



Dietary flavonoid intake and weight maintenance: three prospective cohorts of 124 086 US men and women followed for up to 24 years

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ABSTRACT OBJECTIVE

To examine whether dietary intake of specific flavonoid subclasses (including flavonols, flavones, flavanones, flavan-3-ols, anthocyanins, and flavonoid polymers) is associated with weight change over time.

DESIGN

Three prospective cohort studies.

SETTING

Health professionals in the United States.

PARTICIPANTS

124 086 men and women participating in the Health Professionals Follow-up Study (HPFS), Nurses' Health Study (NHS), and Nurses' Health Study II (NHS II).

MAIN OUTCOME MEASURE

Self reported change in weight over multiple four year time intervals between 1986 and 2011.

RESULTS

Increased consumption of most flavonoid subclasses, including flavonols, flavan-3-ols, anthocyanins, and flavonoid polymers, was inversely associated with weight change over four year time intervals, after adjustment for simultaneous changes in other lifestyle factors including other aspects of diet, smoking status, and physical activity. In the pooled results, the greatest magnitude of association was observed for anthocyanins (−0.23 (95% confidence interval −0.30 to −0.15) lbs per additional standard deviation/day, 10 mg), flavonoid polymers (−0.18 (−0.28 to −0.08) lbs per additional SD/day, 138 mg), and flavonols (−0.16 (−0.26 to −0.06) lbs per additional SD/day, 7 mg). After additional adjustment for fiber intake, associations remained significant for anthocyanins, proanthocyanidins, and total flavonoid polymers but

were attenuated and no longer statistically significant for other subclasses.

CONCLUSIONS

Higher intake of foods rich in flavonols, flavan-3-ols, anthocyanins, and flavonoid polymers may contribute to weight maintenance in adulthood and may help to refine dietary recommendations for the prevention of obesity and its potential consequences.

Introduction

Flavonoids are naturally occurring bioactive compounds that represent a constituent of fruits and vegetables, beyond calorie and macronutrient content, that could potentially influence body weight. In a previous food based analysis, we found wide ranging associations between intake of specific fruits and vegetables and weight maintenance.¹ Among fruits, an increased intake of blueberries, apples, pears, prunes, strawberries, and grapes was inversely associated with weight gain. Increased intake of peppers and celery was also associated with less weight gain. The mechanisms responsible for these benefits and the specific constituents of these fruits and vegetables that may explain these associations are unclear.

The particular fruits and vegetables associated with less weight gain are rich sources of several flavonoid subclasses, particularly flavonols, anthocyanins, and flavones. Animal models and short term human studies provide evidence for underlying mechanisms that relate flavonoids to weight: several flavonoid subclasses have been shown to decrease energy intake,^{2,5} increase glucose uptake in muscle *in vivo*,⁶ and decrease glucose uptake in adipose tissue *in vitro*.^{6,7} Other studies, predominantly focusing on green tea, a rich source of the flavan-3-ol subclass of flavonoids, provide evidence to suggest that flavonoids may decrease fat absorption,^{8,9} increase energy expenditure,⁹⁻¹² and inhibit adipogenesis.^{5,13-15}

Most weight loss studies to date have focused on the flavan-3-ol subclass found in green tea and are limited to small numbers of overweight and obese participants. Here, we examine, to our knowledge for the first time, the associations between habitual consumption of all flavonoid subclasses and weight gain among 124 086 American men and women followed for up to 24 years. The results from this analysis may help to refine previous dietary recommendations and provide people with better guidance on which fruits and vegetables to choose for the prevention of weight gain. Most adults gain weight as they age, and even small increases in weight can have a substantial effect on the risk of

WHAT IS ALREADY KNOWN ON THIS TOPIC

Flavonoids are naturally occurring bioactive compounds that represent a constituent of fruits and vegetables, beyond calorie and macronutrient content, that could potentially influence body weight

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WHAT THIS STUDY ADDS

Increased consumption of most flavonoid subclasses was associated with less weight gain

The greatest magnitude of association was observed for anthocyanins, flavonoid polymers, and flavonols

Choosing high flavonoid fruits and vegetables, such as apples, pears, berries, and peppers, may help with weight control.

hypertension,¹⁶ diabetes,¹⁷ cardiovascular disease,^{18,19} cancer,²⁰ and mortality.²¹ Strategies to help people to maintain a healthy weight from childhood onward are therefore critically needed.

Methods

Study populations

We used three large prospective cohorts of American men and women from 50 states for this analysis. The Health Professionals Follow-up Study (HPFS) began in 1986 and enrolled 51 529 male health professionals aged 40-75 years.²² The Nurses' Health Study (NHS) began in 1976 and enrolled 121 701 female nurses aged 30-55 years.²³ The Nurses' Health Study 2 (NHS II) began in 1989 and enrolled 116 686 younger female nurses aged 25-42 years.²⁴

We excluded men and women with a history of chronic disease at baseline because of potential disease related weight loss. We considered the following to be chronic diseases: obesity (body mass index ≥ 30), diabetes, cancer, cardiovascular disease, renal disease, pulmonary disease, liver disease, ulcerative colitis, lupus, tuberculosis, multiple sclerosis, amyotrophic lateral sclerosis, and Parkinson's disease. With the exception of obesity and cardiovascular disease, we censored people who developed these conditions during follow-up six years before the date of diagnosis. We censored people who developed cardiovascular disease during follow-up at the date of diagnosis. We did not censor people if they became obese during follow-up.

We temporarily excluded women who were pregnant or lactating for one four year interval and censored people who reported gastric bypass surgery. We excluded people over age 65 at baseline and censored the remaining people when they reached age 65 owing to potential age related sarcopenia. Finally, we excluded men and women with missing baseline lifestyle data or who reported implausible energy intake (men < 800 or > 4200 daily calories; women < 600 or > 3500 daily calories). Our final analysis included 20 525 men from the HPFS, 39 423 women from the NHS, and 64 138 women from the NHS II (see supplemental table A for specific exclusions and supplemental table B for a comparison of original versus analysis cohorts).

Patient involvement

No patients were involved in setting the research question or the outcome measures, nor were they involved in the design and implementation of the study. There are no plans to involve patients in the dissemination of the results.

Assessment of weight

All participants self reported their weight in pounds via a questionnaire every two years. We calculated weight change within each four year period as the difference in weight between the beginning and end of each period, so negative differences indicate weight loss and positive differences indicate weight gain. There were a total of six four year periods in the HPFS and NHS (24 years of follow-up) and five in the NHS II (20 years of follow-up).

Pearson correlation coefficients between self reported weight and technician measured weight were 0.97 among a sample of 140 women in the NHS and 123 men in the HPFS.²⁵ We use a conversion factor of 0.454 to convert pounds to kg in the text.

Assessment of flavonoid intakes

All participants self reported their diet every four years using a validated,²⁶ semiquantitative food frequency questionnaire. We used the US Department of Agriculture (USDA) flavonoid content of foods and proanthocyanidin databases to create a database of the flavonoid content per serving of foods and beverages included on the study food frequency questionnaire.^{27,28} We calculated a person's flavonoid intake as the frequency of consumption of specified portions of flavonoid containing foods multiplied by the flavonoid content (aglycone equivalents) per serving of that food, summed across all foods and beverages, as described previously.²⁹

We derived intakes of seven flavonoid subclasses—flavanones (eriodictyol, hesperetin, and naringenin), anthocyanins (cyanidin, delphinidin, malvidin, pelargonidin, petunidin, and peonidin), flavan-3-ols (catechin, gallic acid, epicatechin, epigallocatechin, epicatechin 3-gallate, and epigallocatechin 3-gallate), proanthocyanidins (dimers, trimers, 4-6mers, 7-10mers, and polymers), flavonoid polymers (proanthocyanidins, theaflavins, and thearubigins), flavonols (quercetin, kaempferol, myricetin, and isorhamnetin), and flavones (luteolin and apigenin)—as well as total flavonoids (all subclasses summed). As a sensitivity analysis, we additionally examined flavonoid polymers excluding thearubigins because current quantification methods are somewhat crude. We did not include isoflavones in our analysis because levels are very low (< 3 mg/d) in the habitual US diet and, from the extensive intervention studies, we know that an isoflavone intake of more than 25 mg/d is needed for any biological/clinical effect.^{30,31}

Assessment of covariates

Participants reported their lifestyle habits and any recent physician diagnosed diseases via a questionnaire every two years. We included baseline age and body mass index for that particular four year period as covariates in our main models, as well as change in smoking status, physical activity level,³² hours of sitting or watching TV, and hours of sleep over the same time period and change in intake of the following foods/nutrients: fried potatoes, juice, whole grains, refined grains, fried foods, nuts, whole fat dairy, low fat dairy, sugar sweetened drinks, diet drinks, sweets, processed meats, non-processed meats, trans fat, alcohol, seafood, and caffeine. Because citrus juices are a main source of both flavone and flavanone intake in this population, we used non-citrus juice rather than all fruit juice as a covariate for analyses of these two flavonoid subclasses. In additional models, we further adjusted for change in total fiber intake (g/day).

Statistical methods

We used multivariable generalized linear regression models to examine the association between change in weight (lbs) over four years and change in intake of flavonoid subclasses (standard deviation/day) over the same four year time interval, as described in a previous publication.³³ We used robust variance to account for within person repeated measures because each person could have contributed up to five time periods, and results for all four year time intervals are averaged. We examined each flavonoid subclass individually and did not mutually adjust for other flavonoid subclasses in our models. We included all other lifestyle factors assessable by questionnaire that we and many others have found to be associated with weight change. We did not include biological factors (such as blood pressure) and drugs because these factors could be mediators (in the causal pathway) or direct correlates of mediators of the effect of diet on weight change.

We truncated change in intake of flavonoids and change in weight at the 0.5th and 99.5th centiles to minimize the influence of outliers. The last observation was carried forward for missing values of continuous variables with the exception of diet (main exposure) and weight (main outcome), and indicators were used for missing values of categorical variables. Missing values were carried forward only once for diet and weight,

after which the follow-up was censored. We did several sensitivity analyses, including a complete case analysis in which missing values were not carried forward, models that included obese participants, and models additionally adjusting for change in total fiber, change in total calorie intake, and baseline levels of all covariates.

We pooled results from the three cohorts by using the DerSimonian-Laird estimators and the Q statistic to test for heterogeneity. The three studies are weighted by the inverse of the sum of the study specific variance plus the common between studies variance (random effects pooling). All analyses used SAS version 9.2 and a two tailed α of 0.05.

Results

The average age of participants at baseline ranged from 36.3 years in women in the NHS II to 46.9 years in men in the HPFS and 48.5 years in women in the NHS (table 1). Over each four year period, men gained an average of 2.2 lbs (1 kg), women in the NHS 2.9 lbs (1.3 kg), and women in the NHS II 4.4 lbs (2.0 kg). Men had a median intake of 224 mg/d of total flavonoids, women in the NHS 247 mg/d, and women in the NHS II 236 mg/d. Intake of flavonoid polymers was the highest, ranging from a median of 118 mg/d to 144 mg/d across the three cohorts, and intake of flavones was the lowest, with a median of 1.2 to 2.2 mg/d.

Table 1 | Baseline characteristics and average four year lifestyle changes of men and women in three prospective cohorts

Characteristics	Health Professionals Follow-up Study (n=20 525)		Nurses' Health Study (n=39 423)		Nurses' Health Study II (n=64 138)	
	Baseline* (1986)	Four year change†	Baseline* (1986)	Four year change†	Baseline* (1991)	Four year change†
Age, years	46.9 (2.9)	–	48.5 (2.4)	–	36.3 (3.4)	–
Body mass index, kg/m ²	24.6 (1.4)	–	23.6 (1.4)	–	22.8 (2.2)	–
Weight, lbs	174 (13)	2.2 (–14.0–20.0)	140 (9)	2.9 (–14.0–25.0)	137 (16)	4.4 (–10.0–32.3)
Physical activity, MET-hr/wk	23.5 (19.5)	5.1 (–46.0–100.0)	15.0 (9.9)	1.6 (–34.7–42.6)	21.6 (22.1)	0.4 (–26.1–22.6)
Alcohol, servings/d	0.9 (0.7)	0.0 (–2.1–1.6)	0.5 (0.4)	0.0 (–1.3–0.9)	0.3 (0.4)	0.0 (–0.5–0.8)
Caffeine, mg/d	236 (141)	–11 (–372–306)	305 (112)	–29 (–354–270)	241 (167)	–12 (–280–224)
Flavanols, mg/d	11.7 (7.9–17.0)	2.1 (–14.3–26.9)	12.1 (8.1–18.1)	1.6 (–14.0–27.0)	14.2 (9.3–23.3)	1.3 (–19.5–18.0)
Flavones, mg/d	2.2 (1.2–3.1)	0.1 (–3.4–2.6)	2.1 (1.2–3.1)	0.0 (–2.8–2.2)	1.2 (0.7–2.1)	0.1 (–1.4–2.2)
Flavanones, mg/d	41.7 (16.1–64.9)	–0.1 (–79.3–78.9)	37.6 (14.9–63.7)	–0.7 (–66.1–62.0)	21.7 (9.3–48.6)	–1.3 (–42.5–47.8)
Flavan-3-ols, mg/d	23.9 (13.5–51.0)	0.5 (116.3–115.7)	24.2 (13.2–71.6)	–0.9 (–113.6–133.2)	26.1 (13.0–73.6)	–0.8 (–119.4–79.4)
Anthocyanins, mg/d	7.7 (3.9–16.2)	2.4 (–24.6–28.7)	11.5 (5.2–19.6)	1.8 (–28.4–28.3)	5.7 (3.1–14.6)	4.5 (–17.5–34.6)
Polymers, mg/d	117.6 (63.4–214.4)	7.1 (–344.7–370.8)	134.9 (72.6–254.4)	1.3 (–342.6–396.5)	143.5 (76.5–267.6)	3.9 (–363.6–252.3)
Total flavonoids, mg/d	223.6 (138.1–362.2)	11.9 (–472.6–531.4)	247.1 (149.8–416.0)	2.5 (–480.3–547.0)	235.9 (136.9–410.6)	8.2 (–506.2–360.3)
Total fruit without juice, servings/d	1.5 (0.7)	0.1 (–1.9–2.1)	1.5 (0.5)	0.0 (–1.9–1.9)	1.2 (0.7)	0.0 (–1.3–1.5)
Total vegetables, servings/d	2.9 (1.0)	0.2 (–2.6–3.7)	3.2 (0.8)	0.1 (–2.6–3.4)	3.1 (1.6)	0.0 (–2.7–3.0)
Whole fat dairy, servings/d	1.0 (0.6)	–0.1 (–2.3–1.4)	1.2 (0.5)	–0.1 (–2.3–1.2)	0.8 (0.6)	0.0 (–1.4–1.1)
Low fat dairy, servings/d	0.8 (0.6)	–0.1 (–2.0–1.7)	0.8 (0.4)	0.1 (–1.5–2.0)	1.1 (0.8)	0.0 (–1.4–1.7)
Seafood, servings/d	0.4 (0.2)	0.0 (–0.5–0.6)	0.3 (0.1)	0.0 (–0.5–0.4)	0.3 (0.2)	0.0 (–0.4–0.3)
Whole grains, servings/d	1.5 (0.8)	0.0 (–2.3–2.9)	0.8 (0.4)	0.1 (–1.6–2.2)	1.3 (0.9)	0.0 (–1.5–1.3)
Refined grains, servings/d	1.2 (0.6)	0.0 (–2.1–2.1)	1.2 (0.4)	0.0 (–1.7–1.7)	1.3 (0.7)	–0.1 (–1.3–1.7)
Nuts, servings/d	0.3 (0.3)	0.0 (–0.9–0.8)	0.1 (0.1)	0.0 (–0.6–0.5)	0.1 (0.1)	0.1 (–0.2–0.7)
Sugar sweetened drinks, servings/d	0.3 (0.4)	0.0 (–0.9–0.9)	0.2 (0.2)	0.0 (–0.6–0.7)	0.3 (0.5)	0.0 (–0.9–0.8)
Zero calorie soda, servings/d	0.5 (0.6)	0.0 (–1.5–1.6)	0.5 (0.4)	0.0 (–1.2–1.5)	0.9 (1.1)	–0.1 (–1.8–1.5)
Juice, servings/d	0.8 (0.5)	0.0 (–1.6–1.5)	0.7 (0.4)	0.0 (–1.3–1.3)	0.7 (0.6)	–0.1 (–1.2–1.0)
Sweets, servings/d	1.3 (0.8)	0.0 (–2.4–2.3)	1.1 (0.5)	0.0 (–1.8–2.2)	1.1 (0.8)	–0.1 (–1.5–1.4)
Processed meats, servings/d	0.4 (0.2)	0.0 (–0.8–0.6)	0.3 (0.1)	0.0 (–0.6–0.4)	0.2 (0.2)	0.0 (–0.4–0.4)
Non-processed meats, servings/d	0.6 (0.3)	0.0 (–0.9–0.7)	0.6 (0.2)	0.0 (–0.7–0.6)	0.5 (0.3)	0.0 (–0.5–0.5)
Trans fat, %	1.3 (0.3)	0.0 (–0.7–1.3)	1.7 (0.3)	–0.2 (–1.1–0.9)	1.6 (0.5)	–0.2 (–1.0–0.5)

*Mean (SD) or median (interquartile range).

†Mean (1st–99th centile range).

Table 2 | Four year weight change (lbs, 95% CI) associated with increased daily flavonoid intake (1 SD/day)

Flavonoid	Health Professionals Follow-up Study (n=20525)		Nurses' Health Study (n=39 423)		Nurses' Health Study II (n=64 138)		Pooled* (n=124 086)	
	Model 1	Model 2	Model 1	Model 2	Model 1	Model 2	Model 1	Model 2
Flavonols (SD=7 mg)	-0.16 (-0.21 to -0.12)†	-0.16 (-0.21 to -0.11)†	-0.07 (-0.10 to -0.04)	-0.08 (-0.11 to -0.04)†	-0.26 (-0.30 to -0.24)†	-0.23 (-0.26 to -0.20)†	-0.16 (-0.26 to -0.06)†	-0.16 (-0.26 to -0.06)†
Flavones (SD=0.9 mg)	0.00 (-0.03 to 0.04)	0.00 (-0.04 to 0.03)	-0.03 (-0.04 to 0.03)	-0.01 (-0.04 to 0.02)	-0.02 (-0.05 to 0.02)†	0.00 (-0.03 to -0.03)	0.00 (-0.02 to 0.01)	0.00 (-0.02 to 0.01)
Flavanones (SD=25 mg)	0.01 (-0.04 to 0.05)	0.02 (-0.02 to 0.07)	0.11 (0.07 to 0.15)†	0.12 (0.08 to 0.16)†	0.25 (0.21 to 0.30)†	0.18 (0.14 to 0.22)†	0.11 (0.02 to 0.20)†	0.11 (0.02 to 0.20)†
Flavan-3-ols (SD=45 mg)	-0.13 (-0.20 to -0.06)†	-0.12 (-0.19 to -0.05)†	-0.02 (-0.07 to 0.02)	-0.03 (-0.07 to 0.02)	-0.20 (-0.24 to -0.17)†	-0.14 (-0.18 to -0.10)†	-0.10 (-0.17 to -0.02)†	-0.10 (-0.17 to -0.02)†
Anthocyanins (SD=10 mg)	-0.20 (-0.25 to -0.17)†	-0.20 (-0.24 to -0.16)†	-0.21 (-0.24 to -0.18)†	-0.18 (-0.21 to -0.15)†	-0.37 (-0.39 to -0.35)†	-0.29 (-0.32 to -0.27)†	-0.23 (-0.30 to -0.15)†	-0.23 (-0.30 to -0.15)†
Polymers (SD=138 mg)	-0.23 (-0.30 to -0.17)†	-0.19 (-0.25 to -0.12)†	-0.11 (-0.15 to -0.06)†	-0.10 (-0.15 to -0.06)†	-0.34 (-0.38 to -0.30)†	-0.26 (-0.30 to -0.22)†	-0.18 (-0.28 to -0.08)†	-0.18 (-0.28 to -0.08)†
Proanthocyanidins (SD=72 mg)	-0.45 (-0.52 to -0.38)†	-0.43 (-0.50 to -0.35)†	-0.51 (-0.57 to -0.45)†	-0.55 (-0.62 to -0.49)†	-0.85 (-0.90 to -0.80)†	-0.80 (-0.85 to -0.75)†	-0.60 (-0.82 to -0.37)	-0.60 (-0.82 to -0.37)
Total flavonoids (SD=194 mg)	-0.22 (-0.29 to -0.16)†	-0.22 (-0.28 to -0.15)†	-0.09 (-0.14 to -0.05)†	-0.11 (-0.16 to -0.07)†	-0.33 (-0.37 to -0.29)†	-0.28 (-0.32 to -0.24)†	-0.20 (-0.31 to -0.09)†	-0.20 (-0.31 to -0.09)†

Model 1=adjusted for age.

Model 2=adjusted for age and body mass index at baseline and change in smoking status, physical activity, hours of sitting or watching TV, hours of sleep, fried potatoes, juice (non-citrus for flavanones and flavanones), whole grains, refined grains, fried foods, nuts, whole fat dairy, low fat dairy, sugar sweetened drinks, diet drinks, sweets, processed meats, non-processed meats, trans fat, alcohol, seafood, and caffeine.

*Meta-analysis with random effects.

†95% CI excludes zero (null).

Increased intake of most flavonoid subclasses, with the exception of flavones and flavanones, was inversely associated with weight gain (table 2). The strongest magnitude of association was seen for anthocyanins, flavonoid polymers (including proanthocyanidins alone), and flavonols; each greater standard deviation of daily intake was associated with 0.16 to 0.23 lbs (0.07-0.10 kg) less weight gained over four year intervals. Eighty three per cent of flavanone and 44% of flavone intake was derived from citrus fruits and juices. When we adjusted intakes of flavones and flavanones for total juice intake rather than non-citrus juice, an important contributor to weight gain in these cohorts,³³ the observed associations between intake of these two subclasses and weight gain were stronger, inverse, and statistically significant (table 3, model 4 v model 5).

These associations remained significant for anthocyanins and flavonoid polymers (including proanthocyanidins alone) when we further adjusted for change in total fiber intake but not for flavonols, flavanones, and flavan-3-ols. In these models, increased fiber intake was inversely associated with weight change in all three cohorts. When we excluded thearubigins from total flavonoid polymers, the association with weight change was stronger in fully adjusted models: -0.61 (95% confidence interval -0.83 to -0.38) lbs per increased SD (76 mg).

Models with and without adjustment for change in other aspects of diet were very similar (table 3, model 2 v model 3). All associations were virtually identical when we additionally included change in total energy intake as a covariate in our models. In addition, models that included obese participants (supplemental table C) and models that used a complete case analysis in which missing values were not carried forward (supplemental table D) showed very similar results.

In these cohorts, the main food sources of anthocyanins were blueberries and strawberries, while flavan-3-ols and their oligomers (polymers) were mainly derived from tea and apples. As noted above, orange juice and oranges were the main sources of flavanone and flavone intakes, and tea and onions were the main sources of flavonols in the habitual diet (figure 1). Dietary sources of the different flavonoid subclasses were similar across cohorts, with the exception of beer providing flavan-3-ols in the HPFS, peppers providing flavones in the NHS, and red wine providing flavones in the NHS II and HPFS. The food frequency questionnaire did not include all fruits and vegetables, and some known rich sources of flavonoids, such as blackberries and raspberries, were not specifically included.

Discussion

We found that increased consumption of most flavonoid subclasses was associated with less weight gain among men and women aged 27-65 followed for up to 24 years. The greatest magnitude of association was observed for anthocyanins, flavonoid polymers, and flavonols. These associations remained statistically significant for anthocyanins and flavonoid polymers (including proanthocyanidins alone) even after further adjustment for fiber intake, which was inversely

Table 3 | Four year weight change (lbs, 95% CI) associated with increased daily flavonoid intake (1 SD/day); pooled* associations among 124,086 men and women

Flavonoid	Model 1	Model 2	Model 3	Model 4	Model 5	Model 6	Model 7	Model 8	Model 9
Flavonols (SD=7 mg)	-0.17 (-0.29 to -0.04)†	-0.17 (-0.29 to -0.05)†	-0.14 (-0.23 to -0.04)†	-0.16 (-0.26 to -0.06)†	-0.16 (-0.26 to -0.06)†	-0.04 (-0.10 to 0.02)	-0.15 (-0.26 to -0.03)†	-0.15 (-0.30 to 0.00)	-0.09 (-0.19 to 0.02)
Flavones (SD=0.9 mg)	-0.01 (-0.02 to 0.01)	-0.02 (-0.04 to -0.01)†	0.00 (-0.02 to 0.02)	-0.09 (-0.13 to -0.05)†	0.00 (-0.02 to 0.01)	0.09 (0.04 to 0.14)†	0.01 (-0.01 to 0.03)	0.00 (-0.04 to 0.05)	-0.03 (-0.06 to 0.00)
Flavanones (SD=25 mg)	0.13 (-0.01 to 0.26)	0.12 (-0.01 to 0.26)	0.11 (0.02 to 0.21)†	-0.05 (-0.09 to -0.01)†	0.11 (0.02 to 0.20)†	0.20 (0.08 to 0.32)†	0.13 (0.04 to 0.22)†	0.08 (-0.08 to 0.23)	0.08 (-0.02 to 0.17)
Flavan-3-ols (SD=45 mg)	-0.12 (-0.24 to 0.00)	-0.12 (-0.23 to 0.00)	-0.09 (-0.16 to -0.01)†	-0.10 (-0.17 to -0.02)†	-0.10 (-0.17 to -0.02)†	-0.05 (-0.11 to 0.01)	-0.09 (-0.17 to -0.01)†	-0.07 (-0.23 to 0.09)	-0.03 (-0.10 to 0.04)
Anthocyanins (SD=10 mg)	-0.26 (-0.38 to -0.14)†	-0.28 (-0.38 to -0.17)†	-0.22 (-0.30 to -0.15)†	-0.23 (-0.30 to -0.15)†	-0.23 (-0.30 to -0.15)†	-0.17 (-0.22 to -0.11)†	-0.22 (-0.30 to -0.14)†	-0.24 (-0.40 to -0.09)†	-0.21 (-0.26 to -0.15)†
Polymers (SD=138 mg)	-0.23 (-0.37 to -0.08)†	-0.22 (-0.37 to -0.07)†	-0.17 (-0.27 to -0.07)†	-0.18 (-0.28 to -0.08)†	-0.18 (-0.28 to -0.08)†	-0.10 (-0.16 to -0.03)†	-0.17 (-0.28 to -0.06)†	-0.16 (-0.38 to 0.05)	-0.11 (-0.22 to -0.01)†
Proanthocyanidins (SD=72 mg)	-0.60 (-0.87 to -0.34)†	-0.63 (-0.87 to -0.39)†	-0.55 (-0.75 to -0.35)†	-0.60 (-0.82 to -0.37)†	-0.60 (-0.82 to -0.37)†	-0.43 (-0.61 to -0.26)†	-0.61 (-0.89 to -0.34)†	-0.56 (-0.91 to -0.21)†	-0.52 (-0.72 to -0.36)†
Total flavonoids (SD=194 mg)	-0.22 (-0.37 to -0.06)†	-0.21 (-0.36 to -0.06)†	-0.17 (-0.27 to -0.06)†	-0.20 (-0.31 to -0.09)†	-0.20 (-0.31 to -0.09)†	-0.10 (-0.18 to -0.03)†	-0.19 (-0.31 to -0.07)†	-0.16 (-0.37 to 0.04)	-0.12 (-0.24 to 0.00)

Model 1=adjusted for age.

Model 2=model 1+baseline body mass index and change in smoking status, physical activity, hours of sitting or watching TV, and hours of sleep.

Model 3=model 2+change in fried potatoes, whole grains, refined grains, fried foods, nuts, whole fat dairy, low fat dairy, sugar sweetened drinks, sweets, processed meats, non-processed meats, trans fat, alcohol, seafood, and caffeine.

Model 4=model 3+change in juice.

Model 5=model 4+change in juice (non-citrus for flavones and flavanones).

Model 6=model 5+change in fiber.

Model 7=model 6+change in total energy.

Model 8=model 7+change in baseline age, body mass index, smoking status, physical activity, hours of sitting or watching TV, hours of sleep, fried potatoes, whole grains, refined grains, fried foods, nuts, whole fat dairy, low fat dairy, sugar sweetened drinks, sweets, processed meats, non-processed meats, trans fat, alcohol, seafood, and juice (non-citrus for flavones and flavanones).

Model 9=model 8+model 8.

*Meta-analysis with random effects.

†95% CI excludes zero (null).

associated with weight change in all three cohorts, indicating that high anthocyanin and flavonoid polymer food sources may be associated with less weight gain through mechanisms other than fiber content.

The magnitudes of these associations were small, less than a pound (<0.5 kg) per increased daily standard deviation, but a single serving per day of many fruits can often provide more than one standard deviation (supplemental table E). For example, one half cup of blueberries provides about 121 mg or about 12 standard deviations of anthocyanins. Furthermore, losing even small amounts of weight can improve health: losing just 11-22 lbs (5.0-10 kg) is associated with a decrease in blood pressure,^{16,18} and reducing body mass index by 1-3 kg/m² is associated with a 2-13% lower risk of cardiovascular disease events.¹⁹ Similarly, preventing even small amounts of weight gain could have an important public health impact: gaining 10 lbs (4.5 kg) or more between the ages of 40 and 60 increased the risk of developing diabetes by 40-70%,¹⁷ and a meta-analysis of 221 studies found a 24-59% increased risk of several cancers.²⁰

Comparison with other studies

In a cohort setting, Hughes et al found that women with the highest intake of three flavonoid subclasses, flavones, flavonols, and catechins (a flavan-3-ol), had a smaller increase in body mass index over 14 years than those with lower intakes.³⁴ These associations were not present in men, possibly owing to lower flavonoid intake and smaller increases in body mass index over time compared with women. In our study population, intake of flavonoids was only slightly lower among men than in women, and body mass index also increased more slowly over time among men; however, we detected inverse associations among both men and women. Similar to Hughes et al, however, we saw the strongest inverse associations among younger women (NHS II).

To date, human weight loss trials have examined only supplements containing flavan-3-ols and many have been too short to test meaningful differences in weight loss.^{35,36} Among 80 overweight people with diabetes, no difference was found in weight between the groups randomized to decaffeinated green tea (856 mg/d epigallocatechin gallate) versus placebo for one year.³⁷ On the other hand, weight and subcutaneous fat decreased among 102 overweight/obese people taking 8 g/d oolong tea for six weeks, but there was no placebo comparison group in this study.³⁶ Often, flavan-3-ol interventions are combined with other interventions such as physical activity, caffeine, or insulin, and their effects are hard to tease apart. Comparing results from trials that run for much shorter time periods or use supplements instead of whole foods is also difficult because long term higher intake of specific fruits and vegetables is generally most relevant to public health interventions.

Strengths and limitations of study

Our study has some limitations. Although our validated food frequency questionnaire specifies portion size, measurement error will be present with any assessment

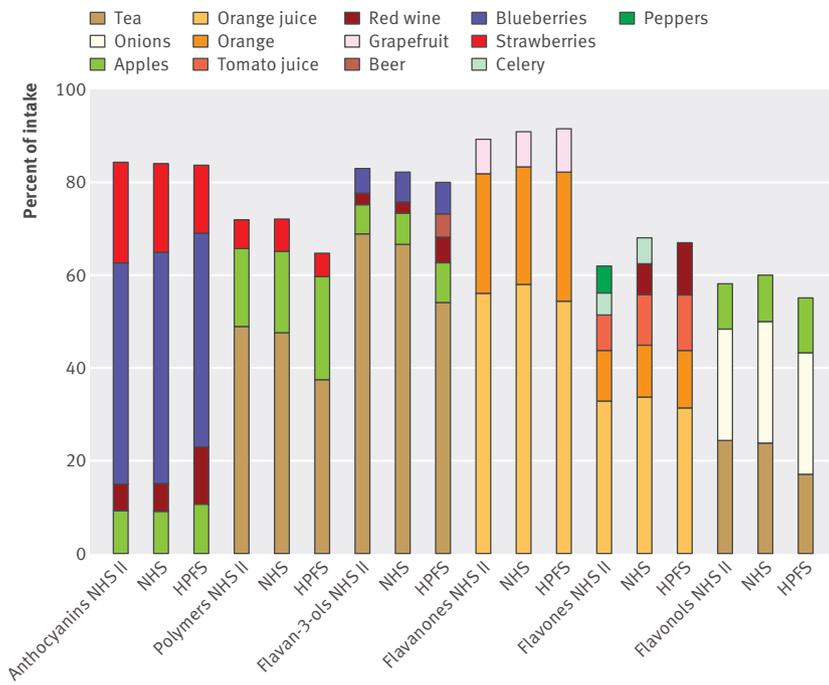


Fig 1 | Major dietary sources of flavonoids by subclass, averaged for 1986-2010 in Health Professionals Follow-up Study (HPFS) and Nurses' Health Study (NHS) and 1991-2007 in NHS II.

of diet and the flavonoid content of foods varies depending on ripeness, storage conditions, food processing, and season, which may introduce additional measurement error. As dietary thearubigin intakes are crudely measured, we did further analysis excluding thearubigins from the flavonoid polymer subclass and found a stronger inverse association with weight change (-0.61 lbs per increased daily SD over four years versus -0.18 lbs).

Furthermore, our food frequency questionnaire did not capture all food sources of flavonoids, but it did capture the main sources present in the habitual diet. Our results could underestimate true associations if people who gained weight in the beginning of a four year period modified their diet in response and began to eat more fruits and vegetables, and hence more flavonoids, in order to lose weight (reverse causality). On the other hand, our results could overestimate true associations if people who gained weight stopped eating fruits and vegetables. In addition, although we incorporated a comprehensive list of confounders, residual confounding is possible owing to imperfect measurement of confounders and potential unmeasured/unknown confounders. The source of confounding most difficult to control is other constituents of the foods that are high in flavonoids; in our analysis, control for fiber intake substantially attenuated the observed associations, except for associations with anthocyanins and flavonoid polymers. Finally, our results may be generalizable only to people of similar age, race, and socioeconomic status.

Strengths of our study include the measurement of all major flavonoid subclasses, the inclusions of more than 100 000 men and women, a wide age range, and up to 24 years of follow-up. Furthermore, each

participant completed a validated, semiquantitative food frequency questionnaire up to seven times during follow-up. We additionally had data on a comprehensive set of potential dietary and lifestyle confounders; this and the restriction of this population to participants with higher education and income reduced residual confounding due to socioeconomic status. Finally, our results were consistent across both sexes and different ages.

Conclusions and policy implications

These data may help to refine previous dietary recommendations for the prevention of obesity and its potential consequences. Our results suggest that choosing high flavonoid fruits and vegetables, such as apples, pears, berries, and peppers, may help with weight control. Most Americans consume less than one cup (less than two servings) of fruits and less than two cups of vegetables daily,³⁸ and juice and potatoes are major contributors to intake.³⁸ Beyond increasing intake to current recommendations of two cups of fruit and 2.5 cups of vegetables per day,³⁹ people may be able to maximize their health benefit by including optimal fruits and vegetables in their daily diets.

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Contributors: MLB, EBR, and AC conceived and designed the experiments. MLB, WCW, FBH, EBR, and AC conducted the experiments. MLB analyzed the data and wrote the first draft of the manuscript. KJM, WCW, FBH, EBR, and AC contributed to the writing of the manuscript. All authors agree with the manuscript's results and conclusions. MLB is the guarantor.

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Ethical approval: The study protocol was approved by the Institutional Review Board of the Brigham and Women's Hospital and by the Harvard T.H. Chan School of Public Health Human Subjects Committee Review Board (ID 2008P000327). All participants provided voluntary responses to mailed questionnaires that served as informed consent.

Transparency declaration: The lead author (the manuscript's guarantor) affirms that the manuscript is an honest, accurate, and transparent account of the study being reported; that no important aspects of the study have been omitted; and that any discrepancies from the study as planned (and, if relevant, registered) have been explained.

Data sharing: Health Professionals Follow-up Study, Nurses' Health Study, and Nurses' Health Study II data may be used in collaboration with a principal investigator. Please see the study websites for more information: www.hsph.harvard.edu/hpfs/hpfs_collaborators.htm and www.channing.harvard.edu/nhs/?page_id=52.

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Supplementary tables