Dietary ratio of animal:plant protein is associated with 24-h urinary iodine excretion in healthy school children

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Abstract

Adequate dietary iodine intake in children is essential for optimal physical and neurological development. Whether lower dietary animal food and salt intake may adversely affect iodine status is under discussion. We examined the association between dietary animal:plant protein ratio with 24-h urinary iodine excretion (24-h UI, μg/d), and whether this is modified by salt intake. A 24-h UI was measured in 1959 24-h urine samples from 516 6- to 12-year-old participants of the Dortmund Nutritional and Anthropometric Longitudinally Designed Study. Parallel 3 d weighed food records were used to estimate dietary intakes. Protein sources were classified as dairy, animal and plant. A repeated-measures regression model (PROC MIXED) was used to analyse the effect of animal:plant protein ratios on 24-h UI. Animal: plant protein ratios ranged from 0.5 (95% CI 0.4, 0.6) to 1.6 (95% CI 1.4, 1.9) (lowest and highest quartile). After adjustment for total energy intake, main dietary iodine sources (dairy and salt intake), and further covariates, the inter-individual variation in animal:plant protein ratio was significantly associated with variation in 24-h UI. One unit higher animal:plant protein ratio predicted 6 μg/d higher 24-h UI (P=0.002) in boys and 5 μg/d (P=0.03) in girls. This relationship was partially mediated by a higher salt intake at higher animal:plant protein ratios. These results suggest that lower consumption of animal protein is associated with a small decline in iodine excretion, partially mediated by decreased salt intake. Because limited salt and increased intake of plant-based foods are part of a preferable healthy food pattern, effective nutrition political strategies will be required in the future to ensure appropriate iodine nutrition in adherent populations.

Key words: Iodine; Children; Dietary protein; Animal protein

The essential role of iodine in thyroid hormone production and in the regulation of metabolism, growth and development has been extensively documented(1–3). In most industrialised countries, the main dietary sources of iodine are iodised salt (approximately 50%) and animal food products; of which, dairy products are the largest contributors, accounting for up to 40% of total iodine intake as has been shown and discussed by different authors(4–8). Despite the global WHO–Universal Salt Iodisation Programme(9), and regional strategies for iodine deficiency prevention and control, which in Europe included initiatives of legislation in the use of iodised salt in industrial food production(10), and more recently in bread(11,12), to mention some; iodine deficiencies in the European region still persist(13–15).

Evidence from studies on the effect of diets containing little to no animal food products on iodine status is limited, but overall, the literature on this topic suggests that lower intakes of animal food can contribute to inadequate iodine intakes(16–18). Although there is still controversy about the impact of reducing dietary protein intake in children and potential health outcomes(19), existing dietary guidelines for healthy eating, for example, the ‘New American Plate’ from the World Cancer Research Fund/American Institute for Cancer Research, advise the limitation in the intake of animal protein and to increase the intake of plant-based foods to prevent chronic diseases(20,21). In addition, reducing Na intake in children is also part of the global health initiatives in the prevention of chronic diseases(22–25). The existing typical Western diets, in contrast, are characterised by high intakes of processed foods, animal food products and salt(26). As studies suggest an association between processed foods as well as meat and increased salt intake(27–29), the question arises whether, diets with lower animal foods (e.g. meats) and thereby possibly also decreased iodised salt intake may contribute to depleted iodine status as reflected in lower iodine excretion. To our knowledge, to date, no studies have assessed

Abbreviations: 24-h UI, 24-h urinary iodine excretion; DONALD, Dortmund Nutritional and Anthropometric Longitudinally Designed Study; EAR, estimated average requirements.

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the association of animal- and plant-based food sources with urinary iodine status in children consuming typical Western diets. To specifically examine the long-term effect of diets predominantly animal or plant based on iodine status, we studied the 24-h urinary iodine excretion (24-h UI) in a sample of German schoolchildren providing multiple collections of both 24-h urine samples (with available Na excretion measurements) and parallel 3d weighed food records, during the years 1993–2010. In Germany, a particularly improved iodine status in the population has been observed since 1993 parallel to legislation amendment, facilitating the use of iodised salt (NaCl) in all processed foods\(^{30,30}\). However, the iodisation of salt used in households or for food production is still on a voluntary basis. During the last years (starting in 2004), the use of iodised salt by the food industry began to decrease in Germany, and by now encompasses only \(< 50\%\) of total added salt, probably contributing to a negative trend in iodine status\(^{4,5}\).

We used the ‘animal-plant protein ratio’ of the children’s habitual diet as an approach to characterise predominantly animal- or plant-based diets. We hypothesised that those children with healthier food patterns (low animal:plant protein ratio) have lower iodine excretion than children with high animal:plant protein ratio; the individual change in dietary sources of protein (animal-related to plant-based) and 24-h UI was evaluated using repeatedly collected dietary records and 24-h urines. As a second aspect, we aimed to investigate whether the potential association between animal:plant protein ratio and urinary iodine excretion may be mediated by an increase in salt consumption (namely iodised salt) with high animal protein diets.

Subjects and methods

Design of the study

The Dortmund Nutritional and Anthropometric Longitudinally Designed (DONALD) Study is an ongoing, observational open cohort study conducted at the DONALD Study Centre of the University of Bonn (formerly Research Institute of Child Nutrition) in Dortmund, Germany. Details on the study protocol have been published elsewhere\(^{31}\). Briefly, since 1985, detailed data on growth, diet and metabolism have been collected from healthy infants between infancy and adulthood. In general, data on growth, diet and metabolism have been collected from children aged 6–12 years participating in the DONALD Study from January 1993 to December 2010 (\(n = 675\) and \(n = 2781\) measurements). To be eligible for the present analysis, children had to be within the age range, and providing at least two complete 24-h urine samples, each with concurrent 3d weighed food record and anthropometrical data during the observational period. The individual number of 24-h urine samples and corresponding dietary records ranged from two to a maximum of seven possible measurements during the 17-year time period (for a 6–12 years age range).

We excluded those observations that met one or more of the following criteria: (1) incompletely collected urine sample, i.e. when body weight-related 24-h creatinine excretion rate was below the cut-off value of \(0.1 \text{mg} / \text{kg body weight per d}\(^{32}\); (2) implausible dietary records, i.e. when the total recorded energy intake was inadequate in relation to the BMR\(^{33}\); (3) estimated total protein intake \(< 0.66 \text{g/kg of body weight, corresponding to the minimum maintenance requirement of protein}\(^{34}\); or (4) use or intake of medication or supplements containing iodine. Hence, the final study sample for the present analysis consisted of 516 children (\(n = 159\) excluded), 51.4\% male and 48.6\% female, providing 1959 collected 24 h urines and related dietary and anthropometrical records (mean 3.8 measurements per subject over the total study period).

Urinary assessment

Annual 24-h urine collections for children older than 3 years were routinely performed in the DONALD Study usually on the third day of the 3d weighed food record under standardised conditions. All urine samples collected during the 24-h collection period were immediately stored frozen at \(-20^\circ\text{C}\) in Extrann (Extran, MA03; Merck) cleaned, preservative-free one litre plastic containers before being transported to the DONALD Study Centre (formerly, Research Institute of Child Nutrition) where they were stored at \(-22^\circ\text{C}\) until analysis\(^\text{35}\). Iodine concentration was determined in the 24-h urine samples by a modified Sandell–Kolthoff method after acidic wet-ashing of the samples\(^\text{36}\). Twenty-four hour Na excretion was measured by flame atomic absorption spectrometry with a Perkin Elmer 1100 Spectrometer (Perkin Elmer).

Nutritional assessment

Food consumption in the DONALD Study was assessed using 3d weighed food records\(^\text{35}\). Energy and nutrient intakes were calculated from the nutrient values of the food ingredients using recipes available in the in-house database LEKETAB, which incorporates information from standard nutrient tables, product labels or recipe simulation based on the labelled ingredients and nutrients\(^\text{37}\). For the present analysis, the data were derived for each participant from the mean of the three dietary recording days. Composite dishes were disaggregated into their corresponding components, e.g. ‘pizza’ into wheat, cheese and water, and assigned to a food group category (e.g. cereals, legumes, fruits, vegetables, milk, cheese, meat and fish). The calculated protein intake was divided into (1) dairy protein including protein content...
from milk and whey-based milk products (yoghurt, butter-milk, cream, condensed milk and quark) and all varieties of cheese; (2) animal protein including protein content from eggs; all type of meats and chicken; fish and seafood and (3) plant protein including protein content from wheat and all wheat products, cereals (breakfast cereals, oats, sprouts); rice, potatoes, fruits, vegetables, mushrooms, soya products (only solid fraction of soya milk), legumes, nuts and pulses. Total protein was estimated as the sum of protein from all sources (dairy + animal protein + plant protein). Based on this classification, dietary animal:plant protein ratio was calculated. A priori, we decided to exclude dairy protein from the ratios for the following reasons: dairy products are often important constituents of the vegetarian food pattern, not just in Germany but also in other industrialised countries such as the USA, as literature suggests (38-39). Milk has a relatively high iodine content (40-43), and recent studies in German children suggest milk to be an important iodine source in the children’s diet (4,5). Due to these reasons, the inclusion of dairy in the animal:plant protein ratio may have introduced a correlation bias, which led to an overestimation of the association between the animal:plant protein ratio and 24-h UI. Milk and milk products whatsoever were included as separate covariate in the PROC MIXED model (see statistical analyses).

Regular use of iodised salt was reported in dietary records or parental interviews by nearly 100% of the participating children. Na (mg) and salt intake (g) were estimated from the urinary Na excretion values (1 g NaCl = 400 mg Na).

**Anthropometry and parental data**

Anthropometric measurements of the DONALD participants were performed at the time of dietary recording and urine collection by trained personnel following standard procedures. Body weight was measured using an electronic scale (Seca 753 E; Seca Weighing and Measuring System) to the nearest 0.1 kg. Height was measured in a standing position to the nearest 0.1 cm using a digital telescopic stadiometer (Harpenden), the children were dressed in underwear only and barefoot. BMI was calculated as body weight (kg) divided by height (m²). Sex- and age-independent BMI standard deviation scores were calculated using German National Reference data (44). Body Surface Area was estimated using the formula of Du Bois & Du Bois (45).

At their child’s admission to the study, parents provided information on educational background and they were also weighed and measured by the same personnel and equipment as the participating children. The weight was taken with light clothes and without shoes. Maternal education and maternal overweight (BMI ≥ 25 kg/m²) were tested additionally as parameters that can potentially influence the dietary preferences in children (46).

**Statistical analyses**

All statistical analyses were carried out by using SAS® (version 9.1.3; SAS, Inc.). In all statistical tests, a P-value < 0.05 was considered as significant. Descriptive data for anthropometric, dietary and urinary parameters are presented as medians and interquartile ranges stratified by (1) sex; and (2) quartiles of animal:plant protein ratio. Quartiles of animal:plant protein ratio (excluding dairy protein) were grouped as low (lowest quartile), medium (middle two quartiles) and high (highest quartile). All observations (i.e. the repeated measurements at different time points from the same individual) were included for the calculations of medians. Sex differences and differences between animal:plant protein ratio quartiles were tested by repeated measures regression models (PROC MIXED, also linear mixed regression model in SAS) to account for the lack of independence that exists between repeated observations on the same person (comparable to a cross-sectional analysis).

PROC MIXED, including both fixed and random effects, was also used for testing the longitudinal association of animal:plant protein ratio with 24-h UI (µg/d). An advantage of the MIXED procedure is that it does not exclude children from the analysis if they have missing data for a particular time point, but it analyses all variables on the assumption that any missing data are missing at random. The random component of the PROC MIXED model accounts among other for the lack of independence that exists between repeated observations on the same person.

The basic longitudinally regression model included animal:plant protein ratio, age, time and time × time as fixed effects. The variable ‘time’ (continuous, 1993–2010) was included as time, and time × time in the model because of the suggested variation of the 24-h UI through the time period (4,50). Potentially confounding factors influencing 24-h UI were considered as covariates. Only variables that substantially modified the coefficient of animal:plant protein ratio by > 10% in the basic models, or significantly predicted the outcome variable (24-h UI), or improved the fit statistic (Akaike’s information criterion) were included in the subsequent multivariate analyses. All final multivariable models (model 1) included animal:plant protein ratio, age (years), time (chronological time of study years 1993–2010 and time × time), energy intake (MJ/d), 24-h urine volume (ml/d) and milk and milk products (g/d) as independent fixed effects. Diets with high animal protein intake may also be characterised by higher salt intake (27), which is one of the most important iodine sources. Because salt intake might mediate the association between animal:plant protein ratio and iodine excretion, 24-h urinary Na excretion (as marker for salt intake) was tested in final multivariable models (model 2). To allow for the distinction between within-person and between-person effects of animal:plant protein ratio on 24-h UI, the data were first centred on the respective person-specific means of the animal:plant protein ratio over time (i.e. individual’s mean), and these means were used to predict the between-person effects on 24-h UI (i.e. the associations of between-person differences in protein consumption with differences in mean 24-h UI between the subjects). After that, time-specific deviations from the person-specific means were calculated (i.e. individual value – the individual’s mean) and used to test whether within-person changes on
protein intakes were associated with within-person changes in 24-h UI.<ref>47, 48</ref>

To estimate the prevalence of children at risk of inadequate iodine intake, 24-h UI excretion levels (µg/d) were compared to the estimated average requirements (EAR, i.e. 55 µg/d for 4–8 years old, 62 µg/d for 9–13 years old)<ref>49</ref>, after considering a mean non-urinary iodine loss of 15%.<ref>60</ref>

**Results**

A general description of the study sample with respect to anthropometry, dietary intakes and 24-h urine parameters is shown in Table 1. In total, median (25th–75th interquartile range) of urinary iodine excretion (24-h UI) among these children was 75 (56–98) µg/d. Both iodine excretion (µg/d) and urinary protein intake were significantly higher in boys than in girls, but no significance remained when corrected by individual energy intake. Regarding dietary intakes, median estimated protein intake was 1.7 (25th–75th interquartile range) g/kg for all children. Animal protein, including dairy, contributed 65.7% to the total protein intake, ranging from 17.3% to 88.8%.

Cross-sectional comparison of anthropometrical, urinary and dietary variables by quartiles of animal:plant protein ratio is shown in Table 2. Both urinary iodine excretion (µg/d) and iodine concentration (µg/l) were lower in the low animal protein diet than in the protein-rich diet (high ratio) in boys and girls; however, they were not significant. Overall, iodine excretion values below the EAR were observed in 14.3% (n 280) of the measurements. The proportion of observations below the EAR was higher in the lower animal protein eaters (12.8% in boys and 19.2% in girls) than in the higher animal protein consumers (10.3% in boys and 14.5% in girls) but statistically was not significant. Children in the high ratio category consumed more protein than children in the lower ratio category (differing by approximately 6 g/d). In the expected direction, Na excretion was higher when more animal protein was consumed (high ratio) but differed significantly only in boys. Proportion of mothers with lower education level was significantly higher in the high animal:plant protein ratio in boys and girls (P<0.0001). Proportion of maternal overweight was higher in the high-ratio group, but was significant only in girls (P=0.0002) (data no shown).

Results of the longitudinal linear mixed effects regression analyses (PROC MIXED) with urinary iodine excretion (24-h UI, µg/d) as dependent variable are shown in Table 3. β-Values for the individual time invariant mean values of animal:plant protein ratios represent the difference of 24-h UI (µg/d) when the dietary animal:plant protein ratio differs by one unit between the individuals across time. β-Values for intra-individual change indicate the within-person changes in the 24-h UI (µg/d) for a unit of change in animal:plant protein ratio in boys and girls (P<0.0001). Proportion of maternal overweight was higher in the high animal:plant protein ratio in boys and girls (P<0.0001). Proportion of maternal overweight was higher in the high-ratio group, but was significant only in girls (P=0.0002) (data no shown).

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**Table 1.** Anthropometric, nutritional and urinary characteristics of 6- to 12-year-old participants of the Dortmund Nutritional and Anthropometric Longitudinally Designed Study (1959 measurements)

<table>
<thead>
<tr>
<th></th>
<th>Boys</th>
<th>Girls</th>
<th>P*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Subjects (n)</td>
<td>265</td>
<td>985</td>
<td></td>
</tr>
<tr>
<td>Measurements (n)</td>
<td>251</td>
<td>974</td>
<td></td>
</tr>
<tr>
<td>Age (years)</td>
<td>9-0</td>
<td>9-0</td>
<td></td>
</tr>
<tr>
<td>Anthropometry</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>BMI</td>
<td>16.4</td>
<td>16.4</td>
<td>0.27</td>
</tr>
<tr>
<td>BMI-SDS</td>
<td>-0.18</td>
<td>-0.20</td>
<td>0.033</td>
</tr>
<tr>
<td>Urinary parameters</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Iodine concentration (µg/l)</td>
<td>109</td>
<td>94</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>24-h Iodine excretion (µg/d)</td>
<td>80</td>
<td>70</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>24-h Iodine excretion (µg/MJ/d)</td>
<td>10.9</td>
<td>10.6</td>
<td>0.22</td>
</tr>
<tr>
<td>Urine volume (ml/d)</td>
<td>730</td>
<td>735</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Urine volume (ml/d/MJ)</td>
<td>98</td>
<td>110</td>
<td>0.86</td>
</tr>
<tr>
<td>Na excretion (mg/d/MJ)</td>
<td>2025</td>
<td>1762</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Na excretion (mg/d)</td>
<td>280</td>
<td>270</td>
<td>0.94</td>
</tr>
<tr>
<td>Dietary parameters</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Estimated salt intake (NaCl, g/d)†</td>
<td>5.1</td>
<td>4.4</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Total energy (MJ/d)</td>
<td>7.4</td>
<td>6.6</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Protein (% of energy)</td>
<td>12.7</td>
<td>12.5</td>
<td>0.21</td>
</tr>
<tr>
<td>Carbohydrates (% of energy)</td>
<td>52</td>
<td>51.9</td>
<td>0.25</td>
</tr>
<tr>
<td>Fat (% of energy)</td>
<td>35.1</td>
<td>35.3</td>
<td>0.38</td>
</tr>
<tr>
<td>Protein per body weight (g/kg)</td>
<td>1.8</td>
<td>1.6</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Animal protein (g/d)‡</td>
<td>15.8</td>
<td>14.4</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Dairy protein (g/d)§</td>
<td>17.2</td>
<td>14.1</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Plant protein (g/d)</td>
<td>17.6</td>
<td>16.0</td>
<td>&lt;0.0001</td>
</tr>
</tbody>
</table>

BMI-SDS, sex- and age-independent BMI standard deviation scores.  
* Sex differences were tested with a linear mixed-effects regression model (PROC MIXED). PROC MIXED was used to account for the dependence of multiple measurements within the same child.  
† Estimated from urinary Na (1 g of NaCl = 0.4 g Na).  
‡ Animal protein includes protein from fish, poultry and all meat products.  
§ Dairy protein includes protein from milk and all milk products.
Table 2. Comparison of anthropometric, nutritional and urinary characteristics between quartiles (Q1 and Q4) of animal:plant protein ratio of 6- to 12-year-old participants of the Dortmund Nutritional and Anthropometric Longitudinally Designed Study (Median values, 25th–75th percentiles)

<table>
<thead>
<tr>
<th>Boys</th>
<th>Girls</th>
</tr>
</thead>
<tbody>
<tr>
<td>Low ratio (Q1)</td>
<td>High ratio (Q4)</td>
</tr>
<tr>
<td><strong>Measurements (n)</strong></td>
<td></td>
</tr>
<tr>
<td>Ratio of animal:plant protein†</td>
<td>0·49</td>
</tr>
<tr>
<td><strong>Anthropometry</strong></td>
<td></td>
</tr>
<tr>
<td>Age</td>
<td>9·0</td>
</tr>
<tr>
<td>Body weight (kg)</td>
<td>30·8</td>
</tr>
<tr>
<td>BSA (m²)</td>
<td>1·09</td>
</tr>
<tr>
<td><strong>Urinary excretion parameters</strong></td>
<td></td>
</tr>
<tr>
<td>Iodine concentration (µg/l)</td>
<td>107</td>
</tr>
<tr>
<td>Iodine per d (µg/d)</td>
<td>76</td>
</tr>
<tr>
<td>% 24-h UI below EAR‡</td>
<td>12·8</td>
</tr>
<tr>
<td>Na per d (mg/d)</td>
<td>1875</td>
</tr>
<tr>
<td>Volume (ml/d)</td>
<td>730</td>
</tr>
<tr>
<td><strong>Dietary parameters</strong></td>
<td></td>
</tr>
<tr>
<td>Total energy (MJ/d)</td>
<td>7·4</td>
</tr>
<tr>
<td>Total protein (g/d)</td>
<td>50·2</td>
</tr>
<tr>
<td>Animal protein (g/d)§</td>
<td>9·0</td>
</tr>
<tr>
<td>Dairy protein (g/d)‖</td>
<td>19·2</td>
</tr>
<tr>
<td>Plant protein (g/d)</td>
<td>20·5</td>
</tr>
<tr>
<td>Dietary fibre (g/d)</td>
<td>20·1</td>
</tr>
</tbody>
</table>

BSA, Body Surface Area; 24-h UI, 24-h urinary iodine excretion; EAR, estimated average requirements.

* $P_{\text{trend}}$ was tested with PROC MIXED using the data of the total sample of 516 subjects with 1959 measurements. PROC MIXED was used to account for the dependence of multiple measurements within the same child.

† Ratios estimated as animal:plant protein (excluding dairy).

‡ Defined by using the EAR<sub>est</sub> corrected for 15% non-urinary losses. Differences were tested with χ² test.

§ Animal protein, include fish, poultry and all meat products.

‖ Dairy protein, include milk and all milk products.
protein ratio across time. A higher animal:plant protein ratio was significantly associated with higher iodine excretion, before (model 1), and after adjustment for Na excretion (model 2); however, β-values diminished after adjustment for Na excretion. Overall, the effect strength was greater between individuals than within individuals (different β-values and significance). Fig. 1 graphically illustrates the proportions of protein sources by animal:plant protein ratio category (low, middle and high) and its effect on 24-h UI, obtained from adjusted means of the 24-h UI from the PROC MIXED model.

In addition, a separate analysis was conducted using the same PROC MIXED model to evaluate the strength of association of the animal:plant protein ratio and 24-h UI, including dairy protein in the animal:plant protein ratios (dairy + animal-/plant-based protein). The results of this analysis were similar to those presented in Table 3; the effect estimates were only slightly enhanced. For instance, in boys at the between-person level, the effect estimate increased from 8.7 μg/d (ratios excluding dairy) to 8.9 μg/d (including dairy) (model 1) and 6.3 to 9.1 μg/d (model 2). In girls, the corresponding values were 7.4 μg/d and 7.4 μg/d (same values) (model 1) and 4.5 to 7.4 μg/d (model 2) respectively.

Discussion

In the present longitudinal analysis of children consuming a diet characterised by relatively high protein intake (typical Western diet), we examined the importance of changes in the proportion of animal to plant protein for changes in iodine status, and the corresponding role of salt intake. Our findings suggest that a high ratio of animal:plant protein is associated with greater iodine excretion (μg/d), even if other relevant dietary sources of iodine (e.g. salt intake) are controlled for.

Suggestive evidence of a higher risk of sub-optimal iodine status has already been proposed when diets lack animal food products. This notion builds on studies mainly conducted in adults, who are vegetarians or vegans, which may lead to deficits in iodine as well as other micronutrients (16,17,50), whereas in children this information is scarce. Most previous cross-sectional studies in children characterising iodine status reported urinary iodine concentration (μg/l), but usually lacked data on the dietary habitual intake (6,7,12,51,52). In our longitudinal analysis, we were able to simulate different dietary protein patterns, and quantify the effect of lower animal food intake, resembling almost a vegetarian diet, on iodine excretion. One example to illustrate the apparent effect of increasing the animal:plant protein ratio on the 24-h UI can be deduced from our model. Assuming a 50 g total protein intake (approximately average protein intake for boys in the low-animal:plant protein ratio), made up by 20 g from dairy, 10 g from animal and 20 g from plant protein, the estimated animal:plant protein ratio is 0.5. Increasing the animal:plant protein ratio by one unit would modify the proportion of the sources, i.e. an intake of 18 g from animal protein, and 12 g of plant-protein, keeping the dairy protein in 20 g. Theoretically, according to our adjusted model (model 2), about 6 μg/d higher 24-h UI would be expected between boys with otherwise similar dietary intakes (including Na intake) and within the same boy, with one unit increase in animal:plant protein ratio. In girls, no association was found between an increase in the ratio and changes at the within person level, possibly due to smaller individual changes that we were not able to fully identify.

One possible mechanism thought to explain the positive association of animal:plant protein ratios with iodine excretion is the suggested association between Na intake and consumption of animal foods. The main dietary sources of Na ingested by children in Europe and Northern America are processed foods, often of animal origin (28,53,54). Longitudinal analysis suggests that meat intake (especially cold cuts) increases 24-h Na excretion as earlier reported in a group of DONALD children (27). These findings were confirmed by our analysis as we observed higher Na intake in the high animal:plant protein ratio category (Table 2). However, the positive association of animal:plant protein ratio with 24-h UI in the
longitudinal PROC MIXED model remained significant after adjustment for Na excretion, albeit with somewhat reduced effect estimates (β-values of the animal:plant protein ratio). Therefore, the association between animal:plant protein ratio and 24-h UI in children seems to be only partially mediated by the amount of Na intake. It is interesting that when including dairy protein in the animal protein fraction from the animal:plant protein ratios predicted a quite similar increase of 24-h UI, compared to the β-values when the dairy is excluded from the ratios. It is possible that the relatively high but constant contribution from the dairy group observed over the whole range of animal and plant protein categories (Fig. 1, pie graphs) explains the observed results.

Traditionally, the iodine status of populations is measured in terms of urinary iodine concentration (µg/d), generally obtained from spot urine samples (55,56) Moreover, recently, it has been acknowledged that whenever feasible, 24-h UI (µg/d) should be the ‘reference standard’ for measuring iodine intake, as it incorporates urine volume, which can strongly affect dilution of the sample and therefore iodine concentration (4,57,58). With our analysis, which comprises a large number of 24-h urine samples (approximately 2000), we were able to confirm the importance of using 24-h UI (µg/d) instead of concentration (µg/l) to define the iodine status. Based on the WHO 100 µg/l cut-off for iodine adequacy, our observed median concentration in boys (109 µg/l) suggests sufficiency, and in girls (94 µg/l) modest insufficiency. Due to an often better hydration status in females compared to males (59,60), such a difference is not unexpected; accordingly, energy-corrected 24h excretion rates of iodine (Table 1) noted that no sex differences exist.

As trends towards alternative nutrition practices, i.e. preferences to consume organic foods, use of no iodised salt, and vegetarian diets, are becoming more popular in European children (61,62), the issue of what constitutes an optimum diet and micronutrient composition has important public health implications. In an effort to achieve healthier diets, even slight changes in food sources, especially in animal products (a decline in dairy, eventually, increasing grain intakes), may potentially negatively affect micronutrients such as iodine as recently observed by Perrine et al. (63) in a group of USA schoolchildren. Accordingly, when the subjects of the present analysis were divided into low (a) high animal:plant protein ratios, the proportion of children with iodine status below EAR was higher in the lower animal foods consumers, although the differences were not significant. Also, the observed effect estimates for a change of animal:plant protein ratio for 24-h UI were small. A change in the animal:plant protein ratio of about one unit predicted a difference in 24-h UI of approximately 6 µg/d, equalling to about 10% of the EAR. Nonetheless, this dietary factor should not be overlooked as a long-term etiological risk factor that might contribute to cause iodine deficiency disorders in children, such as cognitive performance as literature suggests (63,64). In particular, with regard to the planned salt reduction measures that will also reduce the intake of iodised salt (the main iodine source for children), the achievement of an adequate intake in nearly all children is an important goal.

Discrepancies on the adequate amounts and sources of protein intake in children exist. It has been shown in epidemiological and clinical studies that the amounts of dietary protein intake as well as its sources are important in the prevention or risk increase of chronic diseases, as reviewed in the report by Hönnell et al. (19). In the present study population, the median protein intake (1.7 g/kg) was higher than the 0.9 g/kg recommended for German children (65), however, the overall contribution was within the range of representative German schoolchildren’s dietary protein intake (10–15% of energy) (66). Strengths of the present study included the longitudinal design, with two to seven repeated measurements throughout childhood for each individual and the available data in a relatively large number of children, which better allowed us to use models more suitable for the evaluation of the association of
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In conclusion, the present study shows a positive association between higher ratio of animal:plant protein and iodine status in children. Accordingly, children with healthier association between higher ratio of animal:plant protein and 24-h urinary iodine excretion in German schoolchildren and the importance of iodised salt in processed foods. Br J Nutr 106, 1749–1756.


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