#### Reported amount of salt added to food is associated with increased all-cause and

#### cancer-related mortality in older men in a prospective cohort study.

Jonathan Golledge MChir<sup>1,2</sup>

Joseph V. Moxon PhD<sup>1</sup>

Rhondda E. Jones PhD<sup>3</sup>

Kieran McCaul PhD<sup>4</sup>

Graeme J. Hankey PhD<sup>5,6</sup>

Bu B. Yeap PhD<sup>457</sup>

Leon Flicker PhD<sup>4,,5,7</sup>

Paul E. Norman DS<sup>8</sup>

<sup>1</sup> Queensland Research Centre for Peripheral Vascular Disease, School of Medicine and Dentistry, James Cook University, Townsville, Australia. Email: jonathan.golledge@jcu.edu.au;

<sup>2</sup> Department of Vascular and Endovascular Surgery, The Townsville Hospital, Townsville, Australia;

<sup>3</sup> The Australian Institute of Tropical Health and Medicine, James Cook University, Townsville, Australia.

<sup>3</sup> WA Centre for Health & Ageing, Centre for Medical Research, Perth, Australia. Email: kieran.mccaul@uwa.edu.au;

<sup>4</sup> School of Medicine and Pharmacology, University of Western Australia, Perth, Australia. Email: graeme.hankey@uwa.edu.au

<sup>5</sup> Department of Neurology, Sir Charles Gairdner Hospital, Nedlands, Perth, Australia

<sup>6</sup> Department of Endocrinology, Fremantle Hospital, Fremantle, Australia. Email: bu.yeap@uwa.edu.au;

<sup>7</sup> Department of Geriatric Medicine, Royal Perth Hospital, Perth, Australia. Email: leon.flicker@uwa.edu.au;

<sup>8</sup> School of Surgery, University of Western Australia, Perth, Australia. Email: paul.norman@uwa.edu.au.

Correspondence to: Professor Jonathan Golledge, Director, The Vascular Biology Unit,

Queensland Research Centre for Peripheral Vascular Disease, School of Medicine and

Dentistry, School of Medicine and Dentistry, James Cook University Townsville, QLD,

Australia 4811.

Fax +61 7 4433 1401 Telephone +61 7 4433 1417

Email: jonathan.golledge@jcu.edu.au

Abstract word count: 249 Text word count: 2439 Three tables Three figures Running title: Salt intake and mortality.

### Funding

The Health In Men Study has been supported by grants from National Health and Medical Research Council (279408, 379600, 403963, 513823, 540403, 540504, 540405, 634492, 1021416, 1045710 and 1060557). Additional funding from the Queensland Government supported this work. JG holds a Practitioner Fellowship from the National Health and Medical Research Council, Australia (1019921). JG holds a Senior Clinical Research

Fellowship from the Office of Health and Medical Research. The funders had no role in study

design, data collection and analysis, decision to publish, or preparation of the manuscript.

The authors have no other disclaimers relevant to this article.

1 Abstract

2 *Background:* The effect of dietary salt intake on important population outcomes such as

3 mortality is controversial. The aim of this study was to examine the association between the

4 dietary habit of adding salt to food and mortality in older men.

5 Design, participants, setting and measurements: A risk factor questionnaire which contained

6 a question about the dietary habit of adding salt to food was completed by 11742 community

7 recruited older men between 1996 and 1999. The men were followed by means of the

8 Western Australia Data Linkage System until November 30<sup>th</sup> 2010. Deaths due to

9 cardiovascular diseases and cancers were identified using ICD-10 codes in the ranges I00-I99

and C00-D48, respectively. The association between the frequencies of adding salt to food

and mortality was assessed using Kaplan Meier estimates and Cox proportional hazard

12 analysis.

13 *Results:* Median follow-up for survivors was 13.1 years (range 11.8-14.6 years). A total of

14 5399 deaths occurred of which the primary cause registered was cancer and cardiovascular

disease in 1962 (36.3%) and 1835 (34.0%) men, respectively. The reported frequency of

adding salt to food was strongly positively associated with all-cause (p<0.001), cancer-related

17 (p<0.001) but not cardiovascular-related (p=0.649) mortality. Men reporting adding salt to

18 their food always had a 1.12-fold (95% CI 1.05-1.20, p<0.001) and a 1.20-fold (95% CI 1.07-

19 1.34, p=0.001) increased risk of all-cause and cancer-related mortality, respectively, after

adjusting for other risk factors. Men reporting adding salt to their food sometimes had a 1.17-

fold (95% CI 1.05-1.30, p=0.004) increased risk of cancer-related mortality after adjusting for

22 other risk factors.

23 *Conclusion:* A history of adding salt to food is associated with increased cancer-related

24 mortality in older men.

25

*Key words:* Salt; mortality; men.

#### 28 Introduction

29	Good evidence suggests that high salt intake is associated with hypertension and current
30	clinical guidelines and public health policies recommend low salt intake [1-4]. Despite these
31	recommendations the evidence that reducing dietary salt intake decreases mortality is limited
32	and sodium is accepted to be an essential extracellular cation required to maintain
33	hydroelectric balance [5-13]. Findings from a number of studies have associated low salt
34	intake with increased mortality [5-11]. High salt intake has also associated with increased
35	mortality [12, 13]. The previous studies have not been focused on community recruited older
36	men or assessed reported salt added to food which is an aspect easier to assess in community
37	samples. The aim of the current study was to examine the association of the dietary habit of
38	adding salt to food with mortality in a large cohort of older men recruited as part of a
39	community screening study.
40	

#### 41 Methods

42 *Study population:* The Health in Men Study (HIMS) developed from a population-based

43 randomized trial of screening for abdominal aortic aneurysm (AAA) conducted in Perth,

44 Western Australia between 1996 and 1999 which has been previously described in detail [14,

45 15]. Ethics approval for the study was provided by The University of Western Australia

46 Ethics Committee (Project numbers RA/4/1/5765) and all men provided written informed

47 consent.

48

49 Assessment of recruited men: Each man was invited to complete a questionnaire assessing

50 aspects of history and lifestyle relevant to AAA and cardiovascular disease including:

51 smoking history; history of diagnosis of high blood pressure, angina, myocardial infarction,

52 stroke, diabetes and high cholesterol; history of treatment for high blood pressure, angina,

53	diabetes and high cholesterol; frequency of eating meat ( $\geq 6$ times/week, 3–5/week, 1–
54	$2$ /week, <1/week or never); frequency of eating fish ( $\geq 6$ times/week, 3–5/week, 1–2/week,
55	<1/week or never); and hours of non-vigorous exercise (none, $\leq$ 2 hours/ week, >2-4 hours/
56	week, >4-6 hours/ week or >6 hours/ week). Salt addition to food was assessed with the
57	following question 'Do you add salt to your food?' with three possible answers: (a) rarely or
58	never, (b) sometimes, (c) almost always or always. Waist and hip circumference were
59	measured in accordance with guidelines of the International Society for the Advancement of
60	Kinanthropometry [16]. Body mass index was calculated as weight in kilograms divided by
61	height in meters squared as previously described [16]. The greatest transverse and antero-
62	posterior diameter of the infra-renal aorta was measured using a Toshiba Capasee ultrasound
63	machine with a 3.75 MHz probe (Toshiba Australia, North Ryde, NSW). Assessment of
64	intraobserver and interobserver reproducibility in aortic diameter measurement was carried
65	out every 4 months on 10 randomly selected subjects, as previously reported [17]. No
66	significant differences were found between observers with 95% of measurement differences
67	being $<3$ mm [17]. An AAA was defined by infra-renal aortic diameter $\geq$ 30mm.
68	
69	Follow-up and outcome assessment: All men were followed from the time of recruitment
70	until 30th November 2010 by means of the Western Australia Data Linkage System. Deaths
71	due to cardiovascular diseases and cancers were identified from the Death Registry using
72	ICD-10 codes in the ranges I00-I99 and C00-D48, respectively, as previously described [18].
73	The validity of data within the Western Australia linked Death Registry has been previously
74	assessed and found to be good [14].
75	
76	Statistical analyses: All analyses were performed using IBM SSPS Statistics version 22 (St

76 *Statistical analyses:* All analyses were performed using IBM SSPS Statistics version 22 (St.

77 Leonards, New South Wales, Australia), and the publically available R software package.

78	The association of reported salt added to food with all-cause, cancer-related and
79	cardiovascular-related mortality was assessed using Kaplan Meier estimates and Cox
80	proportional hazard analysis. For these analyses men that were still alive were censored at the
81	time of the data linkage. For the cause specific death analyses all men were included and men
82	who died of causes unrelated to the outcome of interest were censored at the date of their
83	death. Initially univariate Cox proportional hazard analysis was performed to assess the
84	association of individual risk factors with: i) all-cause, ii) or cancer-related mortality.
85	Subsequently the association of reported salt added to food with all-cause mortality was
86	adjusted for age (per 5 years), past treatment for hypertension, past treatment for angina, past
87	history of myocardial infarction, past history of stroke, past treatment for diabetes, ever
88	smoking, waist to hip ratio), frequency of eating fish, frequency of non-vigorous exercise and
89	AAA presence, based on significant associations of the risk with all-cause mortality
90	following univariate Cox regression. Similarly, the association of reported salt added to food
91	with cancer-related mortality was adjusted for age (per 5 years), past treatment for
92	dyslipidaemia, ever smoking, waist to hip ratio, body mass index, frequency of eating meat,
93	frequency of non-vigorous exercise and AAA presence. The proportional hazards assumption
94	was assessed for models predicting all-cause or cancer-relating mortality. In order to fulfil the
95	proportional hazards assumption during multivariate analyses, participants were re-
96	categorised into groups with waist to hip ratios of 0.9-1.02, and >1.02. Similarly, participants
97	were re-categorised into groups with BMI of 20-30, 30-49 and >40.
98	
99	Cumulative mortality was compared between men who reported adding salt to their food
100	never, sometimes or always using log rank test.
101	

#### 103 **Results**

#### 104 *Characteristics of the included men*

- 105 Risk factors for the 11742 men reporting the frequencies of adding salt to food at the time of
- recruitment have been previously published [19] [20].
- 107 Median follow-up for survivors was 13.1 years (range 11.8-14.6 years). A total of 5399
- 108 deaths occurred during follow-up of which the primary cause registered was cancer and
- 109 cardiovascular disease in 1962 (36.3%) and 1835 (34.0%) men, respectively. The types of
- 110 cancer registered as the primary cause of death included those of respiratory tract (n=503),
- 111 gastro-intestinal tract (n=473), urogenital (n=373), hematological (n=237) and miscellaneous
- 112 (including skin, soft tissue, muscle, skeletal, brain, thyroid, multiple sites and unknown site;
- n=376) origins. The all-cause mortality rates were 13.2, 32.3 and 45.1% at 5, 10 and 13 years,
- respectively. The cancer-related mortality rates were 5.7, 13.9 and 19.3% at 5, 10 and 13
- 115 years, respectively. The cardiovascular disease-related mortality rates were 5.0, 12.5 and
- 116 18.3% at 5, 10 and 13 years, respectively.
- 117

#### 118 Association of reported frequencies of adding salt to food with mortality

119 Figures 1-3 illustrate the relationship between reported frequencies of adding salt to food and

subsequent all-cause, cancer-related and cardiovascular-related mortalities. Reported

- 121 frequencies of adding salt to food was strongly positively associated with all-cause (Figure 1;
- 122 p<0.001), cancer-related (Figure 2; p<0.001) but not cardiovascular-related (Figure 3;

123 p=0.649) mortality. Men reporting the addition of salt to food never, sometimes or always

- had a cumulative incidence of all-cause mortality of 43.3, 45.0 and 47.6% at 13 years,
- respectively. Men reporting the addition of salt to food never, sometimes or always had a
- 126 cumulative incidence of cancer-related mortality of 16.9, 20.2 and 21.3% at 13 years,
- 127 respectively. Men reporting the addition of salt to food never, sometimes or always had a

cumulative incidence of cardiovascular disease-related mortality of 18.5, 18.1 and 18.1% at

129 13 years, respectively.

130

#### 131 Creating multivariate models to predict all-cause and cancer-related mortality

In order to further assess the association of reported frequencies of adding salt to food with mortality univariate Cox proportional hazard ratios were calculated to assess the association of baseline risk factors with all-cause and cancer-related mortality (Table 1). Risk factors showing significant associations with each outcome via univariate regression were included as covariates in multivariable Cox proportional hazards models to assess the impact of salt consumption on all-cause and cancer-related mortality as appropriate. Ten men with

138 incomplete risk factor data were excluded from multivariate analysis (n for multivariable

139 analyses = 11732).

140

141 Diagnostic statistics demonstrated that the multivariable Cox regression model assessing the 142 relationship of salt consumption with all-cause mortality did not conform with the 143 proportional hazards assumption. To correct this, several variables (previous history of 144 diabetes, ever smoking and frequency of non-vigorous exercise) were stratified prior to 145 entering the model (Table 2). After adjusting for potential confounders, men who reported 146 that they always added salt to their foods had a 1.12-fold (95% CI 1.05-1.20 p<0.001) 147 increased risk of all-cause mortality compared to those who never added salt to their food 148 (Table 2). No significant difference in all-cause mortality was noted for men who sometimes 149 added salt to their food.

150

151 The model assessing the association of salt consumption with cancer-related mortality

152 conformed to the proportional hazards assumption, thus, no further data manipulations were

- 153 performed. Men who reported sometimes or always adding salt to their food had significantly
- increased risk of cancer-related mortality (hazards ratio: 1.16 (95% CI 1.04-1.29), and 1.20
- 155 (95% CI 1.07-1.34) respectively), compared to those who never added salt (Table 3).
- 156
- 157

#### 158 **Discussion**

159 The current study examined the incidence of mortality in a group of community recruited

- 160 older men over a long follow-up of approximately 13 years. Approximately 70% of deaths
- 161 were secondary to cardiovascular and cancer-related causes in keeping with the accepted
- 162 main causes of mortality in Western communities. The main finding from this study was that
- always adding salt to food was associated with increased all-cause and cancer-related

164 mortality in older men. The reliability of this association is supported by the large number of

165 men examined (11742), the long follow-up and the adjustment for potential confounding risk

166 factors. Furthermore the validity of the data is supported by the expected associated of age,

167 cardiovascular risk factors and past history of cardiovascular disease with mortality.

168

Randomized controlled trials suggest that limiting salt intake can reduce resting systolic
blood pressure by approximately 3-4 mmHg during short term follow-up [2, 21]. Randomized
trials have however failed to demonstrate convincingly that limiting dietary salt intake
reduces cardiovascular events or mortality possibly because these studies have been under
powered [22]. Restricting sodium intake has also been associated with some detrimental
effects in experimental studies such as activation of the renin-angiotensin system [23, 24].
Thus the value of dietary salt restriction in improving health is currently controversial [1, 25].

177 A number of prospective studies have examined the association of measures of dietary salt 178 intake, such as dietary questionnaires or 24 hour sodium excretion with mortality with 179 conflicting results [5-13, 26-30].. Three community based studies have associated high salt 180 intake with increased cardiovascular disease or stroke-related mortality in Japan and Europe 181 [12, 14, 26]. In contrast community based studies in the USA and Europe have associated low 182 salt intake with increased mortality [5, 9, 11]. Furthermore studies in patients with diabetes 183 and renal failure have also associated low salt intake with increased mortality [7, 8, 10]. 184 Some studies have suggested that the association between salt intake and cardiovascular 185 death is J-shaped with subjects with low and high sodium excretion having increased 186 mortality[6]. The current study is one of the largest studies to assess the association of 187 adding salt to food with mortality and of note included follow-up for over ten years. While 188 adding salt to food was assessed by a simple question this approach was a very practical way 189 of assessing a large population of older men. It is also possibly a more practical way of 190 advising patients on dietary behavior in that we looked at the specific practice of adding salt 191 to food rather than measures of total salt intake. Data using this approach is also relevant to 192 advising older subjects who may find it very difficult to gauge accurate estimates of sodium 193 intake. Overall we found no association between reported frequency of adding salt to food 194 and cardiovascular mortality.

195

There are a number of possible reasons for this finding. It is possible that high dietary salt intake while predisposing to higher blood pressure in the short term may stimulate other mechanisms in the longer term which correct blood pressure. In support of this theory we previously found no association between reported frequencies of adding salt to food and resting blood pressure [20][19]. Most of the trials examining the effect of modifying salt intake on blood pressure have follow-up limited to weeks [2, 21]. It is also possible that a

single assessment of the frequency of adding salt to food may not be reflective of dietary
behavior over a prolonged follow-up period, and that change in salt consumption during the
period of follow-up might not have been captured. These considerations may have
complicated our assessment of the association of reported frequencies of adding salt to food
with mortality although we adjusted our analyses for cardiovascular risk factors and past
history of cardiovascular disease.

208

209 The association between dietary salt intake and cancer-related mortality has been relatively 210 little studied [27-30]. High dietary salt intake has been positively associated with mortality 211 from stomach cancer in Japanese, Chinese and European populations [26-29]. In the current 212 study men reporting adding salt to their food always had a 1.22-fold increased incidence of 213 cancer-related mortality. This association remained after adjusting for other risk factors that 214 we examined. This finding is in line with experimental and epidemiology data suggesting the 215 role of salt in promoting some cancers such as those within the gastro-intestinal tract [31, 32]. 216 As expected in a cohort of older men the reported cancer types in this series included not only 217 gastro-intestinal but also respiratory, urogenital, hematological and those from other sites. 218 Thus it is possible that the behavior of adding salt always to food may promote cancers at 219 sites other than the gastro-intestinal tract although this requires more specific assessment.

220

A number of possible limitations of this study should be considered including measurement error, reverse causality and residual confounding. Firstly, our assessment of salt added to food was limited to a simple but practical question in which we asked whether salt was added to food never or rarely, sometimes, almost always or always. More sophisticated assessment methods, such as measured of 24-hour urinary sodium excretion, were not used. This approach may have introduced measurement error. It is however accepted that even

227 biochemical methods of estimating salt intake are open to measurement error and self-228 reported dietary intake of salt has been found to be reflective of 24 hour urinary sodium 229 excretion, suggesting that self-report is a valid measure of salt intake[33]. Secondly, we only 230 examined salt added to food on one occasion rather than repeated assessments which would 231 have been ideal. Thirdly, this study is a prospective longitudinal human association study. It 232 is not possible to definitively conclude that the association between always adding salt to 233 food and mortality is causative. The direct role of salt in mortality could only be established 234 by a randomized controlled trial of at risk individuals in which the effect of administering 235 different amounts of salt was compared. Based on data from the current study such a trial 236 would require a large number of subjects and extended follow-up in order to assess the 237 efficacy of salt restriction in limiting mortality. Fourthly, we may have failed to adjust for 238 some confounding factors. The current study included a large number of men and used 239 adjustment for recognized confounding factors such as age, hypertension, high cholesterol, 240 coronary heart disease and stroke. It is possible that other confounding factors which we were 241 not able to assess, such as fruit and vegetable intake, may have contributed to our finding. 242 243 In conclusion the current study suggests that the addition of salt to food always is associated 244 with increased mortality in older men through the promotion of cancer-related deaths. This 245 information supports the concept that dietary salt addition to food should be limited.

246

#### 247 Acknowledgements

The authors thank the men who contributed to the Health in Men Study. We thank the Data
Linkage Unit, Health Department of WA, for their assistance. All individuals who contributed
significantly to this study have been listed as authors.

- 251 *Conflict of interest:* The authors received grant funding which assisted with the completion of
- this work as listed in the funding section on the title page. No other conflicts of interest are
- 253 disclosed.
- 254 *Author roles:*
- 255 Study design: JG; Data collection: GJH, BBY, LF, PEN; Data analysis: JG, JVM, REJ KM;
- 256 Data interpretation: All authors; drafting of manuscript: JG; Critical revision of manuscript:
- 257 All authors.
- 258 *Sponsors role:* The funding bodies had no role in the design, methods, data collection, analysis
- and preparation of this report.
- 260

 Kotchen TA, Cowley AW Jr, Frohlich ED. Salt in health and disease--a delicate balance. N Engl J Med. 2013;368:1229-37.

2. He FJ, Li J, Macgregor GA. Effect of longer term modest salt reduction on blood pressure:

Cochrane systematic review and meta-analysis of randomised trials. BMJ. 2013;346:f1325.

3. Strazzullo P, D'Elia L, Kandala NB, et al. Salt intake, stroke, and cardiovascular disease: metaanalysis of prospective studies. BMJ. 2009;339:b4567.

4. Whelton PK, Appel LJ, Sacco RL, et al. Sodium, blood pressure, and cardiovascular disease: further evidence supporting the American Heart Association sodium reduction recommendations. Circulation. 2012;126:2880-9.

5. Stolarz-Skrzypek K, Kuznetsova T, Thijs L, et al.; European Project on Genes in Hypertension (EPOGH) Investigators. Fatal and nonfatal outcomes, incidence of hypertension, and blood pressure changes in relation to urinary sodium excretion. JAMA. 2011;305:1777-85.

6. O'Donnell MJ, Yusuf S, Mente A, et al. Urinary sodium and potassium excretion and risk of cardiovascular events. JAMA. 2011;306:2229-38.

7. Thomas MC, Moran J, Forsblom C, et al.; FinnDiane Study Group. The association between dietary sodium intake, ESRD, and all-cause mortality in patients with type 1 diabetes. Diabetes Care. 2011;34:861-6.

8. Ekinci EI, Clarke S, Thomas MC, et al. Dietary salt intake and mortality in patients with type 2 diabetes. Diabetes Care. 2011;34:703-9.

9. Cohen HW, Hailpern SM, Fang J, et al. Sodium intake and mortality in the NHANES II followup study. Am J Med. 2006 ;119:275.e7-14.

10. Dong J, Li Y, Yang Z, et al. Low dietary sodium intake increases the death risk in peritoneal dialysis. Clin J Am Soc Nephrol. 2010;5:240-7.

11. Alderman MH, Cohen H, Madhavan S. Dietary sodium intake and mortality: the National Health and Nutrition Examination Survey (NHANES I). Lancet. 1998;351:781-5.

*This version of the paper was accepted for publication in the <u>Journal of Health, Nutrition and Ageing</u> on 08/11/2014 12. Tuomilehto J, Jousilahti P, Rastenyte D, et al. Urinary sodium excretion and cardiovascular* 

mortality in Finland: a prospective study. Lancet. 2001;357:848-51.

13. Nagata C, Takatsuka N, Shimizu N, et al. Sodium intake and risk of death from stroke in Japanese men and women. Stroke. 2004;35:1543-7.

14. Norman PE, Jamrozik K, Lawrence-Brown MM, et al. Population based randomised controlled trial on impact of screening on mortality from abdominal aortic aneurysm. BMJ. 2004;329:1259.

15. Norman PE, Flicker L, Almeida OP, et al. Cohort Profile: The Health In Men Study (HIMS). Int J Epidemiol. 2009;38:48-52.

16. Golledge J, Clancy P, Jamrozik K, et al. Obesity, adipokines, and abdominal aortic aneurysm:Health in Men study. Circulation. 2007;116:2275-9.

17. Norman P, Spencer CA, Lawrence-Brown MM, et al. C-reactive protein levels and the expansion of screen-detected abdominal aortic aneurysms in men. Circulation. 2004;110:862–6.

18. Golledge J, Clancy P, Hankey GJ, et al. Relation between serum thrombospondin-2 and cardiovascular mortality in older men screened for abdominal aortic aneurysm. Am J Cardiol. 2013;111:1800-4.

19. Golledge J, Hankey GJ, Yeap BB, Almeids OP, Flicker L, Norman PE. Reported high salt intake is associated with increased prevalence of abdominal aortic aneurysm and larger aortic diameter in older men. Plos One. In press.

20. Aburto NJ, Ziolkovska A, Hooper L, et al. Effect of lower sodium intake on health: systematic review and meta-analyses. BMJ. 2013;346:f1326.

21. Taylor RS, Ashton KE, Moxham T, et al. Reduced dietary salt for the prevention of cardiovascular disease. Cochrane Database Syst Rev. 2011;7:CD009217.

22. Tikellis C, Pickering RJ, Tsorotes D, et al. Association of dietary sodium intake with atherogenesis in experimental diabetes and with cardiovascular disease in patients with Type 1 diabetes. Clin Sci (Lond). 2013;124:617-26.

# This version of the paper was accepted for publication in the <u>Journal of Health, Nutrition and Ageing</u> on 08/11/2014 23. Sealey JE, Alderman MH, Furberg CD, et al. Renin-angiotensin system blockers may create

more risk than reward for sodium-depleted cardiovascular patients with high plasma renin levels. Am J Hypertens. 2013;26:727-38.

24. Stolarz-Skrzypek K, Liu Y, Thijs L, et al. Blood pressure, cardiovascular outcomes and sodium intake, a critical review of the evidence. Acta Clin Belg. 2012;67:403-10.

25. Tomonari T, Fukuda M, Miura T, et al. Is salt intake an independent risk factor of stroke

mortality? Demographic analysis by regions in Japan. J Am Soc Hypertens. 2011;5:456-62.

26. Murata A, Fujino Y, Pham TM, et al. Prospective cohort study evaluating the relationship between salted food intake and gastrointestinal tract cancer mortality in Japan. Asia Pac J Clin Nutr. 2010;19:564-71.

27. Tsugane S, Akabane M, Inami T, et al. Urinary salt excretion and stomach cancer mortality among four Japanese populations. Cancer Causes Control. 1991;2:165-8.

28. Lu JB, Qin YM. Correlation between high salt intake and mortality rates for oesophageal and gastric cancers in Henan Province, China. Int J Epidemiol. 1987;16:171-6.

29. Joossens JV, Hill MJ, Elliott P, et al. Dietary salt, nitrate and stomach cancer mortality in 24 countries. European Cancer Prevention (ECP) and the INTERSALT Cooperative Research Group. Int J Epidemiol. 1996;25:494-504.

30. Gaddy JA, Radin JN, Loh JT, et al. High Dietary Salt Intake Exacerbates Helicobacter pylori-Induced Gastric Carcinogenesis. Infect Immun. 2013;81:2258-67.

31. D'Elia L, Rossi G, Ippolito R, et al. Habitual salt intake and risk of gastric cancer: a metaanalysis of prospective studies. Clin Nutr. 2012;31:489-98.

32. Rhodes DG, Murayi T, Clemens JC, et al. The USDA Automated Multiple-Pass Method accurately assesses population sodium intakes. Am J Clin Nutr. 2013;97:958-64.

**Figure 1:** Kaplan Meier curves showing the cumulative mortality from all causes in relation to reported frequencies of adding salt to food. Lines represent cumulative mortality for subjects grouped by reported frequencies of adding salt to food. The blue line represents men reporting the addition of salt to food never; the green line represents men reporting the addition of salt to food sometimes; and the brown line represents men reporting the addition of salt to food always. Vertical lines represent subjects censored at loss to follow-up.

**Figure 2:** Kaplan Meier curves showing the cumulative mortality from cancer in relation to reported frequencies of adding salt to food. Lines represent cumulative mortality for subjects grouped by reported frequencies of adding salt to food. The blue line represents men reporting the addition of salt to food never; the green line represents men reporting the addition of salt to food sometimes; and the brown line represents men reporting the addition of salt to food always. Vertical lines represent subjects censored at loss to follow-up.

**Figure 3:** Kaplan Meier curves showing the cumulative mortality from cardiovascular diseases in relation to reported frequencies of adding salt to food. Lines represent cumulative mortality for subjects grouped by reported frequencies of adding salt to food. The blue line represents men reporting the addition of salt to food never; the green line represents men reporting the addition of salt to food never; the green line represents men reporting the addition of salt to food always. Vertical lines represent subjects censored at loss to follow-up.

Numbe	All-cause			Cancer-related		
r						
	Hazar	95% CI	P value	Hazar	95% CI	Р
	d ratio			d ratio		value
4466	1.00	Referenc		1.00	Referenc	
		e			e	
3787	1.06	0.99-1.13	0.078	1.24	1.11-1.38	< 0.00
						1
3489	1.16	1.08-1.23	< 0.001	1.32	1.18-1.47	< 0.00
						1
11742	1.81	1.76-1.87	< 0.001	1.48	1.41-1.56	< 0.00
						1
4202	1.30	1.24-1.38	<0.001	1.05	0.95-1.15	0.343
1120	1.62	1.50-1.76	< 0.001	1.08	0.93-1.26	0.325
1711	1.70	1.59-1.81	< 0.001	1.07	0.94-1.22	0.338
	r 4466 3787 3489 11742 4202 1120	r       Hazar         Hazar       d ratio         4466       1.00         3787       1.06         3489       1.16         11742       1.81         4202       1.30         1120       1.62	rHazar95% CId ratiod ratio44661.00Referenc44661.00Referenc37871.060.99-1.1334891.161.08-1.23117421.811.76-1.8742021.301.24-1.3811201.621.50-1.76	r         Hazar         95% CI         P value           d ratio         -         -         -           d ratio         -         -         -           4466         1.00         Referenc         -           4466         1.00         Referenc         -           3787         1.06         0.99-1.13         0.078           3489         1.16         1.08-1.23         <0.001	rHazar95% CIP valueHazard ratio $-$ P valued ratiod ratio $ -$ d ratio44661.00Referenc $-$ 1.00e $ -$ 37871.060.99-1.130.078 $1.24$ 34891.161.08-1.23 $<0.001$ $1.48$ 117421.81 $1.76-1.87$ $<0.001$ $1.48$ 42021.30 $1.24-1.38$ $<0.001$ $1.05$ 1120 $1.62$ $1.50-1.76$ $<0.001$ $1.08$	rHazar95% CIP valueHazar95% CId ratio $ratio$ $ratio$ $ratio$ $ratio$ $ratio$ 44661.00Referenc1.00Reference1.00Referenc $e$ 37871.060.99-1.130.0781.2434891.161.08-1.23<0.001

Table 1: Univariate association of risk factors with all-cause mortality in 11,742 older men.
---

*This version of the paper was accepted for publication in the <u>Journal of Health, Nutrition and Ageing</u> on 08/11/2014* 

08/11/2014	1	1					
Past history of	903	1.86	1.71-2.02	< 0.001	1.17	0.99-1.39	0.064
stroke							
Past treatment	1333	1.45	1.34-1.56	< 0.001	1.11	0.96-1.28	0.146
for diabetes							
Treatment for	2264	1.01	0.95-1.08	0.733	0.83	0.74-0.94	0.002
high							
cholesterol							
Ever smoker	8337	1.43	1.35-1.53	< 0.001	1.56	1.41-1.74	< 0.00
							1
WHR per	11736	1.07	1.04-1.09	< 0.001	1.05	1.00-1.09	0.046
0.06*							
BMI per 4	11733	0.92	0.89-0.95	< 0.001	0.94	0.90-0.99	0.018
kg/m <sup>2</sup> *							
Eat meat							
(times per							
week)							
≥6	3387	1.00	0.81-1.23	0.998	1.53	1.01-2.32	0.046
3-5	5316	0.94	0.77-1.16	0.557	1.40	0.92-2.11	0.115
1-2	2339	0.92	0.74-1.13	0.413	1.39	0.92-2.12	0.123
<1	503	0.92	0.73-1.18	0.520	1.09	0.68	1.76
Never	197	1.00	Referenc		1.00	Referenc	
			e			e	
Eat Fish (times							
per week)							
≥6	113	0.75	0.55-1.04	0.085	0.72	0.41-1.27	0.255

08/11/2014							
3-5	1141	0.77	0.63-0.93	0.006	0.87	0.63-1.20	0.394
1-2	7337	0.78	0.65-0.92	0.004	0.84	0.62-1.14	0.257
<1	2908	0.76	0.63-0.91	0.002	0.88	0.65-1.20	0.423
Never	243	1.00	Referenc		1.00	Referenc	
			e			e	
Non-vigorous							
exercise							
(hours per							
week)							
None	4122	1.00	Referenc		1.00	Referenc	
			e			e	
≤2	1641	0.93	0.85-1.01	0.075	0.97	0.84-1.11	0.658
>2-4	2221	0.87	0.81-0.94	<0.001	0.92	0.81-1.04	0.190
>4-6	1209	0.79	0.72-0.87	<0.001	0.82	0.70-0.97	0.019
>6	2549	0.88	0.82-0.95	0.001	0.93	0.83-1.05	0.260
AAA	931	1.76	1.62-1.92	<0.001	1.47	1.27-1.71	< 0.00
							1

*This version of the paper was accepted for publication in the <u>Journal of Health, Nutrition and Ageing</u> on 08/11/2014* 

Men with the risk factor were compared to subjects without the risk factor or those with the reference reported level of intake or activity. \*Approximate standard deviation. WHR= Waist to hip ratio; BMI= Body mass index; AAA= Abdominal aortic aneurysm. WHR was missing on 6 men. BMI was missing on 9 men.

Table 2: Multivariate model examining the association of reported frequency of adding salt to

Characteristic	Number	Hazard ratio	95% CI	P value
Reported salt				
addition to food:				
Rare	4462	1.00	Reference	
Sometimes	3784	1.02	0.96-1.09	0.489
Always	3486	1.12	1.05 – 1.2	<0.001
Age per 5 years*	11732	1.76	1.71-1.82	<0.001
Past treatment for	4198	1.13	1.07-1.20	<0.001
hypertension				
Past treatment for	1120	1.18	1.09-1.29	<0.001
angina				
Past history of	1710	1.37	1.28-1.48	<0.001
myocardial				
infarction				
Past history of	902	1.45	1.32-1.58	<0.001
stroke				
WHR <0.9	1854	1.00	Reference	
WHR 0.9-1.02	8110	1.03	0.95-1.11	0.507
WHR >1.02	1768	1.20	1.08-1.33	0.001
Eat Fish (times per				
week)				
≥6	112	0.67	0.49-0.93	0.017
3-5	1141	0.81	0.67-0.98	0.031

### food and all-cause mortality in 11,732 older men.

*This version of the paper was accepted for publication in the <u>Journal of Health, Nutrition and Ageing</u> on 08/11/2014* 

1-2	7330	0.83	0.69-0.98	0.032
<1	2907	0.85	0.71-1.01	0.066
Never	242	1.00	Reference	
AAA	872	1.37	1.25-1.49	<0.001

Men with the risk factor were compared to subjects without the risk factor or those with the reference reported level of intake or activity. All variables shown were included in the multivariate model. Reported levels of non-vigorous exercise, prior treatment for diabetes or ever smoking were included in the model as stratified variables, therefore hazards ratios cannot be calculated.

\*Approximate standard deviation. WHR= Waist to hip ratio; AAA= Abdominal aortic aneurysm.

Table 3: Multivariate model examining the association of reported frequency of adding salt to

Characteristic	Number	Relative risk	95% CI	P value
Reported salt				
addition to food:				
Rare	4462	1.00	Reference	
Sometimes	3784	1.16	1.04-1.29	0.007
Always	3486	1.20	1.07-1.34	0.001
Age per 5 years*	11732	1.46	1.39-1.54	<0.001
Past treatment for	2264	0.84	0.74-0.94	0.004
dyslipidaemia				
Ever smoker	8328	1.48	1.33-1.65	<0.001
BMI<20	276	1.00	Reference	
BMI 20-30	9334	0.59	0.45-0.78	<0.001
BMI 30-39	2093	0.51	0.38-0.69	<0.001
BMI >40	29	0.91	0.39-2.14	0.833
Eat Meat (times				
per week)				
≥6	3382	0.70	0.46-1.07	0.099
3-5	5314	0.78	0.60-1.00	0.052
1-2	2336	0.96	0.85-1.10	0.568
<1	503	0.95	0.86-1.06	0.343
Never	197	1.00	Reference	

### food and cancer-related mortality in 11,732 older men.

Non-vigorous				
exercise (hours pe	er			
week)				
None	4116	1.00	Reference	
≤2	1640	0.99	0.86-1.14	0.856
>2-4	2221	0.93	0.82-1.06	0.283
>4-6	1209	0.84	0.72-0.99	0.039
>6	2546	0.94	0.83-1.06	0.314
AAA	872	1.26	1.08-1.47	0.004

Men with the risk factor were compared to subjects without the risk factor or those with the reference reported level of intake or activity. All variables shown were included in the multivariate model. \*Approximate standard deviation. WHR= Waist to hip ratio; AAA= Abdominal aortic aneurysm.











