



Editor's choice  
Scan to access more  
free content

## ORIGINAL ARTICLE

# Habitual chocolate consumption and risk of cardiovascular disease among healthy men and women

Chun Shing Kwok,<sup>1,2</sup> S Matthijs Boekholdt,<sup>3</sup> Marleen A H Lentjes,<sup>4</sup> Yoon K Loke,<sup>5</sup> Robert N Luben,<sup>4</sup> Jessica K Yeong,<sup>6</sup> Nicholas J Wareham,<sup>7</sup> Phyo K Myint,<sup>1</sup> Kay-Tee Khaw<sup>4</sup>

► Additional material is published online only. To view please visit the journal online (<http://dx.doi.org/10.1136/heartjnl-2014-307050>).

For numbered affiliations see end of article.

## Correspondence to

Dr Chun Shing Kwok, School of Medicine & Dentistry, University of Aberdeen, c/o Professor Phyo Kyaw Myint, Room 4:013, Polwarth Building, Foresterhill, Aberdeen AB25 2ZD, UK; [phyo.myint@abdn.ac.uk](mailto:phyo.myint@abdn.ac.uk)

CSK and SMB are joint first authors.

PKM and K-TK are joint last authors.

Received 22 October 2014  
Revised 25 March 2015  
Accepted 20 April 2015  
Published Online First  
16 June 2015



CrossMark

**To cite:** Kwok CS, Boekholdt SM, Lentjes MAH, et al. *Heart* 2015;**101**:1279–1287.

## ABSTRACT

**Objective** To examine the association between chocolate intake and the risk of future cardiovascular events.

**Methods** We conducted a prospective study using data from the European Prospective Investigation into Cancer (EPIC)-Norfolk cohort. Habitual chocolate intake was quantified using the baseline food frequency questionnaire (1993–1997) and cardiovascular end points were ascertained up to March 2008. A systematic review was performed to evaluate chocolate consumption and cardiovascular outcomes.

**Results** A total of 20 951 men and women were included in EPIC-Norfolk analysis (mean follow-up 11.3 ±2.8 years, median 11.9 years). The percentage of participants with coronary heart disease (CHD) in the highest and lowest quintile of chocolate consumption was 9.7% and 13.8%, and the respective rates for stroke were 3.1% and 5.4%. The multivariate-adjusted HR for CHD was 0.88 (95% CI 0.77 to 1.01) for those in the top quintile of chocolate consumption (16–99 g/day) versus non-consumers of chocolate intake. The corresponding HR for stroke and cardiovascular disease (cardiovascular disease defined by the sum of CHD and stroke) were 0.77 (95% CI 0.62 to 0.97) and 0.86 (95% CI 0.76 to 0.97). The propensity score matched estimates showed a similar trend. A total of nine studies with 157 809 participants were included in the meta-analysis. Higher compared to lower chocolate consumption was associated with significantly lower CHD risk (five studies; pooled RR 0.71, 95% CI 0.56 to 0.92), stroke (five studies; pooled RR 0.79, 95% CI 0.70 to 0.87), composite cardiovascular adverse outcome (two studies; pooled RR 0.75, 95% CI 0.54 to 1.05), and cardiovascular mortality (three studies; pooled RR 0.55, 95% CI 0.36 to 0.83).

**Conclusions** Cumulative evidence suggests that higher chocolate intake is associated with a lower risk of future cardiovascular events, although residual confounding cannot be excluded. There does not appear to be any evidence to say that chocolate should be avoided in those who are concerned about cardiovascular risk.

## INTRODUCTION

Chocolate is an important dietary source of flavonoid antioxidants, which are hypothesised to have a beneficial effect on endothelial function and protect against cardiovascular disease (CVD).<sup>1</sup> Evidence

from a range of small scale intervention trials have reported that intake of chocolate resulted in increased high density lipoprotein (HDL) cholesterol concentrations, decreased low density lipoprotein (LDL) oxidation, and improved endothelial function.<sup>2</sup> Large scale intervention studies have not been performed and therefore the potential benefits of raising chocolate consumption on cardiovascular risk are unknown. However, some evidence on the potential beneficial effects of chocolate can be derived from observational studies.

Many studies have evaluated the risk of cardiovascular outcomes with chocolate consumption.<sup>1–4</sup> One meta-analysis which attempted to quantify systematically the effect of high chocolate consumption was limited by inclusion of a heterogeneous spectrum of outcomes including diabetes and heart failure.<sup>3</sup> A more recent meta-analysis specifically evaluated the effects of flavonols (also found in other foods) and the risk of coronary heart disease (CHD) and found no significant association between flavonol intake and CHD.<sup>4</sup> Many cohort studies which examined the association between chocolate and CVD (defined by CHD and stroke in this study) are limited to sex-specific cohorts.<sup>5–9</sup> Two meta-analyses have been performed which pooled many of these studies and found reduced risk of incident stroke<sup>7</sup> and CVD<sup>10</sup> with higher chocolate consumption. However, these studies are limited because one study only considered stroke while the other study pooled many different cardiovascular events into a single risk estimate.

In order to evaluate any association between habitual chocolate consumption and the risk of cardiovascular events, we analysed data from the European Prospective Investigation into Cancer-Norfolk (EPIC-Norfolk), a large prospective population study in the UK, and incorporated the results from this observational study into the evidence available to date from the literature by conducting a systematic review and meta-analysis.

## METHODS

### EPIC-Norfolk cohort study

EPIC-Norfolk is a prospective population study of over 25 000 men and women, resident in Norfolk, UK (99.6% white Caucasian). The study methods have been described previously in detail.<sup>11</sup> In brief, at the baseline survey between 1993 and 1997,

participants completed a health and lifestyle questionnaire and provided information on physician-diagnosed chronic diseases such as cancer, myocardial infarction, stroke, and diabetes mellitus. We identified cigarette smoking habit (never, former, current)<sup>12</sup> and a validated, self-reported physical activity measure (inactive, moderately inactive, moderately active, active)<sup>13</sup> and assessed reported intake of alcoholic drinks (where units (8 g alcohol)/week were categorised into: zero consumption, >0–7, >7–14, >14–21; >21–28; >28 units/week). At the baseline health examination (N=25 639), a trained nurse measured height, weight, and blood pressure (Acutorr). Total cholesterol, HDL cholesterol and triglycerides were measured in non-fasting blood samples as described previously.<sup>11</sup> Blood samples were stored at –80°C. LDL cholesterol was calculated by the Friedewald formula. Data on lipids were available for 23 074 participants (90%) and for inflammation markers on minimally 18 643 participants (73%).

### Main exposure measure: chocolate consumption

Dietary measurements were obtained by use of the food frequency questionnaire (FFQ), which assessed overall diet in the past year from 24 782 participants (response rate 97%). Every food item or group of food items in the questionnaire had nine answer categories: never or less than once per month, 1–3 per month, once a week, 2–4 per week, 5–6 per week, once a day, 2–3 per day, 4–5 per day, 6+ per day. This instrument and the way it is analysed, using the Compositional Analyses from Frequency Estimates (CAFÉ), are described in detail elsewhere.<sup>14</sup>

Three questions from the FFQ were considered indicative of chocolate consumption, namely ‘Chocolates singles or squares’ (average portion size of 8 g), ‘Chocolate snack bars, for example, Mars, Crunchie’ (average portion size of 50 g), and ‘Cocoa, hot chocolate (cup)’ (average portion size of 12 g powder weight; the liquid to make up the beverage was hence not included). Frequency categories were multiplied by the portion size to derive the amount of chocolate product eaten (g/day). The sum of the weights of these food items consumed, rather than their flavonoid or cocoa content, formed the measure of exposure.

### Clinical outcomes

Participants admitted to hospital were identified using their unique NHS (National Health Service) number by data linkage with ENCORE (East Norfolk Health Authority database). All participants were flagged for death certification at the UK Office of National Statistics, ascertaining vital status for the entire cohort. CHD was defined as ICD-10 (10th Revision of the Internal Classification of Diseases and Related Health Problems) codes I20–25 (which includes the spectrum of CHD including myocardial infarction, unstable angina, and stable angina); stroke was defined as ICD-10 codes I60–69 (which includes ischaemic stroke, haemorrhagic stroke, and stroke of undetermined cause). CVD was defined as the combination of CHD and stroke. A previous validation study in our cohort indicated high specificity for such case ascertainment.<sup>15</sup> We report data with follow-up to 31 March 2008, an average of 11.3 years (2.8 SD, median 11.9 years). The study was approved by the Norwich District Health Authority Ethics Committee. All participants provided signed informed consent.

### Statistical analysis

Participants without a completed FFQ (n=857), with prevalent myocardial infarction or stroke (n=1102), and whose reported chocolate intake was more than 5 SDs above the median

(ie, 100 g/day) were excluded from the analyses (n=86), leaving 23 638 participants eligible; we further excluded 2687 participants who missed data on one or more covariables (n=20 951). Baseline characteristics were calculated by quintiles of chocolate consumption. Since 4195 participants (approximately 20%) reported zero chocolate intake, the lowest quintile corresponds to the non-users only and was used as the reference category in Cox proportional hazards model. HR and corresponding 95% CIs for the risk of future CHD, stroke or CVD were calculated using four models. Regression model 1 adjusted for sex and age. Regression model 2 additionally adjusted for lifestyle factors: smoking, physical activity, energy intake (MJ/day), and alcohol intake (categorical). Regression model 3 adjusted for the variables in model 2 and possible mediators: body mass index (BMI), systolic blood pressure, LDL cholesterol, HDL cholesterol, and prevalent diabetes. In model 4 we additionally adjusted for C-reactive protein (CRP). Analyses were performed using SPSS V.17.0 (Chicago, Illinois, USA).

### Systematic review and meta-analysis

To be eligible for inclusion in our review, studies had to report on the association between chocolate consumption and cardiovascular outcomes. We searched PubMed and EMBASE from inception until June 2013 using the terms described in online data supplement 1, with no language limitations, and we checked bibliographies of included articles. In addition to our search of EMBASE (which already includes unpublished abstracts and conference reports) we also searched conference abstracts from the European Society of Cardiology congresses from 2005 to 2013 inclusive. A further search of ISI Web of Science was conducted in February 2014 to identify additional articles. Two reviewers independently screened abstracts and titles, and then obtained full-text versions of potentially relevant studies to confirm eligibility. Data extraction of included studies was performed by CSK as well as JKY, and checked by YKL. Study validity was evaluated based on ascertainment of chocolate consumption and cardiovascular outcomes as well as steps taken to reduce confounding in the primary studies. We pooled data using the inverse variance method and random effects model in RevMan V.5.2 software (Nordic Cochrane Center, Copenhagen, Denmark). For these comparisons, we used the multivariable adjusted measures of association (HRs, relative risks or ORs) for the highest category of chocolate consumption versus the lowest category of consumption. Heterogeneity was estimated using  $I^2$ , and we considered a value >50% to demonstrate substantial heterogeneity.<sup>16</sup> We planned to evaluate publication bias through asymmetry testing if there were >10 studies in the dataset, and no evidence of significant heterogeneity.<sup>17</sup>

### Sensitivity analyses

We performed additional analysis to determine if a similar direction of effect was observed using propensity score matching on the pre-specified covariates of models 1, 2 and 3. Propensity score matching were performed in STATA using the ‘teffects psmatch’ function which estimates treatment effects from observational data.<sup>18</sup> The results of the propensity score analysis was used as sensitivity analysis for the meta-analysis. We conducted additional analysis excluding the Djousse study because it was a cross-sectional study.

## RESULTS

### EPIC-Norfolk cohort

A complete dataset on relevant baseline characteristics were available for 20 951 study participants; 9214 men and 11 737

women ( $n=16\,162$  when restricted to participants with CRP data). Mean follow-up was  $11.9\pm 2.8$  years, and total person years was 236 942 years. A total of 3013 (14.4%) people experienced a fatal or non-fatal CHD event, stroke or both, referred to as CVD. Among these participants with CVD events, 2434 (11.6%) had a CHD event and 848 (4.0%) had a stroke event. When analyses were limited to people with available CRP at baseline, a total of 2207 (13.7%) study participants experienced a CVD event while 1754 (10.9%) and 648 (4.0%) had a CHD event and/or stroke event, respectively.

### Chocolate consumption in the EPIC-Norfolk cohort

The median daily chocolate consumption was 4.6 g/day (IQR 0.6–12.0); among consumers only, median chocolate intake was 7.0 g/day (IQR 3.5–15.5). Higher chocolate consumption was associated with trends towards a beneficial cardiovascular risk factor profile including lower age, lower BMI, lower waist/hip ratio, lower systolic blood pressure, lower concentrations of apolipoprotein B (apoB) and CRP, a lower prevalence of diabetes mellitus, and more physical inactivity (table 1). In contrast, higher chocolate consumption was more prevalent among men and among current smokers. Higher chocolate intake was associated with a higher energy intake, with lower contributions from protein and alcohol sources and higher contribution from fat and carbohydrates.

### Chocolate consumption and risk of CVD in EPIC-Norfolk

Higher chocolate intake was associated with a statistically significant lower risk of CVD, with stronger associations for CVD mortality than for total CVD (hospitalisation or mortality) (table 2). HRs attenuated after adjustment, but remained borderline significant for total CVD (HR 0.89, 95% CI 0.79 to 1.00) and CVD mortality (HR 0.75, 95% CI 0.62 to 0.92). Adjustment for CRP minimally changed the effect estimates. A significant dose–response association was present for both total incidence and CVD mortality. We assessed the proportionality assumption graphically and the results suggest that proportional hazards were maintained with time.

Higher chocolate consumption was associated with a lower risk of hospitalisation or mortality due to CHD in crude and minimally adjusted models (table 3). This association was attenuated after adjustment for a range of cardiovascular risk factors and after additional adjustment for a set of dietary parameters (HR 0.91, 95% CI 0.80 to 1.04). In the smaller sample of participants with CRP data, we observed a significant association with 18% lower risk in quintile 5 versus quintile 1 (95% CI 0.70 to 0.97). Participants with a high chocolate intake also had a lower risk of stroke. The sex- and age-adjusted HR was 0.77 (95% CI 0.62 to 0.96) for top versus bottom quintile of chocolate consumption (table 3). This association remained statistically significant upon adjustment for smoking, physical activity, and dietary variables. Additional adjustment for mediators did not materially change the estimate (HR 0.78, 95% CI 0.63 to 0.98). Analyses using mortality as outcome, rather than combined mortality and hospitalisations, showed lower risk estimates for CVD, CHD, and stroke, although the number of events were substantially lower and therefore the CIs became wider.

### Propensity score matched analysis for chocolate consumption and risk of CVD

The baseline characteristics of the unmatched and propensity matched cohorts are shown in online supplementary appendix table 1. A Love plot was used to examine the standardised

difference in covariates before and after propensity score adjustment (see online supplementary appendix figure 1). The risk of CHD and stroke by quintiles of chocolate intake considering various levels of adjustments and propensity score matching are shown in online supplementary appendix table 2. After propensity score matching, a trend for benefit with chocolate consumption was apparent for both CHD and stroke, but with lower sample sizes and wider CIs they were not statistically significant.

### Systematic review and meta-analysis

Search results and studies of chocolate and CVD

We screened 392 titles and abstracts and identified eight studies<sup>5–9 19–21</sup> that met eligibility criteria (see online supplementary appendix figure 2). Including the EPIC-Norfolk study, a total of nine studies with 157 809 participants were included in the meta-analysis. These studies included seven cohort studies, one post-hoc analysis of a randomised trial, and one cross-sectional study, and the follow-up duration of the cohort studies ranged from 8–16 years (table 4). Two studies were conducted in the USA and one study was conducted in Australia, but the remainder were conducted in Europe (UK, Netherlands, Germany, Sweden). Three studies included both men and women, but the remainder were sex specific (two studies of men and two studies of women). The mean age of the participants in the included studies ranged from 49–79 years.

Bias assessment for studies of chocolate and CVD

Different methods for evaluating and ascertaining chocolate consumption and cardiovascular outcomes were used across the studies (see online supplementary appendix table 3). One study used patient interviews to ascertain chocolate consumption but the remaining studies used questionnaires. The majority of studies used ICD codes to ascertain cardiovascular diagnoses. Five of the studies used linkage of data to mortality registry/records in order to ascertain mortality. All the included studies were able to use a variety of adjustments to account for the effect of confounders; for the meta-analysis, we used data from model 2 of EPIC-Norfolk. While a key cardiovascular risk factor such as BMI was adjusted for in seven studies, other important risk factors such as cholesterol were only considered in two studies (see online supplementary appendix table 4). We considered the cross-sectional study to have lower validity due to the inability to draw a temporal relationship between chocolate use and adverse cardiovascular outcomes.

Chocolate consumption and risk of cardiovascular events

Overall, chocolate consumption was significantly associated with lower risk of CHD across five studies (pooled RR 0.71, 95% CI 0.56 to 0.92;  $I^2=61\%$ ) (figure 1). The risk of CHD mortality from one study showed no significant difference with and without chocolate consumption (RR 0.98, 95% CI 0.88 to 1.10). For risk of stroke with chocolate consumption there was significantly lower risk of both stroke incidence (pooled RR 0.79, 95% CI 0.70 to 0.87;  $I^2=0\%$ , five studies) (figure 1) and mortality (RR 0.85, 95% CI 0.74 to 0.98; one study). There was a significant lower risk of any cardiovascular event (pooled RR 0.75, 95% CI 0.54 to 1.05;  $I^2=70\%$ , two studies) and cardiovascular mortality (pooled RR 0.55, 95% CI 0.36 to 0.83;  $I^2=67\%$ , three studies) (figure 2). The propensity matched results were incorporated in the meta-analysis and this continued to show a significant association between chocolate consumption and reduction in CHD (pooled RR 0.72, 95% CI 0.55 to 0.93;  $I^2=63$ , five studies) and stroke (pooled RR 0.78, 95% CI 0.70 to 0.88;  $I^2=0\%$ , five studies) (figure 3).

**Table 1** Baseline cardiovascular risk factors by quintiles of chocolate intake in 20 951 men and women of EPIC-Norfolk

Quintiles of daily chocolate intake	Quintile 1	Quintile 2	Quintile 3	Quintile 4	Quintile 5	p Value	Total
Range (g/day)	0	0.6–3.5	4.1–7.0	7.2–15.5	15.6–98.8		
Median (IQR) (g/day)							
Chocolate intake	0	1.1 (0.6, 3.4)	4.6 (4.1, 6.3)	9.2 (8.1, 12.0)	24.9 (22.1, 39.5)		
Chocolate singles	0	0.6 (0.0, 1.1)	0.6 (0.6, 1.1)	1.1 (0.6, 3.4)	1.1 (0.6, 3.4)		
Chocolate bars	0	0.0 (0.0, 0.0)	3.5 (3.5, 3.5)	7.0 (0.0, 7.0)	21.5 (21.5, 21.5)		
Chocolate beverage	0	0.0 (0.0, 0.0)	0.0 (0.0, 0.8)	0.8 (0.0, 5.2)	0.0 (0.0, 5.2)		
N	4195	4437	4035	4106	4178		20 951
Sex, men	44.1 (1852)	43.4 (1927)	40.8 (1648)	43.5 (1786)	47.9 (2001)	<0.001	44.0 (9214)
Age, years	61±9	60±9	58±9	59±9	57±9	<0.001	59±9
Body mass index, kg/m <sup>2</sup>	26.3±4.1	26.1±3.8	26.3±3.7	26.2±3.8	26.0±3.7	<0.001	26.2±3.8
Waist/hip ratio*	0.86±0.09	0.85±0.09	0.84±0.09	0.85±0.09	0.85±0.09	<0.001	0.85±0.09
Smoking							
Current	10.5 (440)	10.7 (473)	11.4 (459)	11.4 (469)	13.0 (543)	<0.001	11.4 (2384)
Previous	44.0 (1847)	43.1 (1914)	40.1 (1618)	40.9 (1678)	38.5 (1610)		41.4 (8667)
Never	45.5 (1908)	46.2 (2050)	48.5 (1958)	47.7 (1959)	48.5 (2025)		47.3 (9900)
Diabetes mellitus	4.6 (193)	1.6 (73)	0.9 (38)	0.9 (38)	0.7 (29)	<0.001	1.8 (371)
Systolic blood pressure, mm Hg	137±19	136±18	134±18	134±18	135±17	<0.001	135±18
Diastolic blood pressure, mm Hg	83±11	83±11	82±11	82±11	82±11	0.010	82±11
Total cholesterol, mmol/L	6.2±1.1	6.2±1.1	6.1±1.1	6.1±1.1	6.1±1.1	0.001	6.1±1.1
LDL cholesterol, mmol/L	4.0±1.0	4.0±1.0	3.9±1.0	4.0±1.0	3.9±1.0	0.07	4.0±1.0
HDL cholesterol, mmol/L	1.43±0.43	1.43±0.42	1.44±0.42	1.43±0.41	1.40±0.41	<0.001	1.43±0.42
Triglycerides, mmol/L	1.5 (1.1–2.2)	1.5 (1.1–2.1)	1.5 (1.0–2.1)	1.5 (1.1–2.1)	1.5 (1.1–2.1)	0.20	1.5 (1.1–2.1)
Apolipoprotein AI, mg/dL*	156±35	157±35	157±35	156±34	155±36	0.29	156±35
Apolipoprotein B, mg/dL*	97±26	97±25	96±25	96±26	94±25	0.001	96±26
CRP, mg/L*	1.6 (0.8–3.2)	1.5 (0.7–3.2)	1.5 (0.7–3.2)	1.4 (0.7–3.1)	1.4 (0.7–2.9)	<0.001	1.5 (0.7–3.1)
Physical activity							
Inactive	32.7 (1373)	28.6 (1267)	26.9 (1084)	27.6 (1134)	27.2 (1138)	<0.001	28.6 (5996)
Moderately inactive	27.5 (1153)	30.2 (1340)	29.4 (1188)	30.2 (1241)	28.1 (1174)		29.1 (6096)
Moderately active	21.8 (914)	23.7 (1051)	24.1 (971)	22.8 (936)	24.3 (1015)		23.3 (4887)
Active	18.0 (755)	17.6 (779)	19.6 (792)	19.4 (795)	20.4 (851)		19.0 (3972)
Energy intake by FFQ, kJ/day	7754±2264	8043±2233	8359±2311	8934±2411	9854±2664	<0.001	8584±2494
Protein intake by FFQ, %en	17.4±3.4	17.3±3.2	17.1±3.0	16.7±2.9	15.5±2.7	<0.001	16.9±3.2
Fat intake by FFQ, %en	32.8±6.7	33.8±6.2	34.3±6.1	34.7±5.7	35.8±5.4	<0.001	34.3±6.1
Carbohydrate intake by FFQ, %en	47.8±7.0	47.3±6.3	47.3±6.1	47.8±5.7	48.0±5.5	<0.001	47.7±6.2
Alcohol intake by FFQ, %en	3.3±5.3	3.4±4.7	3.1±4.2	2.7±3.6	2.5±3.5	<0.001	3.0±4.3
Alcohol intake by HLQ, units/week	3 (1–10)	4 (2–11)	4 (2–10)	4 (1–9)	4 (1–10)	<0.001	4 (1–10)

Data are presented as mean±SD, percentage (number) or median (IQR). Differences between groups were analysed by analysis of variance (ANOVA) for continuous variables, and by  $\chi^2$  test for categorical variables. Continuous variables with a skewed distribution (triglycerides, CRP, alcohol intake) were analysed by the Kruskal-Wallis test.

\*These variables have missing data:  $N_{\text{waist/hip ratio}}=20\ 928$ ;  $N_{\text{ApoAI}}=16\ 035$ ;  $N_{\text{ApoB}}=16\ 022$ ;  $N_{\text{CRP}}=16\ 162$ .

CRP, C-reactive protein; EPIC, European Prospective Investigation into Cancer; FFQ, food frequency questionnaire; HDL, high density lipoprotein; HLQ, Health and Lifestyle Questionnaire; LDL, low density lipoprotein.

**Table 2** Risk of total (fatal and non-fatal) and fatal CVD incidence by quintiles of chocolate intake in EPIC-Norfolk (1993–2008)

Range (g/day)	Total	Quintile 1 0	Quintile 2 0.6–3.5	Quintile 3 4.1–7.0	Quintile 4 7.2–15.5	Quintile 5 15.6–98.8	p Linearity
Total CVD/n (%)	3013/20 951 (14.4)	729/4195 (17.4)	693/4437 (15.6)	528/4035 (13.1)	563/4106 (13.7)	500/4178 (12.0)	
Model 1		1.00	0.94 (0.85 to 1.05)	0.94 (0.84 to 1.05)	0.87 (0.78 to 0.98)	0.84 (0.75 to 0.94)	0.001
Model 2		1.00	0.98 (0.88 to 1.08)	0.97 (0.87 to 1.09)	0.90 (0.80 to 1.00)	0.86 (0.76 to 0.97)	0.006
Model 3		1.00	0.99 (0.89 to 1.10)	0.99 (0.88 to 1.10)	0.91 (0.81 to 1.02)	0.89 (0.79 to 1.00)	0.018
	2207/16 162 (13.7)	548/3251 (16.9)	512/3406 (15.0)	375/3102 (12.1)	422/3220 (13.1)	350/3183 (11.0)	
Model 4a		1.00	1.00 (0.89 to 1.13)	0.96 (0.84 to 1.09)	0.90 (0.79 to 1.03)	0.82 (0.72 to 0.95)	0.003
Model 4b		1.00	1.00 (0.88 to 1.12)	0.95 (0.83 to 1.09)	0.90 (0.79 to 1.02)	0.82 (0.71 to 0.95)	0.003
Fatal CVD/n (%)	1107/20 951 (5.3)	308/4195 (7.3)	238/4437 (5.4)	193/4035 (4.8)	206/4106 (5.0)	162/4178 (3.9)	
Model 1		1.00	0.77 (0.65 to 0.92)	0.90 (0.75 to 1.07)	0.78 (0.66 to 0.94)	0.71 (0.59 to 0.86)	0.001
Model 2		1.00	0.80 (0.68 to 0.95)	0.94 (0.79 to 1.13)	0.81 (0.67 to 0.97)	0.73 (0.60 to 0.89)	0.005
Model 3		1.00	0.82 (0.69 to 0.98)	0.96 (0.80 to 1.15)	0.82 (0.68 to 0.98)	0.75 (0.62 to 0.92)	0.011
	805/16 162 (5.0)	231/3251 (7.1)	174/3406 (5.1)	133/3102 (4.3)	154/3220 (4.8)	113/3183 (3.6)	
Model 4a		1.00	0.82 (0.67 to 1.00)	0.91 (0.73 to 1.13)	0.81 (0.66 to 1.00)	0.71 (0.56 to 0.89)	0.008
Model 4b		1.00	0.81 (0.66 to 0.99)	0.90 (0.73 to 1.12)	0.80 (0.65 to 0.99)	0.70 (0.55 to 0.89)	0.008

Model 1 adjusted for sex and age.

Model 2 adjusted for sex, age, smoking, physical activity, energy intake, and alcohol consumption.

Model 3: as model 2 and diabetes, body mass index, systolic blood pressure, LDL cholesterol, and HDL cholesterol.

Model 4a: as model 3, but restricted to number of participants for whom CRP is available (N=16 162).

Model 4b: as model 3 and CRP (N=16 162).

CRP, C-reactive protein; CVD, cardiovascular disease; EPIC, European Prospective Investigation into Cancer; HDL, high density lipoprotein; LDL, low density lipoprotein.

Publication bias was not assessed because there were fewer than 10 studies included in the analysis.

We conducted additional analysis excluding the Djousse study because it was a cross-sectional study. The lower risk associated with chocolate and CHD was slightly less after exclusion of this study but still remained statistically significant (RR 0.84, 95% CI 0.75 to 0.95 vs RR 0.79, 95% CI 0.70 to 0.87) (data not shown).

## DISCUSSION

In this large prospective population study, higher intake of chocolate up to 100 g/day was associated with a lower risk of CVD and stroke, with stronger associations for mortality than total incidence. We have built on the findings of the previous

meta-analysis<sup>7 10</sup> by including this current study in an updated meta-analysis. The reviews by Larsson *et al*<sup>7</sup> only considered stroke as an outcome while the review by Zhang *et al*<sup>10</sup> considered CVD. The current review included stroke, stroke mortality, CHD, and CHD mortality separately. In addition, we included the EPIC-Norfolk cohort which is a Caucasian population. Our results indicate that chocolate consumption was associated with a lower risk of CVD.

A number of issues have to be taken into account when interpreting the results of the present study. Although FFQs are well-established methods to quantify dietary information in large scale population studies, they have limitations: importantly, recall bias as well as underreporting, particularly among women

**Table 3** Risk of total (fatal and non-fatal) CHD and stroke incidence by quintiles of chocolate intake in EPIC-Norfolk (1993–2008)

Range (g/day)	Total	Quintile 1 0	Quintile 2 0.6–3.5	Quintile 3 4.1–7.0	Quintile 4 7.2–15.5	Quintile 5 15.6–98.8	p Linearity
CHD/n (%)	2434/20 951 (11.6)	577/4195 (13.8)	564/4437 (12.7)	437/4035 (10.8)	449/4106 (10.9)	407/4178 (9.7)	
Model 1		1.00	0.97 (0.86 to 1.09)	0.97 (0.86 to 1.10)	0.87 (0.77 to 0.99)	0.85 (0.75 to 0.96)	0.003
Model 2		1.00	1.01 (0.90 to 1.13)	1.01 (0.89 to 1.15)	0.91 (0.80 to 1.03)	0.88 (0.77 to 1.01)	0.024
Model 3		1.00	1.03 (0.91 to 1.15)	1.03 (0.91 to 1.17)	0.92 (0.81 to 1.05)	0.91 (0.80 to 1.04)	0.06
	1754/16 162 (10.9)	435/3251 (13.4)	407/3406 (11.9)	307/3102 (9.9)	325/3220 (10.1)	280/3183 (8.8)	
Model 4a		1.00	1.00 (0.87 to 1.15)	0.98 (0.85 to 1.14)	0.88 (0.76 to 1.01)	0.83 (0.71 to 0.97)	0.006
Model 4b		1.00	1.00 (0.87 to 1.14)	0.98 (0.84 to 1.13)	0.87 (0.75 to 1.01)	0.82 (0.70 to 0.97)	0.005
Stroke/n (%)	848/20 951 (4.0)	226/4195 (5.4)	187/4437 (4.2)	139/4035 (3.4)	165/4106 (4.0)	131/4178 (3.1)	
Model 1		1.00	0.83 (0.68 to 1.00)	0.84 (0.68 to 1.04)	0.85 (0.70 to 1.04)	0.77 (0.62 to 0.96)	0.036
Model 2		1.00	0.85 (0.70 to 1.03)	0.87 (0.70 to 1.07)	0.86 (0.70 to 1.06)	0.77 (0.62 to 0.97)	0.046
Model 3		1.00	0.86 (0.71 to 1.05)	0.87 (0.71 to 1.08)	0.88 (0.71 to 1.07)	0.78 (0.63 to 0.98)	0.06
	648/16 162 (4.0)	168/3251 (5.2)	149/3406 (4.4)	100/3102 (3.2)	132/3220 (4.1)	99/3183 (3.1)	
Model 4a		1.00	0.95 (0.76 to 1.18)	0.87 (0.67 to 1.12)	0.94 (0.74 to 1.18)	0.81 (0.62 to 1.05)	0.14
Model 4b		1.00	0.95 (0.76 to 1.18)	0.87 (0.67 to 1.11)	0.94 (0.74 to 1.18)	0.80 (0.62 to 1.05)	0.14

Model 1 adjusted for sex and age.

Model 2 adjusted for sex, age, smoking, physical activity, energy intake and alcohol consumption.

Model 3: as model 2 and diabetes, body mass index, systolic blood pressure, LDL cholesterol and HDL cholesterol.

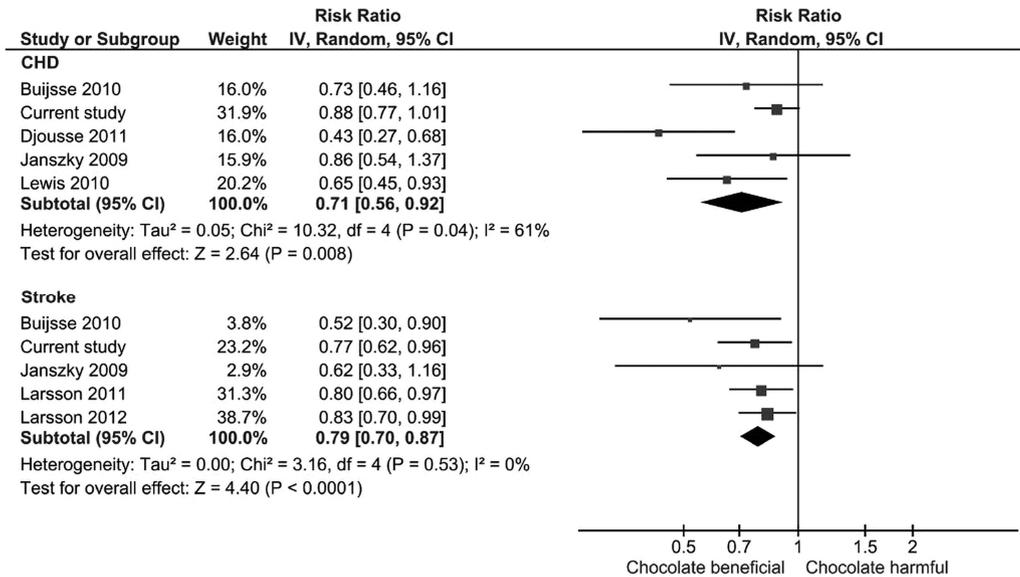
Model 4a: as model 3, but restricted to number of participants for whom CRP is available (N=16 162).

Model 4b: as model 3 and CRP (N=16 162).CHD, coronary heart disease; CRP, C-reactive protein; EPIC, European Prospective Investigation into Cancer; HDL, high density lipoprotein; LDL, low density lipoprotein.

**Table 4** Study design, participants, follow-up and outcomes for studies evaluating chocolate consumption and CVD

Study ID	Dietary exposures	Study design	No in analysis	Characteristics	Follow-up (years)	Outcome measures	Measures of association
Buijsse <i>et al</i> <sup>5</sup>	Cocoa intake	Prospective cohort, Netherlands	470	White men with mean age 72 years	15	Cardiovascular mortality at 15 years	RR: 0.50 (0.32–0.78) for >2.25 g/day vs <0.5 g/day (top vs bottom tertile)
Buijsse <i>et al</i> <sup>20</sup>	Chocolate consumption	Prospective cohort, Germany	19 357	Men and women with mean age of 49 years, 38% were men with no history of myocardial infarction and stroke and not using blood pressure drugs	10	Incident MI, stroke and composite of both at mean of 8.1 years follow-up	RR: MI 0.73 (0.47–1.18), stroke 0.52 (0.30–0.89), composite 0.61 (0.44–0.87) for 4th quartile vs 1st quartile energy-adjusted chocolate consumption
Djousse <i>et al</i> <sup>19</sup>	Chocolate consumption	Cross-sectional, USA	4970	Men and women with mean age of 52 years (range 25–93 years), 45% were men and 5% were of African-American origin	NA	Prevalent coronary heart disease (MI or coronary revascularisation)	OR: 0.43 (0.27–0.68) for >5 times a week vs no consumption (based on some degree of post-hoc categorisation of collected data).
Janszky <i>et al</i> <sup>21</sup>	Chocolate consumption	Prospective cohort, Sweden	1169	Men (71%) and women who survived an acute myocardial infarction with mean age of 59 years	8	Cardiovascular mortality, recurrent MI, stroke and congestive heart failure	HR: Cardiovascular mortality 0.34 (0.17–0.70), recurrent MI 0.86 (0.54–1.37), stroke 0.62 (0.33–1.16) for <2× week vs never (based on some degree of post-hoc categorisation of collected data)
Larsson <i>et al</i> <sup>6</sup>	Chocolate consumption	Prospective cohort, Sweden	33 372	Women with no history of stroke, age range 49–83 years	10.4	Incident stroke	RR: 0.80 (0.66–0.99) >45 g/week vs <8.9 g/week (top vs bottom quartile)
Larsson <i>et al</i> <sup>7</sup>	Chocolate consumption	Prospective cohort, Sweden	37 103	Men with mean age mean of 59 years and no history of stroke	10.2	Incident stroke	RR: 0.83 (0.70–0.99) for >51.6 g/week vs <12 g/week (top vs bottom quartile)
Lewis <i>et al</i> <sup>8</sup>	Chocolate consumption	Post-hoc analysis of a randomised trial of calcium supplementation in women, Australia	1216	Women, mean age 75 years	9.5	Atherosclerotic vascular disease and ischaemic heart disease	HR: Ischaemic heart disease 0.65 (0.46–0.94) for <1 serving/week vs >1 serving/week (based on some degree of post-hoc categorisation of collected data)
Mink <i>et al</i> <sup>9</sup>	Chocolate consumption	Prospective cohort, USA	34 489	Women with mean age 62 years with no history of heart disease	16	CHD mortality and stroke mortality	Rate ratio: CHD mortality 0.98 (0.88–1.10), stroke mortality 0.85 (0.70–1.03) for <1 per week vs >1 per week (based on some degree of post-hoc categorisation of collected data)
Current study	Chocolate consumption	Cohort	25 663	55% women, 99.6% white Caucasians, mean age 59 years with no history of heart disease or stroke	11.8	Incident CHD, stroke and composite of the two, defined as CVD	HR: CHD 0.88 (0.77–1.01), stroke 0.77 (0.62–0.96), CVD 0.86 (0.76–0.97) (top quintile vs bottom quintile)

CHD, coronary heart disease; CVD, cardiovascular disease; MI, myocardial infarction.



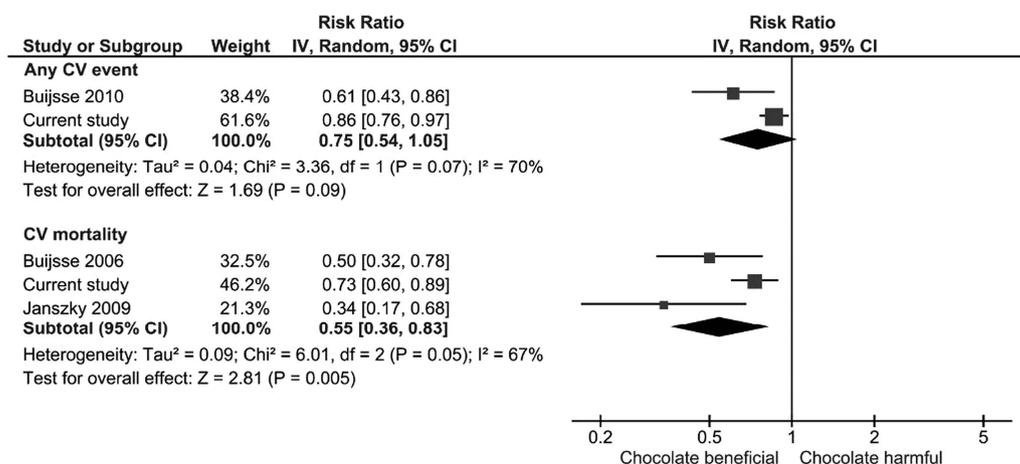
**Figure 1** Meta-analysis risk of the association between chocolate consumption and risk of coronary heart disease (CHD) and stroke.

and participants who are obese.<sup>22</sup> Underreporting is selective, and includes sweets and snacks. It is possible that lower CVD rates among people who report consuming more chocolate might be due to differential underreporting of chocolate intake in those with potentially greater CVD risk such as the obese and physically inactive. It has been suggested that dark chocolate may have more beneficial effects than milk chocolate.<sup>23</sup> Milk chocolate was more frequently consumed than dark chocolate in this cohort (unpublished results); however, we still observed a reduced risk of CVD. This may indicate that not only flavonoids, but also other compounds—possibly related to milk constituents such as calcium and fatty acids—may provide an explanation for the observed association. In addition, the whole dietary pattern may be of relevance, giving concern for unmeasured confounding. Therefore, this observational study cannot provide evidence on the potential causality of the observed association.

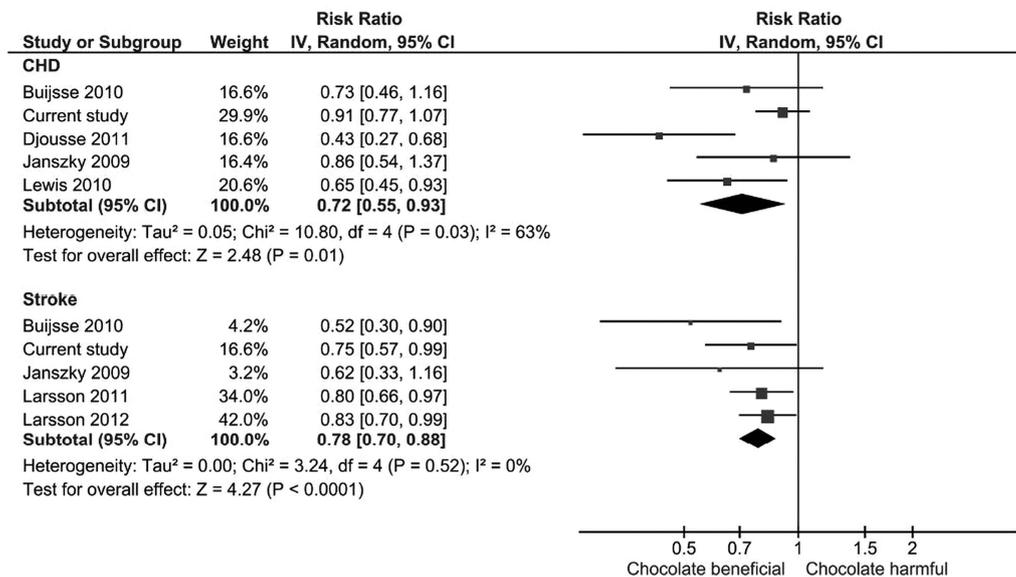
We used the FFQ to assess chocolate intake and this measure of dietary assessment method has been validated in the EPIC-Norfolk cohort.<sup>24</sup> While the FFQ is prone to recall bias and requires the participant to ‘average out’ over a long period

of time, this method has the advantage that it covers a longer time-frame than other methods such as 7-day diet diary (7dDD) and 24 h-diet recall. Chocolate consumption might be a more episodically consumed food for a proportion of the participants and this could have miss-classified the participant as a non-consumer or a high consumer depending in which week the 7dDD or day of 24 h-diet recall would have been completed.

It is possible that part of the observed association could be explained by reverse causality in that people with a higher risk profile, including those with obesity, diabetes mellitus, or prevalent CVD, eat less chocolate-containing foods than people who have a perceived healthy risk profile. However, we excluded people who reported prevalent CVD at the baseline health questionnaire, and analyses were adjusted for diabetes mellitus. We observed that the reference group (non-chocolate consumers) had the highest mean BMI, highest median CRP, highest proportion of participants with diabetes, highest levels of inactivity, and lowest fat intake compared to the other quintiles of chocolate consumers. Alternatively, it may be that higher chocolate consumers have other behaviours that are beneficial for cardiovascular health. Participants with a higher energy intake are also



**Figure 2** Meta-analysis of chocolate consumption and risk of cardiovascular (CV) disease (composite).



**Figure 3** Meta-analysis risk of coronary heart disease (CHD) and stroke with chocolate consumption using propensity matched cohort.

likely to have a higher energy expenditure due to physical activity. They are hence more likely to consume more foods, including chocolate containing foods, which may explain part of the observed associations, although stratified analyses suggested that the association found was homogeneous across levels of physical activity. In addition to multivariable adjustment, we conducted stratified analyses and propensity score matching for baseline differences. Nevertheless we cannot exclude residual confounding from these or other unmeasured factors. The consumption of chocolate as well as its influence on cardiovascular risk may be different depending on the ethnicity of the participants. External validity of our results may be limited to Caucasians.

Our results are somewhat surprising since the expectation was that benefits of chocolate consumption would be mainly associated with dark chocolate rather than the commercially available products generally used in a British population which are high in sugar content and fat. One study has shown reduced incidence of diabetes among men and women with chocolate intake,<sup>25</sup> and other studies have shown that chocolate consumption increases bodyweight.<sup>26 27</sup> This may suggest that there is a balance between benefit and risk with chocolate intake, which is dependent on the risk profile of the individual including baseline weight and dose of chocolate intake.

We did not include heart failure as an outcome for the current analysis for the following reasons. The pathophysiology of CHD and stroke which relates to atherosclerosis differs from that of heart failure. The physiological changes in heart failure are largely driven by neuroendocrine dysregulation and renin-angiotensin-aldosterone system activation. The mechanism by which chocolate might have an impact on these pathways may differ. In addition, the risk factors and confounders in heart failure differ from those in the CVD analysis.

This analysis included clinical biomarkers and anthropometry measured by trained staff. Adjustment for these variables attenuated the association between chocolate consumption and risk of CVD, CHD and stroke, but particularly in the case of stroke and fatal CVD, associations remained significant. CRP was available for a subcohort, in which we observed a better cardiovascular risk profile; however, we observed that CRP did not modify the association between chocolate and CVD, CHD or stroke. Pathways other than anti-inflammatory markers, such as blood

pressure, may be a more likely mechanism to explain the observed association between chocolate and CVD/stroke.

Bias may also affect the systematic review if the original studies made post-hoc decisions on particular outcomes or categories for analysis based on the nature of the findings. The primary studies collected data using different categories of chocolate consumption (eg, frequent vs rare, different quintiles or quartiles) and it seemed that some of the studies made post-hoc decisions on what the cut-offs were in defining categories for analysis (including pooling certain categories while excluding others). This raises the possibility of bias from selective outcome and analysis reporting where categorical cut-offs could have been chosen based on the statistical significance of the findings. Nevertheless, the cumulative evidence reported in this study suggests that high chocolate consumption may be associated with cardiovascular benefit.

### Key messages

#### What is already known on this subject?

- ▶ Chocolate consumption has been associated with a lower risk of cardiovascular disease, but studies have limitations with regard to participant selection and combined outcome measures.

#### What might this study add?

- ▶ We observed evidence that higher intake of chocolate may be associated with lower cardiovascular disease and mortality.
- ▶ Our meta-analysis of eight studies found a lower risk of cardiovascular disease with chocolate consumption.

#### How might this impact clinical practice?

- ▶ Our study adds to the previously found inverse associations between chocolate intake and risk of future cardiovascular events.
- ▶ There does not appear to be any evidence to say that chocolate should be avoided in those who are concerned about cardiovascular risk.

Future research is still needed to explore the association between chocolate and CVD. Studies are needed to understand better the biological mechanism by which chocolate and flavonoids reduce the risk of CVD. It is also possible that some individuals will not benefit from increased chocolate consumption, such as those who are overweight or diabetic. More research is needed to identify individuals who will benefit most.

In conclusion, our findings support the previously reported association between habitual chocolate intake and a lower risk of CHD events in the large EPIC-Norfolk prospective population study, and we have further set this in context with consistent results seen in meta-analysis of current evidence. While randomised controlled trials of chocolate and cardiovascular end points could be conducted, feasibility is uncertain. For the time being, within the general context of existing recommendations for behaviours conducive to cardiovascular health, there does not appear to be evidence that chocolate should be avoided in terms of impact on cardiovascular risk.

#### Author affiliations

<sup>1</sup>Epidemiology Group, Division of Applied Health Sciences, School of Medicine & Dentistry, University of Aberdeen, Aberdeen, UK

<sup>2</sup>Cardiovascular Institute, University of Manchester, Manchester, UK

<sup>3</sup>Department of Cardiology, Academic Medical Center, Amsterdam, The Netherlands

<sup>4</sup>Department of Public Health & Primary Care, University of Cambridge, Cambridge, UK

<sup>5</sup>Department of Public Health & Primary Care, Norwich Medical School, University of East Anglia, Norwich, UK

<sup>6</sup>Lancashire Teaching Hospital NHS Foundation Trust, Preston, UK

<sup>7</sup>Medical Research Council Epidemiology Unit, Cambridge, UK

**Acknowledgements** The authors would like to thank the participants of the EPIC-Norfolk cohort. We thank the nutritionists' team and data management team of the EPIC-Norfolk cohort.

**Contributors** K-TK and NJW are Principal Investigators of EPIC-Norfolk cohort. SMB analysed the EPIC-Norfolk data. RNL is responsible for data management and linkage and contributed to data analysis. MAHL and SMB designed the cohort analysis. CSK and YKL performed the literature search for systematic review. CSK and JKY screened and collected the data for systematic review which was checked by YKL and PKM. CSK and YKL conducted the meta-analysis. SMB, PKM, CSK and YKL drafted the paper and all authors contributed to the writing of the paper.

**Funding** The EPIC-Norfolk study was supported by grants from the Medical Research Council and Cancer Research UK.

**Competing interests** None declared.

**Patient consent** Obtained.

**Ethics approval** Norwich District Health Authority Ethics Committee.

**Provenance and peer review** Not commissioned; externally peer reviewed.

#### REFERENCES

- Corti R, Flammer AJ, Hollenberg NK, *et al.* Cocoa and cardiovascular health. *Circulation* 2009;119:1433–41.
- Ding EL, Hutfless SM, Ding X, *et al.* Chocolate and prevention of cardiovascular disease: a systematic review. *Nutr Metab (Lond)* 2006;3:2.
- Buitrago-Lopez A, Sanderson J, Johnson L, *et al.* Chocolate consumption and cardiometabolic disorders: systematic review and meta-analysis. *BMJ* 2011;343:d4488.
- Wang ZM, Nie ZL, Zhou B, *et al.* Flavonols intake and the risk of coronary heart disease: a meta-analysis of cohort studies. *Atherosclerosis* 2012;222:270–3.
- Buijsse B, Feskens EJ, Kok FJ, *et al.* Cocoa intake, blood pressure, and cardiovascular mortality: the Zutphen Elderly Study. *Arch Intern Med* 2006;166:411–17.
- Larsson SC, Virtamo J, Wolk A. Chocolate consumption and risk of stroke in women. *J Am Coll Cardiol* 2011;58:1828–9.
- Larsson SC, Virtamo J, Wolk A. Chocolate consumption and risk of stroke: a prospective cohort of men and meta-analysis. *Neurology* 2012;79:1223–9.
- Lewis JR, Prince RL, Zhu K, *et al.* Habitual chocolate intake and vascular disease: a prospective study of clinical outcomes in older women. *Arch Intern Med* 2010;170:1857–8.
- Mink PJ, Scrafford CG, Barraj LM, *et al.* Flavonoid intake and cardiovascular disease mortality: a prospective study in postmenopausal women. *Am J Clin Nutr* 2007;85:895–909.
- Zhang Z, Xu G, Liu X. Chocolate intake reduces risk of cardiovascular disease: evidence from 10 observational studies. *Int J Cardiol* 2013;168:5548–50.
- Day N, Oakes S, Luben R, *et al.* EPIC-Norfolk: study design and characteristics of the cohort. European Prospective Investigation of Cancer. *Br J Cancer* 1999;80(Suppl 1):95–103.
- Canoy D, Wareham N, Luben R, *et al.* Cigarette smoking and fat distribution in 21,828 British men and women: a population-based study. *Obes Res* 2005;13:1466–75.
- Wareham NJ, Jakes RW, Rennie KL, *et al.* Validity and repeatability of the EPIC-Norfolk Physical Activity Questionnaire. *Int J Epidemiol* 2002;31:168–74.
- Welch AA, Luben R, Khaw KT, *et al.* The CAFFE computer program for nutritional analysis of the EPIC-Norfolk food frequency questionnaire and identification of extreme nutrient values. *J Hum Nutr Diet* 2005;18:99–116.
- Boekholdt SM, Peters RJ, Day NE, *et al.* Macrophage migration inhibitory factor and the risk of myocardial infarction or death due to coronary artery disease in adults without prior myocardial infarction or stroke: the EPIC-Norfolk Prospective Population study. *Am J Med* 2004;117:390–7.
- Higgins JPT, Green S. *Cochrane Handbook of Systematic Reviews of Interventions*. Version 5.1.0. [Updated March 2011]. The Cochrane Collaboration, 2011. <http://www.cochrane-handbook.org>
- Ioannidis JPA, Trikalinos TA. The appropriateness of asymmetry tests for publication bias in meta-analyses: a large survey. *CMAJ* 2007;176:1091–6. [http://www.ssc.wisc.edu/sscc/pubs/stata\\_psmatch.htm](http://www.ssc.wisc.edu/sscc/pubs/stata_psmatch.htm) (accessed 16 Dec 2014).
- Djousse L, Hopkins PN, North KE, *et al.* Chocolate consumption is inversely associated with prevalent coronary heart disease: The National Heart, Lung, and Blood Institute Family Heart Study. *Clin Nutr* 2011;30:182–7.
- Buijsse B, Weikert C, Drogan D, *et al.* Chocolate consumption in relation to blood pressure and risk of cardiovascular disease in German adults. *Eur Heart J* 2010;31:1616–23.
- Janszky I, Mukamal KJ, Ljung R, *et al.* Chocolate consumption and mortality following a first acute myocardial infarction: the Stockholm Heart Epidemiology Program. *J Intern Med* 2009;266:248–57.
- Subar AF, Kipnis V, Troiano RP, *et al.* Using intake biomarkers to evaluate the extent of dietary misreporting in a large sample of adults: the OPEN study. *Am J Epidemiol* 2003;158:1–13.
- Serafini M, Bugianesi R, Maiani G, *et al.* Plasma antioxidants from chocolate. *Nature* 2003;424:1013.
- Bingham SA, Gill C, Welch A, *et al.* Comparison of dietary assessment methods in nutritional epidemiology: weighted records v. 24 h recalls, food-frequency questionnaires and estimated-diet records. *Br J Nutr* 1994;72:619–43.
- Oba S, Nagata C, Nakamura K, *et al.* Consumption of coffee, green tea, oolong tea, black tea, chocolate snacks and the caffeine content in relation to risk of diabetes in Japanese men and women. *Br J Nutr* 2010;103:453–9.
- Greenberg JA, Buijsse B. Habitual chocolate consumption may increase body weight in a dose-response manner. *PLoS ONE* 2013;8:e70271.
- Greenberg JA, Mason JE, Buijsse B, *et al.* Chocolate-candy consumption and 3-year weight gain among postmenopausal US women. *Obesity* 2015;23:677–83.

## **Correction: *Habitual chocolate consumption and risk of cardiovascular disease among healthy men and women***

---

Kwok CS, Boekholdt SM, Lentjes MAH, *et al.* Habitual chocolate consumption and risk of cardiovascular disease among healthy men and women. *Heart* 2015;101:1279–1287. doi: 10.1136/heartjnl-2014-307050corr1.

In this paper the paragraph Chocolate consumption in the EPIC-Norfolk cohort contains an error. The section which reads ‘Higher chocolate consumption was associated with trends towards a beneficial cardiovascular risk factor profile including lower age, lower BMI, lower waist/hip ratio, lower systolic blood pressure, lower concentrations of apolipoprotein B (apoB) and CRP, a lower prevalence of diabetes mellitus, and more physical inactivity (table 1).’ The final sentence should read ‘and less physical inactivity (table 1)’.

© Article author(s) (or their employer(s) unless otherwise stated in the text of the article) 2018. All rights reserved. No commercial use is permitted unless otherwise expressly granted.

*Heart* 2018;104:532. doi:10.1136/heartjnl-2014-307050corr1

