

Original Article

Whole grain intake, determined by dietary records and plasma alkylresorcinol concentrations, is low among pregnant women in Singapore

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Background and Objectives: To quantify whole grain intake in pregnant women in Singapore in order to provide the first detailed analysis of whole grain intake in an Asian country and in pregnant women. **Methods and Study Design:** Analysis of 24-h diet recalls in a cross-sectional cohort study and analysis of a biomarker of whole grain intake (plasma alkylresorcinols) in a subset of subjects. The Growing Up in Singapore Towards healthy Outcomes-mother offspring cohort study based in Singapore. 998 pregnant mothers with complete 24-h recalls taken during their 26-28th week of gestation. Plasma samples from a randomly select subset of 100 subjects were analysed for plasma alkylresorcinols. **Results:** Median (IQR) whole grain intake for the cohort and the 30% who reported eating whole grains were 0 (IQR 0, 9) and 23.6 (IQR 14.6, 44.2) g/day respectively. Plasma alkylresorcinol concentrations were very low [median (IQR)=9 (3, 15) nmol/L], suggesting low intake of whole grain wheat in this population. Plasma alkylresorcinols were correlated with whole grain wheat intake (Spearman's $r=0.35$; $p<0.01$). **Conclusions:** Whole grain intake among pregnant mothers in Singapore was well below the 2-3 (60-95 g) servings of whole grains per day recommended by the Singapore Health Promotion Board. Efforts to increase whole grain intake should be supported to encourage people to choose whole grains over refined grains in their diet.

Key Words: whole grain, Singapore, pregnant women, alkylresorcinols, dietary intake

INTRODUCTION

Whole grain intake has consistently been associated with a reduced risk of many diet-related diseases, including cardiovascular diseases, metabolic syndrome, diabetes, obesity and some cancers.^{1,2,3} Unlike refined grains, whole grains contain all of their anatomical components i.e. bran, germ and endosperm in the proportions found naturally in the grain.⁴ However, the most commonly

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consumed cereal products are based on refined grains, where the bran and germ are removed, resulting in lower nutrient density (Table 1).

During pregnancy, beyond prevention of related complications such as gestational diabetes,⁵ whole grains may also be important for foetal development due to their high amounts of the methyl donors, folate and betaine.^{6,7} An adequate dietary supply of these compounds may provide an advantageous environment for the programming of the epigenome by meeting the increased need for methyl groups in rapidly dividing cells.^{8,9}

The consistent association between whole grain intake and reduced risk of diseases has led to specific whole grain recommendations in a number of countries, including at least 48 g (3 servings)/d in the United States, to at least 75 g/8.5 MJ in Denmark and Sweden.¹⁰ In the Asia-Pacific region, current recommendations are limited to 'make half your cereal servings whole' in Australia, 'eat plenty of breads and cereals, preferably whole grain' in New Zealand, and 'of 5-7 cereal servings, make 2-3 servings whole grain' in Singapore.¹¹ Where it has been studied, mean whole grain intake has been found to be well below recommended intake, with recent estimates of intakes of 23 g/d in Irish teenagers,¹² 47 g/d in Scandinavian adults,¹³ and 13 g/d among US children and adults.^{14,15}

At present there is little information on whole grain intake in South East Asian countries and no comprehensive studies published in English. The Singapore National Nutrition Survey (SNNS) reported that average whole grain intake among Singaporean adults is low, though around 50% of the population eat whole grain products at least once a month.¹⁶ However there is no quantitative information on whole grain intake among pregnant women, despite whole grains being recommended as part of a healthy diet during pregnancy.¹⁷

In this study, we aimed to gain insight into whole grain intake among pregnant Singaporean women, by performing a retrospective cross sectional study on expectant mothers from the Growing Up in Singapore Towards healthy Outcome (GUSTO) parent-offspring cohort study. As estimating whole grain intake from dietary records is complicated due to a lack of knowledge about identifying whole grain foods, as well as variation in the amount of whole grain ingredients in different foods, we also used plasma alkylresorcinols, a biomarker of whole grain intake,¹⁸ as a non-subjective control of the questionnaire-based estimates of whole grain intake in a subset of 100 mothers. To our knowledge, this work is the first detailed assessment of whole grain intake in an Asian population and in pregnant women.

METHODS

Study design

We used data from the GUSTO study, a longitudinal parent-offspring cohort study in Singapore, which aims to investigate the role of developmental factors in early life to development of metabolic diseases later in life. The study has been described in detail elsewhere.¹⁹ In brief, pregnant women attending antenatal care (<14 weeks' gestation) in KK Women's and Children's Hospital (KKH) and National University Hospital (NUH), which house the major public maternity units in Singapore, were

recruited into the GUSTO study from June 2009 to September 2010. The inclusion criteria included age range between 18-50 years, intention to finally deliver in KKH or NUH, intend to reside in Singapore for the next 5 years, and willingness to donate cord, cord blood, and placenta. Only Chinese, Malay, and Indian women whose parents were from the same ethnic background were included in the study. Women with pre-existing health conditions such as type 1 diabetes mellitus, current depression or mental health-related disorders were excluded. This 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 Institutional Review Boards of the KKH and NUH. Written informed consent was obtained from all subjects on recruitment. The study was approved by both the National Healthcare Group Domain Specific Review Board (reference number D/09/021) and the Sing Health Centralized Institutional Review Board (reference number 2009/280/D).

Maternal characteristics

Data on ethnicity, maternal age and educational level were collected from the participants during recruitment and information about cigarette smoking and alcohol consumption habits during pregnancy was gathered during the 26-28 week pregnancy clinic visits. A 24-h dietary recall was also administered to ascertain participants' diet during pregnancy and for subsequent determination of whole grain intake. At the same clinic visit, maternal weight and height of participants were measured for calculation of body mass index. Fasting blood samples were drawn for subsequent analyses of plasma alkylresorcinols. The participants also underwent oral glucose tolerance testing (OGTT) and subsequently had their gestational diabetes mellitus (GDM) status diagnosed based on World Health Organization criteria.²⁰

Estimation of dietary whole grain and alkylresorcinol intakes

Total whole grain, whole grain wheat, non-wheat whole grain (whole grain rice, oats and corn – other whole grains were not recorded in any of the diet recalls) and alkylresorcinol intake was determined from the 24-h diet recall obtained from the mothers at 26-28 week of gestation. Food pictures were shown during the interview to obtain more accurate measures of portion sizes. Whole grain content in cereal products were converted to grams of whole grain per day based on information supplied on-pack, by manufacturers or estimated from available recipes. The whole grain definition used was that of the Whole Grains Council,⁴ which is used for nutrition labelling guidelines by the Health Promotion Board of Singapore. Alkylresorcinol intake from foods containing whole grain wheat was estimated based on data available for specific foods²¹⁻²³ or estimates based on the average known flour content in each food or food group. Data were corrected for dry weight, based on water content reported in local food composition tables.²⁴ The 24 h recalls were validated against three day food diaries in 400 subjects. While whole grain intake was not determined for the three day food diaries, the Spearman's correlation

Table 1. Comparison of the macronutrient, B-vitamins, and alkylresorcinol content[†] of whole grain (WG) wheat with refined (RG) wheat and white rice with brown rice

	Energy (kJ/100 g)	Carbohydrate (g/100 g)	Protein (g/100 g)	Fat (g/100 g)	Dietary fibre (g/100 g)	Folate (μ g/100 g)	Betaine (mg/100 g)	Choline (mg/100 g)	Thiamine (mg/100 g)	Riboflavin (mg/100 g)	Niacin (mg/100 g)	Vitamin B ₆ (mg/100 g)	Alkylresorcinols (mg/100 g)
Whole grain wheat	1610	70.6	11.6	2.9	13.1	50	90	20	0.47	0.15	4.65	0.41	50
Refined wheat	1697	83.4	9.8	2.2	4.1	26.5	23	10	0.21	0.03	1.04	0.13	5
Difference WG – RG	-87	-12.8	1.8	0.7	9.0	23.5	67	10	0.26	0.12	3.61	0.28	45
Brown rice	1967	79.9	10.4	3.3	4.7	15.6	3	8	0.44	0.06	6.00	0.67	0
White rice	1694	88.4	9.0	0.8	1.1	10.3	3	10	0.05	0.06	0.57	0.64	0
Difference WG – RG	2	8.5	1.5	2.5	3.5	5.2	0	-2	0.40	0	5.43	0.02	0

[†]Data from the Swedish National Food Composition Database²⁴ for all nutrients except for betaine and choline,⁶ and alkylresorcinols.²³

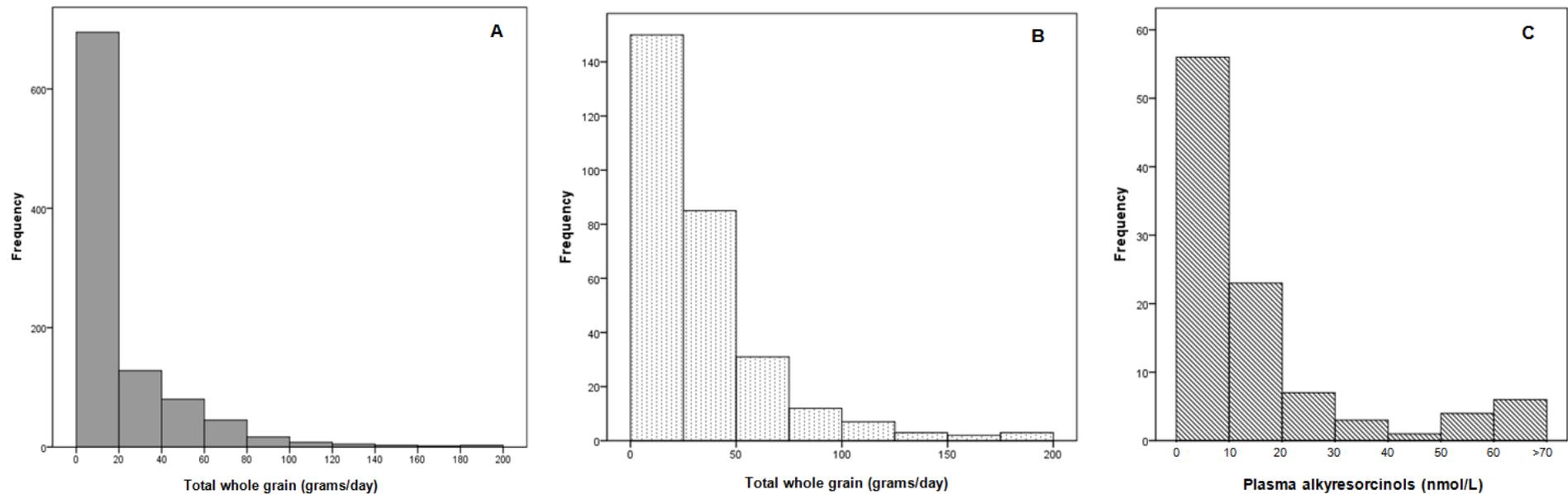


Figure 1. Distribution of measures of whole grain intake in pregnant women in the GUSTO cohort. Whole grain intake in the entire cohort (n=988) was estimated by 24 h recall (A); distribution of whole grain intake among subjects reporting consumption of whole grains (n=293) (B) and distribution of plasma alkylresorcinols among a subset of 100 women from the same cohort (C).

coefficient for dietary fibre was 0.49, and 0.47 for total carbohydrate suggesting agreement between the two methods.

Measurement of plasma alkylresorcinol concentrations

In order to verify estimates of whole grain intake from the 24-hr diet recalls, fasting plasma alkylresorcinols were analysed in a randomly selected subset of 100 subjects from the total cohort of 988 subjects with eligible diet records (see below). Plasma alkylresorcinols were analysed using a modification of a normal phase LC-MS/MS method,²⁵ after extraction with diethyl ether. The modification entailed the use of a shorter column (100 mm x 2 mm, Polaris NH₂, Agilent, Santa Clara, CA, USA, instead of a 250 mm x 2 mm column) and a shortening of the chromatographic run time from 15 minutes to 8 minutes. Control samples analysed using both the original and updated method used here did not differ ($p>0.8$). Matrix matched calibration was used, and two separate control samples were run in triplicate with each batch. Calibration curve standards were run in random order throughout each sequence. Intra-batch repeatability for each control plasma sample was <10% and inter-batch repeatability <15%.

Statistical analyses

Maternal characteristics between the two groups (whole grain vs non-whole grain consumers) were compared using independent sample t-tests for continuous variables and chi-square tests for categorical variables. Dietary whole grain and alkylresorcinol intakes were assessed for normality using the Kolmogorov-Smirnov test, Q-Q plots

and histograms. Normality testing indicated that the data were left-skewed, so non-parametric tests were used for further analyses. Spearman rank correlation coefficients were calculated for the associations between plasma alkylresorcinol and estimates of total dietary whole grain and alkylresorcinol intake ($n=100$). Associations between whole grain intake and plasma alkylresorcinols were analysed with and without adjustment for total energy intake. All statistical calculations were performed using SPSS v. 16.0 (SPSS Inc., Chicago, IL, USA). Two sided p values less than 0.05 were considered statistically significant.

RESULTS

Characteristics of whole grain food consumers and non-consumers

Of the 1162 GUSTO participants recruited, 988 participants had valid 24-hour diet recalls, which were used to determine whole grain intake. Of the 988 subjects, only 293 (29.6%) reported eating any whole grain in the 24-h diet recall (Figure 1A). Of these, mothers of Indian ethnicity had the greatest frequency of whole grain consumption (44%). Participants who were younger ($p=0.011$), had lower educational level ($p=0.0002$), smoked regularly before pregnancy ($p=0.001$), had exposure to smoking before and during pregnancy ($p<0.001$ for both), and had no intake of prenatal supplements ($p=0.06$) were less likely to consume whole grain foods (Table 2).

Whole grain intake did not vary by maternal body mass index (BMI) at 26-28 weeks of gestation, gestational diabetes mellitus, alcohol consumption before and during pregnancy or parity (data not shown).

Table 2. Maternal characteristics according to intake of whole grain in the diet (N=988)

	No whole grain intake		Consumed whole grain		p -value [†]
	N	%	N	%	
Ethnicity					0.0001
Chinese	374	69.3	166	30.7	
Malay	217	83.1	44	16.9	
Indian	104	55.6	83	44.4	
Age, y					0.011
Less than equal 21	35	89.7	4	10.3	
More than 21	660	69.5	289	30.5	
Education [‡]					0.0002
Up to secondary school	231	78.6	63	21.4	
Post-secondary school	273	77.1	81	22.9	
University	186	56.2	145	43.8	
Smoked regularly before pregnancy					0.0005
No	587	68.3	272	31.7	
Yes	108	83.7	21	16.3	
Smoke exposure before pregnancy					0.0003
No	350	61.9	215	38.1	
Yes	345	81.6	78	18.4	
Smoke exposure during pregnancy					0.0003
No	398	64.2	222	35.8	
Yes	297	80.7	71	19.3	
Any prenatal supplements during pregnancy [§]					0.0592
No	181	65.8	94	34.2	
Yes	442	72.3	169	27.7	

[†] p -values were obtained using chi-square tests.

[‡]N=979.

[§]N=886.

Table 3. Spearman's correlations between individual and total plasma alkylresorcinol homologues and estimates of whole grain and alkylresorcinol intakes (n=100). C17:0, C19:0, C21:0, C23:0 and C25:0 are the main alkylresorcinol homologues present in plasma.

	C17:0	C19:0	C21:0	C23:0	C25:0	Total plasma alkylresorcinols
Total whole grain intake (g/day)	0.25*	0.13	0.31**	0.29**	0.13	0.26**
Whole grain wheat (g/dry)	0.24*	0.22*	0.42**	0.36**	0.27**	0.35**
Non-wheat whole grain (g/day)	0.16	0.02	0.05	0.12	-0.05	0.07
Total alkylresorcinol intake (mg/day)	0.26*	0.24*	0.41**	0.37**	0.29**	0.35**

*Spearman's correlation $p < 0.05$; **Spearman's correlation $p < 0.01$.

Table 4. Food sources of whole grain wheat and other whole grains in the cohort and their frequency of intake (% of people consuming the food)

	Freq (%) n=988
Foods made of whole grain wheat	
Wholemeal bread	10.0
Ethnic breads (e.g., chapati, paratha, puri)	5.1
Breakfast cereal with whole grain wheat flour	4.4
Cereal drink (with whole wheat flakes and rice)	2.0
Whole grain wheat biscuits/cookies	2.6
Uppuma (thick porridge made of whole grain wheat)	1.5
Foods made of other whole grain	
Oats and oats products (biscuits, cookies, muesli bar)	5.6
Corn and corn products (sweetcorn cream style, corn chips)	4.3
Brown rice	4.3
Whole grain corn breakfast cereal	2.5
Barley	0.2

Dietary whole grain and alkylresorcinol intakes

In the whole cohort, whole grain intake was highly skewed towards low consumption, with few of those participating in the study eating more than 50 g of whole grain/d (Figure 1A). The median (IQR) whole grain intake of the cohort was 0 (IQR 0, 9) g/day (n=988). Among those who reported consuming any whole grain in their diet (n=293; 29.6%), the median whole grain intake was 23.6 (IQR 14.6, 44.2) g/day (Figure 1B). In this group, 63% (n=185) consumed whole grain wheat, which corresponded to a median intake of 18.1 (IQR 9.4, 35.9) g/d and equivalent to a median of 7.22 (IQR 3.68, 11.7) mg of alkylresorcinols/d. Non-wheat whole grain was consumed by 49.1% (n=144) of the whole grain consumers (median 31.4 g/day; IQR 15.9, 44.2). As alkylresorcinols are only found in high amounts in wheat and rye, the non-wheat sources of whole grain identified from the diet records (rice, corn and oats) do not add to alkylresorcinol intake.

Association of plasma alkylresorcinols with whole grain and alkylresorcinol intakes (n=100)

Of the 100 subjects analysed for plasma alkylresorcinols, only 6 had plasma concentrations above 60 nmol/L (Figure 1C), which corresponds to a suggested threshold for regular intake of at least some whole grain wheat.¹⁸ The median total plasma alkylresorcinol concentration was 9.26 (IQR 2.6, 15.4) nmol/L. The relative percentages of the individual alkylresorcinol homologues was C17:0 (3%), C19:0 (22%), C21:0 (50%), C23:0 (17%) and

C25:0 (8%), indicating that wheat was the source of the alkylresorcinols.²⁶

Plasma alkylresorcinols correlated with total wholegrain, whole grain wheat and alkylresorcinol intake, but not with the intake of non-wheat whole grains. Correlations were significant for all alkylresorcinol homologues measured and strongest for C21:0 and C23:0 (Table 3).

Among subjects who reported consumption of whole grain wheat products (n=24), plasma alkylresorcinol concentrations were higher (median 21.3 nmol/L; IQR 6.73, 59.5) than those not reporting any consumption of whole grain wheat (median 7.37 nmol/L; IQR 2.54, 13.7; $p=0.03$). There was no difference in plasma alkylresorcinol concentration between those reporting eating non-wheat whole grains and those who did not eat any whole grains (data not shown). Adjustment for energy intake did not affect the relationships between whole grain intake and plasma alkylresorcinols.

Key food sources of whole grains in the diets

Of the whole grain foods eaten, whole meal bread was the most commonly eaten, followed by oat products, 'ethnic breads' (e.g. Indian flatbread) and whole grain breakfast cereals (Table 4). Corn-based products were also a relatively commonly consumed source of whole grains. Brown rice was only reported as being eaten by 4.3% of the study participants.

DISCUSSION

The median whole grain intake among pregnant women

in Singapore was low (0 g/d) and below government guidelines for whole grain intake. Among participants who did report eating whole grains, intakes were also well below recommendations (23.6 g/d). Plasma alkylresorcinols in the subpopulation were distributed similarly to the whole grain intake, confirming the findings from the 24-h diet recall data that whole grain intake was indeed low. The low whole grain intake in this cohort are consistent with low intake among the adult Singapore population, as reported in the recent Singapore National Nutrition Survey.¹⁶ Internationally, this is low compared with European countries, where median whole grain intakes among Swedish women of fertile age were 30 g/d,²⁷ 48 g/d in Denmark,²⁸ and around 12 g for men and women between the ages of 19 and 44 in the United Kingdom.²⁹ Data on whole grain consumption during pregnancy is lacking, though extrapolating from Brazilian data¹⁷ would suggest that whole grain intake during pregnancy is also low (mean <5 g/d).

In our cohort of pregnant women, consuming whole grains was associated with a number of healthy behaviours and indicators of higher-socioeconomic status, including not smoking, having a high educational level, and a tendency to consume dietary supplements prior to pregnancy. Associations of greater whole grain intake with older age and higher educational attainment, but not smoking, have also been observed among UK adults.²⁹ Participants of Indian ethnicity had a greater likelihood of eating whole grains, compared with those of Chinese or Malay ethnicity, which we speculate is due to the greater use of whole grains in Indian cuisine. There was no association between eating whole grains and incidence of gestational diabetes or body mass index, though overall intake was so low among whole grain consumers that an effect on these parameters would be unlikely.

In addition to diet recalls to assess whole grain intake, the study results were confirmed using plasma alkylresorcinols, a non-subjective measurement of whole grain intake to confirm the results of the 24-h diet recalls. The advantages of using a biomarker in an observational study is that it does not rely on the population having any background knowledge of what whole grain foods are, nor any bias towards over reporting if participants over-emphasise healthier foods in their diet. However, a limitation of using plasma alkylresorcinols as a biomarker of whole grain intake is that they do not capture brown rice intake, as brown rice does not contain any alkylresorcinols.²¹ This is unlikely to have a large impact on the overall interpretation of our current results as brown rice was not recorded as being commonly eaten by this cohort (Table 4). Moreover, based on the NNS, the mean intake of brown rice intake was also low, estimated at 16 g/d compared with 101 g/d for wholemeal bread and whole grain breakfast cereals.¹⁶ This would suggest that whole grain wheat intake is far greater than brown rice intake and that plasma alkylresorcinols should be able to capture a large proportion of whole grain intake in the Singaporean population.

The median fasting plasma alkylresorcinol concentration in this population was 9 nmol/L, which is low compared with the few population based studies that have been published to date. Median fasting plasma alkylresorcinol concentrations in elderly US citizens was 20

nmol/L,³⁰ while the geometric mean of fasting plasma alkylresorcinol concentrations measured in a subsample of women in a cohort from Northern Sweden was 38 nmol/L.³¹ Based on the mean plasma response to alkylresorcinol intake across 9 studies,¹⁸ a plasma alkylresorcinol concentration of 9 nmol/L would be expected from the intake of 1.3 mg of alkylresorcinols/d, the amount found in approximately 80 g of cooked egg noodles.²³ Plasma alkylresorcinol concentrations from controlled whole grain-free diets (but still containing refined wheat) are generally below 60 nmol/L,¹⁸ confirming that the plasma alkylresorcinol concentrations in this study reflect very low intakes of whole grains, and possibly wheat products in general.

Although the plasma alkylresorcinol concentrations measured were low, they were still significantly correlated with reported whole grain, whole grain wheat and alkylresorcinol intakes. These correlations were in the same range as those previously reported for food frequency questionnaire-based estimates of whole grain intake,¹⁸ and were higher for homologues C21:0 and C23:0. As the association of plasma alkylresorcinols with whole grain intake still holds in this population, we propose that being pregnant does not alter the uptake and metabolism of alkylresorcinols in a way that would confound their use as a biomarker of whole grain intake during pregnancy. Correlations between dietary recall of whole grain intake and plasma alkylresorcinols in the range of 0.25-0.35 appear moderate, and may be seen to raise questions about the relationship between these two. However these correlations are a combination of measurement error for the dietary recall method for whole grains, combined with the many factors that lead to inter-individual differences, including absorption and metabolism of alkylresorcinols, and time since last meal. The deviation (error) from actual intake for both the recall and biomarker are unrelated, so that the use of both methods helps to provide a better picture of actual whole grain intake. In this case, while the overall distribution is similar (Figure 1), the low measured alkylresorcinol concentrations suggested that intake was lower than that reported from the recall measures.

This study represents the first comprehensive analysis of whole grain intake in an Asian population. While some recent reviews have suggested that whole grain may be important for improving health in some Asian populations,³²⁻³⁴ there is little data available on whole grain intakes. Daily whole grain intake in China were estimated to have dropped from 104 g to 24 g from 1982 to 2002 (against a background of 510-402 g/d total grain consumption),³⁵ suggesting that while there may be a tradition of whole grain intake in some Asian populations, economic development or greater availability leads to reduced whole grain intake. The finding that whole grain intake among pregnant Singaporeans is low is a starting point for efforts to encourage greater availability of whole grain foods and increase consumption of whole grains.

The low rates of whole grain consumption among pregnant women in this study are of concern given the substantial evidence based observational data that suggests that whole grains are an important part of a healthy diet.¹ In particular, some whole grains are key sources of

the methyl-donors folate and betaine, which play a role in preventing neural tube defects,^{36,37} as well as a diverse range of vitamins and minerals, and dietary fibre. While there may be efforts to increase whole grain consumption at a government level,³⁸ these data suggest that among pregnant women, few are seeking out whole grain versions of cereal products, or that such products may not be widely available. A public-private partnership to promote eating whole grains first in Denmark resulted in an increase of whole grain intake from 32 g/d to 55 g/d²⁸ demonstrating that whole grain 'campaigns' can lead to substantial increases in intake.

In this observational study, we have found both subjective and non-subjective evidence that whole grain intake among pregnant Singaporeans is very low. This may have implications for long-term disease risk and dietary intake of nutrients important for foetal development. Overall research on whole grain intake and health benefits among South East Asian populations is lacking, and more research is needed to define the potential public health benefits of higher intake of whole grains.

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AUTHOR DISCLOSURES

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REFERENCES

- Ye EQ, Chacko SA, Chou EL, Kugizaki M, Liu S. Greater whole-grain intake is associated with lower risk of type 2 diabetes, cardiovascular disease, and weight gain. *J Nutr.* 2012;142:1304-13. doi: 10.3945/jn.111.155325.
- Aune D, Chan DSM, Lau R, Vieira R, Greenwood DC, Kampman E, Norat T. Dietary fibre, whole grains, and risk of colorectal cancer: systematic review and dose-response meta-analysis of prospective studies. *BMJ.* 2011;343:d6617. doi: 10.1136/bmj.d6617.
- Jonnalagadda SS, Harnack L, Liu RH, McKeown N, Seal C, Liu S, Fahey GC. Putting the whole Grain Puzzle Together: health benefits associated with whole grains-summary of American Society for Nutrition 2010 satellite symposium. *J Nutr.* 2011;141:1011S-22S. doi: 10.3945/jn.110.132944.
- Whole Grains Council. Definition of Whole Grains. 2004 [cited 2014/11/15]; Available from: <http://wholegrainscouncil.org/whole-grains-101/definition-of-whole-grains>.
- Asemi Z, Tabassi Z, Samimi M, Fahiminejad T, Esmailzadeh A. Favourable effects of the dietary approaches to stop hypertension diet on glucose tolerance and lipid profiles in gestational diabetes: a randomised clinical trial. *Br J Nutr.* 2013;109:2024-30. doi: 10.1017/S007114512004242.
- Bruce SJ, Guy PA, Rezzi S, Ross AB. Quantitative measurement of betaine and free choline in plasma, cereals and cereal products by isotope dilution LC-MS/MS. *J Agric Food Chem.* 2010;58:2055-61. doi: 10.1021/jf903930k.
- Patring J, Wandel M, Jägerstad M, Frølich W. Folate content of Norwegian and Swedish flours and bread analysed by use of liquid chromatography-mass spectrometry. *J Food Comp Anal.* 2009;22:649-56. doi: 10.1016/j.jfca.2009.02.007.
- Gluckman PD, Hanson MA, Cooper C, Thornburg KL. Effect of in utero and early-life conditions on adult health and disease. *N Eng J Med.* 2008;359:61-73. doi: 10.1056/NEJMra0708473.
- Waterland RA, Dolinoy DC, Lin JR, Smith CA, Shi X, Tahiliani KG. Maternal methyl supplements increase offspring DNA methylation at Axin Fused. *Genesis.* 2006;44:401-6. doi: 10.1002/dvg.20230.
- Frølich W, Åman P, Tetens I. Whole grain foods and health - a Scandinavian perspective. *Food Nutr Res.* 2013;57. doi: 10.3402/fnr.v57i0.18503.
- Whole Grains Council. Whole grain guidelines worldwide. 2012 [cited 2012/05/12]; Available from: <http://www.wholegrainscouncil.org/whole-grains-101/whole-grainguidelines-worldwide>.
- Devlin NFC, McNulty BA, Gibney MJ, Thielecke F, Smith H, Nugent AP. Whole grain intakes in the diets of Irish children and teenagers. *Br J Nutr.* 2013;110:354-62. doi: 10.1017/S0007114512004989.
- Kyrø C, Skeie G, Loft S, Landberg R, Christensen J, Lund E et al. Intake of whole grains from different cereal and food sources and incidence of colorectal cancer in the Scandinavian HELGA cohort. *Cancer Causes Control.* 2013; 24:1363-74. doi: 10.1007/s10552-013-0215-z.

14. Choumenkovitch SF, McKeown NM, Tovar A, Hyatt RR, Kraak VI, Hastings AV, Herzog JB, Economos CD. Whole grain consumption is inversely associated with BMI Z-score in rural school-aged children. *Public Health Nutr.* 2013;16: 212-8. doi: 10.1017/S1368980012003527.
15. O'Neil CE, Nicklas TA, Zhanovc M, Cho S. Whole-grain consumption is associated with diet quality and nutrient intake in adults: the national health and nutrition examination survey, 1999-2004. *J Am Diet Assoc.* 2010; 110:1461-8. doi: 10.1016/j.jada.2010.07.012.
16. Singapore Health Promotion Board. Report of the National Nutrition Survey 2010. Singapore: Singapore Government Press; 2012.
17. Buss C, Nunes MA, Camey S, Manzolli P, Soares RM, Drehmer M, Giacomello A, Duncan BB, Schmidt MI. Dietary fibre intake of pregnant women attending general practices in southern Brazil - The ECCAGE Study. *Public Health Nutr.* 2009;12:1392-8. doi: 10.1017/S1368980008004096.
18. Ross AB. Present status and perspectives on the use of alkylresorcinols as biomarkers of wholegrain wheat and rye intake. *J Nutr Metab.* 2012;2012:462967. doi: 10.1155/2012/462967.
19. Soh SE, Tint MT, Gluckman PD, Godfrey KM, Rifkin-Graboi A, Chan YH et al. Cohort profile: Growing Up in Singapore Towards healthy Outcomes (GUSTO) birth cohort study. *Int J Epidemiol.* 2014;43:1401-9. doi: 10.1093/ije/dyt125.
20. Wendland EM, Torloni MR, Falavigna M, Trujillo J, Dode MA, Campos MA, Duncan BB, Schmidt MI. Gestational diabetes and pregnancy outcomes - a systematic review of the World Health Organization (WHO) and the International Association of Diabetes in Pregnancy Study Groups (IADPSG) diagnostic criteria. *BMC Pregnancy Childbirth.* 2012;12:13. doi: 10.1186/1471-2393-12-23.
21. Ross AB, Shepherd MJ, Schupphaus M, Sinclair V, Alfaro B, Kamal-Eldin A, Aman P. Alkylresorcinols in cereals and cereal products. *J Agric Food Chem.* 2003;51:4111-8. doi: 10.1021/jf0340456.
22. Ross AB, Kochhar S. Rapid and sensitive analysis of alkylresorcinols from cereal grains and products using HPLC-Coularray-based electrochemical detection. *J Agric Food Chem.* 2009;57:5187-93. doi: 10.1021/jf900239t.
23. Ross AB. Analysis of alkylresorcinols in cereal grains and products using ultrahigh-pressure liquid chromatography with fluorescence, ultraviolet, and coularray electrochemical detection. *J Agric Food Chem.* 2012;60:8954-62. doi: 10.1021/jf301332q.
24. Livsmedelsverket. Livsmedelsdatabasen (Swedish National Food Composition Database). [cited 2013/03/10]; Available from: www.slv.se/naringssock/.
25. Ross AB, Redeuil K, Vigo M, Rezzi S, Nagy K. Quantification of alkylresorcinols in human plasma by liquid chromatography/tandem mass spectrometry. *Rapid Comm Mass Spec.* 2010;24:554-60. doi: 10.1002/rcm.4409.
26. Linko-Parvinen AM, Landberg R, Tikkanen MJ, Adlercreutz H, Penalvo JL. Alkylresorcinols from whole-grain wheat and rye are transported in human plasma lipoproteins. *J Nutr.* 2007;137:1137-42.
27. Amcoff EEAE-B, H; Lindroos, A.K.; Nälén, C.; Pearson, M.; Warensjö-Lemming, E. Riksmaten - vuxna 2010-11: Livsmedels- och näringsintag bland vuxna i Sverige (National Dietary Survey 2010-2011: Food and nutrient intake among adults in Sweden). Stockholm: Livsmedelsverket; 2012.
28. Mejborn HH-Y, K; Fagt, S.; Trolle, E.; Christensen, T. Danskernes fuldkornsindtag 2011-2012 (Wholegrain intake among Danes, 2011-2012). Søborg: DTU Fødevarerinstitutionen; 2013.
29. Thane CW, Jones AR, Stephen AM, Seal CJ, Jebb SA. Comparative whole-grain intake of Br adults in 1986-7 and 2000-1. *Br J Nutr.* 2007;97:987-92.
30. Ma J, Ross AB, Shea MK, Bruce SJ, Jacques PF, Saltzman E, Lichtenstein AH, Booth SL, McKeown NM. Plasma alkylresorcinols, biomarkers of whole-grain intake, are related to lower BMI in older adults. *J Nutr.* 2012;142:1859-64. doi: 10.3945/jn.112.163253.
31. Landberg R, Aman P, Hallmans G, Johansson I. Long-term reproducibility of plasma alkylresorcinols as biomarkers of whole-grain wheat and rye intake within Northern Sweden Health and Disease Study Cohort. *Eur J Clin Nutr.* 2013;67: 259-63. doi: 10.1038/ejcn.2013.10.
32. Misra A, Rastogi K, Joshi SR. Whole grains and health: perspective for Asian Indians. *J Assoc Physicians India.* 2009;57:155-62.
33. Dixit AA, Azar KM, Gardner CD, Palaniappan LP. Incorporation of whole, ancient grains into a modern Asian Indian diet to reduce the burden of chronic disease. *Nutr Rev.* 2011;69:479-88. doi: 10.1111/j.1753-4887.2011.00411.x.
34. Li L, Ying XJ, Sun TT, Yi K, Tian HL, Sun R, Tian JH, Yang KH. Overview of methodological quality of systematic reviews about gastric cancer risk and protective factors. *Asian Pac J Cancer Prev.* 2012;13:2069-79. doi: 10.7314/APJCP.2012.13.5.2069.
35. Ge K. The transition of Chinese dietary guidelines and food guide pagoda. *Asia Pac J Clin Nutr.* 2011;20:439-46. doi: 10.6133/apjcn.2011.20.3.13.
36. Imbard A, Smulders YM, Barto R, Smith DE, Kok RM, Jakobs C, Blom HJ. Plasma choline and betaine correlate with serum folate, plasma S-adenosyl-methionine and S-adenosyl-homocysteine in healthy volunteers. *Clin Chem Lab Med.* 2013;51:683-92. doi: 10.1515/cclm-2012-0302.
37. Benevenga NJ. Consideration of betaine and one-carbon sources of N5-methyltetrahydrofolate for use in homocystinuria and neural tube defects. *Am J Clin Nutr.* 2007;85:946-9.
38. Shiu LK, Loke WM, Vijaya K, Sandhu NK. Nurturing healthy dietary habits among children and youth in Singapore. *Asia Pac J Clin Nutr.* 2012;21:144-50. doi: 10.6133/apjcn.2012.21.1.20.

Original Article

Whole grain intake, determined by dietary records and plasma alkylresorcinol concentrations, is low among pregnant women in Singapore

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由饮食记录和血浆烷基间苯二酚浓度确定的新加坡孕妇全谷物摄入量低

背景与目的：为了给亚洲国家提供一个详细的孕妇全谷物摄入量的分析报告,本文定量分析了新加坡孕妇全谷物的摄入量。**方法与研究设计：**在一个横断面队列研究中分析 24 小时膳食回顾,并分析一个亚组受试者的全谷物摄入量标志物(血浆烷基间苯二酚)。在新加坡长大的母亲子代健康结局队列研究。998 名孕妇在她们怀孕 26-28 周时完成 24 小时膳食回顾。血浆样本来自随机选择的 100 个研究对象组成的亚组,用来分析血浆烷基间苯二酚。**结果：**队列中全部志愿者和队列中 30%报告了摄入全谷物的人群其全谷物摄入的中位数分别为 0 (0, 9) 和 23.6 (14.6, 44.2) 克/天。血浆烷基间苯二酚浓度非常低(中位数=9 (3, 15) nmol/L),提示该人群全谷物摄入量低。血浆烷基间苯二酚与全谷物摄入量相关(Spearman's $r=0.35$; $p<0.01$)。**结论：**新加坡孕妇全谷物摄入量远远低于新加坡健康促进局推荐的每天摄入全谷物 2-3 份(60-95)克。应该鼓励人们在他们的饮食中选择全谷物而不是精制谷物,以增加全谷物的摄入量。

关键词：全谷物、新加坡、孕妇、烷基间苯二酚、膳食摄入量