Comparison of (6S)-5-methyltetrahydrofolic acid v. folic acid as the reference folate in longer-term human dietary intervention studies assessing the relative bioavailability of natural food folates: comparative changes in folate status following a 16-week placebo-controlled study in healthy adults

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Folic acid (pteroylmonoglutamic acid) has historically been used as the reference folate in human intervention studies assessing the relative bioavailability of dietary folate. Recent studies using labelled folates indicated different plasma response kinetics to folic acid than to natural (food) folates, thus obviously precluding its use in single-dose experiments. Since differences in tissue distribution and site of biotransformation were hypothesised, the question is whether folic acid remains suitable as a reference folate for longer-term intervention studies, where the relative bioavailability of natural (food) folate is assessed based on changes in folate status. Healthy adults aged 18–65 years (n 163) completed a 16-week placebo-controlled intervention study in which the relative bioavailability of increased folate intake (453 nmol/d) from folate-rich foods was assessed by comparing changes in plasma and erythrocyte folate concentration with changes induced by an equal reference dose of supplemental (6S)-5-methyltetrahydrofolic acid or folic acid. The relative increase in plasma folate concentration in the food group was 31 % when compared with that induced by folic acid, but 39 % when compared with (6S)-5-methyltetrahydrofolic acid. The relative increase in erythrocyte folate concentration in the food group when compared with that induced by folic acid was 43 %, and 40 % when compared with (6S)-5-methyltetrahydrofolic acid. When recent published observations were additionally taken into account it was concluded that, in principle, folic acid should not be used as the reference folate when attempting to estimate relative natural (food) folate bioavailability in longer-term human intervention studies. Using (6S)-5-methyltetrahydrofolic acid as the reference folate would avoid future results’ validity being questioned.

Human nutrition: (6S)-5-methyltetrahydrofolic acid: Folic acid: Plasma homocysteine: Folate: Bioavailability: Intervention studies

Low folate status is associated with increased risk of neural tube defects in the developing fetus and raising maternal folate status before conception and during pregnancy has been shown to reduce occurrence by approximately 70 % (1). Additionally, low folate status has been associated with increased risk of other disease states in adults, such as CVD (2), dementia (3) and some cancers (4). In the UK, fortification of foods with folate, normally in the form of folic acid, is optional. In some countries, such as the USA, Chile and Canada, a mandatory programme where all flour is fortified with folic acid has been in place for some time (5–7), and is currently under consideration in the UK.

Folate status can be raised either by supplementation or fortification (usually with folic acid), or by increasing consumption of folate-rich foods. Whilst some countries have introduced policies of mandatory fortification, many have not. Those countries without mandatory fortification are thus either dependent mainly on natural food folates to optimise status, or on a mix of natural food folates and voluntary supplementation and/or optional food fortification. Since, for a variety of reasons, folate bioavailability from food is lower than folic acid (8) it is important to be able to assess this in order to make evidence-based recommendations for dietary intake. The assessment of folate bioavailability from whole diets, rather than single foods, has been highlighted as of high priority by a UK Food Standards Agency workshop report (9).

Since, following transfer of folate from the gut mucosal cells, there is extensive but unquantified removal of newly absorbed folate from the hepatic portal vein (liver ‘first-pass’) (10), absolute absorption cannot be estimated. Instead, in an attempt to circumvent this problem, single-dose experimental protocols estimate ‘relative absorption’ from a test dose by comparison with a similar reference dose of folic acid. Whilst short-term single-dose acute protocols involve monitoring the appearance and subsequent clearance of the (6S)-5-methyltetrahydrofolic acid (5-CH3H4PteGlu) response in the fast turnover plasma or serum pool arising from absorbed and biotransformed oral test folates, longer-term chronic dietary intervention protocols assessing ‘relative

Abbreviations: 5-CH3H4PteGlu, (6S)-5-methyltetrahydrofolic acid; RNI, UK reference nutrient intake.

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bioavailability’ involve monitoring changes in markers of folate status (plasma or serum folate, erythrocyte folate)\(^8,11\). Plasma or serum folate is the most responsive indicator, with changes in erythrocyte folate responding more slowly because < 1% of circulating erythrocytes are replaced daily\(^8,11\). Hence, studies wishing to monitor optimal changes in participants’ erythrocyte folate concentrations should have a duration of at least 16 weeks to ensure a turnover of erythrocytes whose average life-span is 120 d.

Our recent studies using labelled folates indicated different plasma response kinetics to folic acid than to natural (food) folates, obviously precluding its use for estimating ‘relative absorption’ in single-dose studies\(^12,13\). Since we hypothesised differences in tissue distribution and site of biotransformation\(^13,14\), the question is whether folic acid remains suitable as a reference folate for long-term intervention studies, where ‘relative bioavailability’ of natural (food) folate is assessed based on changes in folate status.

If, as we hypothesised, absorbed folic acid leads to the more rapid accumulation of liver folate stores compared with other absorbed natural (food) folates, it could be further inferred that the use of folic acid as a reference in long-term feeding studies may at the very least lead to a comparatively accelerated change in plasma or serum folate concentration in fasting blood since this is a function of enterohepatic recirculation (estimated at 60–227 nmol/d\(^15,16\)) which increases in proportion to the magnitude of liver stores\(^17\). Thus, comparison of the change in fasting plasma or serum folate induced by natural food or nature-identical folate to the change in fasting plasma or serum folate induced by folic acid may result in the significant underestimation of ‘relative bioavailability’. We have suggested previously that folic acid could be replaced as a reference folate by 5-CH\(_3\)H\(_4\)PteGlu\(^14\), the natural form of folate found in the systemic circulating blood system.

Here we test this hypothesis using a partially double-blinded, placebo-controlled, nutritional intervention study in apparently healthy adults, which was part of a larger investigation into the effects of folate on biomarkers of cardiovascular risk, in collaboration with the University of Sheffield. In the present paper we assess the relative bioavailability of increased folate intake from naturally folate-rich foods by comparison of changes in plasma and erythrocyte folate concentration with changes induced by an equal reference dose of either supplemental 5-CH\(_3\)H\(_4\)PteGlu or folic acid. The choice of a ‘physiological’ intervention dose is important since it is known that there is a threshold for the postprandial appearance of unmetabolised folic acid in the systemic blood system. When previously using an oral dose of 634 nmol (280 \(\mu\)g) \(^13\)C\(_2\)H\(_4\)H\(_4\)PteGlu with liquid chromatography–MS analysis\(^12,13\) we found no detectable folic acid in the postprandial plasma response; this observation is in line with a previous observation using liquid chromatography–microbiological assay analysis that the threshold was above 603 nmol (266 \(\mu\)g)\(^18\). In contrast, a recent report using a more sensitive version of the liquid chromatography–microbiological assay procedure concluded that the threshold for folic acid is about 453 nmol (200 \(\mu\)g)\(^19\). In the present study we employed the lower dose of 453 nmol (200 \(\mu\)g), thus ensuring results could be interpreted independently of threshold effects.

**Methods**

**Study design**

Ethical approval was obtained from the Norfolk 1 Research Ethics Committee following review by the Institute of Food Research’s Human Research Governance Committee and the East Norfolk and Waveney Research Governance Committee.

Generally healthy males and females aged 18–65 years were recruited initially from the database for volunteers held at the Institute of Food Research’s Human Nutrition Unit based on their eligibility for the study. Volunteers were also recruited through advertisements in the media, recruitment stands, email adverts, posters or word-of-mouth. Exclusion criteria for the study were as follows: females that were pregnant or had been pregnant within 12 months of commencement of the study, females that were breast-feeding, anyone who had donated or intended to donate blood within 16 weeks of the first or last study samples, BMI < 18 or > 40 kg/m\(^2\), receiving vascular disease or anti-hypertensive drugs, those with diabetes, and anyone taking B vitamin or folic acid supplements on a regular basis (including multivitamin supplements). Supplement users willing to stop supplements for the study were able to participate following a minimum 120 d washout period before pre-study screening. Potential participants were invited to a pre-study interview before signing consent. Informed consent was obtained at screening. Volunteers were asked to complete a health questionnaire with a research nurse, who also recorded blood pressure, height and weight (from which BMI was calculated). A first-of-the-day midstream urine sample was provided by the volunteer for a dipstick test (Combur 9 Test\(^8\); Roche Diagnostics Ltd, Burgess Hill, West Sussex, UK); pH, protein, glucose, ketones, urobilinogen, bilirubin, blood, leucocytes, nitrites) and, if satisfactory, a 5 ml fasted blood sample was then taken for full blood count analysis (SPIRE hospital laboratory, Colney, Norwich). Volunteers with satisfactory urinalysis results and full blood counts within the standard reference ranges were accepted on to the study. Blood results outside the normal reference range were assessed by the medical advisor to the Institute of Food Research’s Human Nutrition Unit. Excluded volunteers were notified by telephone of the screening outcomes, and their general practitioners notified by letter.

Once accepted on to the study, volunteers were allocated to one of four intervention groups by a senior staff member unconnected to the study. Volunteers participated in a 16-week dietary intervention 4–6 weeks after screening, during which they consumed in addition to their normal diets either (a) an additional 453 nmol natural food folate per d (from a selection of folate-rich foods), (b) a gelatin capsule containing supplemental 5-CH\(_3\)H\(_4\)PteGlu (Metafolin\(^8\); 453 nmol), (c) a gelatin capsule containing supplemental folic acid (453 nmol; 200 \(\mu\)g), or (d) a gelatin placebo capsule. Naturally folate-rich foods for the food intervention group were selected from the McCance & Widdowson 6th edition\(^20\). High-folate foods with manageable portion sizes were selected and drawn into a choice list for the volunteer. A wide range of foods was available in the choice list, and included: vegetables such as spinach, Brussels sprouts, broccoli, peas, asparagus, cauliflower; cereals such as granary bread and non-fortified muesli; dairy products such as...
camembert, brie and cheddar cheese; and grocery items such as baked beans, orange juice, liver pâté, vegetable pâté and mixed nuts. Each food was given a unit value (where one unit = 113.25 nmol folate) and the volunteers asked to consume four units/d to achieve 453 nmol folate from preweighed food portions, which were delivered to them weekly. The capsules were generously gifted to the project by Merck Eprova AG (Schaffhausen, Switzerland), and were delivered to volunteers every 4th week. Volunteers consumed one capsule within each 24 h period, at their own convenience. Capsules were coded, and the code not broken until the end of the study.

Volunteers provided a 12 h fasted blood sample on day 1 (baseline sample) and a second fasted blood sample the day immediately after the 16-week intervention period for plasma homocysteine and plasma and erythrocyte folate analyses. Samples of blood were also sent for full blood count (SPIRE hospital laboratory, Colney, Norwich); this was principally to obtain packed cell volumes required for the calculation of erythrocyte folate concentrations.

**Dietary assessment**

Dietary folate intakes were assessed by means of a 7 d (every third day) weighed food intake carried out in the month before, and during the last 4 weeks of the intervention phase. Records of foods consumed were coded and analysed by the nutritional software Diet Cruncher for Windows® (WayDownSouth Software, Dunedin, New Zealand) using the UK dataset from McCance and Widdowson, version 6(20).

**Analysis of folate and homocysteine**

Folate concentration in plasma and whole blood was determined using a chloramphenicol-resistant *Lactobacillus rhamnosus* NCIMB 10463 microbiological assay(21) modified by haemolysis of whole blood by 10-fold dilution in 1 % ascorbic acid pre-adjusted to pH 4.25(22). Erythrocyte folate concentration was calculated from whole-blood values using individual packed cell volumes and after correction for plasma folate concentration. Day-to-day precision of the method was closely monitored by including in each analytical run high (plasma mean 48.28 (SD 2.56) nmol/l, CV 5.3 %; erythrocyte folate mean 1049.5 (SD 42.5) nmol/l, CV 4.0 %), medium (plasma mean 1049.5 (SD 42.5) nmol/l, CV 4.0 %), low (plasma mean 872.61 (SD 61) nmol/l, CV 7.0 %; erythrocyte folate mean 379.9 (SD 17.7) nmol/l, CV 4.7 %) and low (plasma mean 8.54 (SD 0.68) nmol/l, CV 8.0 %; erythrocyte folate mean 148.8 (SD 16.2) nmol/l, CV 10.9 %) in-house quality controls from either plasma or whole-blood pools, as appropriate. Additionally, accuracy of the method was monitored monthly with the use of two external quality controls: WHO 1st Serum Folate International Standard IS 03/178 (assigned value 12.1 nmol/l; mean 12.83 (SD 1.07) nmol/l, CV 8.3 %) and WHO 1st Whole Blood International Standard IS 95/528 (assigned value 29.45 nmol/l; mean 30.25 (SD 1.53) nmol/l, CV 5.1 %). Plasma total homocysteine was determined by fluorescence polarisation immunosay at the University of Sheffield, on an Abbott IMX automated analyser together with high, medium and low quality controls containing l-homocysteine in processed human serum (Abbott Diagnostics, Maidenhead, Berks, UK). Commercial quality controls provided with the kit were used to check precision. For the low control (assigned value 7.0 μmol/l) the mean was 7.17 μmol/l, intrabatch CV 2.2 %, interbatch CV 1.9 %. For the median control (assigned value 12.5 μmol/l) the mean was 12.8 μmol/l, intrabatch CV 1.9 %, interbatch CV 1.9 %. For the high control (assigned value 25 μmol/l) the mean was 25.3 μmol/l, intrabatch CV 0.8 %, interbatch CV 2.3 %.

**Statistical analysis**

All analysis was carried out using the statistical package STATISTICA (StatSoft Ltd, Bedford, Beds, UK), with significance levels set at *P*<0.05. Datasets did not deviate significantly from the normal distribution (Kolmogorov–Smirnov test) and are described as mean values with their standard errors.

One-way ANOVA (*F* test) was applied to identify potential differences between treatment groups: where significant for post-treatment changes, anticipated effects (decrease in plasma homocysteine and increase in plasma and erythrocyte folate) were identified by the planned comparison test. The degree of association (\(r^2\)) between baseline parameters was assessed by Pearson \(r\) correlation analysis. Correlations between change and initial baseline values were examined by Bland–Altman correlation plots, where change is correlated against the average of pre- and post-intervention values.

**Table 1.** Baseline descriptive statistics of human volunteers randomised to four intervention treatments (about 200 μg folic acid equivalents per d) for 16 weeks: (i) folate-rich foods; (ii) (6S)-5-methyltetrahydrofolic acid (5-CH₃H₄PteGlu) capsule; (iii) folic acid capsule; (iv) placebo capsule (Mean values with their standard errors)

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Food group</th>
<th>5-CH₃H₄PteGlu</th>
<th>Folic acid</th>
<th>Placebo</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Mean SEM</td>
<td>Mean SEM</td>
<td>Mean SEM</td>
<td>Mean SEM</td>
</tr>
<tr>
<td>Subjects (n)</td>
<td>38  42</td>
<td>40  43</td>
<td>40  43</td>
<td>40  43</td>
</tr>
<tr>
<td>Male</td>
<td>15  18</td>
<td>14  17</td>
<td>14  17</td>
<td>14  17</td>
</tr>
<tr>
<td>Female</td>
<td>23  24</td>
<td>26  26</td>
<td>26  26</td>
<td>26  26</td>
</tr>
<tr>
<td>Age (years)</td>
<td>44.8  1.9</td>
<td>44.5  2.1</td>
<td>48.2  1.9</td>
<td>45.9  2.0</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>25.3  0.6</td>
<td>25.4  0.7</td>
<td>25.3  0.6</td>
<td>26.2  0.7</td>
</tr>
<tr>
<td>Dietary folate intake (nmol/d)</td>
<td>662  45</td>
<td>637  27</td>
<td>711  36</td>
<td>789  52</td>
</tr>
<tr>
<td>Plasma homocysteine (μmol/l)</td>
<td>9.52  0.55</td>
<td>10.35  0.86</td>
<td>9.32  0.41</td>
<td>9.05  0.41</td>
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<tr>
<td>Plasma folate (nmol/l)</td>
<td>24.5  2.3</td>
<td>21.7  1.6</td>
<td>24.1  2.4</td>
<td>25.7  2.1</td>
</tr>
<tr>
<td>Erythrocyte folate (nmol/l)</td>
<td>872  61</td>
<td>793  42</td>
<td>833  56</td>
<td>895  60</td>
</tr>
</tbody>
</table>

\(^\d\text{†}\) One-way ANOVA (*F* test); *P*<0.05 considered to be significant.
Results

Compliance

A total of 163 volunteers successfully completed the study from an initial 181 volunteers who provided a pre-intervention baseline sample. Of those that did not complete, eight volunteers withdrew for personal reasons, and ten volunteers were excluded for subsequently being prescribed medication that fell within the study exclusion criteria. Of those volunteers completing the study, eleven had recently been taking B vitamin or folic acid supplements and were delayed from undergoing pre-study screening for at least a 120d washout period: two in the food group, five in the 5-CH$_3$H$_4$PteGlu group, and four in the folic acid group. Capsule compliance was calculated from counting back the capsules left over at the end of each 4-weekly delivery period and cross-checking with a compliance tick-sheet, and was approximately 96% for each supplement group (folic acid, 5-CH$_3$H$_4$PteGlu and placebo). Food group compliance was assessed by increase in dietary folate intake.

Baseline data

There were no significant differences between any of the treatment groups for age, BMI, plasma homocysteine, plasma folate, erythrocyte folate or dietary folate intake (Table 1), nor in intake of folate metabolism-related vitamins B$_2$, B$_6$, and B$_12$. Average intake of folate was approximately 700 nmol/d, i.e. 154% of the UK reference nutrient intake (RNI) which is defined as the amount above which an intake will almost certainly be adequate$^{(22)}$. Average intakes of other B vitamins were 2 mg/d for vitamin B$_2$ (160% RNI), 2.3 mg/d for vitamin B$_6$ (28.8 µg/g protein; 190% RNI) and 5.18 µg/d for vitamin B$_12$ (345% RNI). Average plasma homocysteine status was approximately 9.5 µmol/l. Although there is no consensus on cut-off points it has been argued, from modelling of folic acid supplementation-induced minimisation of plasma homocysteine, that a normal frequency distribution in folate-replete adults would have a mean of 8.3 (SD 1.7; 95% CI 4.9, 11.7) µmol/l$^{(24)}$. Average folate status (plasma approximately 24 nmol/l, and erythrocyte approximately 850 nmol/l) was within acceptable ranges. The acceptable cut-off for plasma folate is > 14 nmol/l (with deficiency considered to be < 7 nmol/l), and the acceptable cut-off for erythrocyte folates is > 360 nmol/l (with deficiency considered to be < 320 nmol/l)$^{(25)}$. Baseline plasma folate associated positively with baseline erythrocyte folate ($P<0.001$; $r^2$ 0.450).

Effect of intervention on plasma homocysteine, and plasma and erythrocyte folate status

A significant increase in dietary folate intake was confined to the food group (Table 2), indicating that any increase in folate status seen in the 5-CH$_3$H$_4$PteGlu and folic acid capsule intervention groups was due to the effect of the supplements, and not to dietary changes during the course of the study. The mean increase in dietary folate intake in the food group was 431 nmol/d greater that the placebo group but 412 nmol/d greater than the mean increase for all the other three treatment groups combined (approximately equal to 90% of target). When compared with the placebo group, plasma homocysteine was lowered to a similar extent in the three active intervention groups. Changes in both plasma and erythrocyte folate status were significantly greater than the placebo for food, 5-CH$_3$H$_4$PteGlu and folic acid (Table 2), and increases due to 5-CH$_3$H$_4$PteGlu and folic acid were greater than in the food group. The mean plasma increase due to folic acid was 26% higher ($P=0.086$) than the increase due to 5-CH$_3$H$_4$PteGlu. The relative increase in plasma folate concentration in the food group was 31% when compared with that induced by folic acid, but 39% when compared with 5-CH$_3$H$_4$PteGlu. The relative increase in erythrocyte folate concentration in the food group when compared with that induced by folic acid was 43%, and 40% when compared with 5-CH$_3$H$_4$PteGlu. When data for the food, folic acid and 5-CH$_3$H$_4$PteGlu intervention groups were combined, a Bland–Altman correlation plot indicated a positive correlation between the change in plasma folate concentration following intervention and the baseline plasma folate concentration.

### Table 2. Post-supplementation changes in dietary folate intake and fasting folate status of human volunteers randomised to four intervention treatments (about 200 µg folic acid equivalents per d) for 16 weeks: (i) folate-rich foods; (ii) (6S)-5-methyltetrahydrofolic acid (5-CH$_3$H$_4$PteGlu) capsule; (iii) folic acid capsule; (iv) placebo capsule

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Food group</th>
<th>5-CH$_3$H$_4$PteGlu</th>
<th>Folic acid</th>
<th>Placebo</th>
<th>ANOVA† variance ratio (F test)</th>
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</thead>
<tbody>
<tr>
<td>Subjects (n)</td>
<td>38</td>
<td>42</td>
<td>40</td>
<td>43</td>
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<tr>
<td>Male</td>
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<td>17</td>
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<tr>
<td>Female</td>
<td>23</td>
<td>24</td>
<td>26</td>
<td>26</td>
<td></td>
</tr>
<tr>
<td>Dietary folate intake (nmol/d)</td>
<td>370 ± 2$^{a, b}$</td>
<td>36 ± 2</td>
<td>-18.8 ± 2</td>
<td>25.4 ± 4</td>
<td>-44.4 ± 2</td>
</tr>
<tr>
<td>Plasma homocysteine (µmol/l)</td>
<td>-0.61 ± 0</td>
<td>0.37</td>
<td>-1.21 ± 2</td>
<td>0.46</td>
<td>-1.21 ± 2</td>
</tr>
<tr>
<td>Plasma folic acid (nmol/l)</td>
<td>5.91 ± 1</td>
<td>2.17</td>
<td>15.10 ± 1</td>
<td>1.94</td>
<td>18.96 ± 2</td>
</tr>
<tr>
<td>Increase in food group relative to reference folates (ratio)</td>
<td>-0.39</td>
<td>0.31</td>
<td>-0.39</td>
<td>0.31</td>
<td>-0.39</td>
</tr>
<tr>
<td>Erythrocyte folate (nmol/l)</td>
<td>13 ± 2</td>
<td>40</td>
<td>343 ± 2</td>
<td>33</td>
<td>321 ± 2</td>
</tr>
<tr>
<td>Increase in food group relative to reference folates (ratio)</td>
<td>-0.40</td>
<td>0.43</td>
<td>-0.40</td>
<td>0.43</td>
<td>-0.40</td>
</tr>
</tbody>
</table>

$^a, b$ Mean values within a row with unlike superscript letters are significantly different ($P<0.05$; planned comparison test).

$^*$ The folic acid group has a trend to be higher than the 5-CH$_3$H$_4$PteGlu group ($P=0.086$).

† One-way ANOVA (F test): $P<0.05$ considered to be significant.


...largely by the 5-CH₃H₄PteGlu intervention group (P<0.001; r 0.269). Bland—Altman correlation plots between the change in erythrocyte folate concentration and baseline erythrocyte folate concentration indicated positive correlations for the three intervention groups (food P<0.001, r 0.410; 5-CH₃H₄PteGlu P<0.001, r 0.416; folic acid P<0.001, r 0.408). Thus, the higher the initial plasma folate or erythrocyte folate concentration, the greater the tendency for a larger increase in status following the 16-week intervention period.

Discussion

Pre-intervention

There were no significant differences between any of the treatment groups for age, BMI, plasma homocysteine, plasma folate, erythrocyte folate or dietary intakes of folate and folate metabolism-related B vitamins (B₂, B₆ and B₁₂), indicating successful random group allocation. As a whole, the generally healthy volunteers selected for the four study intervention groups had good mean baseline dietary folate intakes, plasma homocysteine status, and plasma and erythrocyte folate status.

Post-intervention

The target intake of an extra 453 nmol/d of natural food folate was chosen because it was considered to be a realistic achievable and maintainable amount for a food intervention protocol. Nevertheless, some participants may have partially replaced existing folate-rich foods in their diet with the provided folate-rich foods. This may explain, in comparison with the other treatment groups combined, why the folate intake of the volunteers in our food intervention group was only approximately 90 % of the target additional intake.

Plasma homocysteine concentrations were lowered moderately but significantly by all three active interventions, with no statistical difference between them in effectiveness. Folate status was raised significantly with folic acid and a nature-identical form of folate (5-CH₃H₄PteGlu), and was raised to a lesser extent by increasing the amount of naturally folate-rich foods in the diet. The increase in fasting plasma folate with folic acid showed a trend (P=0.086) to be significantly greater than the increase induced by 5-CH₃H₄PteGlu. This lowered the estimation of the ‘relative bioavailability’ of natural food folate when based on this group’s change in plasma folate compared with that induced by folic acid (31 %) rather than to 5-CH₃H₄PteGlu (39 %). It is interesting to note previously reported dietary interventions using physiological doses of folic acid or 5-CH₃H₄PteGlu that also suggested a larger change in plasma folate concentration in response to folic acid than to 5-CH₃H₄PteGlu (26,27). Compared with the trend (P=0.086) for a 26 % greater change in plasma folate concentration over 16 weeks with folic acid compared with 5-CH₃H₄PteGlu in the present study, these previous studies (24 weeks with 100 μg/d intervention) indicated a change in plasma folate concentration with folic acid that was either 33 % greater (26), or 52 % greater (27), compared with 5-CH₃H₄PteGlu. Whilst these results also did not reach statistical significance it is nevertheless arguable that these observations, in conjunction with results from the present study, collectively support our hypothesis that folic acid intervention may result in a greater change in fasting plasma folate than that induced by a nature-identical folate (for example, 5-CH₃H₄PteGlu). Folic acid should therefore not be used as the reference folate in longer-term studies where estimates of natural food folate ‘relative bioavailability’ rest on comparative increases in plasma or serum folate concentrations.

When based on changes in erythrocyte folate, the present study indicated little discrepancy when estimating natural food folate ‘relative bioavailability’ using 5-CH₃H₄PteGlu (40 %) rather than folic acid (43 %) as the reference folate. This suggests that results from previous studies estimating food folate bioavailability relative to a reference dose of folic acid are probably valid when using a similar intervention dose and time frame. However, when using a dose of 906 nmol/d, recent studies have concluded that 5-CH₃H₄PteGlu may either maintain (28), or improve (29), erythrocyte folate concentrations to a significantly greater extent than folic acid. Thus, in principle, it would also seem inappropriate to use folic acid as a reference folate when estimates of natural food folate ‘relative bioavailability’ rest on comparative increases in erythrocyte folate concentrations.

It is interesting to note, in particular, that changes in erythrocyte folate concentrations upon active water-soluble folate intervention (food, 5-CH₃H₄PteGlu or folic acid) appeared to be related to initial baseline concentrations such that the higher the initial status, the greater the change upon intervention. Although this is counterintuitive it is a phenomenon that has been reported for fat-soluble compounds such as α-tocopherol, β-carotene and lutein (30). This would mean that folate intervention studies in volunteers with normal baseline status would overestimate the response that is likely to be achieved in any population having a low or deficient status that may subsequently be targeted for folate supplementation.

Conclusions

Relative food folate bioavailability estimates using folic acid as the reference folate are clearly not valid when based on changes in plasma folate concentrations. When recent published observations are additionally taken into account it is also arguable that, in principle, folic acid should not be used as the reference folate even when estimates of food folate bioavailability are based on changes in erythrocyte folate concentrations. Using (6S)-5-CH₃H₄PteGlu as the reference folate in long-term natural food folate bioavailability studies would avoid future results’ validity being questioned.

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