Betaine and Folate Status as Cooperative Determinants of Plasma Homocysteine in Humans

Pål I. Holm, Per M. Ueland, Stein Emil Vollset, Øivind Midttun, Henk J. Blom, Miranda B.A.J. Keijzer, Martin den Heijer

Objective—Two published studies have demonstrated that betaine in the circulation is a determinant of plasma total homocysteine, but none had sufficient power to investigate the possible effect modification by folate status.

Methods and Results—We measured homocysteine, betaine, folate, vitamin B₆, and related compounds in serum/plasma from 500 healthy men and women aged 34 to 69 years before (fasting levels) and 6 hours after a standard methionine loading test. Choline, dimethylglycine, and folate were determinants of plasma betaine in a multiple regression model adjusting for age and sex. The increase in homocysteine after loading showed a strong inverse association with plasma betaine and a weaker inverse association with folate and vitamin B₆. Fasting homocysteine showed a strong inverse relation to folate, a weak relation to plasma betaine, and no relation to vitamin B₆. Notably, adjusted (for age and sex) dose-response curves for the postmethionine increase in homocysteine or fasting homocysteine versus betaine showed that the inverse associations were most pronounced at low serum folate, an observation that was confirmed by analyses of interaction.

Conclusions—Collectively, these results show that plasma betaine is a strong determinant of increase in homocysteine after methionine loading, particularly in subjects with low folate status. (Arterioscler Thromb Vasc Biol. 2005;25:379-385.)

Key Words: betaine ■ choline ■ homocysteine ■ folate ■ methionine
on PML tHcy.9 Notably, low betaine doses in the range of dietary intake have been shown recently to reduce fasting and, in particular, PML tHcy.11

There are few association studies on endogenous betaine in serum/plasma. An inverse association between tHcy and betaine was reported recently in 120 cardiovascular patients,12 and plasma betaine has been shown to be a strong predictor of PML tHcy in 90 patients enrolled in a B vitamin intervention trial.13

We conducted a large study of 500 subjects undergoing methionine loading to investigate the relationship between plasma betaine and basal and PML tHcy and the possible effect modification by folate status.

Methods

Subjects and Protocol
Subjects were recruited through a general practice in The Hague, Holland. These subjects were healthy controls in a study on homocysteine and venous thrombosis. Details on recruitment, inclusion criteria, and data collection have been published previously.14

A 6-hour standard oral methionine loading test (0.1 g L-methionine per kg body weight in 200 mL orange juice) was performed as described.14 The study was approved by the ethics committee of the Leyenburg Hospital, The Hague, Holland. These subjects were healthy controls in a study on homocysteine and venous thrombosis. Details on recruitment, inclusion criteria, and data collection have been published previously.14

Blood Collection and Biochemical Analyses
The EDTA plasma was stored at −20°C until analysis and serum kept at −70°C. Plasma tHcy,15 serum creatinine, serum vitamin B12, serum folate,14 vitamin B6 (sum of pyridoxal 5'-phosphate and pyridoxal),14 betaine, choline, and dimethylglycine (DMG)15 were determined with published methods.

Statistics
Data are presented as medians with 10th to 90th percentiles. Between-group comparisons of continuous variables were done by the Mann–Whitney U test. Spearman rank correlation and multiple regression analyses were used to evaluate associations between individual variables. Multiple linear regression analysis was used to assess the simultaneous relationship between various predictors of tHcy. Plasma tHcy was the dependent variable, whereas the independent variables were presented in the model as quartiles of betaine, folate, cobalamin, and creatinine. Thus, the regression coefficient was used to estimate the difference in mean tHcy between the reference and the other 3 quartiles. tHcy across quartiles was tested for homogeneity of means and for linear trend. Estimates were adjusted for age and sex, or B vitamin levels and creatinine, in addition to age and sex. We investigated the possible interaction between plasma betaine and folate and between betaine and vitamin B6 by including a product term between the 2 variables in multiple linear regression models with tHcy as the dependent variable, retaining betaine and the B vitamin as independent variables. Because tHcy values (fasting and the increase after methionine loading) were not normally distributed, the multiple regression analyses, when appropriate, were also carried with log-transformed tHcy as outcome measures. The dose-response relationships between metabolites were also estimated with Gaussian generalized additive models (GAM),18 as implemented in S-PLUS and R.19 For other analyses, we used SPSS version 11.0 (SPSS).

Results

Subject Characteristics
A total of 500 subjects (292 females and 208 males) with a mean age of 50 years (range 34 to 69) were investigated. Their blood indices before and after methionine loading and according to gender are given in Table 1. Fasting plasma tHcy, betaine, choline, DMG, methionine, creatinine and vitamin B6, and PML betaine, choline, and DMG were all significantly higher in men than in women.

The distribution of plasma betaine in terms of median, 10th to 90th percentiles, and range were 30.3, 18.8 to 45.3, and 9.4 to 94.9 μmol/L. The corresponding values for serum folate were 12.7, 7.0 to 23.6, and 3.0 to 54.4 nmol/L, respectively. Median serum cobalamin was 217 pmol/L.

Loading caused a 20-fold increase in overall median plasma methionine and a 4-fold increase in tHcy. Loading was not associated with any change in median betaine, choline, or DMG (Table 1).

Bivariate Correlations
Spearman rank correlation coefficients are listed in supplemental Table 1 (available online at http://atvb.ahajournals.org). PML ΔtHcy was weakly and inversely related to the folate, cobalamin, and vitamin B6 but showed a moderate inverse relation to fasting (r = −0.27) and PML betaine (r = −0.32; all P < 0.001). Fasting tHcy showed a moderate positive relation to age (r = 0.30) and creatinine (r = 0.37), an inverse relation to folate (r = −0.29) and cobalamin (r = 0.27; all P < 0.001), and a weak but significant positive relation to choline and DMG.

Betaine (fasting and PML) was strongly and positively related to choline and DMG (r = 0.41 to 0.50; P < 0.001) and showed a weak but significant positive relation to folate and vitamin B6 (r = 0.18 to 0.22; P < 0.001).
Folate, Choline, and DMG as Determinants of Betaine

We investigated the relationship between betaine and indices showing a simple correlation with betaine, such as folate, choline, DMG, creatinine, vitamin B₆, and age, using a Gaussian generalized additive regression (GAM), which produces dose-response curves adjusted for age, sex, and other parameters (Figure 2). Serum folate, choline, and DMG showed a positive relation to betaine, but the curve for choline leveled off at 14 μmol/L (99.5 percentile) and that for DMG at 4 μmol/L (80 percentile; Figure 2). Betaine showed no relation to vitamin B₆ and was inversely related to creatinine in this regression model. The associations obtained by GAM were essentially in agreement with those obtained by multiple linear regression (Figure 2, legend).

Estimated Change in PML ΔtHcy and Fasting tHcy by Betaine and Other Determinants

Determinants of ΔtHcy were estimated by multiple regression analysis (Table 2). For all variables, we estimated the difference in mean ΔtHcy between each quartile and the reference quartile. The mean tHcy difference across the extreme quartiles adjusted for age and sex was highest (8.3 μmol/L) for betaine, intermediate (5.7 μmol/L; data not shown) for creatinine, and moderate (4.3 to 4.9 μmol/L) for folate, cobalamin, and vitamin B₆; all P values were <0.001. The betaine–tHcy relationship remained strong (P<0.001) after additional adjustment for all blood indices (including folate and vitamin B₆), whereas multiple adjustments weakened the associations of tHcy with folate, cobalamin, and vitamin B₆; all P values were <0.001. Additional adjustment for interaction between betaine and vitamin B₆ in this regression model demonstrated no interaction between betaine and vitamin B₆ (P for interaction=0.8).

We performed the same calculation of the differences in mean fasting tHcy between quartiles of betaine, folate, cobalamin, vitamin B₆, and creatinine (Table 2). The tHcy difference across the extreme quartiles adjusted for age and sex was moderate (2.2 μmol/L) for betaine, weak (0.6 μmol/L) for vitamin B₆, intermediate (2.6 μmol/L) for cobalamin, and highest (3.7 to 3.9 μmol/L) for folate and creatinine; all P values were <0.001. Additional adjustment for all blood indices reduced the tHcy difference across the extreme betaine quartiles to 0.9 μmol/L, which now became of borderline significance (P=0.07). This adjustment had essentially no effect on the tHcy change across the quartiles.
### TABLE 2. Estimated Change in PML Increase in tHcy and Fasting tHcy According to Quartiles of Betaine and B-Vitamins

<table>
<thead>
<tr>
<th>Blood Indices</th>
<th>Adjusted for Age and Sex</th>
<th>Adjusted for all Parameters*</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Change in PML Increase in tHcy, μmol/L (Mean [95% CI])</td>
<td>Change in Fasting tHcy, μmol/L (Mean [95% CI])</td>
</tr>
<tr>
<td>Betaine (μmol/L)</td>
<td>P&lt;0.001†</td>
<td>P&lt;0.001</td>
</tr>
<tr>
<td>vs &gt;37.0, n=125</td>
<td>30.4–37.0 (n=125)</td>
<td>3.2 (0.2–6.2)</td>
</tr>
<tr>
<td>24.8–30.3 (n=126)</td>
<td>6.4 (3.3–9.4)</td>
<td>6.0 (2.9–9.0)</td>
</tr>
<tr>
<td>&lt;24.8 (n=123)</td>
<td>8.3 (5.0–11.6)</td>
<td>7.2 (3.9–10.6)</td>
</tr>
<tr>
<td>Folate (nmol/L)</td>
<td>P=0.001</td>
<td>P=0.03</td>
</tr>
<tr>
<td>vs &gt;18.1, n=125</td>
<td>12.8–18.1 (n=127)</td>
<td>2.1 (−0.9–5.1)</td>
</tr>
<tr>
<td>9.4–12.7 (n=127)</td>
<td>3.9 (0.9–6.9)</td>
<td>1.8 (−1.2–4.7)</td>
</tr>
<tr>
<td>&lt;9.4 (n=120)</td>
<td>4.9 (1.8–7.9)</td>
<td>3.2 (0.2–6.2)</td>
</tr>
<tr>
<td>Cobalamin (pmol/L)</td>
<td>P&lt;0.001</td>
<td>P&lt;0.001</td>
</tr>
<tr>
<td>vs &gt;291, n=125</td>
<td>218–291 (n=124)</td>
<td>3.6 (0.6–6.6)</td>
</tr>
<tr>
<td>168–217 (n=126)</td>
<td>4.5 (1.5–7.5)</td>
<td>3.7 (0.8–6.6)</td>
</tr>
<tr>
<td>&lt;168 (n=124)</td>
<td>4.3 (1.3–7.3)</td>
<td>3.2 (0.3–6.1)</td>
</tr>
<tr>
<td>Vitamin B6 (nmol/L)</td>
<td>P&lt;0.001</td>
<td>P&lt;0.001</td>
</tr>
<tr>
<td>vs &gt;37.4, n=125</td>
<td>27.7–37.4 (n=125)</td>
<td>1.3 (−1.6–4.3)</td>
</tr>
<tr>
<td>20.8–27.7 (n=125)</td>
<td>4.5 (1.5–7.6)</td>
<td>3.2 (0.2–6.2)</td>
</tr>
<tr>
<td>&lt;20.8 (n=125)</td>
<td>4.9 (1.9–8.0)</td>
<td>3.7 (0.7–6.8)</td>
</tr>
</tbody>
</table>

*All parameters (betaine, folate, cobalamin, vitamin B6, creatinine, age, and sex) are included in the model.
†P for trend.
PML indicates postmethionine load.
of folate, cobalamin, and creatinine but attenuated the tHcy–vitamin B6 relationship (Table 2). Again, essentially the same associations with betaine (P=0.03), folate (P>0.001), cobalamin (P>0.001), vitamin B6 (P=0.8), and creatinine (P>0.001) were obtained after log transformation of tHcy (data not shown).

Additional adjustment for smoking, alcohol, and body weight did not materially change the estimates (data not shown).

**Effect Modification of the PML ΔtHcy: Betaine Relationship by Folate Status**

We used GAM to estimate the dose-response relationship between ΔtHcy and betaine, adjusted for cobalamin, vitamin B6, creatinine, sex, and age. The estimation was done separately in each tertile of plasma folate. In the lowest folate tertile, there was a strong negative relationship between ΔtHcy and betaine. In the top 2 folate tertiles, there was a weaker negative association (Figure 3). Accordingly, we observed an interaction between betaine and folate of borderline significance (P=0.05) by multiple regression (adjusting for cobalamin, vitamin B6, creatinine, sex, and age), with log-transformed ΔtHcy as the outcome parameter.

Adjusted dose-response curves for fasting tHcy versus betaine for each tertile of serum folate were generated by the same procedure. In the lower tertile, there was an inverse association between tHcy and betaine, whereas in the top 2 tertiles, essentially no such association was observed (Figure 3). A significant interaction between betaine and folate (P=0.01) was observed by multiple regression with log-transformed fasting tHcy as the outcome parameter.

**Discussion**

We investigated the role of circulating plasma betaine as a determinant of fasting and PML tHcy in 500 healthy subjects undergoing methionine loading. Plasma betaine was a stronger predictor of the increase in PML tHcy (ΔtHcy) than other parameters investigated, including folate, vitamin B6, and cobalamin, and the inverse association was particularly pronounced in subjects with low serum folate. We also observed that plasma betaine was a predictor of fasting tHcy, but only at low folate status.

**Study Design**

The strength of this study is the number of subjects included, which is high considering the resources and logistics required to perform methionine loading. The logistics are even more complicated for methionine testing, with a sampling interval of 6 hours used in the present study compared with short 2- or 4-hour tests. However, the 6-hour test has been recommended because the homocysteine response has lower within-subject variability than for the short tests. This has been attributed to variable rate of methionine absorption.

**Plasma Levels and Determinants of Betaine, Choline, and DMG**

The median concentrations of betaine (30.3 μmol/L), choline (7.8 μmol/L), and DMG (3.1 μmol/L) determined for this study population of healthy adult men and women were similar to the concentration of betaine,21,23 choline,17,22,23 and DMG17,21,24 reported previously by us and others.

The size of the study population investigated here allowed detailed assessment of predictors of betaine in regression model, including sex, age, and all blood indices (Figure 2). Choline showed a linear relation to betaine and was the strongest metabolic predictor of betaine, which could be explained by choline being the immediate precursor. Folate also showed a linear relation to betaine, suggesting common dietary sources25 or a mutual sparing effect. Finally, the initial linear relationship between DMG and betaine at low DMG may reflect DMG production from betaine, whereas the plateau phase could be attributable to product inhibition of BHMT by DMG.26

**Betaine as Determinant of PML and Fasting tHcy**

This study demonstrated that betaine is a strong determinant of PML ΔtHcy. This effect was only slightly reduced after multiple adjustments (including folate and vitamin B6; Table 2). The observation that betaine is a strong determinant of PML ΔtHcy confirms similar results from a recent small study13 and is in agreement with consistent reports that oral intake of betaine markedly reduces PML tHcy.9,11

We also observed a significant relationship between betaine and fasting tHcy, which became of borderline significance after multiple adjustments, including folate (Table 2). The weak overall association between betaine and fasting tHcy is consistent with such association in 120 cardiovascular patients12 and the observation of moderate reduction in fasting tHcy by betaine supplementation.5,9,11

**Vitamin B6**

We measured vitamin B6 in this study because of the prevailing view that vitamin B6 status is an important determinant of PML tHcy. This idea is based mainly on experiments with vitamin B6-deficient rats27 and studies of subjects with low vitamin B6 status.28 However, in most studies of
humans without overt deficiency, vitamin B_{6} is not related to PML tHcy or is a weaker determinant than folate. In a previous study on cardiovascular patients, we observed no association between vitamin B_{6} and tHcy (fasting and increase after loading). The present study demonstrates that in healthy subjects, vitamin B_{6} is not a predictor of fasting tHcy and is a weaker determinant of the PML increase in tHcy than betaine (Table 2).

**Effect Modification by Folate Status**

We observed that betaine was a strong predictor of PML ΔtHcy in subjects with low serum folate and a weaker predictor at high folate (Figure 3). The effect modification by folate status is the most notable finding in the present study. It elaborates the preliminary observation demonstrating attenuation of the betaine–PML ΔtHcy relationship in subjects supplemented with combinations of B vitamins (cobalamin, folate, and vitamin B_{6}).

We also observed that the weak overall association between betaine and fasting tHcy (Table 2) was attributable to such association in subjects with serum folate in the lower tertile (Figure 3). We are pursuing latter observation in a large ongoing study of the betaine as a determinant of tHcy in folate-deficient subjects.

**Mechanisms**

The strong association between betaine and PML ΔtHcy at low folate suggests increased catalytic activity of the BHMT at low 5-methyltetrahydrofolate, which is supported by measurement of rat liver enzyme activity and by a mathematical model, based on known enzyme kinetics. The increased catalytic activity is accomplished by increasing homocysteine availability and by lowering S-adenosylmethionine, which relieves S-adenosylmethionine-mediated BHMT inhibition.

An important message from this and a previous work, as well as from studies of betaine supplementation, is that betaine reduces tHcy under conditions of high methionine. This seems to be in conflict with the prevailing view based on rat and chick experiments, suggesting that BHMT conserves homocysteine under conditions of methionine deficiency. However, in pigs, BHMT activity in liver increases in response to methionine. Thus, in some species at least, including humans, homocysteine accumulating during folate deficiency may be directed into the BHMT pathway, even in the presence of superfluous methionine.

Betaine is more strongly associated with PML ΔtHcy than with fasting tHcy. This may indicate a role for betaine in regulating postprandial homocysteine status. However, the PML tHcy probably reflects a massive first-pass homocysteine export from the liver after uptake of excess methionine. One may speculate whether high betaine directs the first-pass homocysteine metabolism after loading into betaine-dependent remethylation, thereby reducing homocysteine export and the resulting increase in plasma tHcy. In fasting subjects, betaine may influence homocysteine status in the liver to a larger extent than is reflected by fasting tHcy, which is mainly determined by the activity of ubiquitous methionine synthase, and therefore by overall folate status.

**Implications and Conclusion**

Folate, in addition to choline and DMG, is a major predictor of plasma betaine, which, in turn, is the strongest determinant of PML increase in tHcy hitherto recognized. Notably, the betaine–tHcy relationship is particularly pronounced at low folate status. The plasma betaine varies substantially (ie, 10-fold [from 9.4 to 94.9 μmol/L] between individuals), but a recent study demonstrates that the intraindividual variability is small, with an individual set point that remains stable for years. This suggests that plasma betaine is under strict metabolic control and justifies the concept of betaine status as a component of an individual’s biochemical make-up with ramifications to one carbon metabolism. Betaine status should be investigated in pathologies related to altered metabolism of homocysteine and folate, including cardiovascular disease and cancer.

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**References**


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TABLE I. Bivariate Associations in Terms of Spearman Correlation Coefficients

<table>
<thead>
<tr>
<th></th>
<th>Age</th>
<th>Cr</th>
<th>ΔPML tHcy</th>
<th>Betaine</th>
<th>PML betaine</th>
<th>Choline</th>
<th>DMG</th>
<th>PML Met</th>
<th>Folate</th>
<th>Cbl</th>
<th>B6</th>
</tr>
</thead>
<tbody>
<tr>
<td>tHcy</td>
<td>0.30‡</td>
<td>0.37‡</td>
<td>0.28‡</td>
<td>-0.03</td>
<td>-0.03</td>
<td>0.11*</td>
<td>0.15†</td>
<td>0.05</td>
<td>-0.29‡</td>
<td>-0.27‡</td>
<td>-0.08</td>
</tr>
<tr>
<td>ΔPML tHcy</td>
<td>0.07</td>
<td>0.02</td>
<td>-0.28‡</td>
<td>-0.32‡</td>
<td>-0.08</td>
<td>-0.09*</td>
<td>0.05</td>
<td>-0.13†</td>
<td>-0.15†</td>
<td>-0.17‡</td>
<td>-0.17‡</td>
</tr>
<tr>
<td>Betaine</td>
<td>0.13†</td>
<td>0.27‡</td>
<td>0.88‡</td>
<td>0.50‡</td>
<td>0.44‡</td>
<td>0.17‡</td>
<td>0.21‡</td>
<td>0.06</td>
<td>0.22‡</td>
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</tr>
<tr>
<td>PML betaine</td>
<td>0.13†</td>
<td>0.27‡</td>
<td></td>
<td>0.41‡</td>
<td>0.47‡</td>
<td>0.32‡</td>
<td>0.18‡</td>
<td>0.06</td>
<td>0.21‡</td>
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<tr>
<td>Choline</td>
<td>0.31‡</td>
<td>0.24‡</td>
<td>0.36‡</td>
<td>0.28</td>
<td>0.12†</td>
<td>0.00</td>
<td>0.06</td>
<td></td>
<td></td>
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<tr>
<td>DMG</td>
<td>0.07</td>
<td>0.27‡</td>
<td></td>
<td>0.20‡</td>
<td>-0.09*</td>
<td>0.03</td>
<td>0.03</td>
<td></td>
<td></td>
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</tr>
</tbody>
</table>

*p<0.05
†p<0.01
‡p<0.001

B6, vitamin B6; Cbl, cobalamin; Cr, creatinine; ΔPML, post methionine load increase; DMG, dimethylglycine; Met, methionine; PML, post-methionine load; tHcy, total homocysteine.