood. In addition to the reduced clearance of plasma homocysteine observed, genetic and dialysis-related factors may be involved. Hyperhomocysteinemia is a well established risk factor for accelerated atherogenesis and coronary heart disease. It is also associated with recurrent venous thrombosis in patients without end-stage renal disease.

However, less clear is the relationship between hyperhomocysteinemia and VAT in patients on hemodialysis (Table 2) which is the most common cause of hospitalization among these patients. Mallamaci et al reported a clear association between high homocysteine levels and cardiovascular mortality and atherothrombotic events in hemodialysis patients. In another prospective study of 78 patients, Mallamaci et al found that VAT in dialysis patients is associated with hyperhomocysteinemia.

Table 1 Demographic characteristics of study patients

<table>
<thead>
<tr>
<th></th>
<th>VAT, n = 53</th>
<th>No VAT, n = 72</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, years</td>
<td>61.30 ± 15.7</td>
<td>62.84 ± 13.1</td>
<td>0.55</td>
</tr>
<tr>
<td>Sex, M/F</td>
<td>30/23</td>
<td>51/21</td>
<td>0.09</td>
</tr>
<tr>
<td>Homocysteine level (μmol/L)</td>
<td>25.2 ± 8.38</td>
<td>25.69 ± 16.6</td>
<td>0.27</td>
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<tr>
<td>LDL level</td>
<td>67.1 ± 28.42</td>
<td>65.86 ± 28.14</td>
<td>0.40</td>
</tr>
<tr>
<td>Ferritin level (mg/dL)</td>
<td>761.5 ± 435</td>
<td>766 ± 401</td>
<td>0.47</td>
</tr>
<tr>
<td>Hypertension (μmol/L)</td>
<td>44 (83%)</td>
<td>60 (83.3%)</td>
<td>0.96</td>
</tr>
<tr>
<td>Diabetes mellitus</td>
<td>24 (60.7%)</td>
<td>37 (39.3%)</td>
<td>0.49</td>
</tr>
<tr>
<td>Coronary artery disease</td>
<td>35 (63.6%)</td>
<td>20 (36.4%)</td>
<td>0.22</td>
</tr>
</tbody>
</table>

Abbreviations: LDL, low-density lipoprotein; VAT, vascular access thrombosis.
relationship between homocysteine and VAT.17 Manns et al found no association between homocysteine levels or anti-cardiolipin antibody and VAT in a cross-sectional study involving 118 patients.2 Similarly, the studies reported by Hojs et al and Sirrs et al demonstrated no significant relationship between homocysteine levels and VAT.18,19

Our retrospective study indicates that the total homocysteine level is not a valid marker for VAT. Additionally, we did not discover any trend towards increased total homocysteine levels and VAT. Hypertension and diabetes are associated with premature vascular disease; however, we did not observe this relationship in our study, which is in agreement with the findings of Bowden et al15 and Bednarek-Skublewksa et al.14 Our analysis confirmed the findings of other investigators that the use of antiplatelet agents is associated with a reduced risk of VAT.19–21

In summary, we found no association between total homocysteine concentration and VAT in patients on hemodialysis. Because of the complexity of the relationship between homocysteine and vascular access failure, long-term prospective studies with serial measurement of homocysteine levels may better address the direct metabolic implications of elevated total homocysteine in hemodialysis patients.

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

References