Differential responses to selenomethionine supplementation by sex and genotype in healthy adults

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Abstract

A year-long intervention trial was conducted to characterise the responses of multiple biomarkers of Se status in healthy American adults to supplemental selenomethionine (SeMet) and to identify factors affecting those responses. A total of 261 men and women were randomised to four doses of Se (0, 50, 100 or 200 μg/d as l-SeMet) for 12 months. Responses of several biomarkers of Se status (plasma Se, serum selenoprotein P (SEPP1), plasma glutathione peroxidase activity (GPX3), buccal cell Se, urinary Se) were determined relative to genotype of four selenoproteins (GPX1, GPX3, SEPP1, selenoprotein 15), dietary Se intake and parameters of single-carbon metabolism. Results showed that supplemental SeMet did not affect GPX3 activity or SEPP1 concentration, but produced significant, dose-dependent increases in the Se contents of plasma, urine and buccal cells, each of which plateaued by 9–12 months and was linearly related to effective Se dose (μg/d per kg0.75). The increase in urinary Se excretion was greater for women than men, and for individuals of the GPX1 679 T/T genotype than those of the GPX1 679 C/C genotype. It is concluded that the most responsive Se-biomarkers in this non-deficient cohort were those related to body Se pools: plasma, buccal cell and urinary Se concentrations. Changes in plasma Se resulted from increases in its non-specific component and were affected by both sex and GPX1 genotype. In a cohort of relatively high Se status, the Se intake (as SeMet) required to support plasma Se concentration at a target level (Sepl-target) is: Sein target = [(Sepl-target – Sepl)/18.2 ng d kg0.75 / ml per μg].

Key words: Selenium; Biomarkers; Selenoprotein P; Glutathione peroxidase; Genotype; Supplementation

Se is an essential mineral nutrient required to support the expression of some twenty-five proteins, each of which contains selenocysteine residues as essential constituents(1). These selenoproteins have diverse functions including antioxidant protection(2), thyroid hormone metabolism(3) and Se transport(4). Selenoprotein expression can be reduced by deprivation of Se. SNP can also affect selenoprotein function, as in the case of cytosolic glutathione peroxidase (GPX1)(5). In fact, GPX1 genotype has been associated with cancer risk(6).

High Se status has been associated with reduced cancer risk, and perhaps increased risk of type 2 diabetes. The Nutritional Prevention of Cancer (NPC) Trial showed that increasing Se intake could reduce colon and prostate cancer risk(7), at least for individuals with plasma Se levels <106 ng/ml(8). However, the same study indicated that Se-supplemented subjects whose plasma Se levels had increased to 180–200 ng/ml may have had increased risk of type 2 diabetes(9). Elevated diabetes risk was observed for subjects in the upper quintile of plasma Se (≥137·7 ng/ml) in the third National Health and Examination Survey (NHANES)(10), and among subjects in the upper quartile of plasma Se (≥147 ng/ml) in NHANES 2003–4(11). While increased diabetes risk was not found in response to Se-supplementation in the Se and Vitamin E Cancer Prevention Trial(12), neither was protection against cancer detected in that cohort of men of relatively high baseline Se status (mean plasma Se 156·5 ng/ml). Chiang et al(13) rationalised these apparently discrepant results by suggesting that health risk may be associated with Se status according to a ‘U’ shaped dose–response curve. They demonstrated that at high dietary exposures Se can cause DNA damage in cells and...
induce apoptosis. This may be a mechanism whereby Se exerts its anti-cancer effect; when manifest in non-tumorigenic cells, it may also indicate Se-toxicity.

Such a bi-modal dose–response relationship would suggest that Se may be beneficial for only some individuals, such as those of relatively low Se status and that doses lower than those used previously (200 μg/d(8,12)) may be effective. Should this be the case, then food-based approaches using low Se doses would be facilitated by understanding the quantitative relationship of Se intake, biomarkers of Se status and genetic or sex-based modifying factors.

Projecting the Se intake required to support an Se status target requires understanding the relationship of biomarkers of Se status and level of Se intake. Only a few multi-dose studies have been conducted from which to make such projections. Those studies have used various forms of Se supplements, including selenite which produces only minimal increases (<20 %) in plasma Se(14–16), unless subjects are of low Se status, i.e. plasma Se < 55 ng/ml(17). The only plasma proteins into which selenium-6 can be incorporated, selenoprotein P (SEPP1) and GPX3, are maximally expressed in non-deficient subjects. In contrast, the other major supplemental forms, selenomethionine (SeMet) or SeMet-containing supplements (Se-enriched yeast), increase plasma Se in subjects of both low(17) and higher Se status(14–16,18–21), due to the non-specific incorporation of SeMet into proteins in lieu of methionine (Met). Steady-state plasma Se concentrations are not reached for at least 9 months of SeMet supplementation(21).

We, therefore, conducted a 12-month, randomised, double-blind, multi-dose, placebo-controlled intervention trial to determine the quantitative effects of supplemental SeMet on multiple biomarkers of Se status of healthy, Se-adequate adults. Our hypothesis was that the responses of such individuals could be used to impute the amount of oral Se necessary to raise plasma Se to given target concentrations. We previously presented the baseline findings from that study, comprising a complete assessment using multiple biomarkers of Se status(22). Here, we present the responses to supplementation including subgroup analyses examining the impact of sex and selenoprotein genotype on Se biomarkers.

**Subjects and methods**

This study involved healthy men and women living in the vicinity of Grand Forks, ND, who volunteered and met the eligibility criteria as described previously(22). Volunteers resided in their homes for the duration of the year-long study; they were requested to abstain from Brazil nuts (a significant source of Se) and dietary supplements providing ≥50 μg Se/d for the length of the study.

Sample size was determined from simulations based on results from the NPC trial(7) which were used to estimate the relationship between the observed change in plasma Se after 12 months of supplementation and the effective Se dose (mcg Se/d per kg(17)). For each sample size modelled, anthropometry and experimental design is shown in Fig. 1.

Subjects visited the Research Center 2 weeks before and on the day of initiation of treatment, and at monthly intervals thereafter for 1 year. At each visit anthropometry was performed, a pill count-back was made and subjects received the next calendar pack of supplements. Each volunteer was given a cash honorarium pro-rated for the duration of his/her participation in the study.

This study was conducted according to the guidelines laid down in the Declaration of Helsinki. Oversight was provided by the University of North Dakota Human Subjects Committee (Grand Forks, ND, USA), which reviewed and approved the protocol. The purposes and procedures of the study were explained to the volunteers verbally and in writing, and written informed consent was obtained from each volunteer before his/her participation. The study is registered in the Clinical Trials Registry (ClinicalTrials.gov ID no. NCT00803699).

**Dietary supplement**

Subjects were randomly assigned to treatments consisting of a daily oral supplement containing either 0, 50, 100 or 200 μg Se as l-SeMet and an excipient (dicalcium phosphate) in #2 gelatin capsules (Sabinsa Corporation, Princeton, NJ, USA). The analysed Se contents of these treatments were 0, 56.0, 101.2 and 204.1 μg/capsule, respectively. Supplement capsules were provided in numbered 31-d bubble packs at each monthly visit to the Research Center. Compliance with the study protocol was ascertained in subject interviews and by capsule count-backs. Body weight was also recorded at each of these visits so that Se dose could be adjusted for metabolic body size (kg(175)) in the statistical analysis of the data.

**Anthropometry**

Body weight was measured using an electronic scale. Height was measured at the beginning of the study using a wall-mounted stadiometer.
Sample collection and preparation

Blood and 24 h urine were collected 2 weeks before and on the day of randomisation to treatment as previously described (22), as well as at quarterly intervals. Blood was collected by venepuncture in duplicate 7 ml samples into heparinised, EDTA-treated or non-treated glass tubes. Aliquots of whole blood were subjected to low-speed centrifugation to prepare erythrocyte, buffy coat, plasma and serum fractions. Urine (24 h samples) was collected in sterile polycarbonate bottles. These specimens were held at 4°C pending the completion of screening analyses; excess portions were held at –80°C. Lymphocytes were prepared from whole blood after
lysis of erythrocytes, followed by washing, centrifugation and resuspension in 1% paraformaldehyde (1 h at 2–8°C) to stabilise the cytoplasm, and final resuspension in physiologically buffered saline. Buccal cells were collected using a sterile toothbrush according to Paetau et al. (24); cells were lysed in distilled water, and lysates were held at –80°C for analysis.

**Analytical methods**

Biochemical analyses of methylation status and biomarkers of Se status for the cohort at baseline, before supplementation, were performed as previously described (22). Genotyping was carried out (22) for selenoprotein SNP: GPX1 (rs1050450) (25), GPX4 (rs713041) (26), SEPP1 (rs3877899 and rs7579) (27) and SEPP15 (rs5845) (28). Se status was ascertained on the basis of the activity of GPX and the protein level of SEPP1 in serum, as well as the amounts of Se in plasma, buccal cells and urine. The activity of GPX3 (EC 1.11.1.9) was determined in plasma by the method of Lawrence & Burk (29). The amount of SEPP1 was measured in serum by an enzyme-linked immunosassay (30). Se was determined in plasma, buccal cells and urine by automated electrothermal atomic absorption spectrophotometry using a reduced palladium matrix modifier and an instrument equipped with L’Vov platforms (31). Certified Standards were used (Alfa Aesar, Ward Hill, MA, USA; Perkin Elmer, Waltham, MA, USA and CPI, Santa Rosa, CA, USA) to prepare a calibration set daily with each batch. Calibration validation and calibration blanks were included at the beginning and end of the daily batch and at 10% frequency. Matrix effects for plasma and urine were evaluated using quantitative plasma and urine standards (NIST, Gaithersburg, MD, USA; Seronorm, Billingstad, Norway and Utak, Munich, Germany) to assess the percentage recovery of the analyte in these sample matrices. There is no commercially available quantitative standard for Se in buccal cells; therefore, matrix effects of buccal cell preparations were accounted for by performing spike recoveries using certified calibration standards added directly to one of the samples.

**Selenium distribution in blood**

The components of plasma Se were determined from: (1) total plasma Se; (2) measured plasma GPX3 activities; (3) measured serum SEPP1 levels. To determine the amount of GPX3-derived selenocysteine from the activity of the enzyme, a rate constant of 2.8 × 10^3 mmol/min per mg, molecular weight 92 kDa and a stoichiometry of 4 g-atoms Se per mole GPX3 were assumed (31). For selenocysteine in glycosylated SEPP1, average molecular weight 60 kDa and a stoichiometry of 9.9 g-atoms Se per mole as selenocysteine (32) were assumed. Due to its affinity for heparin, SEPP1 was measured in serum; an inherent assumption was that insignificant quantities of SEPP1 protein precipitated from serum. The difference between the total measured Se and the amounts of Se corresponding to the activity of GPX3 and measured amount of SEPP1 was taken as the amount of Se incorporated non-specifically into plasma proteins. This is presumed to be predominately protein-bound SeMet, as only very low amounts of low molecular weight Se species appear to occur in plasma (<1–2% of total Se) (33).

**Apoptosis and DNA damage assessment**

In order to determine whether this level of Se induced adverse apoptotic responses, that process was evaluated in lymphocytes from a random subset of sixty subjects (thirty women, thirty men) at both 0 and 12 months. Apoptosis was induced by treatment with either H_2O_2 (22 μM), cycloheximide (9.3 mM) or no agent (180 min, 37°C) followed by permeabilisation with 70% ethanol. Cellular DNA was labelled using terminal deoxynucleotidyl transferase and brominated deoxyuridine (BrdU) triphosphate nucleotides, after which cells were

![Graph](https://www.cambridge.org/core/terms).

**Fig. 2.** Time-courses of changes in (a) plasma Se level (diet, P<0.0001; month, P<0.0001; diet × month, P<0.0001), (b) urinary Se (diet, P<0.0001; month, P<0.0001; diet × month, P<0.0001) and (c) glutathione peroxidase 3 (GPX3; diet, P<0.05; month, P<0.0009; diet × month, P<0.034) activity in response to L-selenomethionine supplementation. Values are means, with their standard errors represented by vertical bars.
probed in the dark with fluorescein–BrdU mAb (APO-BrdU™; Phoenix Flow Systems, Inc., San Diego, CA, USA, catalogue no. AU1001). DNA was stained using propidium iodide (PI)/RNase A and within 2 h samples were analysed by flow cytometry using a four-colour instrument (Epics-XL, Beckman-Coulter, Miami, FL, USA) equipped with a 488 nm laser. BrdU and PI were detected by absorbance at 525 and 620 nm, respectively. BrdU incorporation was plotted v. PI incorporation with a logical gate excluding doublets. A total of 15 000 events were collected (flow rate 200–600 events/s) in the region encompassing the main population of intact single leucocytes. Data were analysed using Summit™ Offline flow cytometry analysis software (Cytomation, Inc., Fort Collins, CO, USA).

The extent of DNA damage in peripheral blood lymphocytes was measured by the alkaline comet assay as described by Waters et al. The extent of DNA damage was assessed under basal conditions and after ex vivo exposure of lymphocytes to 22 μM-H2O2. Damage was scored in 200 cells randomly selected from each sample by a single examiner blinded to treatment. SYBR Green 1 stained nucleoids were examined at 200 £ magnification using an epifluorescence microscope. Each cell was scored visually by the method of Duthie & Collins: no damage (type 0); mild to moderate damage (types 1 and 2) and extensive DNA damage (types 3 and 4). Extent of damage was expressed as the percentage of cells of types 3 and 4.

Statistical analyses

All statistical analyses were performed using SAS version 9.1.3 (SAS Institute, Inc., Cary, NC, USA). Data for buccal Se, urine

Fig. 3. Relationships of 1-year changes in (a) plasma Se (Δplasma Se = 12.6 ± 18.2 (effective dose); R² = 0.60, P = 0.0001), (b) glutathione peroxidase 3 (GPX3) activity (Δplasma GPX = 0.06 ± 0.001 (effective dose); R² = 0.0001, P = 0.91), (c) urinary Se (Δurine Se = 17.4 ± 18.6 (effective dose); R² = 0.44, P = 0.0001), (d) selenoprotein P (SEPP1; ΔSEPP1 = 0.15 ± 0.018 (effective dose); R² = 0.02, P = 0.51), (e) buccal cell Se (Δbuccal cell Se = 4.6 ± 2.6 (effective dose); R² = 0.22, P = 0.0001) and (f) non-specific plasma Se (Δnon-specific plasma Se = 8.4 ± 16.9 (effective dose); R² = 0.58, P = 0.0001) with effective Se dose (µg/d per kg⁰.⁷⁵ kg⁰.⁷⁵). For each data set, linear regressions (—) and their 95 % CI (—–) and 95 % prediction intervals (—–—) are shown.
Se, SEPP1, folate, homocysteine (Hcy), vitamin B\textsubscript{12} and thyroid-stimulating hormone were highly skewed for which reason they were logarithmically transformed so that their distributions would more closely approximate normal. For these variables, geometric means with the $^{+}1$ SD confidence limits are reported; other data are expressed as arithmetic means and standard deviations. Nucleic acid data were extracted using Robust Multi-Array Average (RMA) in mAdb (a Multi-Array database tool) (National Cancer Institute, National Institutes of Health) and analysed by paired $t$ tests ($P<0.05$), adjusting for multiplicity using the false discovery rate test.

Repeated-measures ANOVA was used to test for effects of supplemental Se level over time. When appropriate, Tukey’s contrasts were used to compare the supplement levels at each individual time point. Regression analysis was used to model the change in Se status given the effective Se dose, calculated as the Se dose consumed (adjusted for reported compliance) per metabolic body size defined as kg$^{0.75}$. Categorical variables were included in the regressions to test for effects of sex or genotype.

**Results**

A total of 261 subjects (106 men, 155 women) were enrolled in the study and randomised to the treatments. Of these, 243 subjects completed the 12-month study, for an attrition rate of 7%. Of the dropouts (eight men, ten women), none complained of side-effects. Compliance with the treatment protocol, as ascertained by monthly capsule count-backs for subjects completing the study, was 97%.

The study subjects were genotyped for selenoprotein SNPs that have been correlated with cancer incidence or mechanistically implicated in carcinogenesis, and exhibited the following genotype frequencies. The prevalence of dominant alleles in this population was: GPX1 (Pro198Leu; rs1050450: 46% C/C, 43% T/C, 11% T/T), GPX4 (3’-UTR (untranslated region); rs713041: 28% C/C, 52% T/C, 20% T/T), SEPP1 (Ala234Thr; rs3877899: 58% G/G, 38% A/G, 4% A/A and 3’-UTR; rs7579: 44% G/G, 44% A/G, 12% A/A) and SEP15 (3’-UTR; rs5845: 65% C/C, 31% T/C, 4% T/T).

This cohort was of relatively high Se status, as indicated by the baseline values of the biomarkers of Se status, e.g. plasma Se level (142.0 (SD 23.5) ng/ml) and an estimated average Se intake of 109 (SD 44) mg/d. We found no evidence that the background intake of Se from dietary sources, as estimated by a quarterly FFQ, differed significantly between treatment groups or changed significantly during the course of the study. In consideration of the analysed Se contents of the supplements, these treatments provided estimated total Se intakes of approximately 109, 165, 210 and 313 mg/d. Thus, all of the subjects were consuming Se at levels greater than the recommended daily allowance, which was set on the basis of maximal expression of GPX3. There is no recommendation for the concentration of Se in plasma.

![Fig. 4. Relationships of responses of (a, c) plasma and (b, d) urinary Se levels to selenomethionine supplementation for men (M) and women (W): (a, b) dose–response over time and (c, d) 1-year dose–response to effective Se dose (µg/d per kg$^{0.75}$). For each data set, the linear regression is shown for each sex. (d) M: Δurine Se = 20.4 + 12.0 (effective dose) and W: Δurine Se = 20.4 + 20.9 (effective dose).](https://www.cambridge.org/core/terms).
Supplementation with SeMet produced significant increases in some but not all biomarkers of Se status. Dose-dependent increases in the Se contents of plasma (Fig. 2(a)) and urine (Fig. 2(b)) manifested within 3 months, with apparent plateaus being reached by 9–12 months. No significant treatment effect on GPX3 activity was detected, although that parameter showed a small (<2%) but significant increase over the course of the study (Fig. 2(c)).

The 1-year changes in the levels of Se in plasma, urine and buccal cells, and in the non-specific component of plasma Se, each showed significant linear relationships with effective Se dose (µg/d per kg\(^{0.75}\)) (Fig. 3). No changes were observed for GPX3 activity or SEPP1 level. The dose-dependent increase in urinary Se excretion of women was 74% greater \((P=0.001)\) than that of men (Fig. 4), and for individuals of both sexes with the GPX1 T/T genotype was 59% greater \((P<0.006)\) than those with the GPX1 C/C genotype (Fig. 5). These differences were significant within 3 months of SeMet supplementation. Individuals with GPX1 T/T genotype also had significantly lower (7%, \(P<0.05\)) plasma Se levels than those with the GPX1 C/C genotype at baseline\(^{(22)}\). No other differences in responses to SeMet-supplementation were associated with the selenoprotein genotypes tested.

The major components of plasma Se were estimated from the difference between total plasma Se and the measured activities and amounts, respectively, of GPX3 and SEPP1, as described previously\(^{(22)}\). By these estimates, the increase in plasma Se produced by SeMet-supplementation was limited to the non-specific component of that compartment (Table 1).

SeMet-supplementation produced no significant effects on spontaneous or induced apoptosis of peripheral lymphocytes (Table 1). Similarly, no significant difference in spontaneous or induced lymphocyte DNA damage was detected in the alkaline comet assay (Table 1). No significant effects of SeMet-supplementation were detected on the levels of folate, vitamin B\(_{12}\), Hcy, thyroid hormones or buffy coat HbA\(_{1c}\) (Table 1).

Discussion

The baseline plasma Se level of this cohort, 142.0 (SD 23.5) ng/ml, shows that this population was adequately nourished with respect to Se prior to supplementation. Nevertheless, plasma Se increased with SeMet-supplementation, reaching a new steady state within 9–12 months. That change, from the initial steady state to the other, after a year of SeMet-supplementation was described by the following relation:

\[
\Delta S_{\text{pl}} \text{(ng/ml)} = 12.6 \times \text{Se}_{\text{intake}} (P < 0.0001),
\]

where \(\Delta S_{\text{pl}}\) is steady-state change in plasma Se concentration; and \(\text{Se}_{\text{intake}}\) is regular Se intake as SeMet required to raise plasma Se concentration to a particular target level \((S_{\text{pl-target}})\) is:

\[
\text{Se}_{\text{in}} (\mu g/kg^{0.75} \text {d per } g) = \frac{(S_{\text{pl-target}} - S_{\text{pl}})}{18.2 \text{ng d kg}^{0.75}/\text {ml per } g}
\]

where \(S_{\text{pl}}\) is current plasma Se concentration, ng/ml; and \(S_{\text{pl-target}}\) is target plasma Se concentration, ng/ml.

The increase in plasma Se in this cohort did not involve increases in the specific selenoproteins in plasma (GPX3, SEPP1), indicating that the baseline Se intake, estimated to be 109.1 (SD 43.6) µg/d\(^{22}\), was sufficient to support maximal expression of each. Instead, the response of plasma Se was explained by changes solely in the non-specific component of plasma Se, which comprised 46% of plasma Se at baseline and increased, over 1 year of SeMet supplementation, to 46.6, 56.3, 61.8 and 70.5% of plasma total Se with 0, 50, 100 and 200 µg Se/d, respectively. That increase is described by the following relation:

\[
\Delta S_{\text{non-spec}} (\text{ng/ml}) = 7.6 \times \text{Se}_{\text{intake}} (P < 0.0001),
\]

where \(\Delta S_{\text{non-spec}}\) is the steady-state change in non-specific plasma Se level; and \(\text{Se}_{\text{intake}}\) is regular Se intake as SeMet supplied.

Fig. 5. Relationships of 1-year changes in urinary Se levels of subjects of different glutathione peroxidase (GPX1) genotypes and effective Se dose (µg/d per kg\(^{0.75}\)). • GPX1 679 T/T (Leu198Leu); ○ GPX1 679 T/C (Leu198-Pro); ● GPX1 679 C/C (Pro198Pro). For each data set, the linear regression is shown. (a) GPX1 679 T/T: \(\Delta \text{urine Se} = 12.8 + 16.4 \times \text{dose}\) effective dose); GPX1 679 C/C: \(\Delta \text{urine Se} = 12.8 + 19.5 \times \text{dose}\) effective dose). (b) GPX1 679 T/T: \(\Delta \text{urine Se} = 17.3 + 25.4 \times \text{dose}\). GPX1 679 C/C: \(\Delta \text{urine Se} = 17.3 + 19.3 \times \text{dose}\).
### Table 1. Summary of responses to selenium supplementation

(Mean values and standard deviations)

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<th>Characteristics</th>
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<td>BMI (kg/m²)</td>
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<tr>
<td>Plasma Se (ng/ml)</td>
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<td>142.0 ± 23.5</td>
<td>142.0 ± 23.5</td>
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<tr>
<td>Plasma GPX3 (nm/mL mg)</td>
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<td>3.64 ± 0.54</td>
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<td>Plasma SEPP1 (µg/ml)</td>
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<td>Urine Se (ng/mg creatinine)</td>
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<td>Plasma Hcy (µg/mL)</td>
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<td>DNA damage, alkaline comet assay</td>
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<td>Basal (% severe damage)§</td>
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<tr>
<td>H₂O₂-treated (% severe damage)§</td>
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<td><strong>Biomarkers of Se status</strong></td>
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<tr>
<td>Plasma GPX3 (nm/min per mg)</td>
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<tr>
<td>Plasma SEPP1 (µg/ml)</td>
<td>3.70</td>
<td>2.92</td>
<td>4.68</td>
</tr>
<tr>
<td>Non-specific plasma Se† (ng/ml)</td>
<td>137.1b</td>
<td>41.6</td>
<td></td>
</tr>
<tr>
<td>Buccal cell Se (ng/mg protein)</td>
<td>20.59b</td>
<td>13.63</td>
<td>31.09</td>
</tr>
<tr>
<td>Urine Se (ng/mg creatinine)</td>
<td>140.0b</td>
<td>91.6</td>
<td>213.5</td>
</tr>
<tr>
<td>Urine 8OHdG</td>
<td>13.9</td>
<td>5.8</td>
<td>33.0</td>
</tr>
<tr>
<td><strong>Other metabolic markers</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Plasma folate (µg/ml)</td>
<td>23.5</td>
<td>13.7</td>
<td>40.4</td>
</tr>
<tr>
<td>Plasma vitamin B₉ (µg/ml)</td>
<td>461</td>
<td>309</td>
<td>687</td>
</tr>
<tr>
<td>Plasma Hcy (µg/ml)</td>
<td>6.0</td>
<td>4.4</td>
<td>8.1</td>
</tr>
<tr>
<td>TSH (mU/l)</td>
<td>2.61</td>
<td>1.46</td>
<td>4.65</td>
</tr>
<tr>
<td>Free T₃</td>
<td>2.92</td>
<td>0.56</td>
<td></td>
</tr>
<tr>
<td>Free T₄</td>
<td>1.39</td>
<td>0.22</td>
<td></td>
</tr>
<tr>
<td>Glucose</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>HbA1c</td>
<td>4.9</td>
<td>0.8</td>
<td></td>
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<tr>
<td><strong>Apoptosis data‡</strong></td>
<td></td>
<td></td>
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</tr>
<tr>
<td>Spontaneous</td>
<td>1.77</td>
<td>0.98</td>
<td>3.19</td>
</tr>
<tr>
<td>H₂O₂-induced</td>
<td>3.60</td>
<td>1.93</td>
<td>6.69</td>
</tr>
<tr>
<td>CHX-induced</td>
<td>7.54</td>
<td>3.13</td>
<td>18.17</td>
</tr>
<tr>
<td><strong>DNA damage, alkaline comet assay</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Basal (% severe damage)§</td>
<td>14.55</td>
<td>0.03</td>
<td>51.39</td>
</tr>
<tr>
<td>H₂O₂-treated (% severe damage)§</td>
<td>39.70</td>
<td>7.31</td>
<td>78.58</td>
</tr>
</tbody>
</table>

GPX, glutathione peroxidase; SEPP1, selenoprotein P; OHdG, hydroxydeoxyguanosine; TSH, thyroid-stimulating hormone; CHX, cycloheximide.

* Women, n 143; men, n 97.
† See text for calculation.
‡ Women, n 30; men, n 30.
§ Damage scores 3 or 4 (see text for details).
expressed as µg Se/d per kg\(^{0.75}\) (y-intercept is not significantly different from 0 \((P<0.081)\)).

Sexually dimorphic aspects of Se status and response to supplementation have recently been reviewed by Schomburg & Schweizer\(^{36}\). Previously, Méplan et al.\(^{37}\) observed that response of the predominate plasma selenoprotein, SEPP1, to Se supplementation differed by sex. The same group noted that the effect of SNP on the expression of selenoprotein GPX4 depended on sex\(^{38}\). Our present study results show that SeMet-supplementation increased urinary Se output more in women than men, suggesting significant, sex-specific differences in Se metabolism. In rodents, it has been shown that selenoprotein biosynthesis is sexually dimorphic in the liver and kidney\(^{39,40}\) even though total Se concentrations and SEPP levels in serum do not differ. Accordingly, Se supplementation in mice affected selenoprotein biosynthesis in a sex-specific manner, indicating that hepatic metabolism of dietary Se differs in males and females\(^{41}\). It is possible that such an effect may underlie the sexual dimorphic urinary Se excretion that we observed in this study involving human subjects.

We also found increased urinary Se output in individuals with the GPX1 T/T relative to those of the GPX1 C/C genotype. Differential turnover of the GPX1 enzyme, which accounts for some 60% of liver Se\(^{42}\), is likely to affect whole-body Se balance; this may explain the increased urinary output. It is also possible that the urinary response related to the fact that individuals with GPX1 T/T have significantly lower (7%) baseline plasma Se levels than individuals with GPX1 C/C\(^{22}\). Cominetti et al.\(^{43}\) observed a similar difference for morbidly obese women of lower Se status in response to Se from Brazil nuts; however, in their relatively small \((n=37)\) study, they were unable to detect the significant genotype-related differences in baseline plasma Se which we observed in the present cohort\(^{22}\). Hu & Diamond\(^{25}\) found that the GPX1 activity response of cultured breast cancer cells ectopically expressing the GPX1 T/T genotype was less than that of cells expressing the GPX1 C/C genotype. That this difference may be related to cancer risk is suggested by the findings of GPX1 genotype as a determinant of the risk of lung cancer among male smokers in the Finnish Alpha-Tocopherol Beta-Carotene (ATBC) Trial\(^{44}\) and breast cancer in American women\(^{23,45}\), as well as the loss of heterozygosity in lung tumours\(^{46}\).

Gonzalez et al.\(^{47}\) found serum Se and Hcy concentrations to vary inversely, explaining 5-8% of the variance of the latter (2-2% of the variance was accounted for by serum folate concentration). Finding a 63% reduction \((P=0.013)\) in the risk of elevated Hcy concentrations for subjects with serum Se levels in the highest tertile, they concluded that Se should be considered as a potential factor for lowering serum Hcy. The present study, however, detected no effects of SeMet supplementation on circulating levels of folate, vitamin B\(_{12}\) or Hcy.

While Se is required for each of the deiodinases involved in thyroid hormone metabolism, that Se supplementation did not affect thyroid hormone levels suggests that those selenoproteins were maximally expressed in these subjects before supplementation. These findings are consistent with our previous results with another cohort of Se-adults\(^{15}\), although another group\(^{48,49}\) reported such changes.

These results provide no evidence of adverse effects of this level of SeMet supplementation. No effects on lymphocyte DNA damage were observed, although DNA damage was reported for morbidly obese women with the GPX1 679 T/T genotype\(^{43}\) and have been observed in animal models in response to selenite\(^{50}\). Neither did SeMet affect spontaneous or peroxide-induced apoptosis of peripheral lymphocytes.

A limitation of the present study was the use of SeMet as the intervention agent. The biological utilisation of this form of Se is known to be affected by dietary Met\(^{51–55}\), due to the competition of Met and SeMet\(^{53}\) for incorporation into general proteins. This cohort was adequately nourished with respect to total protein, and thus Met. Therefore, Se-enriched yeast, which is mostly SeMet, would be expected to have given comparable results. However, supplementation with inorganic Se, which does not contribute to the non-specific fraction of plasma Se as defined in this study, would be expected to have increased primarily urinary Se and, perhaps, its precursors in the non-protein bound fraction of plasma\(^{55}\).

Conclusions

The present study demonstrates the feasibility of tracking responses to SeMet supplementation in a cohort that is not deficient in the nutrient. In such a cohort, biomarkers with clear functional significance that are limited by Se deficiency, such as GPX3 activity and SEPP1 level, are already maximally expressed. Therefore, they are of no utility as indicators of further increases in Se status. Such tracking demands the use of biomarkers related to tissue/body Se pools. We have demonstrated the use of four such biomarkers: plasma total Se, plasma non-specific Se, buccal cell Se and urinary Se. Each responded to SeMet supplementation; those responses were linear with effective Se dose (i.e. corrected for body mass) over the period of time under investigation. The non-specific component of plasma Se, which we imputed from direct measurements of total plasma Se, SEPP1 and GPX3 activity, was responsive to SeMet supplementation, reflecting the incorporation of SeMet into plasma proteins.

Because SeMet is the dominant form of Se in foods\(^{54,55}\), the algorithms generated in this study can be used to predict the steady-state levels of total and non-specific Se in plasma that will be achieved with any particular regular intake of dietary Se. By this relation, an individual with a plasma Se level of 85 ng/ml would have to consume 1-2 µg Se/d per kg\(^{0.75}\) to support the plasma total Se level (106 ng/ml) associated with prostate cancer risk reduction in the NPC Trial\(^{46}\), whereas an intake of 3-4 µg Se/d per kg\(^{0.75}\) would be required to reach the plasma total Se level (147 ng/ml) associated with increased type 2 diabetes risk in the NHANES 2003–4\(^{14}\). This indicates that Se supplementation targeting cancer risk reduction can be safely achieved.
These results suggest risk categories in which personalised needs for Se can be considered. That women retained ingested Se less well than men suggests that they may have greater dietary Se needs. That the same phenomenon was observed for GPX1 genotype suggests that genotype may also be a determinant of Se need. While the algorithm presented here has application as a general guideline for supplementation, personalised strategies will need to consider the influences of sex and genotype.

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