Whole-grain products and whole-grain types are associated with lower all-cause and cause-specific mortality in the Scandinavian HELGA cohort

Nina F. Johnsen[1]*, Kirsten Frederiksen[1], Jane Christensen[1], Guri Skeie[2], Eiliv Lund[2], Rikard Landberg[3,4], Ingegerd Johansson[5], Lena M. Nilsson[6,7], Jytte Halkjær[1], Anja Olsen[1], Kim Overvad[8,9] and Anne Tjønneland[1]

[1]*Danish Cancer Society Research Center, Strandboulevarden 49, 2100 Copenhagen Ø, Denmark*
[2]*Department of Community Medicine, University of Tromsø – The Arctic University of Norway, Tromsø, Norway*
[3]*Department of Food Science, Uppsala BioCenter, Swedish University of Agricultural Sciences, Uppsala, Sweden*
[4]*Unit of Nutritional Epidemiology, Institute of Environmental Medicine, Karolinska Institutet, Stockholm, Sweden*
[5]*Department of Odontology, Umeå University, Umeå, Sweden*
[6]*Department of Public Health and Clinical Medicine, Umeå University, Umeå, Sweden*
[7]*Arctic Research Centre at Umeå University (Arcum), Umeå, Sweden*
[8]*Section for Epidemiology, Department of Public Health, Aarhus University, Aarhus, Denmark*
[9]*Department of Cardiology, Aalborg University Hospital, Aalborg, Denmark*

Abstract

No study has yet investigated the intake of different types of whole grain (WG) in relation to all-cause and cause-specific mortality in a healthy population. The aim of the present study was to investigate the intake of WG products and WG types in relation to all-cause and cause-specific mortality in a large Scandinavian HELGA cohort that, in 1992–8, included 120 010 cohort members aged 30–64 years from the Norwegian Women and Cancer Study, the Northern Sweden Health and Disease Study, and the Danish Diet Cancer and Health Study. Participants filled in a FFQ from which data on the intake of WG products were extracted. The estimation of daily intake of WG cereal types was based on country-specific products and recipes. Mortality rate ratios (MRR) and 95 % CI were estimated using the Cox proportional hazards model. A total of 3658 women and 4181 men died during the follow-up (end of follow-up was 15 April 2008 in the Danish sub-cohort, 15 December 2009 in the Norwegian sub-cohort and 15 February 2009 in the Swedish sub-cohort). In the analyses of continuous WG variables, we found lower all-cause mortality with higher intake of total WG products (women: MRR 0·89 (95 % CI 0·86, 0·91); men: MRR 0·89 (95 % CI 0·86, 0·91) for a doubling of intake). In particular, intake of breakfast cereals and non-white bread was associated with lower mortality. We also found lower all-cause mortality with total intake of different WG types (women: MRR 0·88 (95 % CI 0·86, 0·92); men: MRR 0·88 (95 % CI 0·86, 0·91) for a doubling of intake). In particular, WG oat, rye and wheat were associated with lower mortality. The associations were found in both women and men and for different causes of deaths. In the analyses of quartiles of WG intake in relation to all-cause mortality, we found lower mortality in the highest quartile compared with the lowest for breakfast cereals, non-white bread, total WG products, oat, rye (only men), wheat and total WG types. The MRR for highest *v.* lowest quartile of intake of total WG products was 0·68 (95 % CI 0·62, 0·75, $P_{\text{trend over quartiles}} < 0.0001$) for women and 0·75 (95 % CI 0·68, 0·81, $P_{\text{trend over quartiles}} < 0.0001$) for men. The MRR for highest *v.* lowest quartile of intake of total WG types was 0·74 (95 % CI 0·67, 0·81, $P_{\text{trend over quartiles}} < 0.0001$) for women and 0·75 (95 % CI 0·68, 0·82, $P_{\text{trend over quartiles}} < 0.0001$) for men. Despite lower statistical power, the analyses of cause-specific mortality according to quartiles of WG intake supported these results. In conclusion, higher intake of WG products and WG types was associated with lower mortality among participants in the HELGA cohort. The study indicates that intake of WG is an important aspect of diet in preventing early death in Scandinavia.

It has long been recognised that a fibre-rich diet is beneficial for health. As early as in the fourth century, the ancient Greek Physician and the 'Father of Western Medicine', Hippocrates, stated that 'To the human body it makes a great difference whether the bread be made of fine flour or coarse, whether with the bran or without the bran'[1]. Early pioneers within the field of nutritional epidemiology suggested that fibre-rich and unrefined foods are particularly

---

healthful and that highly refined foods are the cause of many chronic diseases in the Western part of the world[2-7]. Since then, a number of studies have found a lower mortality with high intake of dietary fibre, particularly from grain[7-9], just as intake of whole grain (WG) and WG products has been shown to be associated with lower risk of obesity/abdominal fatness[10], type 2 diabetes[11], hypertension[12], CVD[11] and colorectal cancer[13]. Clinical trials have shown that increasing the intake of WG lowers body weight[14-17], plasma insulin and insulin resistance[16,18-20], blood pressure[21-24], in addition to lipids, total cholesterol and LDL-cholesterol[14,15,21]. However, in studies with no concomitant effect on weight or waist circumference, intake of WG did not affect cardiovascular risk factors[25-28]. An effect on prostate-specific antigen has also been reported[19], indicating the potential effects on cancer progression as well. Only six studies (two of the references represent the same study) have investigated the intake of WG products in relation to the ultimate end point, mortality[29-35]. Except for one study[35] where intake of WG was not the primary focus, these studies have found beneficial/inverse associations between intake of WG products and all-cause or cause-specific mortality. One study[36] has investigated the intake of quantitative WG, cereal fibre, bran and germ in relation to all-cause and cause-specific mortality among 7827 US women with type 2 diabetes and has found lower all-cause and CVD mortality with higher intake of bran. However, most of the studies have been carried out in low-intake populations. To our knowledge, no study has yet investigated the association between WG types, i.e. WG from different cereal types in relation to mortality.

In Denmark, Norway and Sweden, WG is defined as whole, cracked, flaked or milled kernels of rye, oats, wheat, barley, rice, maize, millet and sorghum, where the bran, germ and endosperm are present in the same proportions as in the intact kernel[37]. This definition is similar to that established by the American Association of Cereal Chemist International[38], except for the inclusion of pseudo-cereals such as amaranth, buckwheat and quinoa in the American definition. WG is not only a rich source of dietary fibre, but also contains a range of vitamins, minerals and phytochemicals present in the outer, nutrient-rich layers[39].

We investigated the intake of different WG products and WG types in relation to all-cause and cause-specific mortality in a large combined Scandinavian cohort.

Materials and methods

Study population

The HELGA cohort is a large Scandinavian cohort combining three prospective studies: the Norwegian Women and Cancer Study[40], the Västerbotten Intervention Programme (VIP) cohort[41], and the Danish Diet Cancer and Health Study[42]. The three cohorts are also part of the European Prospective Investigation into Cancer and Nutrition study (EPIC)[43] and are described in detail elsewhere[41,42,44]. Briefly, participants were all recruited from the general

population in 1992–8, which included 37 231 Norwegian women aged 40–55 years; 13 294 and 12 431 Swedish women and men aged 30, 40, 50 or 60 years, respectively; in addition to 29 875 and 27 178 Danish women and men aged 50–64 years, respectively. In total, 197, 120 and 174 individuals from Denmark, Norway and Sweden, respectively, were excluded due to missing data on vital status, WG intake or potential confounders.

Ethics

The three cohorts were approved by the respective local ethics committees.

Data collection

At baseline, participants filled in a semi-quantitative, country-specific and validated FFQ regarding their habitual diet including WG and alcohol intake[41,44-46]. In Denmark, the FFQ contained 173 items, in Norway eighty-eight, and in Sweden ninety-eight, where the number of items was defined as the number of foods plus the number of standard mixed recipes. In addition, they filled in a comprehensive lifestyle questionnaire covering their lifestyle habits in addition to information on education and smoking intensity. Anthropometrical measurements were carried out by trained professionals at the study centres in Denmark and Sweden; in Norway, these measurements were self-reported.

Assessment of whole-grain intake

In the present study, WG products were assessed from questions referring to the intake of breakfast cereals, non-white bread and crisp bread. WG types refer to subfamilies of the Gramineae family, i.e. oat (Avena), wheat (Triticum), rye (Secale) or other grains, which consist of barley (Hordeum), rice (Oryza) and dried maize (Zea), with rice being the main contributor. Quantification of the intake of different types of WG was based on the intake of breakfast cereals, non-white bread and crisp bread after retrieval of data on the WG content of the three WG products/items from 24-h dietary recalls conducted in a random sample of 8716 participants of the HELGA cohort as part of the calibration study of the EPIC[47]. For each of the three WG food items, the WG content was calculated as a weighted mean from the different more specific foods recorded in the 24-h dietary recalls. These calculations were done country-specific due to differences in the products consumed in the three countries. In Norway and Sweden, the estimation was based on recipes, ingredient lists and data from WG product manufacturers. In Denmark, the estimation was based on a report from the Danish National Food Institute[48]. Values were expressed as the WG content of the unprepared ingredients divided by the weight of the prepared food as it is eaten. When comparing the WG intake measured using 24-h dietary recalls v. FFQ, 68 % of the participants had a WG intake in the same (±1) quintile[49]. In Denmark, two-thirds of WG bread intake was rye bread, whereas the largest contributor to WG

bread intake in Norway and Sweden was mixed-grain bread. In all the three countries, the category crisp bread consisted mostly of products with a high (75 %) WG content, whereas about 20 % was crisp bread with no or a low content of whole grains. Approximately 80 % of breakfast cereals consisted of muesli, porridge, oatmeal or other WG breakfast cereals, and the remaining 20 % were refined grain products such as cornflakes[49].

Ascertainment of mortality

Data on vital status was extracted from the National Central Population Registries and cause of death in the National Cause of Death Registries using the participants' personal identification number given to all citizens in Denmark, Norway and Sweden. End of follow-up was 15 April 2008 in the Danish sub-cohort, 15 December 2009 in the Norwegian sub-cohort and 15 February 2009 in the Swedish sub-cohort. In the Danish sub-cohort, 2173 women and 3333 men died during a median follow-up of 11·9 years. In the Norwegian sub-cohort, 966 women died during a median follow-up of 11·1 years. In the Swedish sub-cohort, 519 women and 848 men died during a median follow-up of 14·2 years.

Concerning cause-specific mortality, 1775 women and 1375 men died from cancer, 298 women and 858 men died from CHD, 137 women and 143 men died from stroke, 125 women and 111 men died from respiratory disease, twenty-four women and seventy men died from diabetes, and 1299 women and 1624 men died from other causes.

Assessment of covariates

Data on alcohol intake (g/d), BMI (kg/m²) and total energy intake (kJ/d) were retrieved from the FFQ. Data on education (none, primary, technical/professional, secondary and longer), smoking intensity (never; former, quit 20+ years; former, quit 11–20 years; former, quit ≤10 years; current, pipe/cigar/occasionally; current, 1–15 cigarettes /d; current, 16–25 cigarettes/d; current, 26+ cigarettes/d, unknown) and the Cambridge physical activity index[50] (a cross-tabulation of occupational activity and time spent on sports and cycling, only Denmark and Sweden) were retrieved from the lifestyle questionnaire.

Statistical analyses

The association between WG intake and sex-specific all-cause mortality was estimated on the basis of Cox proportional hazards models with age being the underlying time scale. Subjects were followed from the age at which they completed the questionnaires to their age at exit, defined as the age of death, emigration, loss to follow-up or end of follow-up, whichever came first. The hazard rate was allowed to change with time under study and was modelled as a linear spline with boundaries at 1, 2 and 3 years after entry to the cohort. Furthermore, the underlying hazard was stratified according to the study centre to control for differences in mortality, questionnaire design, follow-up procedures and

diagnosis or screening procedures. Cause-specific mortality was analysed using a competing risks model.

The WG variables (total WG product intake, different WG products, total intake of WG types or different WG types) were included simultaneously in the three models with different sets of adjusting variables. The basic model included all variables related to WG products or WG types in addition to age (time scale) and follow-up time (linear spline with boundaries at 1, 2 and 3 years after entry). The second model extended the first model by further including the potential confounders, education (categorical: none, primary, technical/professional, secondary, longer and unknown), smoking intensity (categorical: never; former, quit 20+ years; former, quit 11–20 years; former, quit ≤10 years; current, pipe/cigar/occasionally; current, 1–15 cigarettes/d; current, 16–25 cigarettes/d; current, 26+ cigarettes/d, unknown), alcohol intake (linear spline with boundary at 15 g/d), BMI (linear spline with boundaries at 18·5, 25 and 30 kg/m²) and total energy intake (continuous). In the third model, the potentially mediating variables in the causal pathway between WG and mortality, BMI and total energy intake were excluded. In order to allow WG to act via appetite regulation and BMI, the last model was considered the most appropriate.

The intake of 'other types of WG' was very low and was not related to mortality; therefore, 'other types of WG' was omitted from Tables 3–5, although the analyses are still adjusted for this variable, and it is still part of the 'total WG types' variable.

The association between WG variables and mortality seemed to level-off; consequently, a $\log_2$ transformation was applied. Estimates from the Cox regression models, therefore, corresponded to a doubling of the WG intake. Quantitative variables were included linearly in the Cox model after linearity had been evaluated by linear spline models[51,52]. No deviation from linearity was observed for any of the variables after transformation. Heterogeneity of the effects of WG intake on mortality was evaluated among the three countries, among women and men, as well as among WG types and WG products; no significant heterogeneity was found among WG products or types or among countries. Consequently, the amount of WG consumed could be added up across WG products, across types of WG and across countries.

Hazard rate ratios according to quartiles of specific WG products, specific WG types, total WG product intake and total intake of WG types were also estimated. To ensure an equal statistical power in the four quartiles, the definition of quartiles' cut-points/boundaries were based on the distribution of intake among individuals who died during the follow-up; consequently, the number of observations is the same in the four quartiles. Non-consumers of WG (individuals with a WG intake of 0 g/d) were included as part of the first quartile. Specific WG products or types were mutually adjusted and further adjusted for age, follow-up time, education, smoking intensity and alcohol intake.

A forest plot including sex- and country-specific mortality rate ratios for the comparison between the highest and the

lowest quartile of intake of total WG types (adjusted for age, follow-up time, education, smoking intensity and alcohol intake) was produced.

A range of sensitivity analyses were conducted to evaluate the stability of the results in subgroups of individuals (using the model including the intake of total WG types and all-cause mortality, adjusted for age, follow-up time, education, smoking intensity and alcohol intake): (1) the proportional hazards assumption was evaluated by performing separate analyses of deaths within three intervals of follow-up time (boundaries at 1, 2 and 3 years after baseline); (2) a possible interaction with age was investigated (using baseline age 50 years as cut-point) to examine whether older people may particularly benefit from intake of WG; (3) a possible interaction with BMI was investigated to examine whether overweight or obese individuals may benefit more from intake of WG than individuals with low or normal weight; (4) stratifications on physical activity (only Denmark and Sweden) and intake of fruits and nuts, vegetables, meat and alcohol were conducted to ensure that subgroups of individuals identified by their intake of vegetables, fruits and nuts, meat and alcohol (active/inactive or above/below the median intake) had the same mortality; (5) since data on physical activity was only available for Denmark and Sweden, a sensitivity analysis with adjustment for the Cambridge physical activity index[50] was performed in the Danish and Swedish data; (6) correspondingly, as the availability of data on cardiovascular risk factors such as hypertension, hyperlipidaemia and waist circumference as potential confounders was limited in Sweden and Norway, a sensitivity analysis with further adjustment for hypertension, hyperlipidaemia and waist circumference was conducted using the Danish data; (7) to evaluate potential residual confounding by smoking intensity or any diverging results between smokers and non-smokers, a sensitivity analysis was conducted in non-smokers (never smokers and former smokers); (8) to evaluate potential confounding by diet, the analyses were also adjusted for a seven-item diet score that was originally described by Trichopoulou et al.[53] and slightly modified for the present study to include the intake of fruits and nuts, vegetables, fish, meat, milk products, vegetable fat and alcohol. Each individual was assigned one point for each food item where she/he had an intake above the median for foods considered healthy (fruits and nuts, vegetables, fish and vegetable fat) and an intake below the median for foods considered unhealthy (meat, milk products and alcohol); finally (9) in the Danish data, we were able to exclude individuals with a previous diagnosis of CHD, stroke or angina pectoris, as they may not conform to the assumption of proportional hazards.

Two-sided 95 % CI for the mortality rate ratios (MRR) were estimated based on Wald's test of the Cox regression parameter on the log(MRR) scale, and tests of effect modification were performed using the likelihood ratio test. The SAS procedure PHREG on Windows platform was used for the statistical analyses (SAS version 9.3; SAS Institute). Forest plots (based on the results from the PHREG) were generated using the SAS procedure TEMPLATE.

Results

Descriptive analyses

The median age at recruitment was 51 years for women and 54 years for men, and the median BMI was 24 and 26 kg/m², respectively (Table 1). Alcohol intake was 2·4 g/d among women and 12 g/d among men, and total energy intake was 7·1 and 9·8 MJ, respectively. Moreover, 39 % of women and 31 % of men had never smoked, and among both women and men, about 10 % were current smokers and, in addition, smoked sixteen or more cigarettes per d. Regarding education, 27 % of women and 33 % of men had no education or primary school as their highest educational level.

The median intake of breakfast cereals in Swedish women and men was 29 and 31 g/d, respectively, and close to zero in the Danish and Norwegian cohorts (Table 2, upper part). Similarly, Swedish women and men had crisp bread intakes of 34 and 57 g/d, whereas the intakes in the two other cohorts were very low. In contrast, intake of non-white bread was low in the Swedish cohort and of a considerable size in the Danish and Norwegian cohorts (i.e. 103 and 118 g/d among Danish women and men, respectively, and 100 g/d among Norwegian women).

The median intake of WG oats was 2 g/d among both women and men in the combined HELGA cohort, with

Table 1. Baseline characteristics of cohort participants in the HELGA study

(Number of participants and percentages; medians and 5th−95th percentiles)

	Women		Men	
	n	%	n	%
Age (years)				
Median	51		54	
5th−95th percentile	40−63		31−64	
BMI (kg/m²)				
Median	24		26	
5th−95th percentile	20−33		21−33	
Alcohol intake (g/d)				
Median	2·4		12	
5th−95th percentile	0−32		0·4−69	
Energy intake (MJ/d)				
Median	7·1		9·8	
5th−95th percentile	4·3−11		5·8−15	
Smoking intensity				
Never	31 630	39	12 404	31
Former, quit 20+ years	6539	8	3978	10
Former, quit 11−20 years	5067	6	3525	9
Former, quit ≤10 years	7803	10	4281	11
Current, pipe/cigar/occas.	1852	2	5776	15
Current, 1−15 cig/d	15 830	20	3845	10
Current 16−25 cig/d	6898	9	3364	9
Current, 26+ cig/d	636	1	900	2
Unknown	3846	5	1344	2
Education				
None or primary school	21 739	27	13 068	33
Technical/professional	29 848	37	10 806	27
Secondary	17 702	22	5675	14
Longer education	10 812	14	9868	25

occas., Occasionally; cig, cigarettes.

Table 2. Country-specific intakes (g/d) of whole-grain products and whole-grain types of cohort participants in the HELGA study (Medians and 5th–95th percentiles)

| | Denmark | | | | Norway | | Sweden | | | | Total HELGA cohort | | | |
| | Women (n 29 787) | | Men (n 27 069) | | Women (n 37 111) | | Women (n 13 203) | | Men (n 12 348) | | Women (n 80 101) | | Men (n 39 417) | |
	Median	5th–95th percentile	Median	5th–95th percentile	Median	5th–95th percentile	Median	5th–95th percentile	Median	5th–95th percentile	Median	5th–95th percentile	Median	5th–95th percentile
Breakfast cereals	0·8	0–50	0·8	0–50	0	0–72	29	1–165	31	0·4–205	2	0–72	4	0–106
Non-white bread	103	26–213	118	28–263	100	14–240	18	0·3–125	9	0–143	100	3–213	90	0·3–231
Crisp bread	2	0·6–31	2	0·7–14	4	0–31	34	7–71	57	11–113	5	0–48	4	0·8–86
Total whole-grain products	116	38–236	137	44–277	131	35–245	94	26–251	114	30–318	121	34–244	131	35–294
Oat	1	0–30	0·9	0–30	1	0–22	5	0·3–25	5	0·4–31	2	0–30	2	0–31
Rye	21	5–53	33	6–62	10	4–26	28	7–64	50	8–96	17	4–48	37	6–72
Wheat	3	0–13	3	0–13	37	9–76	10	1–31	8	1–35	12	0·4–62	5	0·1–20
Other	0·4	0–2	1	0·1–4	0	0·5–4	3	1–9	5	1–15	0·1	0–0·5	0·1	0–0·9
Total whole-grain types	34	10–76	42	12–87	59	16–104	51	16–103	69	18–140	47	13–95	48	13–111

Sweden having a slightly higher intake than Denmark and Norway (Table 2, lower part). In all the three countries and in both women and men, a considerable intake of rye was observed. Swedish men had the highest intake (50 g/d) compared with Danish women and men (21 and 33 g/d), Norwegian women (10 g/d) and Swedish women (28 g/d). The median intake of WG wheat in the total cohort was 12 g/d among women and 5 g/d among men, and the higher intake among women confer to a rather high intake among Norwegian women (37 g/d). The intake of WG from sources other than oat, rye and wheat was very low (women: median 0·1 (5th–95th percentile 0–0·5) g/d; men: median 0·1 (5th–95th percentile 0–0·9) g/d). The total median intake of different types of WG in the combined HELGA cohort was 47 g/d among women and 48 g/d among men.

All-cause mortality

The MRR and 95 % CI for all-cause mortality in relation to intake of WG are presented in Table 3. For the association between all-cause mortality and intake of WG products, we found lower mortality for each doubling of both breakfast cereals (women: MRR 0·95 (95 % CI 0·94, 0·97); men: MRR 0·96 (95 % CI 0·94, 0·97)) and non-white bread (women: MRR 0·94 (95 % CI 0·92, 0·96); men: MRR 0·97 (95 % CI 0·95, 0·99)). For the intake of total WG products and all-cause mortality, we found lower all-cause mortality for each doubling of total WG product intake (women: MRR 0·89 (95 % CI 0·86, 0·91), men: MRR 0·89 (95 % CI 0·86, 0·91)).

Intake of WG oats, rye and wheat was associated with a statistically significant lower all-cause mortality in both women and men. A doubling of the intake of WG oats was associated with the MRR of 0·98 (95 % CI 0·96, 0·99) for women and 0·98 (95 % CI 0·97, 1·00) for men. A doubling of the intake of WG rye was associated with the MRR of 0·96 (95 % CI 0·93, 0·99) in women and 0·95 (95 % CI 0·92, 0·97) in men. A doubling of the intake of WG wheat was associated with the MRR of 0·93 (95 % CI 0·92, 0·95) for women and 0·96 (95 % CI 0·95, 0·98) for men. A doubling of the total intake of different types of WG cereals was also associated with statistically significant lower all-cause mortality in both women and men (women: 0·88 (95 % CI 0·86, 0·92); men: 0·88 (95 % CI 0·86, 0·91) for a doubling of intake).

In general, adjustment for potential confounders did not change the estimates considerably.

Cancer mortality

A total of 1775 women and 1375 men died from cancer. In the cause-specific associations between WG products and cancer mortality (Table 4), we found lower mortality with higher intake of breakfast cereals (women: MRR 0·97 (95 % CI 0·95, 0·99); men: MRR 0·95 (95 % CI 0·93, 0·97) for a doubling of intake), non-white bread (women: MRR 0·97 (95 % CI 0·94, 1·00); men: MRR 0·96 (95 % CI 0·92, 0·99) for a doubling of intake) and total WG products (women: MRR 0·94 (95 % CI 0·90, 0·99); men: MRR 0·88 (95 % CI 0·84, 0·93) for a doubling of intake). We found a borderline statistically significant

Table 3. The association between intake of whole grain and all-cause mortality of cohort participants in the HELGA study
(Mortality rate ratios (MRR) and 95 % confidence intervals)

	Women (deaths = 3658)						Men (deaths = 4181)					
All-cause mortality	MRR*	95 % CI	MRR†	95 % CI	MRR‡	95 % CI	MRR*	95 % CI	MRR†	95 % CI	MRR‡	95 % CI
Per doubling of whole-grain product intake												
Breakfast cereals	0·95	0·94, 0·96	0·95	0·94, 0·97	0·95	0·94, 0·97	0·95	0·93, 0·96	0·96	0·94, 0·97	0·96	0·94, 0·97
Non-white bread	0·93	0·91, 0·95	0·94	0·92, 0·96	0·94	0·92, 0·96	0·96	0·94, 0·98	0·97	0·95, 0·99	0·97	0·95, 0·99
Crisp bread	0·98	0·96, 0·99	0·98	0·96, 1·00	0·98	0·96, 1·00	0·98	0·95, 1·00	0·98	0·95, 1·00	0·99	0·96, 1·01
Total whole-grain products	0·88	0·85, 0·90	0·88	0·85, 0·91	0·89	0·85, 0·91	0·85	0·83, 0·88	0·87	0·85, 0·90	0·89	0·86, 0·91
Per doubling of whole-grain type intake												
Oat	0·98	0·96, 0·99	0·98	0·96, 0·99	0·98	0·96, 0·99	0·98	0·97, 1·00	0·98	0·97, 1·00	0·98	0·97, 1·00
Rye	0·96	0·93, 0·99	0·95	0·92, 0·98	0·96	0·93, 0·99	0·94	0·91, 0·96	0·93	0·91, 0·96	0·95	0·92, 0·97
Wheat	0·92	0·91, 0·94	0·93	0·92, 0·94	0·93	0·92, 0·95	0·95	0·93, 0·96	0·96	0·95, 0·98	0·96	0·95, 0·98
Total whole-grain types§	0·87	0·85, 0·90	0·88	0·85, 0·91	0·88	0·86, 0·92	0·85	0·82, 0·87	0·87	0·84, 0·90	0·88	0·86, 0·91

* Whole-grain products and whole-grain types (including 'other types' of whole grain) mutually adjusted and adjusted for age (time scale) and follow-up time (linear spline with boundaries at 1, 2 and 3 years after entry).
† First model/column with further adjustment for education (categorical: none, primary, technical/professional, secondary and longer), smoking intensity (categorical: never; former, quit 20 + years; former, quit 11–20 years; former, quit ≤10 years; current; pipe/cigar/occasionally; current, 1–15 cigarettes/d; current, 16–25 cigarettes/d; current, 26+ cigarettes/d; unknown), alcohol intake (linear spline with boundary at 15g/d), BMI (linear spline with boundaries at 18·5, 25 and 30kg/m²) and total energy intake (continuous: kJ/d).
‡ Second model/column without adjustment for the potentially mediating variables, BMI and total energy intake.
§ Sum of intake of oat, rye, wheat and other types of whole grain.

association between the intake of crisp bread and cancer mortality (women: MRR 0·98 (95 % CI 0·95, 1·00); men: MRR 0·96 (95 % CI 0·93, 1·00) for a doubling of intake). When investigating the association between the intake of WG types and cancer mortality, we found lower mortality for each doubling of the WG wheat intake among both women and men (women: MRR 0·95 (95 % CI 0·93, 0·97); men: MRR 0·95 (95 % CI 0·93, 0·97) for a doubling of intake). The intake of total WG types was associated with lower cancer mortality in men (MRR 0·89 (95 % CI 0·84, 0·93) for a doubling of intake).

CHD mortality

A total of 298 women and 858 men died from CHD. In the cause-specific associations between WG products and CHD mortality (Table 4), we found lower CHD mortality with higher intake of breakfast cereals (women: MRR 0·91 (95 % CI 0·86, 0·96); men: MRR 0·96 (95 % CI 0·93, 0·99) for a doubling of intake), total WG products (women: MRR 0·85 (95 % CI 0·77, 0·94); men: MRR 0·92 (95 % CI 0·86, 0·99) for a doubling of intake) and total WG types (women: MRR 0·82 (95 % CI 0·74, 0·91); men: MRR 0·90 (95 % CI 0·84, 0·96) for a doubling of intake). Among women, intake of oat and wheat was associated with a borderline statistically significantly lower CHD mortality (WG oat: MRR 0·95 (95 % CI 0·90, 1·00); WG wheat: MRR 0·95 (95 % CI 0·90, 1·00)).

Stroke mortality

A total of 137 women and 143 men died from stroke. When investigating the cause-specific associations between the intake of WG products and stroke mortality (Table 4), we found lower stroke mortality with higher intake of total WG products (MRR 0·84 (95 % CI 0·72, 0·98) for a doubling of intake) and total WG types (MRR 0·82 (95 % CI 0·71, 0·95) for a doubling of intake) among men.

Respiratory disease mortality

A total of 125 women and 111 men died from respiratory disease. In the cause-specific associations between respiratory disease mortality and intake of WG types or WG products (Table 4), only breakfast cereals, total WG types and WG wheat was associated with lower mortality from respiratory disease in women (breakfast cereals: MRR 0·89 (95 % CI 0·82, 0·98); total WG types: MRR 0·84 (95 % CI 0·71, 0·99); WG wheat: MRR 0·88 (95 % CI 0·83, 0·94) for a doubling of intake).

Diabetes mortality

A total of twenty-four women and seventy men died from diabetes. In the cause-specific associations between diabetes mortality and intake of WG types or WG products (Table 4), we found lower diabetes mortality with higher intake of breakfast cereals (women: MRR 0·79 (95 % CI 0·62, 0·99); men: MRR 0·88 (95 % CI 0·78, 0·98) for a doubling of intake). Among men, higher intake of non-white bread was

Table 4. The association between intake of whole grain and cause-specific mortality of cohort participants in the HELGA study

(Mortality rate ratios (MRR) and 95 % confidence intervals)

	MRR*	95 % CI	MRR†	95 % CI	MRR‡	95 % CI	MRR*	95 % CI	MRR†	95 % CI	MRR‡	95 % CI
Cancer mortality	Women (deaths = 1775)						Men (deaths = 1375)					
Per doubling of whole-grain product intake												
Breakfast cereals	0·97	0·95, 0·99	0·97	0·95, 0·99	0·97	0·95, 0·99	0·94	0·92, 0·96	0·95	0·93, 0·97	0·95	0·93, 0·97
Non-white bread	0·96	0·93, 0·99	0·97	0·93, 1·00	0·97	0·94, 1·00	0·94	0·90, 0·97	0·95	0·92, 0·99	0·96	0·92, 0·99
Crisp bread	0·97	0·95, 1·00	0·98	0·95, 1·00	0·98	0·95, 1·00	0·96	0·92, 1·00	0·96	0·92, 1·00	0·96	0·93, 1·00
Total whole-grain products	0·93	0·89, 0·97	0·94	0·89, 0·99	0·94	0·90, 0·99	0·85	0·80, 0·89	0·88	0·83, 0·93	0·88	0·84, 0·93
Per doubling of whole-grain type intake												
Oat	0·98	0·96, 1·00	0·98	0·96, 1·00	0·98	0·96, 1·00	0·98	0·96, 1·01	0·98	0·96, 1·01	0·99	0·96, 1·01
Rye	1·01	0·96, 1·06	1·00	0·95, 1·05	1·01	0·96, 1·06	0·95	0·90, 0·99	0·94	0·90, 0·99	0·96	0·91, 1·00
Wheat	0·94	0·92, 0·96	0·95	0·92, 0·97	0·95	0·93, 0·97	0·93	0·91, 0·95	0·95	0·93, 0·97	0·95	0·93, 0·97
Total whole-grain types§	0·95	0·90, 1·00	0·97	0·91, 1·02	0·96	0·91, 1·01	0·84	0·80, 0·89	0·88	0·83, 0·93	0·89	0·84, 0·93
CHD mortality	Women (deaths = 298)						Men (deaths = 858)					
Per doubling of whole-grain product intake												
Breakfast cereals	0·91	0·86, 0·95	0·91	0·86, 0·96	0·91	0·86, 0·96	0·95	0·93, 0·98	0·96	0·94, 0·99	0·96	0·93, 0·99
Non-white bread	0·93	0·86, 0·99	0·94	0·87, 1·02	0·94	0·88, 1·01	0·99	0·95, 1·03	1·00	0·95, 1·04	1·00	0·96, 1·05
Crisp bread	0·98	0·92, 1·05	0·98	0·92, 1·05	0·99	0·92, 1·05	1·00	0·95, 1·05	0·99	0·94, 1·05	1·02	0·97, 1·07
Total whole-grain products	0·83	0·76, 0·92	0·86	0·77, 0·95	0·85	0·77, 0·94	0·90	0·84, 0·97	0·91	0·84, 0·98	0·92	0·86, 0·99
Per doubling of whole-grain type intake												
Oat	0·95	0·90, 1·00	0·95	0·90, 1·01	0·95	0·90, 1·00	0·98	0·95, 1·01	0·98	0·95, 1·02	0·98	0·95, 1·01
Rye	0·91	0·82, 1·01	0·90	0·81, 1·00	0·91	0·82, 1·01	0·97	0·91, 1·03	0·95	0·89, 1·01	0·97	0·91, 1·03
Wheat	0·93	0·88, 0·98	0·95	0·90, 1·00	0·95	0·90, 1·00	0·96	0·93, 0·99	0·98	0·95, 1·00	0·98	0·95, 1·01
Total whole-grain types§	0·81	0·73, 0·90	0·82	0·73, 0·92	0·82	0·74, 0·91	0·88	0·82, 0·95	0·88	0·82, 0·95	0·90	0·84, 0·96
Stroke mortality	Women (deaths = 137)						Men (deaths = 143)					
Per doubling of whole-grain product intake												
Breakfast cereals	0·94	0·87, 1·01	0·93	0·86, 1·07	0·94	0·87, 1·02	0·97	0·91, 1·05	0·98	0·91, 1·05	0·98	0·91, 1·05
Non-white bread	0·94	0·84, 1·05	0·92	0·83, 1·03	0·94	0·85, 1·05	0·97	0·87, 1·08	0·97	0·87, 1·09	0·98	0·88, 1·10
Crisp bread	1·05	0·95, 1·16	1·04	0·94, 1·14	1·05	0·95, 1·16	0·95	0·84, 1·08	0·95	0·83, 1·08	0·96	0·85, 1·09
Total whole-grain products	0·89	0·76, 1·05	0·85	0·72, 1·01	0·90	0·76, 1·06	0·81	0·70, 0·95	0·80	0·68, 0·94	0·84	0·72, 0·98
Per doubling of whole-grain type intake												
Oat	0·93	0·86, 1·01	0·93	0·86, 1·00	0·93	0·86, 1·01	1·02	0·94, 1·10	1·02	0·94, 1·10	1·02	0·94, 1·10
Rye	0·97	0·83, 1·14	0·94	0·80, 1·11	0·97	0·83, 1·14	0·87	0·77, 1·00	0·86	0·75, 0·99	0·88	0·77, 1·01
Wheat	1·00	0·91, 1·09	0·99	0·91, 1·08	1·00	0·92, 1·09	0·95	0·88, 1·01	0·96	0·89, 1·03	0·96	0·90, 1·03
Total whole-grain types§	0·85	0·72, 1·01	0·80	0·67, 0·96	0·86	0·72, 1·01	0·79	0·68, 0·92	0·78	0·67, 0·91	0·82	0·71, 0·95
Respiratory disease mortality	Women (deaths = 125)						Men (deaths = 111)					
Per doubling of whole-grain product intake												
Breakfast cereals	0·88	0·81, 0·96	0·89	0·82, 0·97	0·89	0·82, 0·98	0·92	0·84, 1·00	0·92	0·84, 1·00	0·93	0·85, 1·01
Non-white bread	0·89	0·78, 1·01	0·92	0·81, 1·06	0·92	0·80, 1·05	0·98	0·85, 1·12	0·99	0·86, 1·13	1·01	0·88, 1·16
Crisp bread	0·98	0·88, 1·10	1·02	0·92, 1·14	1·00	0·90, 1·11	0·98	0·85, 1·12	1·00	0·87, 1·15	1·00	0·87, 1·15
Total whole-grain products	0·80	0·68, 0·94	0·82	0·69, 0·98	0·84	0·71, 0·99	0·82	0·69, 0·98	0·84	0·70, 1·00	0·88	0·74, 1·05
Per doubling of whole-grain type intake												
Oat	0·96	0·88, 1·05	0·94	0·86, 1·03	0·96	0·87, 1·05	0·92	0·84, 1·01	0·91	0·83, 0·99	0·92	0·84, 1·01
Rye	1·04	0·87, 1·24	1·02	0·86, 1·21	1·05	0·88, 1·24	0·99	0·83, 1·17	0·94	0·80, 1·11	1·00	0·85, 1·18
Wheat	0·86	0·80, 0·91	0·89	0·84, 0·95	0·88	0·83, 0·94	0·98	0·91, 1·05	1·01	0·94, 1·09	1·01	0·94, 1·09
Total whole-grain types§	0·85	0·71, 1·02	0·88	0·72, 1·07	0·89	0·74, 1·08	0·82	0·69, 0·98	0·83	0·70, 0·98	0·88	0·74, 1·04

Table 4. *Continued*

	MRR*	95 % CI	MRR†	95 % CI	MRR‡	95 % CI	MRR*	95 % CI	MRR†	95 % CI	MRR‡	95 % CI
Diabetes mortality	Women (deaths = 24)						Men (deaths = 70)					
Per doubling of whole-grain product intake												
Breakfast cereals	0·78	0·62, 0·99	0·79	0·62, 1·00	0·79	0·62, 0·99	0·87	0·77, 0·97	0·89	0·79, 1·00	0·88	0·78, 0·98
Non-white bread	0·92	0·64, 1·29	0·93	0·65, 1·33	0·95	0·68, 1·34	1·27	1·01, 1·60	1·34	1·05, 1·71	1·30	1·03, 1·64
Crisp bread	0·98	0·77, 1·24	0·96	0·76, 1·23	0·98	0·78, 1·23	1·14	0·97, 1·34	1·19	1·01, 1·40	1·16	0·99, 1·36
Total whole-grain products	0·74	0·53, 1·02	0·77	0·54, 1·09	0·77	0·56, 1·07	1·16	0·87, 1·54	1·27	0·93, 1·74	1·22	0·92, 1·62
Per doubling of whole-grain type intake												
Oat	0·86	0·69, 1·06	0·86	0·69, 1·06	0·85	0·68, 1·05	0·92	0·82, 1·02	0·91	0·82, 1·02	0·92	0·82, 1·03
Rye	0·95	0·66, 1·37	0·95	0·66, 1·37	0·97	0·68, 1·38	1·10	0·87, 1·39	1·06	0·84, 1·34	1·10	0·87, 1·39
Wheat	0·87	0·75, 1·01	0·89	0·77, 1·03	0·89	0·77, 1·04	1·00	0·91, 1·09	1·02	0·93, 1·12	1·02	0·93, 1·12
Total whole-grain types§	0·72	0·51, 1·03	0·74	0·50, 1·08	0·74	0·52, 1·06	1·04	0·80, 1·37	1·11	0·82, 1·48	1·09	0·83, 1·43
Other causes of mortality	Women (deaths = 1299)						Men (deaths = 1624)					
Per doubling of whole-grain product intake												
Breakfast cereals	0·95	0·92, 0·97	0·95	0·93, 0·98	0·95	0·93, 0·98	0·95	0·93, 0·97	0·96	0·94, 0·99	0·96	0·94, 0·98
Non-white bread	0·90	0·87, 0·93	0·90	0·87, 0·94	0·91	0·88, 0·94	0·94	0·91, 0·98	0·96	0·93, 0·99	0·96	0·93, 1·00
Crisp bread	0·97	0·95, 1·00	0·97	0·95, 1·00	0·98	0·95, 1·01	0·98	0·94, 1·01	0·98	0·94, 1·02	0·99	0·95, 1·02
Total whole-grain products	0·82	0·79, 0·86	0·82	0·79, 0·86	0·83	0·80, 0·87	0·83	0·79, 0·87	0·86	0·82, 0·90	0·87	0·83, 0·91
Per doubling of whole-grain type intake												
Oat	0·98	0·96, 1·01	0·98	0·96, 1·01	0·98	0·96, 1·01	0·99	0·96, 1·01	0·99	0·97, 1·01	0·99	0·97, 1·01
Rye	0·91	0·86, 0·95	0·90	0·85, 0·94	0·91	0·87, 0·96	0·91	0·88, 0·95	0·91	0·88, 0·95	0·93	0·90, 0·96
Wheat	0·91	0·89, 0·93	0·91	0·89, 0·94	0·92	0·89, 0·94	0·95	0·93, 0·97	0·97	0·95, 0·99	0·97	0·95, 0·99
Total whole-grain types§	0·82	0·78, 0·86	0·82	0·78, 0·87	0·83	0·79, 0·88	0·83	0·79, 0·87	0·87	0·83, 0·91	0·88	0·84, 0·92

* Whole-grain products and whole-grain types (including 'other types' of whole grain) mutually adjusted and adjusted for age (time scale) and follow-up time (linear spline with boundaries at 1, 2 and 3 years after entry).

† First model/column with further adjustment for education (categorical: none, primary, technical/professional, secondary, longer), smoking intensity (categorical: never; former, quit 20 + years; former, quit 11−20 years; former, quit ≤10 years; current, pipe/cigar/occasionally; current, 1−15 cigarettes/d; current, 16−25 cigarettes/d; current, 26 + cigarettes/d, unknown), alcohol intake (linear spline with boundary at 15 g/d), BMI (linear spline with boundaries at 18·5, 25 and 30 kg/m²) and total energy intake (continuous: kJ/d).

‡ Second model/column without adjustment for the potentially mediating variables, BMI and total energy intake.

§ Sum of intake of oat, rye, wheat and other types of whole grain.

Table 5. The association between quartiles of whole-grain intake and all-cause mortality of cohort participants in the HELGA study

(Mortality rate ratios (MRR) and 95 % confidence intervals)

All-cause mortality	Quartile 1 MRR*	Quartile 2 MRR*	Quartile 2 95 % CI	Quartile 3 MRR*	Quartile 3 95 % CI	Quartile 4 MRR*	Quartile 4 95 % CI	$P_{\text{trend over quartiles}}$
Women (deaths = 3658)†								
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain product intake								
Breakfast cereals	1	0·77	0·67, 0·89	0·79	0·71, 0·88	0·75	0·69, 0·82	<0·0001
Non-white bread	1	0·84	0·76, 0·93	0·74	0·65, 0·84	0·72	0·65, 0·81	<0·0001
Crisp bread	1	0·90	0·82, 0·99	0·86	0·78, 0·94	0·91	0·81, 1·01	0·012
Total whole-grain products	1	0·78	0·71, 0·86	0·77	0·71, 0·85	0·68	0·62, 0·75	<0·0001
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain type intake								
Oat	1	0·85	0·78, 0·93	0·74	0·67, 0·82	0·78	0·70, 0·87	<0·0001
Rye	1	0·92	0·84, 1·01	0·81	0·73, 0·90	0·93	0·83, 1·03	0·134
Wheat	1	0·72	0·65, 0·79	0·65	0·58, 0·72	0·63	0·53, 0·74	<0·0001
Total whole-grain types‡	1	0·80	0·73, 0·87	0·74	0·67, 0·81	0·74	0·67, 0·81	<0·0001
Men (deaths = 4181)†								
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain product intake								
Breakfast cereals	1	0·92	0·82, 1·04	0·82	0·76, 0·89	0·74	0·68, 0·81	<0·0001
Non-white bread	1	0·92	0·83, 1·03	0·85	0·75, 0·95	0·78	0·69, 0·88	<0·0001
Crisp bread	1	0·97	0·89, 1·06	0·94	0·86, 1·02	1·03	0·90, 1·17	0·617
Total whole-grain products	1	0·87	0·80, 0·95	0·74	0·68, 0·81	0·75	0·68, 0·81	<0·0001
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain type intake								
Oat	1	0·87	0·80, 0·95	0·85	0·77, 0·94	0·76	0·69, 0·85	<0·0001
Rye	1	0·91	0·83, 1·00	0·82	0·74, 0·91	0·86	0·78, 0·95	0·001
Wheat	1	0·87	0·80, 0·95	0·76	0·69, 0·84	0·71	0·64, 0·78	<0·0001
Total whole-grain types‡	1	0·82	0·75, 0·90	0·72	0·66, 0·78	0·75	0·68, 0·82	<0·0001

* Whole-grain products and whole-grain types (including 'other types' of whole grain) mutually adjusted and adjusted for age (time scale), follow-up time (linear spline with boundaries at 1, 2 and 3 years after entry), education (categorical: none, primary, technical/professional, secondary, longer), smoking intensity (categorical: never; former, quit 20+ years; former, quit 11–20 years; former, quit ≤10 years; current, pipe/cigar/occasionally; current, 1–15 cigarettes/d; current, 16–25 cigarettes/d; current, 26+ cigarettes/d, unknown), alcohol (linear spline with boundary at 15 g/d), BMI (linear spline with boundaries at 18·5, 25 and 30 kg/m^2), total energy intake (continuous: kJ/d) and alcohol intake (g/d).

† Median intakes of whole grain in each quartile in online Appendix Table.

‡ Sum of oat, rye, wheat and other types of whole grain.

associated with higher diabetes mortality (MRR 1·30 (95 % CI 1·03, 1·64) for a doubling of intake).

Other causes of mortality

A total of 1299 women and 1624 men died from other causes (i.e. causes other than cancer, CHD, stroke, respiratory disease and diabetes). We also investigated other causes of mortality in relation to intake of WG types and WG products (Table 4); here, we found intake of breakfast cereals (women: MRR 0·95 (95 % CI 0·93, 0·97); men: MRR 0·96 (95 % CI 0·94, 0·98) for a doubling of intake), non-white bread (women: MRR 0·91 (95 % CI 0·88, 0·94); men: MRR 0·96 (95 % CI 0·93, 1·00) for a doubling of intake) and total WG products (women: MRR 0·83 (95 % CI 0·80, 0·87); men: MRR 0·87 (95 % CI 0·83, 0·91) for a doubling of intake) to be associated with lower mortality from other causes. In analyses of different types of WG, intake of WG rye and wheat was associated with lower mortality (rye, women: MRR 0·91 (95 % CI 0·87, 0·96), men: MRR 0·93 (95 % CI 0·90, 0·96); wheat, women: MRR 0·92 (95 % CI 0·89, 0·94), men: MRR 0·97 (95 % CI 0·95, 0·99) where all estimates corresponds to a doubling of intake). The total intake of different WG cereal types was also associated with lower mortality from other causes (women: MRR 0·83 (95 % CI 0·79, 0·88); men: MRR 0·88 (95 % CI 0·84, 0·92) for a doubling of intake).

Quartiles of whole-grain intake

In the analyses of quartiles of WG intake in relation to all-cause mortality, we found lower mortality in the highest quartile compared with the lowest quartile for breakfast cereals, non-white bread, total WG products, oat, rye (only men), wheat and total WG types (Table 5).

The MRR for highest *v.* lowest quartile of intake of breakfast cereals was 0·75 (95 % CI 0·69, 0·82, $P_{\text{trend over quartiles}}$ < 0·0001) for women and 0·74 (95 % CI 0·68, 0·81, $P_{\text{trend over quartiles}}$ < 0·0001) for men. The MRR for highest *v.* lowest quartile of intake of non-white bread was 0·72 (95 % CI 0·65, 0·81, $P_{\text{trend over quartiles}}$ < 0·0001) for women and 0·78 (95 % CI 0·69, 0·88, $P_{\text{trend over quartiles}}$ < 0·0001) for men. MRR for highest *v.* lowest quartile of intake of total WG products was 0·68 (95 % CI 0·62, 0·75, $P_{\text{trend over quartiles}}$ < 0·0001) for women and 0·75 (95 % CI 0·68, 0·81, $P_{\text{trend over quartiles}}$ < 0·0001) for men.

The MRR for highest *v.* lowest quartile of intake of oat was 0·78 (95 % CI 0·70, 0·87, $P_{\text{trend over quartiles}}$ < 0·0001) for women and 0·76 (95 % CI 0·69, 0·85, $P_{\text{trend over quartiles}}$ < 0·0001) for men. The MRR for highest *v.* lowest quartile of intake of rye among men was 0·86 (95 % CI 0·78, 0·95, $P_{\text{trend over quartiles}}$ = 0·001). The MRR for highest *v.* lowest quartile of intake of wheat was 0·63 (95 % CI 0·53, 0·74, $P_{\text{trend over quartiles}}$ < 0·0001) for women and 0·71 (95 % CI 0·64,

0·78, $P_{\text{trend over quartiles}} < 0\cdot0001$) for men. The MRR for highest *v.* lowest quartile of intake of total WG types was 0·74 (95 % CI 0·67, 0·81, $P_{\text{trend over quartiles}} < 0\cdot0001$) for women and 0·75 (95 % CI 0·68, 0·82, $P_{\text{trend over quartiles}} < 0\cdot0001$) for men.

In the analyses of quartiles of WG intake in relation to cause-specific mortality, we found lower cancer mortality with higher intake of breakfast cereals, total WG products, oat and wheat (Table 6). Among men, the intake of non-white bread and total WG types was also associated with lower cancer mortality. The MRR for highest *v.* lowest quartile of intake of breakfast cereals was 0·85 (95 % CI 0·74, 0·98, $P_{\text{trend over quartiles}} = 0\cdot003$) for women and 0·75 (95 % CI 0·64, 0·87, $P_{\text{trend over quartiles}} < 0\cdot0001$) for men. The MRR for highest *v.* lowest quartile of intake of total WG products was 0·86 (95 % CI 0·74, 0·99, $P_{\text{trend over quartiles}} = 0\cdot0236$) for women and 0·70 (95 % CI 0·60, 0·81, $P_{\text{trend over quartiles}} < 0\cdot0001$) for men. The MRR for highest *v.* lowest quartile of intake of oat was 0·84 (95 % CI 0·71, 0·99, $P_{\text{trend over quartiles}} = 0\cdot007$) for women and 0·75 (95 % CI 0·64, 0·89, $P_{\text{trend over quartiles}} = 0\cdot003$) for men. The MRR for highest *v.* lowest quartile of intake of wheat was 0·74 (95 % CI 0·61, 0·89, $P_{\text{trend over quartiles}} = 0\cdot006$) for women and 0·58 (95 % CI 0·49, 0·69, $P_{\text{trend over quartiles}} < 0\cdot0001$) for men. Among men, the MRR for highest *v.* lowest quartile of intake of non-white bread was 0·79 (95 % CI 0·64, 0·97, $P_{\text{trend over quartiles}} = 0\cdot019$) and 0·74 (95 % CI 0·63, 0·87, $P_{\text{trend over quartiles}} < 0\cdot0001$) for highest *v.* lowest intake of total WG types.

In the analyses of quartiles of WG intake according to CHD mortality, we found lower CHD mortality with higher intake of breakfast cereals and total WG types. Among women, the intake of total WG products, oat and wheat (borderline) was also associated with lower CHD mortality. The MRR for highest *v.* lowest quartile of intake of breakfast cereals was 0·53 (95 % CI 0·37, 0·77, $P_{\text{trend over quartiles}} = 0\cdot0002$) for women and 0·75 (95 % CI 0·61, 0·91, $P_{\text{trend over quartiles}} = 0\cdot009$) for men. The MRR for highest *v.* lowest quartile of intake of total WG types was 0·65 (95 % CI 0·46, 0·91, $P_{\text{trend over quartiles}} = 0\cdot003$) for women and 0·74 (95 % CI 0·61, 0·91, $P_{\text{trend over quartiles}} = 0\cdot012$) for men. Among women, the MRR for highest *v.* lowest quartile of intake of breakfast cereals was 0·53 (95 % CI 0·37, 0·77, $P_{\text{trend over quartiles}} = 0\cdot0002$). The MRR for highest *v.* lowest quartile of intake of oat was 0·66 (95 % CI 0·45, 0·96, $P_{\text{trend over quartiles}} = 0\cdot048$). The MRR for highest *v.* lowest quartile of intake of wheat was 0·67 (95 % CI 0·45, 1·00, $P_{\text{trend over quartiles}} = 0\cdot062$).

In the analyses of quartiles of WG intake according to stroke mortality, we did not find any statistically significant associations between WG intake and stroke mortality.

In the analyses of quartiles of WG intake according to respiratory disease mortality, we found lower respiratory disease mortality with higher intake of oat. Among women, intake of breakfast cereals and total WG types was also associated with lower respiratory disease mortality. The MRR for highest *v.* lowest quartile of intake of oat was 0·51 (95 % CI 0·28, 0·93, $P_{\text{trend over quartiles}} = 0\cdot006$) for women and 0·52 (95 % CI 0·28, 0·97, $P_{\text{trend over quartiles}} = 0\cdot063$) for men. Among women, the MRR for highest *v.* lowest quartile of intake of breakfast cereals was 0·39 (95 % CI 0·23, 0·65, $P_{\text{trend over quartiles}} < 0\cdot0001$)

and 0·47 (95 % CI 0·28, 0·77, $P_{\text{trend over quartiles}} = 0\cdot005$) for highest *v.* lowest intake of total WG types.

In the analyses of quartiles of WG intake according to diabetes mortality, we found lower diabetes mortality with higher intake of breakfast cereals and oat among men. The MRR for highest *v.* lowest quartile of intake of breakfast cereals was 0·45 (95 % CI 0·23, 0·90, $P_{\text{trend over quartiles}} = 0\cdot004$), and the MRR for highest *v.* lowest quartile of intake of oat was 0·40 (95 % CI 0·20, 0·81, $P_{\text{trend over quartiles}} = 0\cdot003$).

In the analyses of quartiles of WG intake according to other causes of mortality, we found lower mortality from other causes with higher intake of breakfast cereals, non-white bread, total WG products, oat (only borderline among men), rye, wheat and total WG types. The MRR for highest *v.* lowest quartile of intake of breakfast cereals was 0·74 (95 % CI 0·65, 0·84, $P_{\text{trend over quartiles}} < 0\cdot0001$) for women and 0·78 (95 % CI 0·68, 0·90, $P_{\text{trend over quartiles}} = 0\cdot0002$) for men. The MRR for highest *v.* lowest quartile of intake of non-white bread was 0·54 (95 % CI 0·45, 0·64, $P_{\text{trend over quartiles}} < 0\cdot0001$) for women and 0·65 (95 % CI 0·54, 0·79, $P_{\text{trend over quartiles}} < 0\cdot0001$) for men. The MRR for highest *v.* lowest quartile of intake of total WG products was 0·57 (95 % CI 0·47, 0·68, $P_{\text{trend over quartiles}} < 0\cdot0001$) for women and 0·70 (95 % CI 0·61, 0·80, $P_{\text{trend over quartiles}} < 0\cdot0001$) for men. The MRR for highest *v.* lowest quartile of intake of oat was 0·81 (95 % CI 0·67, 0·98, $P_{\text{trend over quartiles}} = 0\cdot0006$) for women and 0·85 (95 % CI 0·73, 1·00, $P = 0\cdot107$) for men. The MRR for highest *v.* lowest quartile of intake of rye was 0·81 (95 % CI 0·68, 0·97, $P_{\text{trend over quartiles}} = 0\cdot007$) for women and 0·76 (95 % CI 0·65, 0·89, $P_{\text{trend over quartiles}} = 0\cdot0003$) for men. The MRR for highest *v.* lowest quartile of intake of wheat was 0·51 (95 % CI 0·39, 0·67, $P_{\text{trend over quartiles}} < 0\cdot0001$) for women and 0·75 (95 % CI 0·64, 0·88, $P_{\text{trend over quartiles}} = 0\cdot0009$) for men. The MRR for highest *v.* lowest quartile of intake of total WG types was 0·61 (95 % CI 0·51, 0·73, $P_{\text{trend over quartiles}} < 0\cdot0001$) for women and 0·71 (95 % CI 0·62, 0·82, $P_{\text{trend over quartiles}} < 0\cdot0001$) for men.

Country-specific results

The forest plots in Fig. 1(a) and (b) illustrate the sex- and country-specific MRR (95 % CI) for the association between intake of total WG types and all-cause mortality. The forest plots illustrate the homogeneity in results between countries.

Sensitivity analyses

Totally, nine sensitivity analyses were conducted (see sections on Methods and Statistical analyses) with the analysis of total intake of WG types and all-cause mortality as the basis model (Table 3, third and seventh column: MRR of 0·88 (95 % CI 0·86, 0·92) for women and 0·88 (95 % CI 0·86, 0·91) for men, for each doubling in the intake of total WG types): (1) the assumption of proportional hazards were not violated; (2) and (3) tests for effect modification by age or BMI showed no statistically significant effect modification by age or BMI on the association between WG intake and all-cause mortality; (4) stratification by physical activity (only Denmark

Table 6. The association between quartiles of whole-grain intake and cause-specific mortality of cohort participants in the HELGA study

(Mortality rate ratios (MRR) and 95 % confidence intervals)

	Quartile 1	Quartile 2		Quartile 3		Quartile 4		
	MRR*	MRR*	95 % CI	MRR*	95 % CI	MRR*	95 % CI	$P_{trend\ over\ quartiles}$
Women†								
Cancer mortality (deaths = 1775)								
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain product intake								
Breakfast cereals	1	0·98	0·83, 1·15	0·84	0·74, 0·95	0·85	0·74, 0·98	0·003
Non-white bread	1	1·01	0·86, 1·18	0·93	0·79, 1·08	0·89	0·75, 1·05	0·085
Crisp bread	1	0·89	0·77, 1·03	0·84	0·75, 0·95	0·87	0·74, 1·02	0·012
Total whole-grain products	1	0·86	0·76, 0·98	0·85	0·75, 0·97	0·86	0·74, 0·99	0·0236
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain type intake								
Oat	1	0·89	0·78, 1·01	0·77	0·67, 0·90	0·84	0·71, 0·99	0·007
Rye	1	0·98	0·87, 1·11	1·00	0·86, 1·17	0·99	0·79, 1·25	0·957
Wheat	1	0·76	0·65, 0·88	0·74	0·63, 0·87	0·74	0·61, 0·89	0·006
Total whole-grain types‡	1	0·99	0·87, 1·13	0·94	0·82, 1·07	0·88	0·77, 1·02	0·069
CHD mortality (deaths = 298)								
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain product intake								
Breakfast cereals	1	0·71	0·43, 1·16	0·64	0·47, 0·87	0·53	0·37, 0·77	0·0002
Non-white bread	1	0·72	0·52, 0·99	0·54	0·37, 0·79	0·72	0·47, 1·09	0·023
Crisp bread	1	0·81	0·56, 1·18	0·95	0·68, 1·33	0·94	0·66, 1·34	0·864
Total whole-grain products	1	0·72	0·54, 0·97	0·64	0·46, 0·89	0·56	0·40, 0·78	0·0004
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain type intake								
Oat	1	0·74	0·52, 1·05	0·85	0·59, 1·21	0·66	0·45, 0·96	0·048
Rye	1	0·79	0·57, 1·11	1·02	0·75, 1·38	0·69	0·46, 1·05	0·249
Wheat	1	0·83	0·57, 1·21	0·87	0·60, 1·27	0·67	0·45, 1·00	0·062
Total whole-grain types‡	1	0·83	0·60, 1·14	0·59	0·42, 0·82	0·65	0·46, 0·91	0·003
Stroke mortality (deaths = 137)								
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain product intake								
Breakfast cereals	1	0·70	0·33, 1·47	0·73	0·43, 1·23	0·66	0·43, 1·02	0·062
Non-white bread	1	0·86	0·48, 1·54	1·02	0·55, 1·87	0·75	0·41, 1·37	0·470
Crisp bread	1	1·03	0·62, 1·74	0·91	0·54, 1·56	1·22	0·68, 2·18	0·655
Total whole-grain products	1	0·78	0·50, 1·22	0·57	0·35, 0·94	0·85	0·53, 1·37	0·264
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain type intake								
Oat	1	1·03	0·64, 1·67	0·66	0·38, 1·16	0·63	0·34, 1·15	0·095
Rye	1	0·71	0·41, 1·22	0·81	0·47, 1·37	0·91	0·53, 1·57	0·934
Wheat	1	0·72	0·44, 1·18	0·83	0·50, 1·38	0·55	0·23, 1·30	0·323
Total whole-grain types‡	1	0·55	0·34, 0·88	0·50	0·31, 0·82	0·80	0·48, 1·33	0·319
Respiratory disease mortality (deaths = 125)								
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain product intake								
Breakfast cereals	1	0·28	0·11, 0·69	0·48	0·28, 0·83	0·39	0·23, 0·65	<0·0001
Non-white bread	1	0·87	0·50, 1·53	0·47	0·24, 0·93	0·74	0·41, 1·35	0·228
Crisp bread	1	0·80	0·51, 1·25	0·59	0·32, 1·07	1·03	0·59, 1·79	0·801
Total whole-grain products	1	0·56	0·35, 0·89	0·53	0·33, 0·86	0·82	0·49, 1·37	0·113
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain type intake								
Oat	1	0·56	0·36, 0·88	0·48	0·27, 0·85	0·51	0·28, 0·93	0·006
Rye	1	1·62	0·86, 3·04	1·12	0·61, 2·05	1·12	0·60, 2·10	0·658
Wheat	1	0·63	0·41, 0·99	0·57	0·34, 0·94	0·44	0·09, 2·05	0·064
Total whole-grain types‡	1	0·65	0·40, 1·08	0·64	0·39, 1·04	0·47	0·28, 0·77	0·005
Diabetes mortality (deaths = 24)								
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain product intake								
Breakfast cereals	1	0·80	0·23, 2·77	0·45	0·13, 1·59	0·28	0·07, 1·04	0·036
Non-white bread	1	0·47	0·15, 1·48	0·40	0·11, 1·50	0·55	0·17, 1·73	0·522
Crisp bread	1	1·21	0·41, 3·54	1·25	0·36, 4·35	1·09	0·27, 4·43	0·860
Total whole-grain products	1	0·41	0·13, 1·29	0·41	0·13, 1·29	0·90	0·31, 2·62	0·494
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain type intake								
Oat	1	0·88	0·34, 2·27	0·36	0·09, 1·51	0·25	0·05, 1·31	0·028
Rye	1	0·93	0·28, 3·09	0·46	0·14, 1·53	0·70	0·22, 2·25	0·390
Wheat	1	0·69	0·26, 1·85	0·61	0·20, 1·88	3·17	0·27, 37·59	0·627
Total whole-grain types‡	1	0·77	0·25, 2·41	0·44	0·14, 1·38	0·64	0·20, 2·04	0·308
Mortality from other causes (deaths = 1299)								
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain product intake								
Breakfast cereals	1	0·68	0·52, 0·88	0·75	0·62, 0·91	0·74	0·65, 0·84	<0·0001
Non-white bread	1	0·70	0·59, 0·82	0·60	0·49, 0·74	0·54	0·45, 0·64	<0·0001
Crisp bread	1	0·85	0·72, 1·01	0·87	0·75, 1·01	0·86	0·72, 1·04	0·059
Total whole-grain products	1	0·80	0·69, 0·92	0·69	0·59, 0·80	0·57	0·47, 0·68	<0·0001
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain type intake								
Oat	1	0·82	0·70, 0·95	0·75	0·64, 0·89	0·81	0·67, 0·98	0·0006
Rye	1	0·84	0·72, 0·98	0·68	0·57, 0·81	0·81	0·68, 0·97	0·007
Wheat	1	0·66	0·56, 0·78	0·55	0·46, 0·66	0·51	0·39, 0·67	<0·0001
Total whole-grain types‡	1	0·67	0·58, 0·77	0·67	0·57, 0·79	0·61	0·51, 0·73	<0·0001

Table 6. *Continued*

	Quartile 1	Quartile 2		Quartile 3		Quartile 4		
	MRR*	MRR*	95 % CI	MRR*	95 % CI	MRR*	95 % CI	$P_{\text{trend over quartiles}}$
Men†								
Cancer mortality (deaths = 1375)								
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain product intake								
Breakfast cereals	1	0·99	0·81, 1·21	0·73	0·63, 0·85	0·75	0·64, 0·87	<0·0001
Non-white bread	1	0·92	0·76, 1·12	0·97	0·79, 1·19	0·79	0·64, 0·97	0·019
Crisp bread	1	0·99	0·85, 1·15	0·92	0·79, 1·07	0·83	0·67, 1·03	0·080
Total whole-grain products	1	0·84	0·72, 0·98	0·74	0·64, 0·86	0·70	0·60, 0·81	<0·0001
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain type intake								
Oat	1	0·80	0·69, 0·94	0·75	0·64, 0·89	0·75	0·64, 0·89	0·003
Rye	1	0·97	0·82, 1·14	0·80	0·65, 0·98	0·93	0·78, 1·11	0·261
Wheat	1	0·84	0·72, 0·98	0·69	0·58, 0·82	0·58	0·49, 0·69	<0·0001
Total whole-grain types‡	1	0·85	0·73, 0·99	0·68	0·58, 0·79	0·74	0·63, 0·87	<0·0001
CHD mortality (deaths = 858)								
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain product intake								
Breakfast cereals	1	0·88	0·69, 1·11	0·87	0·72, 1·05	0·75	0·61, 0·91	0·009
Non-white bread	1	1·13	0·89, 1·42	0·95	0·74, 1·22	0·87	0·65, 1·17	0·128
Crisp bread	1	0·98	0·81, 1·19	0·95	0·78, 1·16	1·23	0·88, 1·71	0·704
Total whole-grain products	1	0·94	0·78, 1·15	0·80	0·66, 0·97	0·85	0·70, 1·04	0·041
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain type intake								
Oat	1	1·01	0·83, 1·23	1·01	0·80, 1·28	0·82	0·65, 1·03	0·052
Rye	1	0·77	0·62, 0·95	0·81	0·67, 0·97	0·88	0·73, 1·07	0·216
Wheat	1	0·80	0·66, 0·97	0·70	0·55, 0·88	0·84	0·65, 1·08	0·273
Total whole-grain types‡	1	0·79	0·65, 0·96	0·80	0·66, 0·97	0·74	0·61, 0·91	0·012
Stroke mortality (deaths = 143)								
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain product intake								
Breakfast cereals	1	0·92	0·59, 1·45	0·86	0·56, 1·32	0·87	0·47, 1·28	0·488
Non-white bread	1	0·74	0·44, 1·22	0·56	0·35, 0·91	0·86	0·50, 1·46	0·181
Crisp bread	1	1·10	0·67, 1·79	0·97	0·60, 1·57	1·30	0·71, 2·39	0·590
Total whole-grain products	1	1·01	0·66, 1·56	0·55	0·35, 0·87	0·86	0·54, 1·37	0·101
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain type intake								
Oat	1	0·67	0·42, 1·07	1·03	0·60, 1·76	0·71	0·41, 1·21	0·200
Rye	1	0·77	0·48, 1·23	0·72	0·39, 1·31	0·61	0·36, 1·05	0·097
Wheat	1	0·59	0·35, 0·99	0·72	0·41, 1·27	0·74	0·41, 1·32	0·946
Total whole-grain types‡	1	0·70	0·44, 1·10	0·69	0·43, 1·11	0·71	0·44, 1·15	0·179
Respiratory disease mortality (deaths = 111)								
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain product intake								
Breakfast cereals	1	1·50	0·79, 2·82	1·23	0·76, 1·98	0·66	0·37, 1·16	0·350
Non-white bread	1	1·04	0·52, 2·08	0·62	0·28, 1·35	0·94	0·46, 1·94	0·560
Crisp bread	1	0·77	0·45, 1·32	0·98	0·59, 1·62	1·00	0·50, 2·01	0·947
Total whole-grain products	1	1·03	0·61, 1·76	1·05	0·63, 1·77	0·60	0·33, 1·07	0·098
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain type intake								
Oat	1	0·99	0·59, 1·68	0·87	0·50, 1·51	0·52	0·28, 0·97	0·063
Rye	1	1·16	0·65, 2·10	0·70	0·32, 1·57	1·02	0·53, 1·97	0·624
Wheat	1	1·46	0·83, 2·55	1·40	0·76, 2·56	1·06	0·57, 1·95	0·999
Total whole-grain types‡	1	0·96	0·58, 1·61	0·65	0·38, 1·11	0·74	0·42, 1·30	0·139
Diabetes mortality (deaths = 70)								
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain product intake								
Breakfast cereals	1	0·59	0·24, 1·50	0·31	0·14, 0·68	0·45	0·23, 0·90	0·004
Non-white bread	1	1·89	0·66, 5·41	1·59	0·52, 4·82	2·30	0·79, 6·72	0·188
Crisp bread	1	1·33	0·68, 2·61	1·07	0·53, 2·15	2·35	1·10, 5·03	0·085
Total whole-grain products	1	1·76	0·83, 3·74	1·09	0·49, 2·42	1·67	0·79, 3·51	0·418
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain type intake								
Oat	1	0·60	0·32, 1·12	0·33	0·16, 0·72	0·40	0·20, 0·81	0·003
Rye	1	1·14	0·53, 2·48	0·40	0·12, 1·34	1·80	0·80, 4·05	0·175
Wheat	1	0·75	0·37, 1·56	1·07	0·51, 2·22	1·18	0·60, 2·34	0·245
Total whole-grain types‡	1	0·92	0·45, 1·87	0·95	0·49, 1·86	1·18	0·59, 2·38	0·636
Other causes of mortality (deaths = 1624)								
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain product intake								
Breakfast cereals	1	0·86	0·71, 1·05	0·83	0·72, 0·94	0·78	0·68, 0·90	0·0002
Non-white bread	1	0·83	0·70, 0·98	0·75	0·63, 0·91	0·65	0·54, 0·79	<0·0001
Crisp bread	1	0·95	0·82, 1·09	0·91	0·79, 1·05	1·06	0·88, 1·28	0·928
Total whole-grain products	1	0·83	0·72, 0·95	0·70	0·61, 0·80	0·70	0·61, 0·80	<0·0001
MRR in quartile 2, 3 and 4 compared to quartile 1 (reference) of whole-grain type intake								
Oat	1	0·87	0·75, 1·01	0·94	0·81, 1·09	0·85	0·73, 1·00	0·107
Rye	1	0·83	0·73, 0·96	0·66	0·55, 0·80	0·76	0·65, 0·89	0·0003
Wheat	1	0·88	0·76, 1·02	0·82	0·70, 0·96	0·75	0·64, 0·88	0·0009
Total whole-grain types‡	1	0·81	0·70, 0·93	0·71	0·62, 0·81	0·71	0·62, 0·82	<0·0001

* Whole-grain products and whole-grain types (including 'other types' of whole-grain) mutually adjusted and adjusted for age (time scale), follow-up time (linear spline with boundaries at 1, 2 and 3 years after entry), education (categorical: none, primary, technical/professional, secondary, longer), smoking intensity (categorical: never; former, quit 20+ years; former, quit 11–20 years; former, quit ≤10 years; current, pipe/cigar/occasionally; current, 1–15 cigarettes/d; current, 16–25 cigarettes/d; current, 26 + cigarettes/d, unknown), alcohol (linear spline with boundary at 15 g/d), BMI (linear spline with boundaries at 18·5, 25 and 30 kg/m²), total energy intake (continuous: kJ/d) and alcohol intake (g/d).

† Median intakes of whole-grain in each quartile in online Appendix Table.
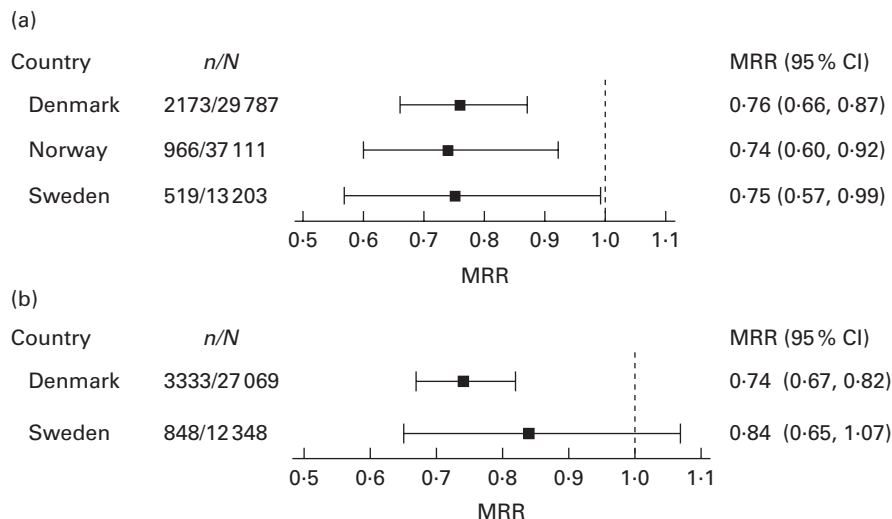
‡ Sum of oat, rye, wheat and other types of whole grain.

(a)

Country	n/N		MRR (95 % CI)
Denmark	2173/29 787		0·76 (0·66, 0·87)
Norway	966/37 111		0·74 (0·60, 0·92)
Sweden	519/13 203		0·75 (0·57, 0·99)

0·5   0·6   0·7   0·8   0·9   1·0   1·1
MRR

(b)

Country	n/N		MRR (95 % CI)
Denmark	3333/27 069		0·74 (0·67, 0·82)
Sweden	848/12 348		0·84 (0·65, 1·07)

0·5   0·6   0·7   0·8   0·9   1·0   1·1
MRR

Fig. 1. (a) Forest plot of the mortality rate ratios (MRR) and 95 % CI for the association between intake of total whole-grain types and all-cause mortality of female participants in the HELGA study. (b) Forest plot of the MRR and 95 % CI for the association between intake of total whole-grain types and all-cause mortality of male participants in the HELGA study.

and Sweden) and intake of fruits and nuts, vegetables, meat and alcohol did not reveal any inconsistent results below/above median activity or intake; (5) adjustment for the physical activity index (only Denmark and Sweden) revealed MRR of 0·89 (95 % CI 0·86, 0·93) for women and 0·90 (95 % CI 0·87, 0·93) for men; (6) further adjustment for hypertension, hyperlipidaemia and waist circumference (using only the Danish data) did not change the results the least; (7) results in non-smokers (never-smokers and former smokers) revealed MRR of 0·91 (95 % CI 0·87, 0·96) for women and 0·94 (95 % CI 0·90, 0·99) for men; (8) the analyses adjusted for the diet index gave a MRR of 0·90 (95 % CI 0·87, 0·93) for women and 0·89 (95 % CI 0·87, 0·92) for men; (9) exclusion of individuals with a previous diagnosis of CHD, stroke or angina pectoris did not change the results (change in MRR < 1 %).

Discussion

In this large Scandinavian population, intake of WG products was associated with lower all-cause mortality. In particular, in the analyses of continuous WG variables, intake of breakfast cereals and non-white bread was associated with lower mortality. Also, the total intake of different cereal types was associated with lower all-cause mortality, and this association was due to lower mortality associated with higher intake of WG oat, wheat and rye. The estimates were, with few exceptions, all in the same direction and of the same magnitude, and many were also highly statistically significant. Adjustment for potential confounders generally did not change the strength of associations, and the many sensitivity analyses further supported the overall conclusion of the study.

In the analyses of cause-specific mortality, intake of total WG products and total WG types was associated with lower mortality (except for mortality from respiratory disease, and total WG product intake and cancer mortality in women). The large part of the remaining associations had MRR below

unity, in the range of 1–12 % lower mortality for each doubling in the intake of specific WG products or specific WG types (Table 4) and in the order of 3–17 % for total WG products or total WG types. Crisp bread was not associated with mortality after adjustment for potential confounders.

In the analyses of quartiles of WG intake in relation to all-cause mortality, we found lower mortality in the highest quartile compared with the lowest quartile for breakfast cereals, non-white bread, total WG products, oat, rye (only men), wheat and total intake of different WG types. The analyses on cause-specific mortality according to quartiles of WG intake were characterised by lower power, but still supported the overall results of the study.

The present study has a major advantage in being carried out in a population with a considerable and large variation in the intake of different types of WG. Furthermore, the combination of three large, prospective, population-based cohorts with linkage to National registries ensures a complete and valid follow-up of deaths and causes of death in addition to high statistical power. Consequently, the results were consistent and highly statistically significant. The study does, however, also have some limitations: there may be some degree of misclassification in the assessment of WG intake due to the self-reported data from the FFQ, to a large variation in WG contents of WG products, and to the fact that the questionnaires were not designed to measure the intake of WG. However, the questionnaires were validated locally and performed well[45,46,54], and the most likely consequence of this non-differential misclassification is attenuation of associations. Diet was assessed only at baseline and may not reflect diet earlier in life nor changes during the relatively long follow-up; consequently, the relevance of a single measure of exposure depends on the degree to which it tracks over time. When comparing the first 5, 5–10 and 10–15 years with the rest of the follow-up, the associations were not weakened with longer follow-up. If study participation was

associated with a risk factor for mortality, there is a chance of selection bias. However, as disease status was unknown at baseline, selection bias is less likely to have affected the results, but cannot be completely ruled out. To evaluate possible bias from prevalent disease (non-proportionality), we allowed the mortality rate ratios to change with time under study. Furthermore, the results from analyses of the first, second and third year of follow-up were compared with results from the remaining part of follow-up; no differences were found. There is always a risk of residual confounding from unknown factors or measurement error in known confounders. In particular, individuals with a high intake of WG may, in general, have a healthier lifestyle and a better health than non-consumers, also regarding factors not included in our three sets of potential confounders. Last, as the three population samples do not completely reflect the general population, the external validity and generalisability is limited to a population slightly healthier than the general population.

The present findings of lower mortality with higher intake of WG products are in line with all, except one, studies on WG and mortality. The single study[35] that did not find an association did not have intake of WG as the primary focus (but the Mediterranean diet) and had been characterised by a study population with an exceptionally low intake of WG. A study of US male health professionals[32] has investigated the frequency of intake of whole-grain and refined-grain breakfast cereals in relation to total and cause-specific mortality. Consistent with our study, they found a strong association between intake WG breakfast cereals and all-cause mortality. In the Iowa Women's Health Study, consisting of 41836 postmenopausal women aged 55–69 years[29,31], they have found a strong association between total WG product intake and all-cause and cause-specific mortality. This association was confined to a lower mortality with higher intake of dark bread, but not to intake of WG breakfast cereals and cause-specific mortality. Another study of Norwegian women and men has investigated a WG bread score in relation to total and cause-specific mortality[30], and consistent with our analyses on WG product intake and mortality, they found lower all-cause mortality in women and men with a high WG bread score. The remaining two, rather small studies with 187 and 867 deaths[29,31,33,34], were conducted in American populations where the intakes of WG were extremely low and associated with other healthy behaviours. Still, after adjustment for potential confounders, they found lower mortality with higher intake of WG. In the study by Sahyoun et al.[33], the lower mortality associated with higher number of WG servings per day was confined to cardiovascular mortality. This is in line with a recent and combined study of the Nurses Health Study and the Health Professionals Follow-up Study[55] and with the present results on CHD where intake of both WG products and WG types was associated with lower CHD mortality.

The sub-study of the Nurses Health Study, conducted on 7822 women with type 2 diabetes[37], has found a borderline significant trend for the relative risks over quintiles of WG intake. The lowest intake in the highest quintile was only 25·5 g/d, corresponding to a considerably lower intake than in the present study and only one-third of the official recommendation. However, the relative risks were of the same size as in the present study (relative risk 0·89 (95 % CI 0·71, 1·11) in the highest quintile compared with the lowest), and it is most likely that with more deaths and a larger range of intake, the study would find stronger associations between WG intake and mortality.

The present study indicates that particularly WG wheat, but also WG oats and WG rye, are associated with lower mortality. Rye is characterised by the highest amount of dietary fibre among cereals, and the proportion of soluble fibre components is much higher in rye than in wheat[56]. Minimally processed oat products are rich in viscous forming β-glucans, which have well-established cholesterol-lowering properties[57]. The analyses on different WG types and mortality are justified by the possible differential effects of different types of grain, but in the present study, there were no large differences between the different types of (whole) grain cereals.

The remarkable consistency of the present results between women and men, between the cause-specific analyses (disregarded the lower statistical power in split analyses), between the type and the product measures of WG and the fact that the estimates with few exceptions are all below unity increases our belief that the findings reflect true associations and not chance or bias. The size of the estimates may seem small (1–17 % lower mortality for each doubling in WG intake), but considering the fact that WG is just one single dietary component, and diet is just one of a range of lifestyle factors predicting mortality, it is still a valuable and achievable goal for the Scandinavian populations. In the Scandinavian countries, an official recommendation of 75 g of WG/10 MJ/d has been published[58–60]. Between 16 % (Danish men) and 35 % (Norwegian women) reached the recommended daily intake of WG[49], and although the intake of WG is part of the traditional diet in the Scandinavian countries, effort has to be made to increase and keep the intake high.

In conclusion, the intake of WG (products or cereal types) was associated with lower mortality in the HELGA cohort. These results were found quite consistently for different causes of death and across categories of sex and types or products of WG. The present study indicates that intake of WG is an important aspect of diet in preventing early death, and there were no indications that the associations between WG and mortality were caused by one specific type or product of WG.

Supplementary material

To view supplementary material for this article, please visit http://dx.doi.org/10.1017/S0007114515001701

Acknowledgements

British Journal of Nutrition

References

1. Burkitt D (1987) Dietary fibre – historical aspects. *Scand J Gastroenterol Suppl* **129**, 10–13.
2. Burkitt D, Morley D & Walker A (1980) Dietary fibre in under- and overnutrition in childhood. *Arch Dis Child* **55**, 803–807.
3. Burkitt DP (1952) Acute abdomens – British and Baganda compared. *East Afr Med J* **29**, 189–194.
4. Cleave TL (1956) The neglect of natural principles in current medical practice. *J R Nav Med Serv* **42**, 54–83.
5. Trowell H (1972) Dietary fibre and coronary heart disease. *Rev Eur Etud Clin Biol* **17**, 345–349.
6. Walker AR (1947) The effect of recent changes of food habits on bowel motility. *S Afr Med J* **21**, 590–596.
7. Walker AR (1977) Health implications of fibre-depleted diets. *S Afr Med J* **52**, 767–770.
8. Chuang SC, Norat T, Murphy N, *et al.* (2012) Fiber intake and total and cause-specific mortality in the European Prospective Investigation into Cancer and Nutrition cohort. *Am J Clin Nutr* **96**, 164–174.
9. Park Y, Subar AF, Hollenbeck A, *et al.* (2011) Dietary fiber intake and mortality in the NIH-AARP diet and health study. *Arch Intern Med* **171**, 1061–1068.
10. Karl JP & Saltzman E (2012) The role of whole grains in body weight regulation. *Adv Nutr* **3**, 697–707.
11. Ye EQ, Chacko SA, Chou EL, *et al.* (2012) Greater whole-grain intake is associated with lower risk of type 2 diabetes, cardiovascular disease, and weight gain. *J Nutr* **142**, 1304–1313.
12. Lillioja S, Neal AL, Tapsell L, *et al.* (2013) Whole grains, type 2 diabetes, coronary heart disease, and hypertension: links to the aleurone preferred over indigestible fiber. *Biofactors* **39**, 242–258.
13. Aune D, Chan DS, Lau R, *et al.* (2011) Dietary fibre, whole grains, and risk of colorectal cancer: systematic review and dose–response meta-analysis of prospective studies. *BMJ* **343**, d6617.
14. Katcher HI, Legro RS, Kunselman AR, *et al.* (2008) The effects of a whole grain-enriched hypocaloric diet on cardio-vascular disease risk factors in men and women with metabolic syndrome. *Am J Clin Nutr* **87**, 79–90.
15. Poppitt SD, Keogh GF, Prentice AM, *et al.* (2002) Long-term effects of *ad libitum* low-fat, high-carbohydrate diets on body weight and serum lipids in overweight subjects with metabolic syndrome. *Am J Clin Nutr* **75**, 11–20.
16. Rave K, Roggen K, Dellweg S, *et al.* (2007) Improvement of insulin resistance after diet with a whole-grain based dietary product: results of a randomized, controlled cross-over study in obese subjects with elevated fasting blood glucose. *Br J Nutr* **98**, 929–936.
17. Pol K, Christensen R, Bartels EM, *et al.* (2013) Whole grain and body weight changes in apparently healthy adults: a systematic review and meta-analysis of randomized controlled studies. *Am J Clin Nutr* **98**, 872–884.
18. Juntunen KS, Niskanen LK, Liukkonen KH, *et al.* (2002) Postprandial glucose, insulin, and incretin responses to grain products in healthy subjects. *Am J Clin Nutr* **75**, 254–262.
19. Landberg R, Andersson SO, Zhang JX, *et al.* (2010) Rye whole grain and bran intake compared with refined wheat decreases urinary C-peptide, plasma insulin, and prostate specific antigen in men with prostate cancer. *J Nutr* **140**, 2180–2186.
20. Pereira MA, Jacobs DR Jr, Pins JJ, *et al.* (2002) Effect of whole grains on insulin sensitivity in overweight hyperinsulinemic adults. *Am J Clin Nutr* **75**, 848–855.
21. Kelly SA, Summerbell CD, Brynes A, *et al.* (2007) Wholegrain cereals for coronary heart disease. *The Cochrane Database of Systematic Reviews,* CD005051.
22. Keenan JM, Pins JJ, Frazel C, *et al.* (2002) Oat ingestion reduces systolic and diastolic blood pressure in patients with mild or borderline hypertension: a pilot trial. *J Fam Pract* **51**, 369.
23. Pins JJ, Geleva D, Keenan JM, *et al.* (2002) Do whole-grain oat cereals reduce the need for antihypertensive medications and improve blood pressure control? *J Fam Pract* **51**, 353–359.
24. Saltzman E, Das SK, Lichtenstein AH, *et al.* (2001) An oat-containing hypocaloric diet reduces systolic blood pressure and improves lipid profile beyond effects of weight loss in men and women. *J Nutr* **131**, 1465–1470.
25. Brownlee IA, Moore C, Chatfield M, *et al.* (2010) Markers of cardiovascular risk are not changed by increased whole-grain intake: the WHOLEheart study, a randomised, controlled dietary intervention. *Br J Nutr* **104**, 125–134.
26. Giacco R, Lappi J, Costabile G, *et al.* (2013) Effects of rye and whole wheat versus refined cereal foods on metabolic risk factors: a randomised controlled two-centre intervention study. *Clin Nutr* **32**, 941–949.
27. Kristensen M, Toubro S, Jensen MG, *et al.* (2012) Whole grain compared with refined wheat decreases the percentage of body fat following a 12-week, energy-restricted dietary intervention in postmenopausal women. *J Nutr* **142**, 710–716.
28. Leinonen KS, Poutanen KS & Mykkanen HM (2000) Rye bread decreases serum total and LDL cholesterol in men with moderately elevated serum cholesterol. *J Nutr* **130**, 164–170.
29. Jacobs DR Jr, Meyer KA, Kushi LH, *et al.* (1999) Is whole grain intake associated with reduced total and cause-specific death rates in older women? The Iowa Women's Health Study. *Am J Public Health* **89**, 322–329.
30. Jacobs DR Jr, Meyer HE & Solvoll K (2001) Reduced mortality among whole grain bread eaters in men and women in the Norwegian County Study. *Eur J Clin Nutr* **55**, 137–143.
31. Jacobs DR Jr, Andersen LF & Blomhoff R (2007) Whole-grain consumption is associated with a reduced risk of noncardio-vascular, noncancer death attributed to inflammatory diseases in the Iowa Women's Health Study. *Am J Clin Nutr* **85**, 1606–1614.
32. Liu S, Sesso HD, Manson JE, *et al.* (2003) Is intake of breakfast cereals related to total and cause-specific mortality in men? *Am J Clin Nutr* **77**, 594–599.
33. Sahyoun NR, Jacques PF, Zhang XL, *et al.* (2006) Whole-grain intake is inversely associated with the metabolic syndrome and mortality in older adults. *Am J Clin Nutr* **83**, 124–131.

34. Steffen LM, Jacobs DR Jr, Stevens J, et al. (2003) Associations of whole-grain, refined-grain, and fruit and vegetable consumption with risks of all-cause mortality and incident coronary artery disease and ischemic stroke: the Atherosclerosis Risk in Communities (ARIC) Study. Am J Clin Nutr 78, 383–390.

35. van den Brandt PA (2011) The impact of a Mediterranean diet and healthy lifestyle on premature mortality in men and women. Am J Clin Nutr 94, 913–920.

36. He M, van Dam RM, Rimm E, et al. (2010) Whole-grain, cereal fiber, bran, and germ intake and the risks of all-cause and cardiovascular disease-specific mortality among women with type 2 diabetes mellitus. Circulation 121, 2162–2168.

37. Frolich W, Aman P & Tetens I (2013) Whole grain foods and health – a Scandinavian perspective. Food Nutr Res 57, 18503.

38. AACC International Whole Grains Definition. St Paul, MN: AACC International. http://www.aaccnet org/initiatives/definitions/Pages/WholeGrain aspx 2013 (accessed June 2013).

39. Okarter N & Liu RH (2010) Health benefits of whole grain phytochemicals. Crit Rev Food Sci Nutr 50, 193–208.

40. Lund E, Dumeaux V, Braaten T, et al. (2008) Cohort profile: The Norwegian Women and Cancer Study – NOWAC – Kvinner og kreft. Int J Epidemiol 37, 36–41.

41. Winkvist A, Hornell A, Hallmans G, et al. (2009) More distinct food intake patterns among women than men in northern Sweden: a population-based survey. Nutr J 8, 12.

42. Tjonneland A, Olsen A, Boll K, et al. (2007) Study design, exposure variables, and socioeconomic determinants of participation in Diet, Cancer and Health: a population-based prospective cohort study of 57,053 men and women in Denmark. Scand J Public Health 35, 432–441.

43. Riboli E, Hunt KJ, Slimani N, et al. (2002) European Prospective Investigation into Cancer and Nutrition (EPIC): study populations and data collection. Public Health Nutr 5, 1113–1124.

44. Hjartaker A, Andersen LF & Lund E (2007) Comparison of diet measures from a food-frequency questionnaire with measures from repeated 24-hour dietary recalls. The Norwegian Women and Cancer Study. Public Health Nutr 10, 1094–1103.

45. Johansson I, Hallmans G, Wikman A, et al. (2002) Validation and calibration of food-frequency questionnaire measurements in the Northern Sweden Health and Disease cohort. Public Health Nutr 5, 487–496.

46. Tjonneland A, Overvad K, Haraldsdottir J, et al. (1991) Validation of a semiquantitative food frequency questionnaire developed in Denmark. Int J Epidemiol 20, 906–912.

47. Slimani N, Kaaks R, Ferrari P, et al. (2002) European Prospective Investigation into Cancer and Nutrition (EPIC) calibration study: rationale, design and population characteristics. Public Health Nutr 5, 1125–1145.

48. Mejborn H, Biltoft-Jensen A, Trolle E, et al. (2012) Fuldkorn – Definition og vidensgrundlag for anbefaling af fuldkornsindtag i Danmark (Wholegrain – Definition and Scientific Background for Recommendations of Wholegrain Intake in Denmark). Søborg: National Food Institute, Technical University of Denmark. http://www.fuldkorn.dk/files/Rapporter/Fuldkorn%20definition%20og%20vidensgrundlag.pdf (accessed June 2012).

49. Kyro C, Skeie G, Dragsted LO, et al. (2012) Intake of whole grain in Scandinavia: intake, sources and compliance with new national recommendations. Scand J Public Health 40, 76–84.

50. Wareham NJ, Jakes RW, Rennie KL, et al. (2003) Validity and repeatability of a simple index derived from the short physical activity questionnaire used in the European Prospective Investigation into Cancer and Nutrition (EPIC) study. Public Health Nutr 6, 407–413.

51. Greenland S (1995) Avoiding power loss associated with categorization and ordinal scores in dose–response and trend analysis. Epidemiology 6, 450–454.

52. Greenland S (1995) Dose–response and trend analysis in epidemiology: alternatives to categorical analysis. Epidemiology 6, 356–365.

53. Trichopoulou A, Costacou T, Bamia C, et al. (2003) Adherence to a Mediterranean diet and survival in a Greek population. N Engl J Med 348, 2599–2608.

54. Parr CL, Veierod MB, Laake P, et al. (2006) Test–retest reproducibility of a food frequency questionnaire (FFQ) and estimated effects on disease risk in the Norwegian Women and Cancer Study (NOWAC). Nutr J 5, 4.

55. Wu H, Flint A, Qi Q, et al. (2015) Association between dietary whole grain intake and risk of mortality. JAMA Intern Med. 175, 373–384.

56. Andersson SC, Olsson ME, Johansson E, et al. (2009) Carotenoids in sea buckthorn (Hippophae rhamnoides L.) berries during ripening and use of pheophytin a as a maturity marker. J Agric Food Chem 57, 250–258.

57. Wolever TM, Tosh SM, Gibbs AL, et al. (2010) Physicochemical properties of oat β-glucan influence its ability to reduce serum LDL cholesterol in humans: a randomized clinical trial. Am J Clin Nutr 92, 723–732.

58. Danish Veterinary and Food Administration (2012) De 8 kostråd (The Danish Food-based Dietary Guidelines). Mørkhøj, Denmark: The Danish Food Institute. http://www altomkost dk/Anbefalinger/De_8_kostraad/forside htm

59. Norwegian Nutrition Council (2011) Kostråd for å fremme folkehelsen og forebygge kroniske sykdommer (Dietary Guidelines to Improve Public Health and Prevent Chronic Diseases). Oslo, Norway: Norwegian Nutrition Council. http://www helsedirektoratet no/publikasjoner/kostrad-for-a-fremme-folkehelsen-og-forebygge-kroniske-sykdommer/Sider/default aspx

60. The Swedish Food Administration (2011) Kostråd vuxna (Dietary Recommendations – Adults). Uppsala, Sweden: Swedish National Food Agency. http://www slv se/sv/grupp1/Mat-och-naring/Kostrad/Vuxna/