# Potato consumption and risk of pancreatic cancer in the HELGA cohort

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# Abstract

Potatoes have been a staple food in many countries throughout the years. Potatoes have a high glycaemic index (GI) score, and high GI has been associated with several chronic diseases and cancers. Still, the research on potatoes and health is scarce and contradictive, and we identified no prospective studies that had investigated the association between potatoes as a single food and the risk of pancreatic cancer. The aim of this study was to prospectively investigate the association between potato consumption and pancreatic cancer among 114 240 men and women in the prospective HELGA cohort, using Cox proportional hazard models. Information on diet (validated FFQ's), lifestyle and health was collected by means of a questionnaire, and 221 pancreatic cancer cases were identified through cancer registries. The mean follow-up time was 11.4 (95% CI 0.3, 16.9) years. High consumption of potatoes showed a non-significantly higher risk of pancreatic cancer in the adjusted model (hazard ratio (HR) 1.44; 95% CI 0.93, 2.22,  $P_{\text{for trend}}$  0.030) when comparing the highest *v*. the lowest quartile of potato consumption. In the sex-specific analyses, significant associations were found for females (HR 2.00; 95% CI 1.07, 3.72,  $P_{\text{for trend}}$  0.020), but not for males (HR 1.01; 95% CI 0.56, 1.84,  $P_{\text{for trend}}$  0.34). In addition, we explored the associations by spline regression, and the absence of dose–response effects was confirmed. In this study, high potato consumption was not consistently associated with a higher risk of pancreatic cancer. Further studies with larger populations are needed to explore the possible sex difference.

#### Key words: Cohort studies: Epidemiology: Potatoes: Cancer

Pancreatic cancer is the eighth most common cause of cancer death worldwide, and 227 000 individuals were diagnosed with pancreatic cancer in  $2008^{(1)}$ . In the Nordic countries, pancreatic cancer contributes to 2.4% of the cancer cases in men and 2.7% of the cancer cases in women<sup>(2)</sup>. Pancreatic cancer is hard to detect early<sup>(3)</sup>, and the cancer is generally advanced, and usually fatal, by the time it is diagnosed<sup>(4)</sup>. The 5-year prevalence of people living with pancreatic cancer worldwide is estimated to be 4.1 per  $100\,000^{(3)}$ . There is limited evidence suggesting consumption of red and processed meat, foods and beverages containing fructose and heavy alcohol drinking (more than 3 drinks/d) as risk factors of pancreatic cancer. The evidence for fruits and physical activity is too limited and inconsistent to draw any conclusion<sup>(5)</sup>. Other non-dietary risk factors for pancreatic cancer are more established.

Approximately 25% of cases of pancreatic cancer are attributable to tobacco smoking<sup>(5)</sup>, and the risk also increases with age. Factors such as body fatness and greater childhood growth have been identified as, respectively, convincing and probable causes of pancreatic cancer<sup>(5)</sup>. Chronic pancreatitis, insulin resistance, type 2 diabetes<sup>(4,5)</sup> and infection with *Helicobacter pylori* have also been identified as risk factors of this cancer<sup>(4)</sup>. In addition, pancreatic cancer seems to run in some families, and some of these are caused by an inherited syndrome<sup>(6)</sup>. About 10% of the pancreatic cancer cases are owing to genetic changes, indicating that most people who are diagnosed with pancreatic cancer do not have a family history of it. Potatoes are the fourth most plentiful food crop in the world after wheat, rice and maize<sup>(7)</sup>, and are a good source of fibre, carbohydrates (starches), niacin, folate, vitamin C and minerals (e.g. K, Mg

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Abbreviations: GI, glycaemic index; HR, hazard ratio.

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and Fe)<sup>(8,9)</sup>. According to World Cancer Research Fund/American Institute of Cancer Research report (WCRF/AICR), the evidence for associations between folate, vitamin C, carbohydrates and pancreatic cancer risk is weak or inconclusive<sup>(5)</sup>. Generally, the research on health effects of potato consumption is limited and contradictory, especially regarding the long-term health effects of potato in diets worldwide<sup>(10,11)</sup>. Yet, some studies have found a high potato consumption to be associated with higher risk of pancreatic cancer<sup>(12)</sup>, colon cancer (tendency)<sup>(13)</sup> and colorectal cancer<sup>(14)</sup>. Other studies have found associations between high potato consumption and risk of gastric cancer among women but not among men<sup>(15)</sup>, and risk of rectal cancer among Whites but not among African-Americans<sup>(16)</sup>. An association between potato consumption and risk of oral and pharyngeal cancer has also been observed<sup>(17)</sup>. On the other hand, some short-term studies have implicated that potatoes contain anti-tumour agents<sup>(11)</sup>. Moreover, a casecontrol study found that potato consumption had a protective effect on rectal cancer among women, but no associations were found for men and, in addition, no associations between potato consumption and risk of colon cancer was found<sup>(18)</sup>. A prospective study of intake of fibre from different sources and risk of colon cancer found that fibre from potatoes was inversely related to colon cancer among men; however, for women the intake of potato fibre was associated with higher risk<sup>(19)</sup>. In addition, two case-control studies found potato consumption to be associated with decreased risk of bladder cancer<sup>(20,21)</sup>. Potatoes have a high glycaemic index (GI) and glycaemic load (GL)<sup>(22-24)</sup>. Even though two earlier meta-analyses concluded that there was no association between GI/GL and pancreatic cancer<sup>(25,26)</sup>, recent studies have found the opposite<sup>(27,28)</sup>. Potatoes are usually eaten as part of a meal<sup>(9)</sup>; therefore, the effect potatoes have on disease risk alone is difficult to determine. A typical Western dietary pattern with high consumption of red and processed meat, potato chips, high fat, dairy products and eggs has been associated with a higher risk of pancreatic cancer among men<sup>(29,30)</sup>, although there are other studies in which this association has not been found<sup>(31)</sup>. Preparation methods of potatoes vary highly across the world, and this is relevant to consider when studying potato consumption and risk of disease. Boiling is assumed to be the healthiest way of preparing, as frying and roasting adds other components, such as fats and salt, to the meal. Acrylamide formation may also be a problem when potatoes are cooked at high temperatures<sup>(22,23)</sup>. However, in the Scandinavian countries, boiling is the most common preparation method<sup>(32-34)</sup>. Information from 24-h dietary recalls performed as part of the European Prospective Investigation into Cancer and Nutrition (EPIC) study showed that boiling was preferred in 70-88% (standard error of the estimates for Norway, Sweden and Denmark: 0.01) of the meals containing potatoes in the Danish, Swedish and Norwegian populations used for the current study<sup>(14)</sup>. To our knowledge, no prospective study has investigated the association between potatoes as a single food, rather than potatoes being part of a dietary pattern, and the risk of pancreatic cancer. The aim of the present study was to investigate the associations between potato consumption and the risk of pancreatic cancer in a prospective cohort of women and men in the Scandinavian countries.

#### Methods

# The HELGA study

The HELGA Study is a large population-based Scandinavian cohort consisting of 119978 participants from The Norwegian Women and Cancer study<sup>(35)</sup>, the Northern Sweden Health and Disease Study Cohort<sup>(36)</sup> and the Danish Diet, Cancer and Health Study<sup>(37)</sup>. The three cohorts are also part of the EPIC study<sup>(38)</sup>. Data collection was carried out in 1998-1999 (Norway), 1992-1996 (Sweden) and in 1993-1997 (Denmark). In the Norwegian cohort, women aged 40-55 years, from all over Norway, were selected randomly and invited to participate. In the Swedish cohort, the participants consisted of men and women, aged 30, 40, 50 or 60 years, who all were invited to attend a general health screening within the northern county of Västerbotten. The Danish participants were inhabitants in the Copenhagen and Aarhus areas, aged 50-64 years, free of cancer and born in the country who were invited to participate in the Diet, Cancer, and Health study. The participants provided informed consent, and medical ethics review boards in each of the three countries approved the studies.

## Dietary assessment

Semi-quantitative FFO, filled in at baseline, were used for dietary assessment. The country-specific FFO reflected the habitual diet during the previous year, and they have all been validated and described in detail elsewhere<sup>(39-41)</sup>. The Norwegian FFQ consisted of eighty-five frequency questions, the Danish consisted of 192 frequency questions and the Swedish consisted of eighty-four frequency questions. The questions on potato consumption varied between the three countries. In Norway, the participants were asked one general question on how many potatoes they ate (never/seldom, 1-4/week, 5-6/ week, 1/d, 2/d, 3/d or 4+/d). No questions on preparation method were asked. In Denmark, the questionnaire contained seven different questions, where the participants chose which preparation method (boiled, baked, roasted, mashed, stewed, potato salad or French fries) they had used, and they were asked how many times during a specific period of time they ate potatoes prepared in these various ways: never, during a month (<1, 1, 2-3 times), per week (1, 2-4, 5-6 times) or per day (1, 2-3, 4-5, 6-7, 8+ times). For boiled and baked potatoes, the portion size was specified as one potato, and the participants were asked how many times during a specific period of time they ate one boiled or baked potato (e.g. 1 potato 5-6 times/ week would be 5-6 potatoes/week). In Sweden, the questionnaire contained five questions where the participants chose which preparation method (boiled/baked, roasted, French fries, mashed or potato salad) they had used, and how many times during a specific period of time they ate potatoes prepared in various ways: never, sometimes during a year, per month (1-3 times), per week (1, 2-3, 4-6 times) or per day (1, 2-3, 4+). The amount of food consumed every time was assessed using pictures (a plate with meat, vegetables and potatoes) illustrating four different portion sizes the participants could choose between. In each country, frequencies of all preparation methods were combined with quantities (in g/d) and summed

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into a general variable on potato consumption. Both potatoes as a single food and potato as an ingredient in dishes (pots, soups and hash) were included. The dishes were broken down to the ingredients levels and the ingredients were then added to their respective food groups. As information on preparation methods was unavailable in the Norwegian cohort, and variables for specific preparation methods were not available in the standardised data set, we have focused on total potato consumption. On the basis of the high proportion of boiled potatoes consumed in Norway, Sweden and Denmark<sup>(14,42)</sup>, it is likely that the exposure variable in the present study mainly reflects intake of boiled potatoes. The questionnaires also contained questions on lifestyle and health. The weight and height of the Danish and Swedish participants were measured, whereas in Norway this information was self-reported.

# Identification of pancreatic cancer cases

Over 95% of pancreatic cancers are adenocarcinomas of the exocrine pancreas<sup>(5)</sup>. We included malignant, primary pancreatic cancer of the exocrine pancreas (carcinomas and adenocarcinomas) as defined by the International Classification of Diseases 10th revision as C25 (C25·0–C25·4 and C25·7–C25·9). Information on cancer incidence and vital status was obtained from national cancer registries, and cause of death registries.

#### Exclusions

We excluded 2597 participants with prevalent cancer. The preliminary number of pancreatic cancer cases was 268. We excluded 539 participants, including one case, owing to implausible reported daily energy intake (lower than 2500 kJ for both sexes, and higher than 18000 kJ for women and 21 000 kJ for men) and another twenty-seven (no cases) owing to implausible potato intake (>1 kg/d). As we included only adenocarcinomas from the exocrine pancreas, forty-four pancreatic cancer cases were excluded because they were neuroendocrine pancreatic tumours, lymphoma, carcinoid, malignant cell and malignant tumour. One case was excluded owing to missing information on cancer morphology. Further, we excluded 2494 participants, including one case, owing to missing information on smoking. Finally, thirty-six participants with follow-up time registered as zero were excluded, as they did not contribute to follow-up. Hence, 114240 participants (38766 men and 75474 women) (Norwegian cohort: 33 690, Swedish cohort: 24 305 and Danish cohort: 56 245) were included in the final analyses. Of these, 221 (121 men and 100 women) were diagnosed with pancreatic cancer during follow-up.

# Statistical analysis

Participant characteristics are presented as medians and 5th–95th percentiles and frequency distributions as appropriate. We used Cox proportional hazard models with hazard ratios (HR) and 95% CI to estimate the association between potato consumption and the risk of pancreatic cancer. Follow-up time was used as the time variable, and the participants were followed up from date of entry into the study until the date of

cancer diagnosis, date of death, date of emigration or the end of follow-up (Denmark, 31 December 2007: Sweden and Norway, 31 December 2008). We present one model stratified by sex, and adjusted for age at recruitment, interaction between age and follow-up time and total energy (kI), and one fully adjusted model, additionally adjusted for BMI, height and smoking (see below for more details). Owing to differences in the question formulation and general differences in procedures and measurements between the three cohorts, all analyses were stratified by country. Variables classified as 'probably' or 'convincingly' associated with risk of pancreatic cancer in the WCRF/AICR<sup>(5)</sup> were tested as possible confounders or risk factors: BMI (weight in kg divided by height in m squared, <25 kg/m<sup>2</sup>: underweight/normal-weight, 25–29.9 kg/m<sup>2</sup>: overweight,  $\geq 30 \text{ kg/m}^2$ : obese) and smoking status (never, former and current). We also made a finer categorisation of smoking, dichotomising former smokers into those who quit  $\leq 10$  years ago or ≥11 years ago, and current smokers into those who smoke 1–15 cigarettes/d or  $\geq$ 16 cigarettes/d. However, as this categorisation did not have any material effect on the results, we used the more robust adjustment for smoking status in the final models. Greater childhood growth measured as adult attained height and BMI at aged approximately 20 years is also a probable risk factor, and hence we adjusted for height at baseline. All continuous variables (potato consumption, total energy and height) were tested for linear associations with the outcome. Potato consumption was divided into quartiles in order to fully cover the non-linearity in pancreatic cancer risk, whereas total energy intake was applied in tertiles. Additional variables associated with potato consumption in the Norwegian cohort(42) were assessed for confounding effects: red and processed meat, vegetables, education and diabetes (selfreported). In addition, we did separate analyses by sex. As the number of cases was relatively small, stratification by diabetes was not possible, but in addition to adjust for diabetes we performed sensitivity analyses excluding diabetics. Owing to the small number of cases, we did not perform separate analyses by country, but we repeated all analyses in the Danish cohort only, as this was the largest sub-cohort containing most of the cases. In addition, we performed analyses adjusting for preparation methods in the Danish cohort. We also tested a multivariable adjusted model additionally including dietary factors (red and processed meat, alcohol, carbonated/soft/ isotonic drinks and diluted syrups, and fat) that are characteristic of an obesogenic environment and the metabolic syndrome<sup>(43,44)</sup> (results not shown). As these variables did not materially change the risk estimates, they were not included in the final models. Adjustment for GI was not done, as these estimates were not available from the food composition database, but we constructed a model (data not shown) where we adjusted for intake of pasta and rice to rule out associations with other main sources of dietary carbohydrate used at dinner meals. Variables were included in the final models if they were significantly associated with pancreatic cancer, or if they influenced the HR by more than 10%. We also adjusted for total energy intake, as this adjustment is usually appropriate to control for confounding in studies on disease and diet. In addition, as our study consisted of three sub-cohorts in different

countries, this adjustment attenuated possible differences in the information obtained from the FFQ. Tests for trend were performed for all regression analyses, and quartile medians of potato consumption were used in the tests. We performed sensitivity analyses that excluded participants with a pancreatic cancer diagnosis <1 year  $(n \ 12)$  and 3 years  $(n \ 34)$  after having completed the questionnaire, owing to the possibility that preclinical symptoms affected eating habits. Proportional hazard (PH) assumptions were checked using Schoenfeld residuals, which showed sign of deviation from proportionality. By including an interaction term between age at entry and follow-up time in the models, the PH assumption was fulfilled for all variables. A  $\chi^2$  test was performed to check for heterogeneity between sexes. A restricted cubic spline regression model with four knots at fixed percentiles (5, 35, 65 and 95, as suggested by Harrell<sup>(45)</sup>) was used to further explore dose-response effects between potato consumption and risk of pancreatic cancer. All variables were checked for multi-collinearity using variance inflation factor, and the results showed no violation of this assumption. Data were analysed using STATA version 14. Calculations showed that the sample size and number of incident pancreatic cancer cases in the HELGA study were sufficiently large to detect a HR of 1.5 for the highest quartile of potato intake compared with the lowest, with a statistical power of 85% and significance level of 5%.

# Results

The mean follow-up time was 11.4 (95% CI 0.3, 16.9) years. Median age of participants in the quartiles of potato consumption was 52–53 years (Table 1). The cohort consisted of more women than men (66.1% females) (data not shown), whereas slightly more cases were men (54.8%). The Swedish sub-cohort had the longest follow-up (mean: 13.3 years), whereas the Danish sub-cohort was the largest (49% of the participants), provided most of the cases (68.8%) and had, on average, the oldest participants (results not shown in table).

Participants in the highest quartiles of potato consumption tended to be lower educated, taller and to consume more processed meat, fat and total energy (Table 1). Consumption of red meat and alcohol were noticeably lowest in the lowest quartile, and consumption of carbonated/soft/isotonic drinks and diluted syrups was highest in the lowest and highest quartile. In addition, potato consumption was highest in the Swedish participants, and among men. Smoking showed only minor differences across the quartiles of potato consumption. Minor differences were also seen for intake of vegetables, fruits, pasta and rice. Owing to the small differences between the adjusted models (Table 2), we only present HR and CI from the adjusted model 2. Higher consumption of potatoes showed tendencies of higher risk of pancreatic cancer (HR 1.44; 95% CI 0.93, 2.22,  $P_{\text{for trend}}$  0.03) when comparing the highest v. the lowest quartile of consumption in the multivariable model (Table 2). However, compared with the second quartile, both upper quartiles showed a significantly increased risk (data not shown). In the sex-specific analyses, higher consumption was associated with a higher risk of pancreatic cancer for females (HR 2.00; 95% CI 1.07, 3.72, Pfor trend 0.02). No significant associations were found for men (HR 1-01; 95% CI 0-56, 1-84). Results from the spline regression models confirmed the absence of dose–response effects between potato consumption and risk of pancreatic cancer (Fig. 1). In sensitivity analyses, excluding participants diagnosed with pancreatic cancer <1 (twelve cases) or 3 years (thirty-four cases) after completing the questionnaire did not influence the results. Analysis with a finer categorisation of smoking (including intensity and time since quitting) also did not alter the conclusions. Excluding participants with diabetes (twenty-two cases) did not influence the results. When we repeated all analyses in the Danish cohort (152 cases) only, the overall results were the same (data not shown). Repeating the sex-specific analysis or excluding participants with diabetes in the Danish cohort was not possible owing to the small number of cases.

## Discussion

In this study, we found that participants reporting higher consumption of potatoes showed tendencies at higher risk of pancreatic cancer. In the sex-specific analyses, we found that higher consumption of potatoes was associated with a higher risk of pancreatic cancer for women only. When we further explored the associations by spline regression, the absence of dose-response effects was confirmed. The strengths of this study include its prospective design, which minimises the influence of recall bias. In addition, the high consumption of particularly boiled potatoes in these countries gave us an opportunity to study a population with high potato consumption, where it is not likely that, for example, acrylamide or fat will influence the results. Nevertheless, potatoes are not eaten isolated, but as part of a meal. Further, owing to the use of cancer registries and population registries in the Scandinavian countries, the case coverage was excellent and loss to follow-up was minimal. Another strength of the study is that we, in addition to using standard Cox proportional hazard models, repeated all analyses with spline regression to further explore a possible dose-response effect between high potato consumption and pancreatic cancer risk. It has been discussed that there are other regression methods that could be better suited for analysis than the standard categorical analysis we used at first<sup>(46)</sup>, and one of them is spline regression as standard categorical analysis does not make efficient use of within-category information. As the standard categorical analysis showed relatively inconsistent and non-linear results, with high potato consumption being associated with higher risk of pancreatic cancer in some subgroups, it was important to explore this further. As the results from the spline regression supported the results from the Cox regression, and there was no apparent biological evidence to support the results, we cannot, on the basis of our results, conclude that there is an association between potato consumption and pancreatic cancer. One limitation of the study was that even though the study had a large sample size, the number of cases was relatively small, as expected with a relatively rare cancer type. This restricted our possibility to perform subgroup and stratified analyses, as this affected the statistical power. We tried solving this by, for

#### Table 1. Demographic, lifestyle and dietary characteristics by potato consumption in the HELGA cohort (Medians and ranges and 5th–95th percentiles)

	1st quartile (0–73⋅3 g)		2nd quartile (73·3–126·3 g)		3rd quartile (126·3–185·9 g)		4th quartile (185·9–999·5 g)	
Characteristics	Median	5th–95th percentile	Median	5th–95th percentile	Median	5th–95th percentile	Median	5th–95th percentile
n		28 103		28 800		28748		28 589
Cases of pancreatic cancer		35		43		67		76
Age at recruitment								
Median		51.0		53.0		52.8		52.7
Range	40	).3–61.7	40	0.0-63.3	39.9-63.5		30.6-63.6	
Sex (%)								
Men		16·9		29.7		36.5		52.3
Women		83.1		70.3		63·5		47.7
Country (%)								
Denmark		34.9		58.3		53.3		50·1
Sweden		12.9		17.8		22.2		32.0
Norway		52.2		23.9		24.5		17.9
Smoking status		022		200		240		17.5
Never		39.2		42.8		41.2		39.1
Former		30.0		28.0		28.1		27.6
Current		30.8		29·2		30.7		33.3
		30.0		29.2		30.7		33.3
Highest level of education completed (%)								
None/primary school		22.8		27.4		30.3		35.6
Technical/professional school	34.9		36.9		34.1		30.6	
Secondary school Higher education (including	24·4 17·9		18·3 17·4		17·9 17·7		16·8 17·0	
university)								
Diabetes (%)								
Diabetics		1.8		1.7		1.7		1.8
Non-diabetics	96.7		95.8		96.0		95.7	
Unknown		1.6		2.5		2.2		2.5
BMI (kg/m <sup>2</sup> )	24.3	19.7-32.3	24.8	20.0-32.7	24.9	20.0-32.6	25.0	20.0-32.7
Height (cm)	167	157-181	168	156-183	169	157-184	171	158-186
Consumption								
Vegetables (g/d)	122	25–309	133	28–335	139	30–340	127	25–343
Fruits (g/d)	133	17-425	137	22-433	141	23-433	131	20-433
Red meat (g/d)	25	4-119	42	6-119	43	7–130	43	8-145
Processed meat (g/d)	25	4–73	25	6–68	30	7-76	37	9–94
Pasta (g/d)	19	0-79	19	1-84	18	0-84	16	0-84
Rice (g/d)	16	0-64	16	1-64	16	1–75	14	1–75
Carbonated/soft/isotonic drinks, diluted syrups(q/d)	39	0-454	29	0–500	33	0–500	39	0–507
Alcohol (g/d)	2.6	0-37.3	6.0	0-46.2	5.3	0–50.3	5.3	0-56.4
	2·0 58	31–104	67	0–46·2 36–117	5·3 73	40–127	5.3 82	0-36-4 44-146
Total fats (g/d)	58 6666	31–104 3912–10709		4616-12060	-	40-127		44-146 5674-14912
Total energy (kJ/d)	0000	5912-10709	/011	+010-12000	0200	3037-13038	900Z	5074-1491

instance, excluding those with diabetes instead of performing separate analysis on diabetics. The results remained unchanged. Adjustments for diabetes did not have any material effect on the results either. We applied a finer categorisation for smoking status, which did not have any effect on the results. Nevertheless, the study will probably suffer from some residual confounding. Our results were based on data from selfadministered questionnaires, which is a potential source of measurement error. However, the food data were standardised by common standardisation guidelines. The questions on potato consumption varied across the three countries, and the questionnaire in the Norwegian cohort did not have any questions on preparation method. Therefore, total potato consumption was summed up into a general variable in g/d. Ideally, questions on preparation method should have been included; however, information from 24-h dietary recalls performed as part of the EPIC study showed that boiling was

the most common preparation method in meals containing potatoes in the Danish, Swedish and Norwegian populations included in the current study<sup>(14)</sup>. Adjustment for GI was not done, as these estimates were not available from the food composition database. In addition, information on H. pylori, insulin resistance and chronic pancreatitis, which have been associated with pancreatic cancer, was not available. In addition, it is important to keep in mind that the results of the present study were based on only one measurement, and dietary changes could have occurred during the follow-up period. As potatoes are usually eaten as part of a meal, it is also important to keep in mind that the effect potatoes have on disease risk alone is difficult to determine. Foods that are part of a meal interact with other dietary components and nutrients, and thus the impact of potato consumption on disease risk may depend on which other foods they are grouped with in a dietary pattern<sup>(9)</sup>. These issues are also present regarding GI, as the

Table 2. Risk\* for pancreatic cancer according to potato consumption in the HELGA study (n 114240)

(Hazard ratios and 95% confidence intervals)

		Adjust	ed model 1†	Adjusted model 2‡		
Potatoes (g/d)	Cases	HR	95 % CI	HR	95 % CI	
Both sexes						
Q1	35	Ref.		Ref.		
Q2	43	0.87	0.55, 1.38	0.87	0.55, 1.38	
Q3	67	1.33	0.87, 2.05	1.32	0.86, 2.03	
Q4	76	1.45	0.94, 2.25	1.44	0.93, 2.22	
P§			0.026	0.0301		
Females ( <i>n</i> 75 474)						
Q1: 0–73·3	16		Ref.	Ref.		
Q2: 73·3–126·2	17	1.14	0·61, 2·11	1.15	0.62, 2.12	
Q3: 126·3–185·9	39	1.51	0.82, 2.77	1.49	0.81, 2.73	
Q4: 185·9–910·2	49	2.05	1.10, 3.81∥	2.00	1.07, 3.72∥	
P§			0.016	0.020		
Males ( <i>n</i> 38766)						
Q1: 0–73·3	19		Ref.	Ref.		
Q2: 73·3–126·3	26	0.58	0.29, 1.17	0.58	0.29, 1.17	
Q3: 126·3–185·9	28	1.06	0.58, 1.94	1.06	0.58, 1.94	
Q4: 185·9–999·5	27	1.01	0.56, 1.84	1.01	0.56, 1.84	
P§			0.349	0.344		

Q, quartile; Ref., referent values.

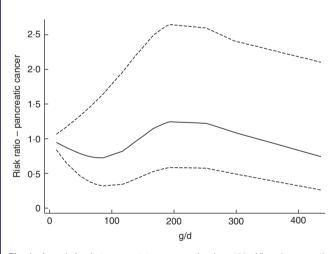
\* As estimated by Cox proportional hazards regression.

† Stratified by country and sex, and adjusted for age at recruitment, interaction between age and time, and total energy (kJ).

‡ Additionally adjusted for BMI, height and smoking.

§ Test for trend, based on median potato consumption in each quartile.

Il Values are significant.



**Fig. 1.** Association between potato consumption (per 100 g/d) and pancreatic cancer risk. As estimated by a restricted cubic Spline regression model with four knots at fixed percentiles (5, 35, 65, 95, suggested by Harrell<sup>(45)</sup>). Adjusted for sex, total energy (kJ), BMI, height, smoking status and stratified by country.

postprandial glycaemic response is influenced by several factors in a meal, such as the source and amount of carbohydrates and the type and amount of fibre present<sup>(9)</sup>. In addition, the ingredients or other foods eaten together with potatoes will influence the GI and the postprandial glycaemic response. The variety of potatoes and the preparation method will also play a part<sup>(9)</sup>. The research on pancreatic cancer and GI is inconsistent; two earlier meta-analyses concluded that there was no association<sup>(25,26)</sup>, whereas two recent studies found that diets with high GI and GL increased the risk of pancreatic cancer<sup>(27,28)</sup>. As mentioned, adjustment for GI was not possible in the present study, but we constructed a model (data not shown) where we adjusted for intake of pasta and rice to rule out associations with other main sources of dietary carbohydrate used at dinner meals, and the results were still the same. Adjusting for preparation methods (boiled, baked, fried, roasted, mashed, potato salad and stewed) in the Danish cohort did not change the original results (results not shown). Participants in the highest quartile of potato consumption showed some characteristics that are known to be associated with pancreatic cancer, like they tended to have a higher BMI. The participants in the highest quartile were also slightly taller. Regarding dietary factors, they consumed more processed meat, and had a generally higher intake of total fat and total daily energy intake. The evidence regarding these dietary factors in association with pancreatic cancer is limited<sup>(5)</sup>; however, these factors are characteristics of an obesogenic environment and the metabolic syndrome<sup>(43,44)</sup>. The metabolic syndrome is a cluster of risk factors related to risk for stroke, heart disease, diabetes type 2, overweight, obesity and insulin resistance<sup>(43)</sup>, and the incidence of this syndrome increases with age<sup>(47)</sup>. However, multivariable adjusted models including these variables did not materially change the risk estimated, and they were not included in the final models. The main reason for not including them was the limited number of pancreatic cancer cases in our study sample, and we aimed to retain power in our statistical analyses. The metabolic syndrome has become more common owing to increasing obesity rates among adults<sup>(43)</sup>. In addition, several studies have found the metabolic syndrome to be associated with a higher risk of pancreatic cancer<sup>(48,49)</sup>. The key component of the metabolic syndrome in pancreatic carcinogenesis was suggested to be diabetes<sup>(49)</sup>. As mentioned, exclusion of participants with diabetes (twenty-two cases) did not influence the results in the present study. Even though there is a lack of research regarding long-term cancer-related health effects of potatoes<sup>(11)</sup>, there are several studies that have found an association between potato consumption and cancer: in particular, one case-control study found that a high potato consumption was associated with a higher risk of pancreatic cancer<sup>(12)</sup>. It is also interesting that some studies on dietary patterns, rather than studies on potatoes per se, show that a western dietary pattern that includes potatoes has been associated with a higher risk of pancreatic cancer among men<sup>(29,30)</sup>. However, these findings are not consistent<sup>(31)</sup>. In addition, some studies have found potato consumption to be associated with a higher risk of other cancers of the digestive system, such as rectal cancer<sup>(16)</sup>, colon cancer (tendencies)<sup>(13)</sup> and colorectal cancer<sup>(14)</sup>. Other studies have shown contradictory or inconclusive results regarding gastric, colon and rectal cancer<sup>(15,18,19)</sup>.

# Conclusion

In this study, we cannot conclude that there is an association between potato consumption and risk of pancreatic cancer. However, potatoes are a staple food in many countries, and there are generally few and inconsistent studies within this field. In addition, the inconsistent findings in the present study emphasise the need for more and larger studies investigating the association between potato consumption and pancreatic cancer, and potatoes and health in general.

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