Correspondence by TEI, other substances such as As. We are currently testing this hypothesis in a randomized controlled trial of creatine supplementation. In addition, as Basu et al. (2011) noted, and as we have previously reported (Gamble and Liu 2005), one implication of the observed association between uCr and As methylation capacity is that urinary As should not be expressed per gram creatinine to correct for urine concentration. Rather, uCr should be included as a covariate in regression models.

One concerning aspect of the study by Basu et al. (2011) is the handling of blood samples used for nutrient measurements. As noted by Basu et al. and in a previous publication on these same participants (Chung et al. 2006), the blood samples were stored in an ice chest in the field for up to 24 hr before processing. This 24-hr delay can be problematic for some nutrients, especially folate, which is extremely sensitive to oxidative degradation (Drammeh et al. 2008). Basu et al. (2011) reported that in univariate analyses, they observed higher urinary percentages of InAs in individuals with higher serum folate concentrations. This finding is contrary to our previous findings that folate facilitates As methylation (Gamble et al. 2005, 2006, 2007; Hall et al. 2007, 2009). This discrepancy might be explained by differences in sample processing.

Basu et al. (2011) also reported associations between dietary intake of several nutrients (assessed using a modified 24-hr recall) and As methylation capacity. One of the most critical and widely discussed issues in nutritional epidemiology is the method used to adjust for total energy intake (TEI) (Willett et al. 1997). The main reasons to adjust for TEI are to a) adjust for potential confounding by TEI, b) remove extraneous variation in nutrient intakes that is due only to their correlation with TEI, and c) simulate a dietary intervention. What is often most relevant is diet composition, or nutrient intake in relation to TEI (Willett et al. 1997). Several methods are available to adjust for TEI, and the best approach can vary depending on the nutrient and question of interest. Basu et al. (2011) adjusted for TEI by dividing each nutrient intake by TEI (nutrient density method). While this approach is appealing because of its simplicity, in reality it can create a complex variable (Willett and Stampfer 1998). For example, when TEI is related to the outcome of interest, the use of nutrient densities can actually induce confounding in the opposite direction. Although we cannot determine from Basu et al.’s article whether TEI measured by the 24-hr recall was associated with As methylation, in theory, an association seems plausible. Also, because their statistical analysis tested for associations between multiple nutrients and urinary As metabolites, it is best to acknowledge that some of the statistically significant associations might be due to chance alone. The authors declare that they have no actual or potential competing financial interests.

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We thank Hall and Gamble for their interest in our findings. In our article (Basu et al. 2011), we cited their earlier results concerning urinary dimethylarsinic acid (DMA) and creatine concentrations, noting that in 2005 they reported a strong correlation between urinary creatine and the percentage of DMA (DMA%) (Gamble et al. 2005). We also noted rather similar findings from another group working in Bangladesh (Nermell et al. 2008), so together with our findings in India, there are three separate studies that have reported this association (Basu et al. 2011; Gamble et al. 2005; Nermell et al. 2008). Our results highlight the strength of the relationship between urinary creatine and urinary DMA%, which was stronger than the relationship between DMA% and any of the 19 dietary factors and 16 blood micronutrients that we investigated.

Gamble and Hall state that they had previously reported (Gamble and Liu 2005) that one implication of the observed association between urinary creatine and arsenic methylation was that urinary As should not be expressed as per gram creatinine to correct for urine concentrations. This is not quite what they said, although they did point out risks in expressing urinary arsenic per gram of creatinine. However, in that letter (Gamble and Liu 2005) and again in this one, they state that urinary creatine should be included as a covariate in regression models. We do not agree with this recommendation. Steinmaus et al. (2009) reported a spurious relationship between low-level arsenic concentrations and diabetes in the United States after including urinary creatine in a regression model. They noted that urinary creatine concentrations may change as a consequence of diabetes, which is known to affect renal function, and that it is not appropriate to adjust for a factor that is a consequence of the disease being studied.

We also do not agree with Gamble and Hall’s criticism of our blood sample handling, which they appear to have raised because our folate findings were different from theirs (e.g., Gamble et al. 2005). For our study (Basu et al. 2011), blood samples were collected in remote villages, and on occasion they had to be stored on ice for up to 24 hr. Gamble and Hall state that “this 24-hr delay can be problematic for some nutrients, especially folate, which is extremely sensitive to oxidative degradation.” In support of this statement they cite Drammeh et al. (2008), who found reductions in serum folate concentrations.
(about 30%) from whole blood stored for 1 day at 32°C. However, storing blood samples on ice at about 0°C is not the same as storing at 32°C, so this result is not relevant. In addition, Drammeh et al. cited a study that reported stable plasma folate levels during storage of up to 7 days at 4°C (Kubasik et al. 1979). We therefore do not think our findings should be criticized based on sample storage.

However, this does not mean we suggest that folate is not related to DMA%. Findings may vary from study to study, and we accept that Gamble et al. have reported several studies suggesting that folate is associated with small increases in arsenic methylation, including a randomized trial with folate supplements (Gamble et al. 2006). In a study of pregnant women in Bangladesh, a small association between folate and arsenic methylation was reported (Gardner et al. 2011; Li et al. 2008); the authors concluded that nutritional status has a marginal influence on the metabolism of arsenic in pregnant women. Although the overall evidence suggests that folate may increase arsenic methylation, the magnitude of the effect is small and in our opinion of little consequence.

In their letter Gamble et al. report they are now conducting a randomized trial with creatine. Well-run randomized trials of nutritional supplements with sufficient statistical power typically require a large amount of public health funds and resources. We suggest that randomized trials should be conducted only on people whose exposure to arsenic has ceased. However, it is clear that exposures did not cease in the randomized trial of folate they conducted, but that exposures remained markedly elevated based on urine arsenic concentrations (Gamble et al. 2006). The purpose of a randomized trial of folate or creatine is to determine whether these agents might affect arsenic methylation to help reduce arsenic toxicity and increase excretion. However, the relevance of this goal is questionable because we already know the best and most appropriate way to reduce arsenic toxicity—by reducing exposure. The first priority should be to stop exposure to arsenic. If exposure to arsenic ceased, there would be very little arsenic to methylate and enhancement of methylation with supplements would not be necessary. The authors declare that they have no actual or potential competing financial interests.

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