

**Nut intake and 5-year changes in body weight and obesity risk in adults: results from the EPIC-PANACEA study**

Heinz Freisling, Hwayoung Noh, Nadia Slimani, Véronique Chajes, Anne M. May, Petra H. Peeters, Elisabete Weiderpass, Amanda J. Cross, Guri Skeie, Mazda Jenab, Francesca R. Mancini, Marie-Christine Boutron-Ruault, Guy Fagherazzi, Verena A. Katzke, Tilman Kühn, Annika Steffen, Heiner Boeing, Anne Tjønneland, Cecilie Kyrø, Camilla P. Hansen, Kim Overvad, Eric J. Duell, Daniel Redondo-Sánchez, Pilar Amiano, Carmen Navarro, Aurelio Barricarte, Aurora Perez-Cornago, Konstantinos K. Tsilidis, Dagfinn Aune, Heather Ward, Antonia Trichopoulou, Androniki Naska, Philippos Orfanos, Giovanna Masala, Claudia Agnoli, Franco Berrino, Rosario Tumino, Carlotta Sacerdote, Amalia Mattiello, H.B(as). Bueno-de-Mesquita, Ulrika Ericson, Emily Sonestedt, Anna Winkvist, Tonje Braaten, Isabelle Romieu, Joan Sabaté

H. Freisling (*corresponding author*), H. Noh

Nutritional Methodology and Biostatistics Group, Section of Nutrition and Metabolism, International Agency for Research on Cancer (IARC-WHO), 150, cours Albert Thomas, 69372 Lyon Cedex 08, France; Tel +33(0)47273 8664, Fax +33(0)47273 8361; e-mail: freislingh@iarc.fr

N. Slimani, V. Chajes, M. Jenab, I. Romieu

Nutritional Epidemiology Group, Section of Nutrition and Metabolism, International Agency for Research on Cancer (IARC-WHO), Lyon, France

A. M. May, P. H. Peeters

Julius Centre for Health Sciences and Primary Care, University Medical Centre Utrecht, Utrecht, The Netherlands

P. H. Peeters, A. J. Cross, K. K. Tsilidis, D. Aune, H. Ward, B. H. Bueno-de-Mesquita

Department of Epidemiology & Biostatistics, School of Public Health, Imperial College London, London, United Kingdom

E. Weiderpass, G. Skeie, T. Braaten

Department of Community Medicine, Faculty of Health Sciences, University of Tromsø, The Arctic University of Norway, Tromsø, Norway

E. Weiderpass

Department of Research, Cancer Registry of Norway, Oslo, Norway

Department of Medical Epidemiology and Biostatistics, Karolinska Institutet, Stockholm, Sweden

Genetic Epidemiology Group, Folkhälsan Research Center, Helsinki, Finland

F. R. Mancini, M.-C. Boutron-Ruault, G. Fagherazzi  
Inserm U1018, Gustave Roussy Institute, CESP, Villejuif, France  
University Paris-Saclay, University Paris-Sud, Villejuif, France

V. Katzke, T. Kühn  
German Cancer Research Center (DKFZ), Division of Cancer Epidemiology, Heidelberg, Germany

A. Steffen, H. Boeing  
Department of Epidemiology, German Institute of Human Nutrition Potsdam-Rehbrücke, Nuthetal, Germany

A. Tjønneland, C. Kyrø  
Danish Cancer Society Research Center, Copenhagen, Denmark

C. P. Hansen, K. Overvad  
Department of Public Health, Section for Epidemiology, Aarhus University, Aarhus, Denmark

E. J. Duell  
Unit of Nutrition and Cancer, IDIBELL, Catalan Institute of Oncology, Barcelona, Spain

D. Redondo-Sánchez  
Escuela Andaluza de Salud Pública, Instituto de Investigación Biosanitaria IBS GRANADA, Hospitales  
Universitarios de Granada/Universidad de Granada, Granada, Spain  
CIBER de Epidemiología y Salud Pública (CIBERESP), Spain

P. Amiano  
Public Health Division of Gipuzkoa, BioDonostia Research Institute, San Sebastian, Spain  
CIBER Epidemiology and Health Public, Madrid, Spain

C. Navarro  
Department of Epidemiology, Murcia Regional Health Council, IMIB-Arrixaca, Murcia, Spain  
CIBER Epidemiología y Salud Pública (CIBERESP), Spain  
Department of Health and Social Sciences, Universidad de Murcia, Murcia, Spain

A. Barricarte  
Navarra Public Health Institute, Pamplona, Spain  
Navarra Institute for Health Research (IdiSNA) Pamplona, Spain  
CIBER Epidemiology and Public Health CIBERESP, Spain

A. Perez-Cornago  
Cancer Epidemiology Unit, Nuffield Department of Population Health University of Oxford, United Kingdom

K. K. Tsilidis

Department of Hygiene and Epidemiology, School of Medicine, University of Ioannina, Ioannina, Greece

D. Aune

Björknes University College, Oslo, Norway

A. Trichopoulou, A. Naska, P. Orfanos

Hellenic Health Foundation, Athens, Greece

WHO Collaborating Center for Nutrition and Health, Unit of Nutritional Epidemiology and Nutrition in Public Health, Dept. of Hygiene, Epidemiology and Medical Statistics, School of Medicine, National and Kapodistrian University of Athens, Greece

G. Masala

Cancer Risk Factors and Life-Style Epidemiology Unit, Cancer Research and Prevention Institute – ISPO, Florence, Italy

F. Berrino, C. Agnoli

Epidemiology and Prevention Unit, Fondazione IRCCS Istituto Nazionale dei Tumori, Milan

R. Tumino

Cancer Registry and Histopathology Unit, "Civic- M.P.Arezzo" Hospital, ASP Ragusa, Italy

C. Sacerdote

Unit of Cancer Epidemiology, Città della Salute e della Scienza University-Hospital and Center for Cancer Prevention (CPO), Turin, Italy

A. Mattiello

Dipartimento di Medicina Clinica E Chirurgia Federico II University, Naples, Italy

B. H. Bueno-de-Mesquita

Dept. for Determinants of Chronic Diseases, National Institute for Public Health and the Environment (RIVM), Bilthoven, The Netherlands

Dept. of Social & Preventive Medicine, Faculty of Medicine, University of Malaya, Kuala Lumpur, Malaysia

U. Ericson, E. Sonestedt

Department of Clinical Sciences Malmö, Lund University, Malmö, Sweden

A. Winkvist

Department of Internal Medicine and Clinical Nutrition, The Sahlgrenska Academy, University of Gothenburg, Gothenburg, Sweden

J. Sabaté

Center for Nutrition, Healthy Lifestyle and Disease Prevention, Loma Linda University, School of Public Health, Loma Linda, USA

### **Electronic supplementary material**

The online version of this article contains supplementary material, which is available to authorized users.

### **Acknowledgments**

**Funding/Support:** This publication arises from the project PANACEA, which has received funding from the European Union, in the framework of the Public Health Programme (project number: 2005328).

The coordination of EPIC is financially supported by the European Commission (DG-SANCO) and the International Agency for Research on Cancer. The national cohorts are supported by Danish Cancer Society (Denmark); Ligue Contre le Cancer, Institut Gustave Roussy, Mutuelle Générale de l'Éducation Nationale, Institut National de la Santé et de la Recherche Médicale (INSERM) (France); German Cancer Aid, German Cancer Research Center (DKFZ), Federal Ministry of Education and Research (BMBF), Deutsche Krebshilfe, Deutsches Krebsforschungszentrum and Federal Ministry of Education and Research (Germany); the Hellenic Health Foundation (Greece); Associazione Italiana per la Ricerca sul Cancro-AIRC-Italy and National Research Council (Italy); Dutch Ministry of Public Health, Welfare and Sports (VWS), Netherlands Cancer Registry (NKR), LK Research Funds, Dutch Prevention Funds, Dutch ZON (Zorg Onderzoek Nederland), World Cancer Research Fund (WCRF), Statistics Netherlands (The Netherlands); Nordic Centre of Excellence programme on Food, Nutrition and Health (Norway); Health Research Fund (FIS), PI13/00061 to Granada; , PI13/01162 to EPIC-Murcia), Regional Governments of Andalucía, Asturias, Basque Country, Murcia and Navarra, ISCIII RETIC (RD06/0020) (Spain); Swedish Cancer Society, Swedish Research Council and County Councils of Skåne and Västerbotten (Sweden); Cancer Research UK (14136 to EPIC-Norfolk; C570/A16491 and C8221/A19170 to EPIC-Oxford), Medical Research Council (1000143 to EPIC-Norfolk, MR/M012190/1 to EPIC-Oxford) (United Kingdom). The current study was financially supported by Loma Linda University (LLU contract No.: 2150183). JS received funding from the INC International Nut and Dried Fruit Council.

The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

### **Additional Contributions:**

We thank all EPIC participants and staff for their contribution to the study.

For information on how to submit an application for gaining access to EPIC data and/or biospecimens, please follow the instructions at <http://epic.iarc.fr/access/index.php>.

1 **Abstract**

2 *Purpose* There is inconsistent evidence regarding the relationship between higher intake of  
3 nuts, being an energy-dense food, and weight gain. We investigated the relationship between nut  
4 intake and changes in weight over 5 years.

5 *Methods* This study includes 373,293 men and women, 25-70 years old, recruited between  
6 1992 and 2000 from 10 European countries in the European Prospective Investigation into Cancer and  
7 Nutrition (EPIC) study. Habitual intake of nuts including peanuts, together defined as nut intake, was  
8 estimated from country-specific validated dietary questionnaires. Body weight was measured at  
9 recruitment and self-reported 5 years later. The association between nut intake and body weight  
10 change was estimated using multilevel mixed linear regression models with center/country as random  
11 effect and nut intake and relevant confounders as fixed effects. The relative risk (RR) of becoming  
12 overweight or obese after 5 years was investigated using multivariate Poisson regressions stratified  
13 according to baseline body mass index (BMI).

14 *Results* On average, study participants gained 2.1 kg (SD 5.0 kg) over 5 years. Compared to  
15 non-consumers, subjects in the highest quartile of nut intake had less weight gain over 5 years (−0.07  
16 kg; 95% CI, −0.12- −0.02) ( $P$ -trend=0.025) and had 5% lower risk of becoming overweight (RR, 0.95;  
17 95% CI, 0.92-0.98) or obese (RR, 0.95; 95% CI, 0.90-0.99) (both  $P$ -trend <0.008).

18 *Conclusions* Higher intake of nuts is associated with reduced weight gain and a lower risk of  
19 becoming overweight or obese.

20 **Keywords** Nut intake, weight gain, obesity, energy balance, adults, Europe

## 21 **Introduction**

22 Observational studies and clinical trials, including the recent PREDIMED trial [1], have provided  
23 evidence that high nut consumption has beneficial effects on the occurrence of chronic diseases such  
24 as cardiovascular disease and type 2 diabetes [2–5], and a possible role in cancer prevention [5–8].

25 Nuts can provide 160–200 kcal per serving (30 g) and thus have energy-density similar to  
26 foods such as crackers, chocolate candies, and cookies. Therefore, concerns persist that high nut intake  
27 may lead to weight gain and increased long-term risk of obesity [9]. Whether frequent nut  
28 consumption promotes weight gain is not yet conclusive. Weight gain may not occur if nuts are  
29 incorporated into an isocaloric diet in which they are substitute for other foods such as red meat or  
30 processed meat or refined carbohydrates, as opposed to being added to an existing diet [10].

31 Randomized nut-feeding trials showed that compared with control diets, isocaloric diets  
32 enriched with nuts did not increase body weight, body mass index (BMI), or waist circumference [11,  
33 12]. However, these trials were limited by small numbers of volunteers, consuming a controlled diet  
34 over relatively short periods, with one notable exception, where median follow-up time was 4.8 years  
35 [12], and were not primarily designed to evaluate body weight changes. In addition, such trials are  
36 expensive to conduct across populations and may not test real-life settings.

37 There are little existing data from prospective observational studies [13–17], and these are  
38 limited because they were based on homogeneous populations and with one exception [14], did not  
39 account for overall dietary patterns. Dietary patterns may confound findings associated with nut intake  
40 because individuals who eat higher quantities of nuts usually also have a better overall diet quality  
41 [18], and other favourable lifestyle habits such as higher physical activity levels. Thus, it is important  
42 to account for dietary quality and other lifestyle behaviours in prospective observational settings.

43 We propose to address these knowledge gaps utilizing data of the EPIC-PANACEA study;  
44 PANACEA (Physical Activity, Nutrition, Alcohol, Cessation of smoking, Eating out of home in  
45 relation to Anthropometry) is the sub-cohort of the EPIC (European Prospective Investigation into  
46 Cancer and nutrition) study, where repeated assessments of weight are available making it possible to  
47 study weight changes.

48 The main objective of the present study was to investigate the relationship between nut intake and  
49 subsequent changes in weight after an average of 5 years of follow-up accounting for dietary patterns  
50 and other lifestyle factors that may co-vary with nut intake. A secondary objective was to estimate  
51 risks of becoming overweight or obese associated with higher nut intake.

52

## 53 **Methods**

### 54 *Study population.*

55 The EPIC study is an ongoing prospective cohort study across 23 centers in 10 European countries:  
56 Denmark, France, Germany, Greece, Italy, the Netherlands, Norway, Spain, Sweden, and the United  
57 Kingdom (UK). The cohort of 521,448 men and women recruited from 1992 to 2000 (age range: 25 to  
58 70 years) was enrolled from the general population with exceptions for France (national health  
59 insurance scheme members), Utrecht and Florence (breast cancer screening participants), Oxford  
60 (health conscious, mainly vegetarian, volunteers), and some centres from Italy and Spain (blood  
61 donors). The rationale for EPIC, study design, and methods have been described in detail elsewhere  
62 [19, 20]. The EPIC study was approved by the Ethical Review Board of the IARC and the Institutional  
63 Review Board of each participating EPIC centers.

64 For the present study, we excluded pregnant women, participants with missing dietary or  
65 lifestyle information, missing data on weight and height or with implausible anthropometric values at  
66 baseline (n=23,713). We further excluded 122,154 individuals with missing weight at follow-up and  
67 2,288 individuals with outlying anthropometry at follow-up: weight change  $< -5$  or  $> 5$  kg/year and  
68 BMI at follow-up  $< 16$  kg/m<sup>2</sup>. More details on follow-up exclusions are given in **Figure S1** (Online  
69 Resource) and have been previously detailed [21, 22]. The final analyses included 103,303 men and  
70 269,990 women with complete and plausible body weight data.

71

### 72 *Anthropometric measures and weight change.*

73 Two body weight measures were available for each participant: at baseline and after a median follow-  
74 up time of 5 years (min.: 2 years for Heidelberg (Germany); max.: 11 years for Varese (Italy)). At  
75 baseline, body weight and height were measured in most centres using comparable, standardized

76 procedures with the exception of those taken in France, Norway and the health conscious group of the  
77 Oxford centre in which subjects self-reported their weight. As for the follow-up weight assessments,  
78 all values were self-reported, except in Norfolk (UK) and Doetinchem (The Netherlands) where  
79 weight was measured [21, 22]. The accuracy of self-reported anthropometric measures – at baseline  
80 and at follow-up – was improved with the use of prediction equations derived from subjects with both  
81 measured and self-reported weight at baseline [23]. Our main outcome was weight change in kg per 5  
82 years, calculated as weight at follow-up – weight at baseline divided by the follow-up time in years  
83 and multiplied by 5 years.

84

#### 85 *Dietary assessment.*

86 Habitual food consumption during the previous 12 months was assessed at baseline for each individual  
87 with center-specific methods; in most cases food-frequency questionnaires (FFQs) [20]. These  
88 questionnaires were developed and validated in each country/center to capture country-specific dietary  
89 habits. In most centers FFQs were self-administered, with the exception of Greece, Ragusa (Italy),  
90 Naples (Italy) and Spain where face-to-face interviews were performed. Extensive quantitative FFQs  
91 were used in northern Italy, the Netherlands, Germany and Greece. Questionnaires structured by meals  
92 were used in Spain, France and Ragusa (Italy). Semi-quantitative FFQs were used in Denmark,  
93 Norway, Naples (Italy) and Umea (Sweden). In the UK, both a semi-quantitative FFQ and a 7-day  
94 record were used, whereas a method combining a FFQ with a 7-day record on lunch and dinner was  
95 used in Malmö (Sweden) [20]. Details of the questionnaire items regarding nut intake for each center  
96 or country, have been described previously [8]. In brief, the respective questionnaire food item(s) in  
97 France, Germany, Greece, Ragusa (Italy), the Netherlands, Spain, and the UK asked non-specifically  
98 for intake of any kind of nuts incl. peanuts; in Denmark and Norway specifically for peanuts, and in  
99 Umea (Sweden) specifically for “peanuts, salted”; in northern Italy specifically for “walnuts,  
100 hazelnuts, almonds, and peanuts”, and in Naples (Italy) for “walnuts”; in Spain for an exhaustive list  
101 of different types of nuts incl. peanuts and seeds; in Malmö (Sweden), the FFQ included peanuts as  
102 snacks, whereas other nuts had to be added to an open-ended question or recorded at lunch and dinner  
103 meals; finally, in Germany, the Netherlands, and the UK separate items on peanut butter intake were



104 asked for and we included this item in our overall nut intake variable. Here we define the combined  
105 intake of any of the items described above as “nut intake”; because nut intake was assessed in these  
106 broad categories, a stratified analysis by specific types of nuts was not possible. Non-consumers were  
107 determined from the FFQs and defined as those with an intake of nuts equal to zero.

108         In order to account for healthy diet, which may confound nut intake, we used the modified  
109 relative Mediterranean Diet Score (mrMDS) [24]. This score included the nutritional components that  
110 characterize the Mediterranean diet: i.e. higher intake of vegetables, legumes, fruit and nuts, cereals,  
111 fish and seafood, plant oils, and moderate alcohol consumption; and lower intakes of meat/products,  
112 and dairy products. Each mrMDS component (apart from alcohol) was measured in grams per 1000  
113 kcal (to express intake as energy density) and higher scores (range: 0-18) characterizing a  
114 Mediterranean diet [24]. In order to avoid over-adjustment, we used the mrMDS after subtracting the  
115 “fruit and nuts” component.

116

#### 117 *Assessment of other covariates.*

118 Data on objectively validated physical activity [25], smoking status, and education were collected at  
119 baseline through questionnaires [20]. Information on smoking status was also collected at follow-up at  
120 the same time as anthropometric data collection. Thus, we could account for smoking status  
121 modification during follow-up (stable current smoker, stable former smoker, stable never smoker, quit  
122 smoking, started smoking).

123

#### 124 *Statistical analyses.*

125 Habitual nut intake as estimated from the dietary questionnaires was analysed both on a continuous  
126 scale per 15 g/day increment, which corresponds to the mean intake of nut consumers in the highest  
127 cohort category of intake, and by categories with all non-consumers (~25%) placed in the first  
128 (reference) category and the consumers divided by quartiles into the remaining four categories of  
129 intake (categories 2 to 5), similar as in Jenab et al. [8]. As a secondary analysis, we also modelled  
130 frequency of nut intake using the following categories: “never/almost never”, “0.5-2 times/month”,  
131 “0.5-≤1 times/week”, “more than 1 times/week”, which is similar to Bes-Rastrollo et al.[14].

132 Frequency data for the centers Cambridge (UK) (n=14,535) and Malmö (Sweden) (n=21,566) were  
133 not available because open-ended dietary methodologies were used.

134 The association between nut intake and body weight change (kg/5 years) was estimated using  
135 multilevel mixed linear regression models with center as random effect and nut intake and relevant  
136 confounders as fixed effects. Models with three different sets of adjustment were fit (see footnotes of  
137 Table 2 for complete list). Participants with missing values for physical activity (1.5%), education  
138 (2.1%), and smoking status at follow-up (0.4% after replacing missing values at follow-up [10.5%] by  
139 smoking status at baseline) were classified as a separate category and included in the models. Model  
140 assumptions and fit were checked visually by plotting the residuals against each of the categorical  
141 covariates. The linearity of the associations for each continuous covariate was evaluated by three-knot  
142 restricted cubic spline models at Harrell's default percentiles (i.e. 10<sup>th</sup>, 50<sup>th</sup>, and 90<sup>th</sup>) in combination  
143 with a Wald-type test [26]. Because baseline BMI and follow-up time in years (both *P* non-linear <  
144 0.001) showed a non-linear relationship with weight change, splines with 3 knots for these two  
145 variables were included as covariates.

146 In order to evaluate heterogeneity across countries/centers, we performed country/center-  
147 specific analyses using generalized linear models and pooled results by random-effect meta-analysis  
148 and calculated *I* squared and respective *P* values for heterogeneity [27].

149 We performed a range of sensitivity analyses such as excluding participants with chronic  
150 diseases at baseline or missing values in covariates, excluding countries where nut intake included  
151 peanuts only or adjusting for main food groups instead of the mrMDS (**Table S1**, Online Resource).

152 We tested *a priori* for effect modification by age (categorised as younger than median age <51  
153 and ≥ 51 years), sex, BMI categories at baseline (<25, 25-≤30, >30kg/m<sup>2</sup>), and change of smoking  
154 status (never, current, start smoking, quitter, former) by including interaction terms between each  
155 variable and nut intake (continuous per 15 g/d) in the models. *P* values for the interaction term were  
156 calculated by using *F* tests.

157 We used a modified Poisson regression approach [28] to estimate the relative risks (RR) and  
158 95% confidence intervals (CI) of becoming overweight or obese according to nut intake (in categories  
159 of absolute intakes and frequency of intake). Analyses were stratified by initial BMI categories (<25:

160 normal weight,  $25 \leq \text{BMI} < 30$ : overweight and  $\geq 30 \text{ kg/m}^2$ : obese). RRs were adjusted as in our model 3  
161 described above. The BMI after 5 years was calculated from the 5 year follow-up weight and baseline  
162 height.

163 Differences were considered statistically significant at  $P < 0.05$ . All statistical analyses were  
164 performed with STATA 12.1 (College Station TX).

165

## 166 **Results**

167 The main characteristics of the study population at baseline by categories of nut intake are shown in  
168 **Table 1**. Higher intake of nuts was associated with younger age, a lower BMI, a higher educational  
169 level, never smoking, and being more physically active. Participants in the highest category of nut  
170 intake also had higher intakes of vegetables, fruit, cereals/cereal products, non-alcoholic and alcoholic  
171 beverages, but also of sugar/confectionary, and cakes/biscuits; they also had a slightly higher mrMED  
172 score. In contrast, they had lower intakes of meat/products, dairy, fish, and potatoes. On average, study  
173 participants gained 2.1 kg of weight between baseline and the 2<sup>nd</sup> weight assessment with considerable  
174 variation between subjects (SD 5.0 kg).

175 Body weight changes (kg) over 5 years according to baseline nut intake are shown in **Table 2**.  
176 After adjustment for potential confounders, each 15g/day increase in nut intake was associated with  
177 less weight gain ( $-0.04 \text{ kg/5-years}$ , 95% CI,  $-0.071$ -  $-0.012$ ). The observed effects were small and  
178 corresponded to  $\sim 2.5\%$ -reduction in body weight increase. Associations remained virtually unchanged  
179 after further adjustment for Mediterranean diet using the mrMDS (Model 3, Table 2). Estimated  
180 results were consistent across countries/centers with low heterogeneity ( $I$ -squared=21%,  $P$   
181 heterogeneity = 0.22) (**Figure S2**, Online Resource). Analyses by categories of nut intake confirmed  
182 the findings using intake on a continuous scale, where participants in the highest category of nut intake  
183 gained 0.07 kg/5-years less weight as compared to non-consumers ( $P$  trend = 0.025) (Table 2).  
184 Furthermore, when we analyzed frequency of nut intake without accounting for amounts of intake,  
185 strengths of associations increased, where subjects consuming nuts more than once per week gained  
186 0.1 kg/5-years less weight as compared to non-consumers ( $P$  trend  $< 0.001$ ) (Table 2).

187 Our main findings were also robust to a range of sensitivity analyses (**Table S1**, Online  
188 Resource). For example, excluding participants who started or quit smoking during follow-up (Model  
189 S4), with missing values in any of the covariates (Model S8), , or in non-smokers only (to exclude  
190 residual confounding in smokers) (Model S16) resulted in virtually similar effect estimates. Similarly,  
191 excluding participants from Denmark, Norway, and Umea (Sweden), where the country/center-  
192 specific FFQ only included peanuts, did not alter the estimates (Model S9). In contrast, when we  
193 excluded France (Model S11), where the FFQ item on nuts was asked only in relation to “aperitif”  
194 before lunch or dinner, which in France is typically consumed with an alcoholic beverage, effect  
195 estimates per 15g/day nut intake doubled from  $-0.042$  (95% CI,  $-0.071$ - $-0.012$ ) to  $-0.083$  kg/5-years  
196 (95% CI,  $-0.114$ -  $-0.051$ ). Another important finding in our sensitivity analysis was that adjustment  
197 for main food groups as indicated in Table 1, instead of the mrMDS, resulted in similar effect  
198 estimates (Model S12), but only when intake of meat/products was excluded. Inclusion of intake of  
199 meat/products completely attenuated associations between intake of nuts and peanuts (15g/day) and 5-  
200 y weight change ( $0.004$  kg/5-y; 95% CI,  $-0.027$ -  $0.034$ ) (Model S13).

201 No effect modification was found with regard to baseline age ( $P$  interaction = 0.54), sex ( $P$   
202 interaction = 0.62), baseline weight status ( $P$  interaction = 0.18) or change in smoking status ( $P$   
203 interaction = 0.95).

204 Adjusted relative risks (95% CI) of becoming overweight or obese after 5 years according to  
205 categories of nut intake and initial BMI are presented in **Table 3**. At baseline, 197,291 subjects were  
206 normal weight, 127,445 were overweight and 48,557 were obese. After 5 years, 31,215 (15.8%)  
207 normal weight subjects became overweight or obese and 14,913 (13.2%) overweight subjects became  
208 obese. Compared to non-consumers of nuts, normal weight subjects at baseline in the highest category  
209 of nut intake had a 5% (95% CI, 2%-8%) lower risk of becoming overweight or obese. Similarly,  
210 overweight subjects at baseline had a 5% (95% CI, 1%-10%) lower risk of becoming obese.  
211 Frequency of nut intake was also associated with 5% (95% CI, 1%-10%) lower risk of becoming  
212 overweight or obese in subjects that were normal weight at baseline. However, no association was  
213 observed for risk of becoming obese in subjects that were already overweight at baseline ( $P$  trend =  
214 0.39).

215 **Discussion**

216 Gradual age-related body weight increase during adulthood is a well observed phenomenon in many  
217 non-obese populations — in our study, about 0.4 kg per year. Using baseline and follow-up data from  
218 a large European multi-center cohort study, EPIC-PANACEA, we found that long-term weight gain  
219 was significantly less in individuals consuming higher levels of nuts. These inverse associations were  
220 modest for absolute intake of nuts, but were more pronounced for the frequency of consumption –  
221 possibly reflecting different dietary habits or difficulties in reporting portion size accurately – where  
222 >1 serving of nuts per week was associated with a 10% lower body weight increase. Importantly, our  
223 findings are not likely to be confounded by a better overall diet quality, which is often observed in  
224 high consumers of nuts, because we adjusted for dietary patterns and other lifestyle factors notably  
225 physical activity and smoking.

226 In a post hoc analysis, we found that habitual high intake of meat and processed meat appears  
227 to attenuate associations. We believe that the observed effects of nut intake on body weight change are  
228 at least partly mediated via a reduced intake of meat/products shown to be positively associated with  
229 weight gain [22, 29]. This has been hypothesized earlier as being one of the potential pathways of  
230 weight stabilizing effects of nuts [10] and confirmed in our sensitivity analysis (Table S1, Online  
231 Resource).

232 Our findings are in line with the few other prospective observational studies [13–17]. Women  
233 in the Nurses' Health Study II (NHS II), who reported eating nuts  $\geq 2$  times/wk, experienced 0.5 kg  
234 less weight gain (95% CI, -0.8- -0.2) after a mean 8 years of follow-up compared with those who  
235 rarely ate nuts [14]. Similar results were observed in the Seguimiento Universidad de Navarra (SUN)  
236 study, a prospective cohort in Spain, where weight change in men and women was assessed after a  
237 median of 28 months [13] and after 6 years [16]. In the Nurses' Health Study (NHS), no differences in  
238 weight gain over 16 years of follow-up across categories of nut consumption were observed [15]. A  
239 pooled analysis of the NHS, the NHS II, and the Health Professionals Follow-up Study, where the  
240 relationship of dietary changes over 4-year periods was related to changes in body weight, found that  
241 per serving increase in nut intake, study participants gained 0.57 lb (~0.3 kg) less weight per 4-year  
242 period [17]. The observed differences in effect sizes across these studies can most likely be explained

243 by a combination of factors including differences in length of weight follow-up, confounder  
244 adjustment, accuracy of dietary assessment instruments used, but also differences in terms of  
245 frequency and amount of consumed nuts, underlying dietary habits and other lifestyle factors that are  
246 specific to a population. Interestingly, the only randomized controlled nut-feeding trial (PREDIMED)  
247 that had a comparably long follow-up as in our study reported very similar results with regard to  
248 adjusted difference in 5 year changes in bodyweight in the nut group as compared with the control  
249 group ( $-0.08$  kg) though not statistically significant (95% CI,  $-0.50-0.35$  kg) and only in the context  
250 of a Mediterranean diet [12]. We specifically accounted for Mediterranean dietary patterns in our  
251 analysis in order to evaluate associations of nut intake with weight change in the context of other diets.  
252 Romaguera et al. showed previously in the same study population that high adherence to a  
253 Mediterranean diet was associated with a 5-year weight change of  $-0.16$  kg (95% CI,  $-0.24- -0.07$  kg)  
254 and were 10% (95% CI, 4%-18%) less likely to develop overweight or obesity compared to  
255 individuals with a low adherence [30].

256         Several mechanistic hypotheses have been proposed that could explain the association  
257 between nut consumption and lessened weight gain, despite a potentially higher total energy intake in  
258 nut consumers [10, 31]. These include increased satiety/supressed hunger due to the high dietary fibre  
259 and plant protein content of nuts; the high content of unsaturated fat, which together with the high  
260 protein content can lead to an increase in resting energy expenditure and diet-induced thermogenesis,  
261 both of which can reduce body weight and weight gain; and incomplete mastication of nuts may cause  
262 a low level of fat absorption that could result in the loss of available energy [10, 31]. In addition,  
263 individuals who consume nuts regularly tend to consume less red and processed meat [10]. As already  
264 mentioned above, such a replacement is likely to be beneficial for the prevention of weight gain  
265 because red and processed meat intake have been associated with weight gain, risk of obesity and  
266 higher BMI [17, 22, 29].

267         Our study has limitations. First, only self-reported weight at follow-up was available in most  
268 centers. To mitigate this potential source of bias, we used a prediction equation to improve self-  
269 reported weight estimates [23]. Furthermore, in the EPIC-Norfolk study, a sub-cohort of EPIC, a high  
270 correlation between self-reported and measured weight data has been shown ( $r=0.97$  in men and

271  $r=0.98$  in women), which means that ranking of participants according to self-reported weight was  
272 adequate [32]. Second, we were not able to accurately measure changes in body composition (e.g.,  
273 using dual-energy x-ray absorptiometry, DXA); therefore we had to assume that observed weight  
274 changes are largely due to changes in body fat mass and not in lean body mass. Third, we were not  
275 able to account for potential changes in diet during follow-up; yet, magnitudes of changes in weight  
276 appear to be more pronounced and more robust if changes in diet can be accounted for [33].  
277 Nevertheless, mean dietary changes at the population level are often small; for example, in the NHS,  
278 the mean 4-year change in nut intake corresponded to a 5% increase of the baseline intake [17].  
279 Fourth, we were not able to stratify our analysis by specific types of nuts because nut intake was  
280 assessed in broad categories of nut intake across the EPIC centers/countries. Finally, measurement  
281 error is a limitation inherent to all epidemiological studies using self-reported dietary data. We  
282 attempted to minimize this bias by adjusting for total energy intake and for plausibility of dietary  
283 energy reporting; the latter has been recently shown in the EPIC-Potsdam sub-study to improve  
284 expected associations between intakes of energy-dense foods and BMI [34].

285         Strengths of our study include its prospective design with a reasonably long follow-up, the  
286 very large sample size, which provided sufficient power to also detect smaller associations, despite the  
287 large variability of weight change, and to perform a number of sensitivity analyses. In order to  
288 improve dietary intake assessment of nuts, like for many other food groups, it is important to continue  
289 the search for and validation of biomarkers of nut intake in the future and metabolomics approaches  
290 may offer new opportunities in this regard [35]. Future research may also assess the mediating role of  
291 plasma fatty acid changes in the association between nuts and weight change.

292         We conclude that in this prospective study of middle-aged adults from 10 European countries  
293 representing populations with heterogeneous diets, higher nut intake is associated with slightly less  
294 weight gain after 5 years of follow-up. Higher nut consumers also demonstrated a lower risk of  
295 becoming overweight or obese. Our findings are thus in line with short-term randomized nut-feeding  
296 trials and support dietary recommendations to increase nut consumption to reduce chronic disease risk  
297 and mortality.

**Ethical standards** The study has been performed in accordance with the ethical standards laid down in the 1964 Declaration of Helsinki and its later amendments and obtained ethical approval from participating centres and IARC ethics committees. Informed consent was given by all study participants.

**Conflicts of interest** The authors declare that they have no conflict of interest.



## References

1. Estruch R, Ros E, Salas-Salvado J, et al (2013) Primary prevention of cardiovascular disease with a Mediterranean diet. *N Engl J Med* 368:1279–1290. doi: 10.1056/NEJMoa1200303
2. Bao Y, Han J, Hu FB, et al (2013) Association of nut consumption with total and cause-specific mortality. *N Engl J Med* 369:2001–2011. doi: 10.1056/NEJMoa1307352
3. Nash SD, Nash DT (2008) Nuts as part of a healthy cardiovascular diet. *Curr Atheroscler Rep* 10:529–535.
4. Sabate J, Ang Y (2009) Nuts and health outcomes: new epidemiologic evidence. *Am J Clin Nutr* 89:1643S–1648S. doi: 10.3945/ajcn.2009.26736Q
5. Aune D, Keum N, Giovannucci E, et al (2016) Nut consumption and risk of cardiovascular disease, cancer, all-cause and cause-specific mortality: a systematic review and dose-response meta-analysis of prospective studies. *BMC Med* 1–14. doi: 10.1186/s12916-016-0730-3
6. Wang W, Yang M, Kenfield SA, et al (2016) Nut consumption and prostate cancer risk and mortality. *Br J Cancer* 1–4. doi: 10.1038/bjc.2016.181
7. Nakanishi M, Chen Y, Qendro V, et al (2016) Effects of walnut consumption on colon carcinogenesis and microbial community structure. *Cancer Prev Res (Phila)*. doi: 10.1158/1940-6207.CAPR-16-0026
8. Jenab M, Ferrari P, Slimani N, et al (2004) Association of nut and seed intake with colorectal cancer risk in the European Prospective Investigation into Cancer and Nutrition. *Cancer Epidemiol Biomarkers Prev* 13:1595–1603.
9. Appel LJ, Van Horn L (2013) Did the PREDIMED trial test a Mediterranean diet? *N Engl J Med* 368:1353–1354. doi: 10.1056/NEJMe1301582
10. Jackson CL, Hu FB (2014) Long-term associations of nut consumption with body weight and obesity. *Am J Clin Nutr* 100:408–411. doi: 10.3945/ajcn.113.071332
11. Flores-Mateo G, Rojas-Rueda D, Basora J, et al (2013) Nut intake and adiposity: Meta-analysis of clinical trials. *Am J Clin Nutr* 97:1346–1355. doi: 10.3945/ajcn.111.031484
12. Estruch R, Martínez-González MA, Corella D, et al (2016) Effect of a high-fat Mediterranean diet on bodyweight and waist circumference: A prespecified secondary outcomes analysis of the PREDIMED randomised controlled trial. *Lancet Diabetes Endocrinol* 4:666–676. doi: 10.1016/S2213-8587(16)30085-7
13. Bes-Rastrollo M, Sabate J, Gomez-Gracia E, et al (2007) Nut consumption and weight gain in a Mediterranean cohort: The SUN study. *Obesity (Silver Spring)* 15:107–116. doi: 10.1038/oby.2007.507
14. Bes-Rastrollo M, Wedick NM, Martinez-Gonzalez M a, et al (2009) Prospective study of nut consumption, long-term weight change, and obesity risk in women. *Am J Clin Nutr* 89:1913–1919.
15. Jiang R, Manson JE, Stampfer MJ, et al (2002) Nut and peanut butter consumption and risk of

- type 2 diabetes in women. *JAMA* 288:2554–2560.
16. Martínez-González M a., Bes-Rastrollo M (2011) Nut consumption, weight gain and obesity: Epidemiological evidence. *Nutr Metab Cardiovasc Dis* 21:40–45. doi: 10.1016/j.numecd.2010.11.005
  17. Mozaffarian D, Hao T, Rimm E, et al (2011) Changes in Diet and Lifestyle and Long- Term Weight Gain in Women and Men. *N Engl J Med* 364:2392–404.
  18. O’Neil CE, Keast DR, Fulgoni VL, Nicklas T a. (2010) Tree nut consumption improves nutrient intake and diet quality in US adults: An analysis of national health and nutrition examination survey (NHANES) 1999-2004. *Asia Pac J Clin Nutr* 19:142–150.
  19. Riboli E, Kaaks R (1997) The EPIC Project: rationale and study design. *European Prospective Investigation into Cancer and Nutrition. Int J Epidemiol* 26 Suppl 1:S6-14.
  20. Riboli E, Hunt K, Slimani N, et al (2002) European Prospective Investigation into Cancer and Nutrition (EPIC): study populations and data collection. *Public Health Nutr* 5:1113-24. doi: 10.1079/PHN2002394
  21. Vergnaud AC, Norat T, Romaguera D, et al (2012) Fruit and vegetable consumption and prospective weight change in participants of the European prospective investigation into cancer and nutrition-physical activity, nutrition, alcohol, cessation of smoking, eating out of home, and obesity study. *Am J Clin Nutr* 95:184–193. doi: 10.3945/ajcn.111.019968
  22. Vergnaud AC, Norat T, Romaguera D, et al (2010) Meat consumption and prospective weight change in participants of the EPIC-PANACEA study. *Am J Clin Nutr* 92:398–407. doi: 10.3945/ajcn.2009.28713
  23. Spencer E a, Appleby PN, Davey GK, Key TJ (2002) Validity of self-reported height and weight in 4808 EPIC-Oxford participants. *Public Health Nutr* 5:561–565. doi: 10.1079/PHN2001322
  24. Buckland G, González CA, Agudo A, et al (2009) Adherence to the mediterranean diet and risk of coronary heart disease in the spanish EPIC cohort study. *Am J Epidemiol* 170:1518–1529. doi: 10.1093/aje/kwp282
  25. Peters T, Brage S, Westgate K, et al (2012) Validity of a short questionnaire to assess physical activity in 10 European countries. *Eur J Epidemiol* 27:15–25. doi: 10.1007/s10654-011-9625-y
  26. Orsini N, Greenland S (2011) A procedure to tabulate and plot results after flexible modeling of a quantitative covariate. *Stata J* 11:1–29.
  27. Higgins JPT, Thompson SG (2002) Quantifying heterogeneity in a meta-analysis. *Stat Med* 1539–1558. doi: 10.1002/sim.1186
  28. Zou G (2004) A Modified Poisson Regression Approach to Prospective Studies with Binary Data. *Am J Epidemiol* 159:702–706. doi: 10.1093/aje/kwh090
  29. Rouhani MH, Salehi-Abargouei A, Surkan PJ, Azadbakht L (2014) Is there a relationship between red or processed meat intake and obesity? A systematic review and meta-analysis of

- observational studies. *Obes Rev* 15:740–8. doi: 10.1111/obr.12172
30. Romaguera D, Norat T, Vergnaud A-C, et al (2010) Mediterranean dietary patterns and prospective weight change in participants of the EPIC-PANACEA project. *Am J Clin Nutr* 92:912–921. doi: 10.3945/ajcn.2010.29482
  31. Vadivel V, Kunyanga CN, Biesalski HK (2012) Health benefits of nut consumption with special reference to body weight control. *Nutrition* 28:1089–1097. doi: 10.1016/j.nut.2012.01.004
  32. Park JY, Mitrou PN, Keogh RH, et al (2012) Self-reported and measured anthropometric data and risk of colorectal cancer in the EPIC-Norfolk study. *Int J Obes (Lond)* 36:107–18. doi: 10.1038/ijo.2011.61
  33. Smith JD, Hou T, Hu FB, et al (2015) A Comparison of Different Methods for Evaluating Diet, Physical Activity, and Long-Term Weight Gain in 3 Prospective Cohort Studies. *J Nutr* 145:2527–2534. doi: 10.3945/jn.115.214171
  34. Gottschald M, Knüppel S, Boeing H, Buijsse B (2016) The influence of adjustment for energy misreporting on relations of cake and cookie intake with cardiometabolic disease risk factors. *Eur J Clin Nutr* 70:1318–1324. doi: 10.1038/ejcn.2016.131
  35. Mora-Cubillos X, Tulipani S, Garcia-Aloy M, et al (2015) Plasma metabolomic biomarkers of mixed nuts exposure inversely correlate with severity of metabolic syndrome. *Mol Nutr Food Res* 59:2480–90. doi: 10.1002/mnfr.201500549

**Table 1** Main characteristics of the study population according to categories<sup>a</sup> of nut intake (n = 373,293)

	Non-consumers (n = 97,852)	>0-0.8 g/d (n = 85,470)	>0.8-2.8 g/d (n = 55,335)	>2.8-6.0 g/d (n = 65,815)	>6.0 g/d (n = 68,821)
Nut intake, g/d, median [IQR]	0.0	0.5 [0.2-0.7]	1.7 [1.5-2.3]	4.1 [3.3-4.9]	12.4 [8.1-18.8]
Follow-up time, y	4.6 ± 1.7	7.0 ± 2.7	5.0 ± 2.2	5.0 ± 2.2	4.7 ± 2.0
Weight change, kg/5y <sup>b</sup>	1.7 ± 5.3	2.1 ± 4.4	2.2 ± 5.0	2.2 ± 4.9	2.3 ± 5.1
Women %	73.7	66.0	72.9	77.7	72.7
Age, y	53.8 ± 8.3	51.5 ± 9.8	52.3 ± 9.6	50.7 ± 9.1	49.9 ± 9.7
BMI at inclusion, kg/m <sup>2</sup>	25.8 ± 4.4	25.7 ± 4.2	25.0 ± 4.1	24.9 ± 4.1	24.8 ± 4.0
BMI categories, %					
<25 kg/m <sup>2</sup>	47.8	48.1	55.6	58.1	58.7
25-<30 kg/m <sup>2</sup>	36.3	37.7	33.0	31.1	30.5
30-≤35 kg/m <sup>2</sup>	12.5	11.2	9.1	8.6	8.7
>35 kg/m <sup>2</sup>	3.4	3.0	2.3	2.2	2.1
University degree or higher, %	17.4	22.1	28.4	28.5	31.3
Missing	1.5	0.6	1.5	1.6	1.5
Physically inactive, %	25.1	20.7	19.5	16.9	17.2
Missing	1.4	0.4	1.5	1.7	2.6
Smoking status at follow-up, %					
Never	49.9	40.0	46.4	45.2	43.9
Former	27.8	27.3	28.6	28.1	29.6
Current	19.1	15.4	14.0	14.7	16.2
Missing	3.3	17.3	11.0	12.0	10.3
Previous illness, % <sup>c</sup>	9.3	6.8	8.3	7.0	7.1
Missing	12.7	5.8	10.1	7.1	4.9
Dietary intake					
Total energy intake, kcal/d	1,980 ± 594	2,015 ± 598	2,061 ± 573	2,071 ± 576	2,297 ± 626
Vegetables, g/d	208 ± 136	185 ± 139	231 ± 147	236 ± 152	255 ± 167
Fruits, g/d	233 ± 184	218 ± 169	236 ± 171	235 ± 171	252 ± 185
Legumes, g/d	19 ± 31	8 ± 14	14 ± 20	15 ± 21	20 ± 25
Meat/products, g/d	106 ± 59	99 ± 56	99 ± 59	96 ± 58	100 ± 65
Dairy, g/d	332 ± 232	329 ± 249	337 ± 231	308 ± 214	325 ± 226
Fish, g/d	50 ± 42	29 ± 25	32 ± 27	40 ± 38	36 ± 36
Egg/egg products, g/d	21 ± 19	15 ± 15	18 ± 16	19 ± 17	20 ± 18
Potatoes, g/d	94 ± 70	102 ± 87	88 ± 65	84 ± 58	85 ± 58
Cereals/cereal products, g/d	198 ± 99	224 ± 112	210 ± 103	212 ± 95	225 ± 103
Sugar/confectionary, g/d	38 ± 48	44 ± 55	44 ± 46	40 ± 41	42 ± 39
Cakes/biscuits, g/d	37 ± 42	41 ± 43	41 ± 42	42 ± 40	45 ± 43
Added fat, g/d	27 ± 18	30 ± 18	27 ± 18	26 ± 17	28 ± 19
Nonalcoholic beverages, g/d	983 ± 792	1,086 ± 804	1,225 ± 731	1,100 ± 719	1,136 ± 735
Alcoholic beverages, g/d	145 ± 265	182 ± 293	172 ± 262	165 ± 253	192 ± 270
mrMED score units/d	8.7 ± 3.0	8.4 ± 3.1	9.0 ± 3.0	9.2 ± 2.9	9.4 ± 3.0

Data are expressed as arithmetic mean ± SD if not stated otherwise.

<sup>a</sup> First category corresponds to non-consumers of nut intake based on food-frequency questionnaires; categories 2-5 are quartiles of consumers; note that proportion of subjects in categories 2-5 is unequal because observations with the same value were categorised in the same band ('xtile' command in Stata).

<sup>b</sup> Calculated as weight at follow-up minus weight at baseline divided by the follow-up time in years and multiplied by 5 years.

<sup>c</sup> Type 2 diabetes, cardiovascular disease, cancer.

Abbreviations: BMI, body mass index (calculated as weight in kilograms divided by height in meters squared); IQR, interquartile range; mrMED, modified relative Mediterranean diet score (range: 0-18; higher scores characterizing a Mediterranean diet).

**Table 2** Difference in body weight gain (kg) over 5 years according to baseline nut intake in 373,293 men and women

	N (%)	Median nut intake [IQR] g/d	Model 1 <i>beta</i> (95% CI)	Model 2 <i>beta</i> (95% CI)	Model 3 <i>beta</i> (95% CI)
<i>Beta</i> per 15g/d	373,293 (100)	0.9 [0.0-4.3]	-0.046 (-0.075, -0.018)	-0.046 (-0.075, -0.017)	-0.042 (-0.071, -0.012)
----- <i>Categories of absolute nut intake</i> -----					
Non-consumer	97,852 (26)	0.0	Reference	Reference	Reference
>0-0.8 g/d	85,470 (23)	0.5 [0.2-07]	-0.039 (-0.095, 0.018)	-0.038 (-0.094, 0.019)	-0.035 (-0.092, 0.021)
>0.8-2.8 g/d	55,335 (15)	1.7 [1.5-2.3]	-0.04 (-0.096, 0.015)	-0.022 (-0.077, 0.034)	-0.014 (-0.070, 0.041)
>2.8-6.0 g/d	65,815 (18)	4.1 [3.3-4.9]	-0.059 (-0.112, -0.007)	-0.047 (-0.099, 0.006)	-0.037 (-0.089, 0.016)
>6.0 g/d	68,821 (18)	12.4 [8.1-18.8]	-0.089 (-0.142, -0.036)	-0.082 (-0.135, -0.028)	-0.069 (-0.123, -0.015)
<i>P</i> trend (linear)			0.001	0.006	0.025
----- <i>Frequency of nut intake</i> <sup>a</sup> -----					
Never/almost never	87,520 (26)	-	Reference	Reference	Reference
0.5-2 times/mo	93,221 (28)	-	-0.03 (-0.083, 0.023)	-0.022 (-0.075, 0.03)	-0.018 (-0.071, 0.034)
0.5-≤1 times/wk	72,760 (21)	-	-0.077 (-0.128, -0.026)	-0.065 (-0.117, -0.014)	-0.058 (-0.110, -0.006)
>1 times/wk	83,691 (25)	-	-0.124 (-0.177, -0.071)	-0.115 (-0.169, -0.061)	-0.102 (-0.156, -0.047)
<i>P</i> trend (linear)			<0.001	<0.001	<0.001

Multilevel linear mixed models with random effect on the intercept and slope according to center.

Overall mean 5-year weight gain corresponded to 2.1 kg (SD 5.0) and negative beta-values indicate less weight gain (kg) over the same period.

Model 1 adjusted for age, sex, and body mass index (BMI) at baseline (3-knot restricted cubic spline); Model 2 was further adjusted for follow-up time in years (3-knot restricted cubic spline), total energy intake (kcal/day), educational level, levels of physical activity, smoking status at follow-up, and plausibility of dietary energy reporting; Model 3 was further adjusted for the modified relative Mediterranean diet score (without fruit and nut component). IQR, interquartile range.

<sup>a</sup> Frequency data for the centers Cambridge (UK) (n=14,535) and Malmö (Sweden) (n=21,566) were not available.

**Table 3** Adjusted relative risks (RR) (95% CI) of becoming overweight or obese over 5 years according to baseline nut intake and baseline body mass index (BMI) in men and women

	BMI <25 kg/m <sup>2</sup> at baseline n=197,291			BMI ≥25 to <30 kg/m <sup>2</sup> at baseline n=127,445		
	N (%)	N overweight or obese (%)	RR of becoming overweight or obese (95% CI)	N (%)	N obese (%)	RR of becoming obese (95% CI)
<i>Categories of absolute nut intake</i>						
Non-consumer	46,784 (24)	7,082 (23)	Reference	31,495 (28)	3,637 (25)	Reference
>0-0.8 g/d	41,148 (21)	8,374 (27)	0.97 (0.94, 1.00)	28,283 (25)	4,353 (29)	0.96 (0.92, 1.00)
>0.8-2.8 g/d	30,786 (16)	4,360 (14)	0.94 (0.91, 0.97)	16,244 (14)	2,110 (14)	0.98 (0.93, 1.03)
>2.8-6.0 g/d	38,206 (19)	5,629 (18)	0.95 (0.93, 0.98)	18,337 (16)	2,432 (16)	0.93 (0.89, 0.98)
>6.0 g/d	40,367 (20)	5,770 (18)	0.95 (0.92, 0.98)	18,771 (17)	2,381 (16)	0.95 (0.90, 0.99)
<i>P</i> trend (linear)			0.002			0.018
<i>Frequency of nut intake<sup>a</sup></i>						
Never/almost never	40,688 (23)	6,678 (24)	Reference	27,825 (28)	3,776 (28)	Reference
0.5-2 times/mo	50,523 (28)	8,100 (29)	0.98 (0.95, 1.01)	28,250 (28)	3,802 (28)	0.98 (0.94, 1.03)
0.5-≤1 times/wk	39,836 (22)	6,644 (23)	0.96 (0.94, 0.99)	21,443 (21)	3,121 (23)	0.94 (0.90, 0.98)
>1 times/wk	48,416 (27)	6,822 (24)	0.95 (0.92, 0.98)	22,859 (23)	2,924 (21)	0.99 (0.95, 1.04)
<i>P</i> trend (linear)			0.001			0.385

A modified Poisson regression approach (Zou 2004) was used to calculate the RR and 95% CI.

Adjusted for age, sex, country/center, BMI at baseline (3-knot restricted cubic spline), follow-up time in years (3-knot restricted cubic spline), total energy intake (kcal/day), educational level, levels of physical activity, smoking status at follow-up, and plausibility of dietary energy reporting, and for the modified relative Mediterranean diet score (without fruit and nut component).

<sup>a</sup> Frequency data for the centers Cambridge (UK) (n=14,535) and Malmö (Sweden) (n=21,566) were not available.