Short Communication

Validation of a FFQ for estimating whole-grain cereal food intake

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Estimation of whole-grain (WG) food intake in epidemiological and nutritional studies is normally based on general diet FFQ, which are not designed to specifically capture WG intake. To estimate WG cereal intake, we developed a forty-three-item FFQ focused on cereal product intake over the past month. We validated this questionnaire against a 3-d-weighed food record (3DWFR) in thirty-one subjects living in the French-speaking part of Switzerland (nineteen female and twelve male). Subjects completed the FFQ on day 1 (FFQ1), the 3DWFR between days 2 and 13 and the FFQ again on day 14 (FFQ2). The subjects provided a fasting blood sample within 1 week of FFQ2. Total cereal intake, total WG intake, intake of individual cereals, intake of different groups of cereal products and alkylresorcinol (AR) intake were calculated from both FFQ and the 3DWFR. Plasma AR, possible biomarkers for WG wheat and rye intake were also analysed. The total WG intake for the 3DWFR, FFQ1, FFQ2 was 26 (SD 22), 28 (SD 25) and 21 (SD 16) g/d, respectively. Mean plasma AR concentration was 55·8 (SD 26·8) nmol/l. FFQ1, FFQ2 and plasma AR were correlated with the 3DWFR (r 0·72, 0·81 and 0·57, respectively). Adjustment for age, sex, BMI and total energy intake did not affect the results. This FFQ appears to give a rapid and adequate estimate of WG cereal intake in free-living subjects.

Whole-grain cereals: Food-frequency questionnaires: Alkylresorcinol: Dietary assessment

Whole-grain (WG) cereal intake has been strongly associated with a decreased risk of heart disease, diabetes and some cancers(1,2). WG intake is usually estimated based on answers to three to fourteen foods in total diet FFQ(3–5). Since the United States Department of Agriculture recommendations in 2004 that people should eat at least three servings of WG cereals per day(6), many new products based on different amounts of WG have become available, making accurate estimation/ranking on the basis of a limited number of questions potentially difficult. There is a need for new approaches for estimating WG intake, including FFQ that focus on cereal intake, and using biomarkers of WG intake, such as plasma alkylresorcinols (AR)(7–11).

Currently, there are no validated FFQ focused on WG cereal intake, nor information about the correlation between plasma AR and habitual WG intake. With the growing interest in WG cereals and the general importance of cereal-based foods for human nutrition, it is important to develop tools that improve our understanding of their intake and role in nutrition. In this present study, we aimed to develop and validate a FFQ that can rapidly estimate WG cereal intake in free-living subjects in the French-speaking area of Switzerland.

Subjects and methods

Questionnaire design

The FFQ was designed to capture information about intake over the past month from four general categories of cereal products: bread, breakfast cereals, snacks and desserts and cooked cereals. Questions were asked about the frequency of specified portions of food eaten (never/0 time/month, 1–3 times/month, 1 time/week, 2–4 times/week, 5–6 times/week, 1 time/d, 2–4 times/d and 5 or more times/d). Individual portion sizes were corrected using photos for food categories with large variation in portion size (breakfast cereal and cooked cereals). For each food category, space was available for subjects to include products that were not included in the category. Frequencies were converted into g/d for each product. For each food category, space was available for subjects to include products that were not included in the category. Frequencies were converted into g/d for each product. For each food, an average value for each variable studied (total cereal content, type of cereal ingredients, percentage of WG, fibre and AR concentration) was estimated by collecting data for up to six different products available at local supermarkets (Lausanne area, Switzerland). All products were converted to dry weight. Where direct analysis data on AR concentration for a food did not exist, it was estimated using previously published values for WG wheat, rye and barley flours(12–14).

Abbreviations: AR, alkylresorcinol; FFQ1, FFQ on day 1; FFQ2, FFQ on day 14; WG, whole grain; 3DWFR, 3-d-weighed food record.

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The relative validity of the FFQ was determined against a 3-d-weighed food record (3DWFR). Cereal food intake from the 3DWFR was calculated using Genesis R&D nutrient composition database software (ESHA, Salem, OR, USA). Nutrient data were from the Swiss nutrient database (http://www.swissfir.ethz.ch/index_EN), or, if not available, from the German, French or American nutrient databases. The following data were calculated from the FFQ and 3DWFR: total WG intake (g/d), total cereal intake (g/d), cereal fibre (g/d), WG wheat (g/d), WG rye (g/d), WG oats (g/d), WG maize (g/d), other WG (g/d, including barley and millet), AR intake (mg/d), intake of breads and biscuits (g/d), breakfast cereals (g/d), snacks and sweets (g/d) and cooked cereals (rice, pasta etc; g/d). Data were handled using Excel (Microsoft, Denver, CO, USA).

Subjects and study design

Thirty-three subjects from the Nestlé Research Centre volunteered for the trial (filled out the first FFQ), and a total of thirty-one completed the entire study (nineteen females and twelve males; Table 1). All subjects were of European or Caucasian cultural origin.

Subjects were given a first coded FFQ (FFQ1), and asked to fill this out and then commence a 3DWFR over the next 14 d. The 3DWFR was filled out in booklets divided into days and meal times, including space to include recipes. Subjects were asked to weigh and complete the record on two weekdays and one weekend day, and recommended to include as many details about the foods they ate as possible. On completion of the 3DWFR, the subjects were asked to complete the FFQ a second time (FFQ2), and then over the next week provided a fasting blood sample. The present study was conducted according to the guidelines laid down in the Declaration of Helsinki, and all procedures involving human subjects/patients were approved by the Clinical Trial Ethical Committee of the Lausanne region. Written informed consent was obtained from all subjects.

Laboratory analyses

Blood was collected and plasma separated by centrifuging the blood at 1000 rpm for 10 min. Plasma was stored at 2°C until analysis. Plasma glucose, TAG, LDL and HDL cholesterols were measured using enzymatic kits and an XPAND autoanalyzer (kits and instrument from Dade Behring, Düb gen, Switzerland).

AR were analysed according to the method of Landberg et al. (15), except that 1 ml of 50 % ethanol was used for deproteinisation before extraction, and N-methyl-N-(trimethylsilyl) trifluoroacetamide + 1 % trimethylchlorosilane was used as the silylating reagent.

Statistical analyses

FFQ1, FFQ2, 3DWFR and plasma AR data did not follow a normal distribution, so were normalised using log transformation. Data from FFQ1, FFQ2 and plasma AR concentrations were compared with the 3DWFR using several different methods. Pearson correlation coefficients were computed to measure the strength of the relationship between the variables and Bland–Altman plots (16), were used to assess the homogeneity of the individual data, and FFQ and plasma AR concentrations were classified according to tertiles of intake to assess the ability of the methods to correctly group subjects.

Correlations were determined with and without weighting for sex, BMI and age. Different means of tertiles (17) were analysed using ANOVA with the Tukey–Kramer multiple-comparison test. Differences and correlations were considered significant at \( P<0.05 \).

NCSS for Windows 2007 (Kaysville, UT, USA) and Microsoft Excel were used for statistical calculations.

Results and discussion

Thirty-three subjects were recruited onto the study and thirty-one completed the three dietary records. Clinical chemistry values for subjects were within normal ranges except for two, who had not fasted before sampling and were removed from analyses for plasma AR (Table 1). The mean consumption of WG cereals estimated by the 3DWFR, FFQ1 and FFQ2 was 26, 29 and 21 g/d, respectively (Supplementary Table S1, available online only at http://www.journals.cambridge.org/bjn). Results from both FFQ1 and FFQ2 were correlated with WG intake estimated by the 3DWFR (Fig. 1 and Supplementary Table S2, available online only at http://www.journals.cambridge.org/bjn). Globally, FFQ1 slightly overestimated WG intake (3 g) and FFQ2 underestimated intake (−5 g; Supplementary Fig. S1, available online only at http://www.journals.cambridge.org/bjn). A possible explanation for the difference between the FFQ and the 3DWFR is that recording diet over 3 d may not be sufficient to capture cereal intake in the case of main meal cereal foods (such as rice, pasta etc), as these may only be eaten 1–2 times/week, but contribute a large amount to overall cereal intake. The FFQ overestimated for cooked cereal intake while providing an adequate estimate for the other important sources of cereals (bread and breakfast cereals; Supplementary Fig. S2, available online only at http://www.journals.cambridge.org/bjn). A possible explanation for the difference between the FFQ and the 3DWFR is that recording diet over 3 d may not be sufficient to capture cereal intake in the case of main meal cereal foods (such as rice, pasta etc), as these may only be eaten 1–2 times/week, but contribute a large amount to overall cereal intake. The FFQ overestimated for cooked cereal intake while providing an adequate estimate for the other important sources of cereals (bread and breakfast cereals; Supplementary Fig. S2, available online only at http://www.journals.cambridge.org/bjn).

Few studies have attempted to validate total WG cereal intake determined by FFQ against dietary records. Hu et al. (18) found that while cold breakfast cereal was well correlated (Pearson correlation of log transformed data unless otherwise stated) between a general diet FFQ and a dietary record (0.56 and 0.77 for two repeats of the same FFQ), other food sources of WG were poorly correlated (0.31 and

### Table 1. Subject biometric characteristics and fasting plasma values*

<table>
<thead>
<tr>
<th>Mean</th>
<th>Standard Deviation</th>
<th>Range</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>38.12</td>
<td>23–64</td>
</tr>
<tr>
<td>Weight (kg)</td>
<td>67.12</td>
<td>50–93</td>
</tr>
<tr>
<td>Height (cm)</td>
<td>171</td>
<td>154–199</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>23.3</td>
<td>19–30</td>
</tr>
<tr>
<td>Total energy intake (MJ)†</td>
<td>8.35</td>
<td>4.52–14.57</td>
</tr>
<tr>
<td>Total AR (nmol/l)</td>
<td>55.80</td>
<td>16.70–108.38‡</td>
</tr>
<tr>
<td>AR ratio C17 : 0:C21 : 0</td>
<td>0.18</td>
<td>0.05–0.66‡</td>
</tr>
</tbody>
</table>

AR, alkylresorcinol.

*There were a total of thirty-one subjects of which nineteen were females.

†Measured using the 3-d-weighed food record.

‡Twenty-nine subjects (two subjects removed – see text for details).
0.27). Salvini et al. (19) obtained correlation coefficients of 0.43–0.66 for bread and 0.69–0.75 for cold breakfast cereal, although the intake of WG from these products was not estimated. Earlier validation work on FFQ, later used to assess WG intake, reports only correlations for nutrients rather than individual foods. Crude fibre intake determined by 1 week food record and FFQ had unadjusted Pearson correlations of 0.40–0.52 (20,21). This specific WG FFQ with correlation coefficients of 0.72–0.81 with the 3DWFR appears to be on par or an improvement on the previous general diet FFQ estimation of WG intake, although previous estimations have been based on intake over 1 year, while this FFQ focused on intake over the past month.

Total WG consumption as estimated by FFQ1 and FFQ2 was correlated (r 0.75, P < 0.0001) and was not different (95% CI for slope: 0.58, 1.17) indicating that overall, the two questionnaires gave similar results. At the individual level, repeatability was not as good as indicated by the overall correlation, as ten subjects out of thirty-one stated that they ate either twice as much or less WG for FFQ1 than for FFQ2, suggesting that there was a learning effect of the 3DWFR on FFQ2. This can be partly accounted for by misconceptions about what constitutes a WG cereal product, especially for bread, the most frequently consumed cereal-based food in the present study.

The average plasma AR concentration was 55.8 (SD 26.8) nmol/l, with a range from 16.7 to 108.4 nmol/l (Table 1). Plasma AR were correlated with total WG cereal intake estimated by the 3DWFR (r 0.57), FFQ1 (r 0.54) and FFQ2 (r 0.55). Plasma AR were also correlated with total AR intake.

Table 2. Correlations and correct classification into tertiles for the 3-d-weighed food record (3DWFR), FFQ on day 1 (FFQ1) and plasma alkylresorcinol (AR) concentrations (n 29)

<table>
<thead>
<tr>
<th></th>
<th>3DWFR</th>
<th>FFQ1</th>
<th>FFQ2</th>
<th>AR</th>
</tr>
</thead>
<tbody>
<tr>
<td>3DWFR</td>
<td>–</td>
<td>0.72</td>
<td>0.81</td>
<td>0.57</td>
</tr>
<tr>
<td>FFQ1</td>
<td>0.72</td>
<td>–</td>
<td>0.74</td>
<td>0.54</td>
</tr>
<tr>
<td>FFQ2</td>
<td>0.81</td>
<td>0.74</td>
<td>–</td>
<td>0.55</td>
</tr>
<tr>
<td>AR</td>
<td>0.57</td>
<td>0.54</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Correctly classified (%)</td>
<td>–</td>
<td>77</td>
<td>52</td>
<td>52</td>
</tr>
</tbody>
</table>

*Classification into the same tertile of WG cereal intake determined by the 3DWFR.
methods cannot be(22). In the present study, AR had a
of subject recall and bias in a way that recall and diary
by FFQ, 24 h recall or weighed records (in the order of
generally have a weak correlation with dietary intake assessed
attractive way of obtaining a non-subjective measurement of
intake biomarker such as plasma AR concentration is an
sample size also prevented meaningful stratification of data
and the upper and lower tertiles, which may in part be due to
the low sample size used in the present study. The low
sample size also prevented meaningful stratification of data
by sex and age.

Because of the difficulties in assessing WG intake, using an
intake biomarker such as plasma AR concentration is an
attractive way of obtaining a non-subjective measurement of
WG wheat and rye intakes. While biomarkers of food intake
generally have a weak correlation with dietary intake assessed
by FFQ, 24 h recall or weighed records (in the order of
r 0.3–0.5), their true benefit is that they are truly independent
of subject recall and bias in a way that recall and diary
methods cannot be(22). In the present study, AR had a
correlation with WG intake estimated by the 3DWFR and
the two FFQ of between 0.54 and 0.57, which is in the
upper range of correlations from other biomarkers of dietary
intake such as serum α-carotene for vegetable intake (diet
record r 0.52, FFQ r 0.35)(23) and fatty acid 15:0 for total
dairy intake (DR r 0.43, FFQ r 0.28)(24). Intervention studies
have found correlations of between 0.21 and 0.58 for plasma
AR and AR intakes(9), 0.34 for plasma AR and rye bread
intakes(7) and 0.41 for plasma AR and cereal fibre intakes(10).
The AR C17:0:21:0 ratio, which indicates whether the AR
mostly came from WG wheat (approximately 0.1) or rye
(approximately 1)(8,13), was 0.17 (sd 0.14), indicating that
wheat was the most common source of AR in the diet. Two
subjects had ratios over 0.5, but both had low total concen-
trations of AR (<35 nmol/l). The mean and range observed
(Table 1) are lower than baseline samples reported in Swedish
(103 nmol/l)(19) and Finnish studies (65–98 nmol/l)(7,10).

Subjects with a low intake of WG (<16 g/d) still had low–
moderate amounts of circulating AR (average for the lowest
tertile of WG intake = 40.1 nmol/l). These low concentrations
are probably due to relatively high intake of refined cereal
products that can still contain low amounts of AR(12), which
could lead to measurable plasma AR concentrations. The results
from the present study support the hypothesis that AR are acceptable
biomarkers of WG cereal intake. Analysis of other plasma par-
ameters and WG intake did not find any associations, except
for a negative correlation between plasma glucose and WG
intake measured by 3DWFR (r = −0.40, P = 0.031). This trend
was NS for WG cereal food intake estimated by FFQ. While
this finding is based on a low number of subjects, it does lend
support to the idea that WG cereal intake may have an effect on glucose metabolism(25).

The FFQ designed for the present study could be further
refined, as some foods that were included in the questionnaire
were not eaten by any subjects – most notably popcorn and
porridge, which are important sources of WG in other
countries. The current time taken to fill out the questionnaire
(5–10 min) is acceptable for the goal of a rapid questionnaire.
As WG cereal intake estimated using this FFQ correlates with
WG cereal intake estimated using a 3DWFR, and proposed
biomarkers of WG wheat and rye intakes (plasma AR), it
appears to be suitable for estimating WG cereal intake in
free-living subjects over the past month.

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A. B. R. designed the questionnaire and wrote the manuscript.
A. B. R., D. M. B. and S. K. designed the study. A. B. R.,
B. D. and M. B. collected the data. A. B. analysed the
samples. N. P. and A. B. R. performed the data analysis. All
authors contributed to the final revision of the manuscript.
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alkylresorcinols and increases their concentration compared
Alkylresorcinols from whole-grain wheat and rye are trans-
resorcinols as biomarkers of whole-grain wheat and rye intake:
plasma concentration and intake estimated from dietary records.
Plasma alkylresorcinols and urinary alkylresorcinol metabolites

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