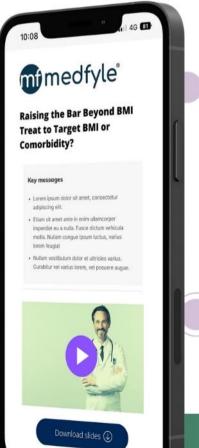


# Medfyle Conference Coverage from ObesityWeek® 2022

Discover a new way to catch up with the latest advances in obesity research and care presented at ObesityWeek® 2022.

- Medfyle summaries
- Expert interviews
- Expert presentations
- Posters

### **CLICK TO ACCESS NOW**



This activity is supported by an educational grant from Lilly

#### ORIGINAL ARTICLE

Epidemiology/Genetics

Revised: 3 November 2022

## Association between classes and subclasses of polyphenol intake and 5-year body weight changes in the EPIC-PANACEA study

Jazmin Castañeda <sup>1</sup> 💿   Mercedes Gil-Lespinard <sup>1</sup> 💿   Enrique Almanza-Aguilera <sup>1</sup> 💿
Fjorida Llaha <sup>1</sup> 💿   Jesús-Humberto Gómez <sup>2,3</sup> 💿   Nicola Bondonno <sup>4,5</sup> 💿
Anne Tjønneland <sup>4,6</sup> 💿   Kim Overvad <sup>7</sup> 💿   Verena Katzke <sup>8</sup> 💿
Matthias B. Schulze <sup>9,10</sup> 💿 📔 Giovanna Masala <sup>11</sup> 💿 📔 Claudia Agnoli <sup>12</sup> 💿 📔
Maria Santucci de Magistris 13   Rosario Tumino 14 💿   Carlotta Sacerdote 15 💿
Guri Skeie <sup>16</sup>   Magritt Brustad <sup>16,17</sup>   Cristina Lasheras <sup>18</sup>
Esther Molina-Montes <sup>3,19,20,21</sup> María-Dolores Chirlaque <sup>2,3</sup>
Aurelio Barricarte <sup>3,22,23</sup>   Emily Sonestedt <sup>24</sup>   Marisa da Silva <sup>25</sup>
Ingegerd Johansson <sup>26</sup> 💿   Johan Hultdin <sup>27</sup> 💿   Anne M. May <sup>28</sup>
Nita G. Forouhi <sup>29</sup> 💿   Alicia K. Heath <sup>30</sup> 💿   Heinz Freisling <sup>31</sup> 💿
Elisabete Weiderpass <sup>31</sup> 💿   Augustin Scalbert <sup>31</sup> 💿   Raul Zamora-Ros <sup>1,32</sup> 💿

#### Correspondence

Raul Zamora-Ros, Department of Nutrition, Food Sciences, and Gastronomy, Faculty of Pharmacy and Food Sciences, University of Barcelona, Avinguda Joan XXIII, 27-31, 08028 Barcelona, Spain. Email: rzamora@ub.edu and rzamora@ idibell.cat

#### Funding information

Consejo Nacional de Ciencia y Tecnología, Grant/Award Number: ID 693636; Instituto de Salud Carlos III, Grant/Award Numbers: CD20/ 00071, CPII20/00009, FI19/00185, PI18/ 00191

#### Abstract

**Objective:** The aim of this study was to evaluate the associations among the intake of total polyphenols, polyphenol classes, and polyphenol subclasses and body weight change over 5 years.

**Methods:** A total of 349,165 men and women aged 25 to 70 years were recruited in the Physical Activity, Nutrition, Alcohol, Cessation of Smoking, Eating Out of Home and Obesity (PANACEA) project of the European Prospective Investigation into Cancer and Nutrition (EPIC) cohort from nine European countries. Body weight was measured at baseline and at follow-up after a median time of 5 years. Polyphenol intake, including four main polyphenol classes and eighteen subclasses, was estimated using validated dietary questionnaires and Phenol-Explorer. Multilevel mixed linear regression models were used to estimate the associations.

**Results:** Participants gained, on average, 2.6 kg ( $\pm$ 5.0 kg) over 5 years. Total flavonoids intake was inversely associated with body weight change (-0.195 kg/5 years, 95% CI: -0.262 to -0.128). However, the intake of total polyphenols (0.205 kg/ 5 years, 95% CI: 0.138 to 0.272) and intake of hydroxycinnamic acids (0.324 kg/

Jazmin Castañeda and Mercedes Gil-Lespinard contributed equally to this study.

For affiliations, refer to page 10.

This is an open access article under the terms of the Creative Commons Attribution-NonCommercial License, which permits use, distribution and reproduction in any medium, provided the original work is properly cited and is not used for commercial purposes. © 2023 The Authors. *Obesity* published by Wiley Periodicals LLC on behalf of The Obesity Society. 1930739x, 0, Downloaded from https

5 years, 95% CI: 0.267 to 0.381) were positively associated with body weight gain. In analyses stratified by coffee consumption, hydroxycinnamic acid intake was positively associated with body weight gain in coffee consumers (0.379 kg/5 years, 95% CI: 0.319 to 0.440), but not in coffee nonconsumers (-0.179 kg/5 years, 95% CI: -0.490 to 0.133).

**Conclusions:** Higher intakes of flavonoids and their subclasses are inversely associated with a modest body weight change. Results regarding hydroxycinnamic acids in coffee consumers require further investigation.

#### INTRODUCTION

Overweight and obesity are defined as abnormal or excessive body fat accumulation. Obesity is one of the principal contributors to the global burden of chronic diseases, particularly cardiovascular disease, type 2 diabetes, and certain types of cancers [1]. The prevalence has increased rapidly: the World Health Organization reports that more than 2 billion adults have overweight and obesity worldwide, whereas, in Europe, nearly 60% of adults are classified as having overweight or obesity [2, 3]. Obesity results from a complex interaction of several factors such as diet, lifestyle, socioeconomics, genetics, and environment [4].

Even a moderate weight loss of 5% to 10% has been shown to lead to the significant improvement of several cardiometabolic parameters (e.g., triglycerides, blood pressure, waist circumference, insulin sensitivity,  $\beta$ -cell function) [5] and a lower risk of comorbidities, including cardiovascular disease [6], diabetes mellitus [6], hypertension [7], dyslipidemia [8], and overall mortality [9].

Polyphenols are bioactive phytochemicals found in plant foods and beverages such as fruit, vegetables, seeds, herbs, spices, whole grains, tea, coffee, and wine [10]. They comprise a large variety of chemical structures and they are divided into four main classes: flavonoids, phenolic acids, stilbenes, and lignans [10, 11]. Increasing preclinical and clinical evidence has suggested a role of polyphenols as antiobesity compounds. Indeed, several *in vitro* and *in vivo* studies have shown that polyphenols may stimulate thermogenesis and energy expenditure, inhibit adipocyte differentiation and growth, increase lipolysis, induce  $\beta$  oxidation, and decrease appetite [12, 13]. The antiobesity effects of polyphenols, particularly flavonoids, have also been supported by numerous clinical trials [11, 14].

In contrast, few observational epidemiological studies have examined the role of polyphenol intake in body weight control. The Supplementation with Antioxidant Vitamins and Minerals (Supplementation en Vitamines et Mineraux Antioxydants [SU.VI.MAX]) study observed that a high intake of total polyphenols was associated with lower waist circumference and body mass index (BMI) after 6 years of follow-up [15]. Similarly, an inverse association among total flavonoid intake and body weight, BMI, and waist circumference was observed in the National Health and Nutrition Examination Survey (NHANES) [16]. Likewise, results from the Prevención con Dieta Mediterránea (REDIMED) study indicated that total urinary polyphenol excretion was inversely associated with changes in body weight,

#### Study Importance

#### What is already known?

- Several experimental studies have reported that polyphenols can stimulate different mechanisms in body weight loss, such as thermogenesis, energy expenditure, and induced β oxidation, among others.
- Some clinical trials have described the antiobesity effects of pharmacological doses of some polyphenols.
- There is scarce epidemiological research investigating the associations between classes and subclasses of polyphenols and body weight change, especially with nonflavonoids.

#### What does this study add?

- This study provides evidence suggesting that the intake of flavonoids is associated with the maintenance of body weight in both men and women.
- Hydroxycinnamic acid intake from coffee is associated with an increase in body weight in coffee consumers, but not in coffee nonconsumers.

## How might these results change the direction of research?

- For a better understanding on the influence of polyphenols on body weight loss, future randomized controlled trials using combinations of polyphenols or plant extracts mimicking polyphenol-rich diets are needed.
- Further research evaluating the effect of hydroxycinnamic acids in body weight change is also warranted.

BMI, waist circumference, and waist to height ratio over 5 years [17]. Furthermore, the Health, Alcohol and Psychosocial factors in Eastern Europe (HAPPIEE) study reported an inverse association among high total polyphenol intake, particularly from stilbenes and lignans, and BMI and waist circumference [18]. However, these epidemiological studies have some limitations, including a limited number of classes and/or subclasses of polyphenols investigated and a low variability of polyphenol intake due to small geographic variations. Therefore, the present study aimed to examine the associations among intakes of total polyphenols and all polyphenol classes and subclasses estimated using the Phenol-Explorer database and body weight change over 5 years in the large, multicountry European Prospective Investigation into Cancer and Nutrition-Physical Activity, Nutrition, Alcohol, Cessation of Smoking, Eating Out of Home and Obesity (EPIC-PANACEA) cohort.

#### **METHODS**

#### Study population

EPIC is a prospective cohort study with 521,448 participants, aged from 25 to 70 years, recruited between 1992 and 2000 in 23 centers from 10 Western European countries: Denmark, France, Germany, Greece, Italy, the Netherlands, Norway, Spain, Sweden, and the UK. The EPIC Study was approved by the local ethics committees from the participating centers and the ethical review board of the International Agency for Research on Cancer. All participants signed an informed consent. Further details on the study design and methods have been described previously [19].

The EPIC-PANACEA study is a subcohort of EPIC with follow-up data of anthropometric measurements, and it was originally designed to investigate the determinants of obesity and weight change in EPIC countries. In the present study, we excluded pregnant women and participants with missing diet or lifestyle questionnaires, missing data on weight and height, or unreliable anthropometric values at baseline (n = 23,713). In addition, we excluded participants with missing data on weight at followup (n = 122,154) and those with unrealistic body weight changes (<-5 or >5 kg/y over several years) or implausible/unusual anthropometry at follow-up (BMI at follow-up < 16 kg/m<sup>2</sup>; n = 2288). More details regarding the EPIC-PANACEA study design have been described elsewhere [20]. Finally, participants from Greece (n = 24,128) did not provide data for the present study; therefore, they were also excluded. The final analysis included 93,435 men and 255,730 women.

#### Anthropometric measures and body weight change

Body weight in EPIC-PANACEA was measured at baseline and at follow-up. The time between the first and second measurements ranged from 2 years (Germany) to 11 years (Italy), with an overall median of 5 years. Standardized methods were used to take anthropometric measurements (body weight and height), except for the centers of France, Norway, and Oxford (UK), where participants self-reported their anthropometric values at baseline. In the followup, body weight was self-reported in all centers except in Cambridge (UK) and Doetinchem (the Netherlands), where weight was measured via standardized methods [20]. For the accuracy of self-reported anthropometric measurements (at baseline and followup), Oxford correction equations were used to predict measured

weight and to calculate the corrected weight change [21]. Our primary outcome was weight change in kilograms per 5 years, calculated as weight at follow-up minus weight at baseline divided by the follow-up time in years to obtain the annual weight change and then multiplied by 5 years.

#### Dietary assessment and other covariates

Habitual diet was recorded at baseline by validated country- or center-specific dietary questionnaires that captured food and beverage intakes over the previous 12 months [19]. In most centers, these were self-administered food frequency questionnaires, except for Ragusa (Italy), Naples (Italy), and Spain, where face-to-face interviews were conducted and meal-structured questionnaires were used. A combined method of a semiguantitative food frequency questionnaire and a 7-day record was used in the UK and Malmö (Sweden) [19]. Nutrients and total energy intakes were estimated using the standardized EPIC Nutrient Database [22]. Dietary polyphenol intake was estimated using the Phenol-Explorer database, which contains content values for 502 polyphenols in 452 foods and beverages, together with retention factors for cooked and processed food [23]. Dietary polyphenols were divided into four major classes: total flavonoids, phenolic acids, stilbenes, and lignans, plus a minority class of polyphenols and 18 subclasses, specified in Table 2; all classes and subclasses were then summed to calculate total polyphenol intake.

Moreover, validated questionnaires were used at baseline to collect data on tobacco use, education level, menstrual history, physical activity (inactive, moderately inactive, moderately active, active), and clinical data. In addition, information on smoking status (never, former, current) was also collected at follow-up.

#### Statistical analyses

Polyphenol intake was assessed as a categorical variable based on quintiles distributed throughout the entire EPIC-PANACEA study. In addition, linear trend tests were calculated assigning the median of each quintile as scores. Polyphenol intake was also analyzed as a continuous variable after log<sub>2</sub> transformation to reduce the skewness of intake distributions. Before log<sub>2</sub> transformation, zero values were replaced with half of the nonzero minimum of the polyphenol class or subclass. One-unit increase corresponded to the absolute body weight change (kilograms per 5 years) associated with doubling in intake.

Multilevel mixed linear regression models were used to estimate the association among total, classes, and subclasses of polyphenol intake and body weight change over 5 years, using the EPIC center as a random effect and polyphenol intake and relevant confounders as fixed effects. Missing values (3.5% for educational level, 1.5% for physical activity, 2.1% for education, 4.7% for smoking status) were classified into a separate category (unknown) and included in the models. BMI at baseline and follow-up time in years had a nonlinear association with weight change. Consequently, they were included in the models as restricted cubic splines with three knots (10th, 50th,

and 90th percentiles) according to Harrell [24]. We fitted multivariable models adjusting for potential confounders (as fixed effects) selected a priori. Model 1 was adjusted for sex (male and female), age at baseline (years), and BMI (kilograms per meters squared). In model 2, we also adjusted for lifestyle characteristics; follow-up time (years), alcohol consumption (grams per day), education level (none, primary education, technical or professional school, secondary school, higher education, and unknown), physical activity (inactive, moderately inactive, moderately active, active, and unknown), smoking status at follow-up (never, former, and current), and menopausal status (pre-, post-, and peri-surgically postmenopausal and unknown). Model 3 was further adjusted for variables related to energy: total energy intake (kilocalories per day) and plausibility of energy intake reporting (yes and no) [19]. In addition, for model 3, we replaced total energy (kilocalories per day) with the all-components model (adjusting for all individual components providing energy of the diet) [25]: however, the results remained similar to the previous model; therefore, we did not present them. Finally, model 4 was model 3 additionally adjusted for other dietary factors: vitamin C (milligrams per day) and fiber (grams per day) intake. Furthermore, polyphenol intake was included in the statistical models as energy density (2000 mg/kcal/d). Results from both methods were almost identical, and energy density results were not reported.

The main food source of phenolic acids is coffee, which plays a role in body weight change [26]. Therefore, we performed analyses separately for hydroxycinnamic acid (HCA) class intake in coffee consumers and nonconsumers. Also, because caffeine has been associated with body weight loss [26], we evaluated the associations between total coffee and the type of coffee (caffeinated vs. decaffeinated) and body weight in order to investigate the effect of caffeine in body weight and differentiate the relationships between HCAs and caffeine from coffee intake.

Interaction analyses were performed between classes and subclasses of polyphenol intake (continuous, milligrams per day) and sex, age (<50 and ≥50 years), menopause status (peri-, post-, and premenopause), physical activity (inactive, moderately inactive, moderately active, and active), smoking status at follow-up (never, former, and current smokers), and BMI at baseline (underweight, normal weight, overweight, and obesity) in relation to body weight change. P values for the interaction were calculated using the likelihood ratio test. Sex, menopause, smoking status, and BMI had a statistically significant interaction; therefore, models were further fitted separately for each category of the variables. Finally, to assess the robustness of the results, we conducted a sensitivity analysis excluding participants with chronic diseases at baseline (type 2 diabetes, cardiovascular disease, and/or cancer) and participants who either quit smoking or started smoking or had missing data on smoking during follow-up. All p values presented were considered statistically significant at p < 0.05. Statistical analyses were performed using SAS software version 9.3 (SAS Institute Inc., Cary, North Carolina).

#### RESULTS

Participants in the highest quintile of polyphenol intake were more likely to be men and older and they had, on average, a lower BMI and

a higher education, alcohol consumption, and total energy intake (Table 1). In addition, they were more likely to be more physically active and current smokers. Women with higher total polyphenol intakes were less likely to be premenopausal than those with lower intakes. In all quintiles of total polyphenol intake, phenolic acids were the main contributors, followed by flavonoids, whereas lignans and stilbenes were consumed in low amounts (between 1 and 3 mg/d).

On average, study participants gained 2.6 kg (±5.0 kg) per 5 years of body weight during the follow-up. Body weight changes over 5 years by quintile of total, classes, and subclasses of polyphenol intakes are shown in Table 2 and Supporting Information Table S1. Total polyphenol intakes were positively associated with body weight gain; participants in the highest intake quintile had a 0.205-kg (95% confidence interval [CI]: 0.138 to 0.272) greater 5-year weight gain compared with those in the lowest quintile after multivariable adjustments (model 4; Table 2). Analysis by polyphenol classes showed that higher intakes of flavonoids ( $\beta_{O5}$  vs.  $_{O1}$  –0.195 kg/5 years, 95% CI: -0.262 to -0.128) and stilbenes (only when modeled continuously,  $\beta$ log<sub>2</sub> -0.032 kg/5 years, 95% CI: -0.039 to -0.024) were inversely associated with body weight change. Similarly, intakes of individual subclasses of flavonoids, except for isoflavonoids, were statistically, significantly, and inversely associated with body weight change (Table 2). A body weight gain was observed comparing participants in the extreme quintiles of total phenolic acid intake ( $\beta$  <sub>Q5</sub> vs. <sub>Q1</sub> 0.328 kg/5 years, 95% CI: 0.268 to 0.386) and its subclass HCAs ( $\beta$  <sub>0.5</sub> vs. <sub>0.1</sub> 0.324 kg/5 years, 95% CI: 0.267 to 0.381). However, other subclasses of phenolic acids showed an inverse association with body weight change, such as hydroxybenzoic acid ( $\beta$  <sub>Q5</sub> vs. <sub>Q1</sub>= -0.244 kg/5 years, 95% CI: -0.317 to -0.170) and hydroxyphenylacetic acid ( $\beta$   $_{0.5}$ vs.  $_{0.1}$  = -0.204 kg/5 years, 95% CI: -0.275 to -0.132). Minor polyphenol classes such as tyrosols and hydroxycoumarins showed an inverse association with body weight change. Analyses by quintiles of classes, subclasses, and total polyphenol intake were supported by the results using the continuous variable after the log<sub>2</sub> transformation (Figure 1).

Women presented a slightly stronger positive association between total polyphenol intake and body weight gain compared with men (p value interaction < 0.001; Supporting Information Table S2). Total polyphenol and phenolic acid intakes were strongly associated with body weight gain among women in perimenopause (Supporting Information Table S3). For smoking status at follow-up, we found that total polyphenol and phenolic acid class intakes were more strongly associated with weight gain in former smokers, whereas the opposite occurred with total flavonoid intake (Supporting Information Table S4). For BMI, there was an inverse trend between total flavonoid intake and body weight change in participants with underweight, normal weight, and overweight, but a positive trend was detected among participants with obesity (Supporting Information Table S5).

Additional analyses for HCAs were performed by dividing the analysis by coffee consumption, in which we observed a positive association with weight gain in coffee consumers, but not in coffee nonconsumers (Supporting Information Table S6). Subsequently, we separately analyzed the association for total coffee and type of coffee



**TABLE 1** Baseline characteristics of the population according to quintiles of total polyphenol intake in the EPIC-PANACEA study (*n* = 349,165)

	Quintile of intake				
Category	Quintile 1 (n = 69,832)	Quintile 2 (n = 69,826)	Quintile 3 (n = 69,826)	Quintile 4 (n = 69,841)	Quintile 5 (n = 69,840)
Total polyphenol intake (mg/d)	558.2 (441.2-706.0)	854.8 (788.8-970.7)	1117.5 (1049.4-1248.4)	1427 (1342-1607.3)	1923.8 (1759.7-2856.3
Follow-up time (y)	5.1 ± 2.4	5.5 ± 2.7	5.3 ± 2.4	4.9 ± 2.0	4.6 ± 1.5
Weight change (kg/5 years) <sup>a</sup>	2.4 ± 5.2	2.5 ± 4.9	2.6 ± 5.0	2.7 ± 4.9	2.8 ± 5.0
Age (y)	50.4 ± 8.9	50.8 ± 9.2	51.7 ± 9.3	52.6 ± 9.2	52.9 ± 8.6
BMI (kg/m <sup>2</sup> )	25.6 ± 4.4	25.1 ± 4.1	25.0 ± 4.0	24.8 ± 3.9	24.7 ± 3.8
Alcohol use (g/d)	6.4 ± 12.2	9.5 ± 13.7	12.3 ± 15.6	14.3 ± 17.6	16.7 ± 20.8
Energy intake (kcal/d)	1777 ± 517	1964 ± 544	2092 ± 567	2182 ± 592	2366 ± 634
Women (%)	79.4	74.1	72.0	70.9	69.7
Education level (%)					
None	11.6	4.5	2.4	1.3	0.8
Primary school	28.4	26.9	22.8	19.6	18.3
Technical school	20.2	23.0	23.0	23.2	22.1
Secondary school	20.7	22.1	22.1	21.9	21.9
Higher education	17.5	26.2	26.2	28.3	31.0
Missing	1.4	3.2	3.2	5.4	5.7
Physical activity level (%)					
Inactive	23.6	19.3	17.3	16.8	16.0
Moderately inactive	32.6	35.0	34.9	34.6	33.6
Moderately active	29.5	27.7	26.1	26.1	26.9
Active	12.5	16.2	19.3	20.4	22.3
Missing	1.5	1.6	1.8	1.8	1.0
Smoking at follow-up (%)					
Never	54.1	47.8	47.6	47.8	44.2
Former	25.5	28.4	30.1	32.2	34.3
Current	15.7	17.2	16.8	16.7	20.0
Missing	4.5	6.4	5.3	3.1	1.3
Prevalent diseases at baseline	(%) <sup>b</sup>				
No	85.6	86.7	85.3	81	78.7
Yes	8.0	7.5	7.7	7.5	7.2
Missing	6.3	5.6	6.9	11.3	14.0
Menopause (%)					
Premenopausal	29.1	26.3	23.1	21.0	19.9
Postmenopausal	30.6	30.1	32.4	33.8	32.9
Perimenopausal	17.4	15.6	14.1	13.5	14.5
Surgery	2.1	2.0	2.3	2.4	2.3
Classes of polyphenol intake (					
Phenolic acids	249.7 (142.4-500.7)	462.0 (328.0-704.5)	585.6 (430.3-917.6)	708.0 (525.7-1215.6)	1025.7 (675.9-1918.3)
Flavonoids	224.8 (150.6-449.6)	340.7 (234.7-649.6)	468.2 (325.5-856.3)	647.8 (424.7-1132.2)	896.6 (548.0-1633.7)
Other polyphenol classes	32.2 (19.9-72.0)	40.1 (26.5-93.5)	45 (28.3-106.4)	48.8 (29.5-117.5)	57 (34.8-134.3)
					(Continu

(Continues)

#### **TABLE 1** (Continued)

	Quintile of intake				
Category	Quintile 1	Quintile 2	Quintile 3	Quintile 4	Quintile 5
	(n = 69,832)	(n = 69,826)	(n = 69,826)	(n = 69,841)	(n = 69,840)
Lignans	1.2	1.3	1.4	1.5	1.7
	(0.9-3.1)	(1-4.6.0)	(1.1-4.5)	(1.2-3.8)	(1.4-3.7)
Stilbenes	0.2	0.4	0.7	1.1	1.4
	(0.05-3.5)	(0.1-6.5)	(0.2-8.4)	(0.2-9.9)	(0.3-11.2)

*Notes*: Means ± SD are presented for continuous variables, and percentages are presented for categorical variables. Medians and percentiles (25th to 95th percentile) are presented for polyphenol intake.

Abbreviations: EPIC, European Prospective Investigation into Cancer and Nutrition; PANACEA, Physical Activity, Nutrition, Alcohol, Cessation of Smoking, Eating Out of Home and Obesity.

<sup>a</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>b</sup>Type 2 diabetes, cardiovascular disease, and cancer.

(caffeinated vs. decaffeinated) as exposure variables with body weight changes. Decaffeinated coffee intake was associated with a slightly greater body weight gain than caffeinated coffee (when modeled continuously; Supporting Information Table S6).

In the sensitivity analysis, after excluding participants with chronic disease at baseline (n = 57,617) and participants who quit smoking or started smoking during follow-up or with missing values (n = 35,489), we observed that the associations among total polyphenol intake and polyphenol classes and body weight change were similar to our main results (Supporting Information Tables S7 and S8).

#### DISCUSSION

In this large prospective study, a mean body weight gain of 2.6 kg during the 5 years of follow-up was observed. Progressive age-related weight gain in adulthood is a well-observed phenomenon in many nonobese populations such as in the Nurses' Health Study II cohort, which showed a weight change of 4.4 lb (2.0 kg) per 4.4 years of follow-up [27], and in NHANES, the weight change of which was 2.5 kg per 9.8 years of follow-up [28]. In the current study, a positive association among total polyphenol and phenolic acid intakes and body weight gain was observed. Conversely, higher intakes of flavonoids, including anthocyanins, dihydrochalcones, and dihydroflavonols, and other minor polyphenol classes such as tyrosols and stilbenes were inversely associated with body weight change, supporting the evidence from experimental studies that some polyphenol classes and subclasses may play a role in obesity prevention; several mechanisms have been proposed, such as activation of  $\beta$  oxidation processes, stimulation of energy expenditure, and inhibition of adipocyte differentiation [12]. Recently, it has been discussed that polyphenols may modulate type 2 taste receptors responsible for bitter taste recognition, which may play a role in energy/body weight homeostasis [29]. Similar to our findings, a cohort study by Adriouch and colleagues reported that high intakes of flavanones, flavones, and lignans were associated with lower waist circumference and lower BMI. In their study, total polyphenol and phenolic acid intakes were associated with both a lower body weight gain and a lower increase in

adiposity over 6 years [15]. Our study showed that total polyphenols were positively related to body weight gain; however, after excluding phenolic acids, we observed an inverse association with body weight change. Such changes were driven by phenolic acids (specifically HCAs), the main contributors to total polyphenols; therefore, the results with total polyphenols need to be interpreted with caution.

Flavonoids are the most-studied polyphenol class in relation to their effects on body weight [13]. In our study, we observed that a higher intake of flavonoids, particularly anthocyanins, flavan-3-ol monomers, theaflavins, flavones, and flavonols, was strongly and inversely associated with body weight change over 5 years. Similarly, in a large prospective cohort, statistically significant inverse associations among subclasses of anthocyanins, flavanols (including proanthocyanidins), and flavonols and body weight change were observed after a 24-year follow-up [27]. Another cohort from the Netherlands observed an association among a higher intake of flavonol/flavone and catechin and a lower increase in BMI in women, but not in men [30]. Similarly, the Mediterranean healthy Eating, Aging and Lifestyle (MEAL) cohort study, with a follow-up over 14 years, reported that women with a high intake of total flavonoids were less likely to have obesity and that flavonol intake was inversely associated with obesity [31]. Although the magnitudes of body weight loss were small, they may contribute to body weight maintenance, which has been reported as a protective factor for diseases such as type 2 diabetes, hypertension, and cardiovascular disease [32]. Some trials have also investigated these body weight effects showing, in general, a body weight reduction after the intervention with supplements rich in polyphenols [11, 14].

Contrary to our expectations, this study found an inverse association among the minor subclasses of phenolic acids (i.e., hydroxybenzoic acids and hydroxyphenylacetic acids) and body weight change, whereas total phenolic acid and HCA intakes were associated with an increase in body weight. Their main food source is coffee [10] and, because other compounds of coffee such as caffeine, trigonelline, and magnesium may possess antiobesogenic properties [33], after we stratified by coffee consumption, non-coffee consumers showed a null association between HCA intake and body weight change, whereas, in coffee consumers, phenolic acid intake

nenol classes, and subclasses (milligrams per day) and body weight change over 5 years in the EPIC-PANACEA study	
<b>BLE 2</b> Associations between intakes of total polyphenols, polyphenol clast	: 349,165)
TAB	0 = U

HENC	DL CL	ASSI	ES AI	ND V	VEIG	GHT	СНА	NGE	IN E	PIC-	PAN	ACE	A					_	O A Res	bearch	es	sit	ty	K		HE BESI OCIE	TY TY	.W	/11	LE	ΞY	
	p value for trend		<0.001		<0.001		<0.001		<0.001		0.081		<0.001		<0.001		<0.001		<0.001		<0.001		<0.001		<0.001		0.004		<0.001		<0.001	(Continues)
	Quintile 5, β (95% CI)	>1630.5	0.205 (0.138 to 0.272)	>786.7	-0.195 (-0.262 to -0.128)	>595.7	-0.141 (-0.207 to -0.075)	>246.9	-0.200 (-0.265 to -0.134)	>345.9	-0.071 (-0.136 to -0.005)	>59.1	-0.170 (-0.233 to -0.106)	>63.5	-0.178 (-0.246 to -0.109)	>64.7	-0.148 (-0.214 to -0.082)	>64.7	-0.270 (-0.336 to -0.204)	>15.3	-0.201 (-0.269 to -0.133)	>3.3	-0.187 (-0.240 to -0.134)	>3.4	-0.258 (-0.335 to -0.181)	>0.10	0.058 (-0.044 to 0.160)	>881.2	0.328 (0.268 to 0.386)	>838.2	0.324 (0.267 to 0.381)	
	Quintile 4, $\beta$ (95% CI)	1263.2-1630.5	0.095 (0.034 to 0.157)	525.4-786.7	-0.183 (-0.244 to -0.122)	369.8-595.7	-0.100 (-0.16 to -0.041)	69.8-246.9	-0.114 (-0.175 to -0.053)	243.3-345.9	-0.124 (-0.183 to -0.065)	24.9-59.1	-0.197 (-0.261 to -0.134)	36.4- 63.5	-0.118 (-0.181 to -0.055)	35.8-64.7	-0.159 (-0.216 to -0.102)	32.1-64.7	-0.167 (-0.223 to -0.111)	10.6-15.3	-0.200 (-0.258 to -0.141)	1.9-3.3	-0.124 (-0.172 to -0.075)	1.1-3.4	-0.142 (-0.211 to -0.072)	0.04-0.10	0.001 (-0.095 to 0.097)	619.1-881.2	0.102 (0.047 to 0.156)	570.7-838.2	0.094 (0.04 to 0.147)	
	Quintile 3, ß (95% CI)	983.6-1263.1	0.011 (-0.046 to 0.069)	359.5-525.3	-0.136 (-0.194 to -0.079)	238.5-369.7	-0.072 (-0.128 to -0.015)	30.7-69.7	-0.069 (-0.127 to -0.01)	175.8-243.2	-0.086 (-0.141 to -0.030)	3.5-24.8	-0.068 (-0.125 to -0.011)	23.7-36.3	-0.107 (-0.166 to -0.047)	20.3-35.7	-0.104 (-0.158 to -0.051)	17.1-32.0	-0.065 (-0.118 to -0.012)	7.6-10.5	-0.143 (-0.198 to -0.088)	0.8-1.8	-0.085 (-0.133 to -0.037)	0.16-1.0	-0.025 (-0.092 to 0.043)	>0.01-0.03	-0.038 (-0.134 to 0.057)	458.9-618.9	0.006 (-0.047 to 0.059)	416.3-570.6	-0.015 (-0.067 to 0.038)	
	Quintile 2, β (95% CI)	720.1-983.5	0.009 (-0.044 to 0.063)	231.6-359.4	-0.069 (-0.122 to -0.016)	142.8-238.4	-0.023 (-0.076 to 0.03)	16.0-30.6	-0.044 (-0.098 to 0.009)	113.5-175.7	-0.060 (-0.112 to -0.007)	<3.5	-0.011 (-0.071 to 0.050)	14.3-23.6	-0.070 (-0.124 to -0.015)	10.8-20.2	-0.087 (-0.139 to -0.036)	7.7-17.0	-0.079 (-0.131 to -0.028)	4.9-7.5	-0.125 (-0.177 to -0.073)	<0.8	-0.048 (-0.182 to 0.086)	<0.16	0.050 (-0.018 to 0.119)	>0-0.01	-0.019 (-0.115 to 0.078)	298.3-458.8	-0.048 (-0.100 to 0.004)	240.6-416.2	-0.063 (-0.115 to -0.012)	
Quintile of intake	Quintile 1, $\beta$ (95% CI)	<720.1	0 (ref)	<231.6	0 (ref)	<142.8	0 (ref)	<16.0	0 (ref)	<113.5	0 (ref)	0	0 (ref)	<14.3	0 (ref)	<10.8	0 (ref)	<7.7	0 (ref)	<4.9	0 (ref)	0	0 (ref)	0	0 (ref)	0	0 (ref)	<298.3	0 (ref)	<240.6	0 (ref)	
	Polyphenol (mg/d)	Total polyphenols	Model 4	Flavonoids	Model 4	Total flavanols	Model 4	Flavan-3-ol monomers	Model 4	Proanthocyanidins	Model 4	Theaflavins	Model 4	Flavonols	Model 4	Anthocyanins	Model 4	Flavanones	Model 4	Flavones	Model 4	Dihydrochalcones	Model 4	Dihydroflavonols	Model 4	Isoflavonoids	Model 4	Total phenolic acids	Model 4	Hydroxycinnamic	Model 4	

7

(Continued)	
~ ~	1
4	ļ
2	ב נ
H	

	Quintile of intake					
Polyphenol (mg/d)	Quintile 1, $eta$ (95% CI)	Quintile 2, ß (95% CI)	Quintile 3, ß (95% CI)	Quintile 4, ß (95% CI)	Quintile 5, β (95% Cl)	p value for trend
Hydroxybenzoic	<5.4	5.4-14.9	15.0-33.0	33.1-83.5	>83.5	
Model 4	0 (ref)	-0.083 (-0.141 to -0.025)	-0.104 (-0.171 to -0.037)	-0.147 (-0.218 to -0.077)	-0.244 (-0.317 to -0.170)	<0.001
Hydroxyphenyl acetic	<0.01	0.01-0.05	0.06-0.14	0.15-0.32	>0.32	
Model 4	0 (ref)	-0.037 (-0.088 to 0.014)	-0.176 (-0.228 to -0.124)	-0.219 (-0.274 to -0.163)	-0.204 (-0.275 to -0.132)	<0.001
Lignans	<1.0	1.0-1.2	1.3-1.6	1.7-2.1	>2.1	
Model 4	0 (ref)	-0.0003 (-0.054 to 0.053)	-0.070 (-0.128 to -0.012)	-0.010 (-0.077 to 0.056)	0.093 (0.013 to 0.174)	<0.001
Stilbenes	<0.09	0.09-0.33	0.34-1.1	1.2-2.6	>2.6	
Model 4	0 (ref)	-0.082 (-0.495 to 0.331)	-0.120 (-0.533 to 0.294)	-0.238 (-0.652 to 0.175)	-0.373 (-0.788 to 0.042)	<0.001
Other polyphenol classes	<23.6	23.6-36.3	36.4-50.8	50.9-70.9	>70.9	
Model 4	0 (ref)	0.053 (0.0004 to 0.106)	-0.003 (-0.059 to 0.053)	-0.074 (-0.134 to -0.013)	-0.073 (-0.145 to -0.002)	0.002
Alkylphenols	<5.3	5.3-19.7	19.8-35.2	>35.3-54.7	>54.7	
Model 4	0 (ref)	0.095 (0.037 to 0.152)	-0.005 (-0.071 to 0.060)	-0.112 (-0.183 to -0.041)	-0.167 (-0.247 to -0.086)	<0.001
Hydroxycoumarins	0	>0-0.01	>0.01-0.57	0.58-0.18	>0.18	
Model 4	0 (ref)	-0.052 (-0.134 to 0.030)	-0.083 (-0.172 to 0.006)	-0.214 (-0.303 to -0.124)	-0.270 (-0.367 to -0.174)	<0.001
Tyrosol	<1.1	1.1-2.5	2.6-5.1	5.2-11.4	>11.4	
Model 4	0 (ref)	-0.126 (-0.178 to -0.075)	-0.222 (-0.276 to -0.168)	-0.331 (-0.393 to -0.268)	-0.343 (-0.422 to -0.264)	<0.001
Alkymethoxyphenols	<0.9	0.9-1.9	2.0-2.8	2.9-4.2	>4.2	
Model 4	0 (ref)	-0.146 (-0.356 to 0.064)	-0.201 (-0.412 to 0.011)	-0.071 (-0.284 to 0.141)	0.135 (-0.079 to 0.348)	<0.001
Note: Overall mean 5-year we	ight gain corresponded to 2	Note: Overall mean 5-year weight gain corresponded to 2.6 kg (SD 5.0), and negative $\beta$ values indicate less weight gain (kilograms) over the same period.	ues indicate less weight gain (kilo	grams) over the same period.		

■\_\_\_WILEY\_ Obesity ●

Model 4: generalized linear mixed models with random effect on the intercept and slope according to center and adjusted for age, sex, and BMI at baseline (three-knot restricted cubic spline), follow-up time in years (three-knot restricted cubic spline), educational level, smoking status, physical activity, alcohol consumption, menopausal status, energy intake, plausibility of dietary energy reporting, vitamin C, and total fiber intakes.

Abbreviations: EPIC, European Prospective Investigation into Cancer and Nutrition; PANACEA, Physical Activity, Nutrition, Alcohol, Cessation of Smoking, Eating Out of Home and Obesity; ref, reference.

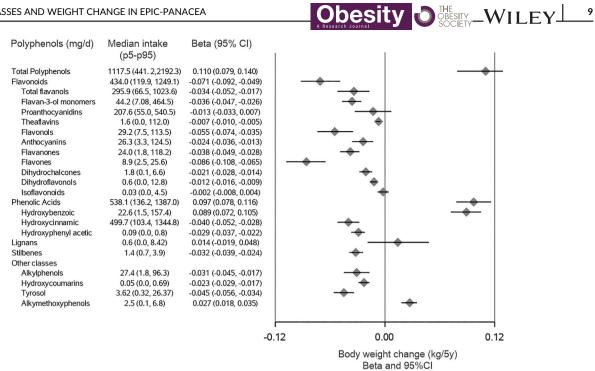


FIGURE 1 The association between intakes of total polyphenols, polyphenol classes, and subclasses (milligrams per day) as continuous variables (after log<sub>2</sub> transformation) and body weight change (kilograms) over 5 years in the EPIC-PANACEA study. Model 4: generalized linear mixed models with random effect on the intercept and slope according to center and adjusted for age, sex, and BMI at baseline (three-knot restricted cubic spline), follow-up time in years (three-knot restricted cubic spline), educational level, smoking status, physical activity, alcohol consumption, menopausal status, total energy intake, plausibility of dietary energy reporting, vitamin C, and total fiber intakes

was positively associated with body weight gain. Some in vivo studies have suggested potential mechanisms of HCAs from coffee in weight loss through regulation of lipid and glucose metabolism, e.g., via sterol regulatory element binding transcription factor 1 (SREBP-1C), peroxisome proliferator activated receptor  $\alpha$  (PPAR- $\alpha$ ), increased fatty acid oxidation, and increased insulin secretion; however, there is not information reported about possible mechanisms of HCAs associated with body weight gain [34]. The direct role of coffee consumption on body composition has been studied in previous cohorts; however, the results are still inconclusive. For instance, two previous cohorts have concluded that higher coffee consumption was associated with lower weight gain, BMI, and waist circumference [26, 35], whereas consumption of more than six cups of coffee per day was associated with higher BMI compared with the group consuming less than two cups per day among Swedish women [35]. Similarly, other studies conducted in Asia have reported that higher coffee consumption (>3 cups/d) was associated with higher risk of obesity compared with lower coffee consumption [36, 37]. However, these results need to be taken with caution, as they considered instant coffee blends that contain sugar and cream. In this sense, the method of coffee preparation, the types of coffee varieties, roasting degree, the size of the serving, and use of milk or cream and sugar added to coffee are factors that may influence the coffee-obesity relationship [38]. For example, the types of coffee have been evaluated (caffeinated vs. decaffeinated), and it was found that decaffeinated coffee was associated with higher body weight gain compared with caffeinated coffee [39]. In our study,

we observed an increase in body weight for both types of coffee; however, it is important to highlight that the consumption of caffeinated coffee was much higher (mean = 222.1 mL/d) compared with decaffeinated coffee (mean = 32.3 mL/d). Therefore, phenolic acids and caffeine seem to not have harmful effects on body weight, but it is unclear which potential coffee compounds, if any, might have an obesogenic effect.

The current study found an inverse association of stilbenes with body weight change, which is consistent with that reported in the HAPIEE study for change in waist circumference [18]. However, their results were not statistically significant. Furthermore, it has been previously observed that consumption of stilbenes from berries, red grapes, and wine has antiadipogenic and antilipogenic effects, improving changes in BMI and waist circumference [12, 40]. Future studies on the current findings are still warranted.

Our results also showed that tyrosols are inversely associated with body weight change. The main food sources of tyrosols include olives and olive oil, of which the principal component is hydroxytyrosol [10]. It has been largely studied for its protective effects on lowdensity lipoprotein oxidation and reduction of oxidative stress [41], and it may be responsible for the antiobesogenic properties of olive oil [41]. Moreover, extra-virgin olive oil is the main source of dressing and cooking fats in the Mediterranean diet, which has been reported to have a protective effect against weight gain, particularly in younger people [42]. The PREDIMED study evaluated the long-term effects of an unrestricted-calorie Mediterranean diet rich in extra-virgin olive oil on adiposity measures and observed a small reduction in body weight in participants given the Mediterranean diet interventions compared with the control groups [23]. However, more clinical and epidemiological evidence is needed in order to clarify the effect of tyrosols on body weight.

Results of the interaction analyses should be interpreted with caution because even very small body weight differences among subgroups are statistically significant because of our large sample size. We observed that women with perimenopause had a slightly greater weight gain associated with total polyphenol intake. Several polyphenols have estrogenic effects [12], and they would help to prevent body weight gain related to menopause. However, our results pointed out the opposite, probably because of reverse causality. Women with perimenopause might improve their diets and lifestyles [43], but, when they become menopausal, they tend to return to their regular habits [43]. Moreover, analyses by smoking status at follow-up showed that current smokers benefitted more from higher intakes of total polyphenols and phenolic acids, probably because of their ability to reduce tobacco oxidative stress [44]. Finally, interactions by BMI showed that flavonoid intakes were associated with a lower body weight in all groups, except those with obesity. Our hypothesis was that participants with obesity would benefit more from the intake of potential antiobesity compounds [12]. However, our contradictory results could be because participants with obesity tend to underestimate unhealthy foods and overestimate healthy foods more than those without obesity [45].

Strengths of our study include the multicenter prospective design, two measurements of body weight (to calculate body weight change), and a large sample size, which provided sufficient statistical power to perform multiple subgroup and sensitivity analyses. Another strength is the use of Phenol-Explorer, the most comprehensive database on polyphenol content in foods, to measure polyphenol intake. However, our study also had some limitations. First, weight was self-reported at follow-up in most centers; however, we improved the accuracy of these data by using a prediction equation, and the results in the two EPIC centers with measured weight (Cambridge and Doetinchem) were consistent with the rest of the cohort [21]. Second, the use of both selfreport diet and lifestyle questionnaires with a single measurement at baseline did not allow us to consider dietary or lifestyle (except tobacco consumption) changes during follow-up. Third, participants diagnosed with severe diseases during follow-up might have changed their dietary and lifestyle habits. However, we performed sensitivity analyses excluding participants with preexisting conditions, and the results remained robust to those of the entire cohort. Fourth, the information regarding the method of preparation and type of coffee was limited, and the quantification of HCAs in coffee was probably underestimated [10]. Fifth, breastfeeding, a part of pregnancy, can also interfere in the standard body weight trajectory. Pregnant women were excluded from our analysis; therefore, most of the breastfeeding women were consequently excluded, except those lactating only at baseline. We assume that this number is very low because the mean average was 50 years old, and, in some centers (such as Spain and Italy), participants were

mostly blood donors, and it is not possible to do a blood donation until 6 to 9 months after giving birth. For this reason, we recommend complementing the results from dietary questionnaires with nutritional biomarkers in future studies. Although validated center- and country-specific questionnaires were used to collect polyphenol-rich food data [19], we cannot exclude some measurement error leading to a potential underestimation of any true association. Finally, all models were adjusted for potential confounders; however, some potential residual bias cannot be ruled out.

In conclusion, this study identified a small inverse association between flavonoid intake and body weight change, specifically for anthocyanin, flavan-3-ol monomer, flavanone, flavone, and flavonol subclasses. These results suggest that flavonoids from foods may be promising compounds for weight control. Future randomized controlled trials using combinations of polyphenols or plant extracts mimicking polyphenol-rich diets may provide more definitive evidence to validate these results. In addition, HCAs from coffee showed a positive association with weight gain in coffee consumers. Future research related to coffee constituents, including HCAs, and weight change is warranted.O

#### AFFILIATIONS

<sup>1</sup>Unit of Nutrition and Cancer, Cancer Epidemiology Research Programme, Catalan Institute of Oncology (ICO), Bellvitge Biomedical Research Institute (IDIBELL), L'Hospitalet del Llobregat, Spain <sup>2</sup>Department of Epidemiology, Regional Health Council, IMIB-Arrixaca, Murcia, Spain

<sup>3</sup>CIBER in Epidemiology and Public Health (CIBERESP), Madrid, Spain <sup>4</sup>Unit of Diet, Genes and Environment, Danish Cancer Society Research Center, Copenhagen, Denmark

<sup>5</sup>Institute for Nutrition Research, School of Medical and Health Sciences, Edith Cowan University, Perth, Western Australia, Australia <sup>6</sup>Department of Public Health, Section of Environmental Health, Faculty of Health and Medical Sciences, University of Copenhagen, Copenhagen, Denmark

<sup>7</sup>Department of Public Health, Aarhus University, Aarhus, Denmark
 <sup>8</sup>Department of Cancer Epidemiology, German Cancer research
 Center (DKFZ), Heidelberg, Germany

<sup>9</sup>Department of Molecular Epidemiology, German Institute of Human Nutrition Potsdam-Rehbruecke, Nuthetal, Germany

<sup>10</sup>Institute of Nutritional Science, University of Potsdam, Potsdam, Germany

<sup>11</sup>Cancer Risk Factors and Life-Style Epidemiology Unit, Institute for Cancer Research Prevention and Clinical Network (ISPRO), Florence, Italy

<sup>12</sup>Epidemiology and Prevention Unit, Department of Research, IRCCS National Cancer Institute Foundation, Milan, Italy

<sup>13</sup>Department of Clinical and Experimental Medicine, Federico II University, Naples, Italy

<sup>14</sup>Hyblean Association for Epidemiological Research (AIRE-ONLUS), Ragusa, Italy

<sup>15</sup>Unit of Cancer Epidemiology, Città della Salute e della Scienza University-Hospital, Turin, Italy <sup>16</sup>Department of Community Medicine, Faculty of Health Sciences, University of Tromsø, The Arctic University of Norway, Tromsø, Norway

<sup>17</sup>The Public Dental Health Service Competence Center of Northern Norway, Tromsø, Norway

<sup>18</sup>Department of Functional Biology, University of Oviedo, Oviedo, Spain

<sup>19</sup>Department of Nutrition and Food Science, Campus of Cartuja, University of Granada, Granada, Spain

<sup>20</sup>Biosanitary Research Institute of Granada - ibs.Granada, Granada, Spain

<sup>21</sup>Institute of Nutrition and Food Technology (INYTA) 'José Mataix', Biomedical Research Centre, University of Granada, Granada, Spain
<sup>22</sup>Navarra Public Health Institute, Pamplona, Spain

<sup>23</sup>Navarra Institute for Health Research (IdiSNA), Pamplona, Spain

<sup>24</sup>Nutritional Epidemiology, Department of Clinical Sciences Malmö, Lund University, Malmö, Sweden

<sup>25</sup>Register-based Epidemiology, Department of Clinical Sciences Lund, Lund University, Lund, Sweden

<sup>26</sup>Department of Odontology, Umeå University, Umeå, Sweden
<sup>27</sup>Department of Medical Biosciences, Umeå University, Umeå, Sweden

<sup>28</sup>Julius Center for Health Sciences and Primary Care, University Medical Center Utrecht, Utrecht University, Utrecht, The Netherlands <sup>29</sup>MRC Epidemiology Unit, University of Cambridge School of Clinical Medicine, Institute of Metabolic Science, Cambridge Biomedical Campus, Cambridge, UK

<sup>30</sup>Department of Epidemiology and Biostatistics, School of Public Health, Imperial College London, London, UK

<sup>31</sup>International Agency for Research on Cancer Nutrition (IARC-WHO), Lyon, France

<sup>32</sup>Department of Nutrition, Food Sciences, and Gastronomy, Food Innovation Network (XIA), Institute for Research on Nutrition and Food Safety (INSA), Faculty of Pharmacy and Food Sciences, University of Barcelona, Barcelona, Spain

#### ACKNOWLEDGMENTS

We thank all EPIC participants and staff for their contribution to the study. We also thank Bertrand Hémon and Catalina Bonet for their valuable help with the EPIC database. We also thank the National Institute for Public Health and the Environment (RIVM; Bilthoven, the Netherlands); the University of Paris-Saclay, the University of Versailles Saint-Quentin-en-Yvelines (UVSQ), National Institute of Health and Medical Research (INSERM), Gustave Roussy Institute, and the Center for Research in Epidemiology and Population Health (CESP; Villejuif, France); the Ministry of Health of the Basque Government, Sub-Directorate for Public Health and Addictions of Gipuzkoa, the Public Health Division of Gipuzkoa, and BioDonostia Research Institute (San Sebastian, Spain); and Oxford University (UK) for their contribution and ongoing support to the EPIC Study.

#### FUNDING INFORMATION

This study has been funded by Carlos III Health Institute through the project PI18/00191 (co-funded by European Regional Development Fund [ERDF], a way to build Europe). The Physical Activity, Nutrition, Alcohol, Cessation of Smoking, Eating Out of Home and Obesity (PANACEA) project was funded by the European Union in the framework of the Public Health Programme (project number: 2005328). The coordination of European Prospective Investigation into Cancer and Nutrition (EPIC) is financially supported by International Agency for Research on Cancer (IARC) and also by the Department of Epidemiology and Biostatistics. School of Public Health. Imperial College London. which has additional infrastructure support provided by the National Institute for Health and Care Research (NIHR) Imperial Biomedical Research Centre (BRC). The national cohorts are supported by the following: Danish Cancer Society (Denmark): League Against Cancer, Gustave Roussy Institute, General Mutual of National Education, and the National Institute of Health and Medical Research (INSERM: France): German Cancer Aid, German Cancer Research Center (DKFZ), German Institute of Human Nutrition Potsdam-Rehbruecke (DIfE), and the Federal Ministry of Education and Research (BMBF) (Germany): Italian Association for Cancer Research (AIRC), Company of Saint Paul, and the 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), and Statistics Netherlands (the Netherlands); Health Research Fund (FIS), Carlos III Institute of Health (ISCIII), Regional Governments of Andalucía, Asturias, Basque Country, Murcia, and Navarra, and the Catalan Institute of Oncology (ICO; Spain); Swedish Cancer Society, Swedish Research Council, and the County Councils of Skåne and Västerbotten (Sweden); Cancer Research UK (14136 to EPIC-Norfolk; C8221/A29017 to EPIC-Oxford) and Medical Research Council (MRC: 1000143 to EPIC-Norfolk: MR/M012190/1 to EPIC-Oxford; the UK). Nita G. Forouhi acknowledges support from MRC Epidemiology Unit (MC\_UU\_0006/3) and from the NIHR Cambridge Biomedical Research Centre diet, nutrition, and lifestyle theme (IS-BRC-1215-20014). She is an NIHR Senior Investigator. The authors thank the Research Centers of Catalonia (CERCA) Program/Government of Catalonia for the institutional support to Bellvitge Biomedical Research Institute (IDIBELL). Jazmin Castañeda thanks the National Council for Science and Technology (CONACYT) predoctoral fellowship (identifier 693636). Mercedes Gil-Lespinard, Enrique Almanza-Aguilera, and Raul Zamora-Ros were supported by the predoctoral contracts for training in research into health (PFIS) (FI19/ 00185), the "Sara Borrell" (CD20/00071), and the "Miguel Servet II" (CPII20/00009) programs from the Carlos III Institute of Health (cofunded by European Social Fund [ESF], investing in your future).

#### DATA AVAILABILITY STATEMENT

For information on how to apply for getting access to EPIC data and/or biospecimens, please follow the instructions at the following link: http://epic.iarc.fr/access/index.php.

#### CONFLICT OF INTEREST

The authors declared no conflict of interest.

#### DISCLAIMER

Where authors are identified as personnel of the International Agency for Research on Cancer/World Health Organization, the authors alone are responsible for the views expressed in this article, and they do not necessarily represent the decisions, policy, or views of the International Agency for Research on Cancer/World Health Organization.

#### ORCID

Jazmin Castañeda 🕩 https://orcid.org/0000-0002-8521-9946 Mercedes Gil-Lespinard D https://orcid.org/0000-0002-7387-2139 Enrique Almanza-Aguilera 🕩 https://orcid.org/0000-0002-4805-0774 Fiorida Llaha D https://orcid.org/0000-0003-0534-6484 Jesús-Humberto Gómez 🕩 https://orcid.org/0000-0001-8442-8327 Nicola Bondonno D https://orcid.org/0000-0001-5905-444X Anne Tjønneland 🕩 https://orcid.org/0000-0003-4385-2097 Kim Overvad https://orcid.org/0000-0001-6429-7921 Verena Katzke () https://orcid.org/0000-0002-6509-6555 Matthias B. Schulze D https://orcid.org/0000-0002-0830-5277 Giovanna Masala D https://orcid.org/0000-0002-5758-9069 Claudia Agnoli D https://orcid.org/0000-0003-4472-1179 Rosario Tumino 🕩 https://orcid.org/0000-0003-2666-414X Carlotta Sacerdote D https://orcid.org/0000-0002-8008-5096 Guri Skeie D https://orcid.org/0000-0003-2476-4251 Magritt Brustad D https://orcid.org/0000-0003-0114-5271 Cristina Lasheras b https://orcid.org/0000-0003-0482-4229 Esther Molina-Montes b https://orcid.org/0000-0002-0428-2426 María-Dolores Chirlague D https://orcid.org/0000-0001-9242-3040 Aurelio Barricarte 🕩 https://orcid.org/0000-0001-6750-1270 Emily Sonestedt D https://orcid.org/0000-0002-0747-4562 Marisa da Silva b https://orcid.org/0000-0003-1215-8625 Ingegerd Johansson 💿 https://orcid.org/0000-0002-9227-8434 Johan Hultdin 🝺 https://orcid.org/0000-0002-9599-0961 Nita G. Forouhi D https://orcid.org/0000-0002-5041-248X Alicia K. Heath 🗅 https://orcid.org/0000-0001-6517-1300 Heinz Freisling D https://orcid.org/0000-0001-8648-4998 Elisabete Weiderpass D https://orcid.org/0000-0003-2237-0128 Augustin Scalbert b https://orcid.org/0000-0001-6651-6710 Raul Zamora-Ros 🕩 https://orcid.org/0000-0002-6236-6804

#### REFERENCES

- Dai H, Alsalhe TA, Chalghaf N, Riccò M, Bragazzi NL, Wu J. The global burden of disease attributable to high body mass index in 195 countries and territories, 1990–2017: an analysis of the Global Burden of Disease Study. *PLoS Med.* 2020;17:e1003198. doi:10. 1371/journal.pmed.1003198
- World Obesity. Prevalence of obesity. Accessed July 15, 2022. https://www.worldobesity.org/about/about-obesity/prevalence-ofobesity.
- World Health Organization. WHO European Regional Obesity Report. WHO Regional Office for Europe; 2022.
- Albuquerque D, Nóbrega C, Manco L, Padez C. The contribution of genetics and environment to obesity. *Br Med Bull.* 2017;123: 159-173.

- Semlitsch T, Stigler FL, Jeitler K, Horvath K, Siebenhofer A. Management of overweight and obesity in primary care—a systematic overview of international evidence-based guidelines. *Obes Rev.* 2019;20: 1218-1230.
- Wilding JPH, Jacob S. Cardiovascular outcome trials in obesity: a review. Obes Rev. 2021;22:e13112. doi:10.1111/obr.13112
- Fantin F, Giani A, Zoico E, Rossi AP, Mazzali G, Zamboni M. Weight loss and hypertension in obese subjects. *Nutrients*. 2019;11:1667. doi:10.3390/nu11071667
- Pedersen LR, Olsen RH, Anholm C, et al. Weight loss is superior to exercise in improving the atherogenic lipid profile in a sedentary, overweight population with stable coronary artery disease: a randomized trial. *Atherosclerosis*. 2016;246:221-228.
- Ma C, Avenell A, Bolland M, et al. Effects of weight loss interventions for adults who are obese on mortality, cardiovascular disease, and cancer: systematic review and meta-analysis. *BMJ*. 2017;359:j4849. doi:10.1136/bmj.j4849
- Zamora-Ros R, Knaze V, Rothwell JA, et al. Dietary polyphenol intake in Europe: the European Prospective Investigation into Cancer and Nutrition (EPIC) study. *Eur J Nutr.* 2016;55:1359-1375.
- 11. Farhat G, Drummond S, Al-Dujaili EAS. Polyphenols and their role in obesity management: a systematic review of randomized clinical trials. *Phytother Res.* 2017;31:1005-1018.
- Boccellino M, D'Angelo S. Anti-obesity effects of polyphenol intake: current status and future possibilities. Int J Mol Sci. 2020;21:5642. doi:10.3390/ijms21165642
- 13. Most J, Goossens GH, Jocken JW, Blaak EE. Short-term supplementation with a specific combination of dietary polyphenols increases energy expenditure and alters substrate metabolism in overweight subjects. *Int J Obes (Lond)*. 2014;38:698-706.
- 14. Llaha F, Zamora-Ros R. The effects of polyphenol supplementation in addition to calorie restricted diets and/or physical activity on body composition parameters: a systematic review of randomized trials. *Front Nutr.* 2020;7:84. doi:10.3389/fnut.2020.00084
- Adriouch S, Kesse-Guyot E, Feuillet T, et al. Total and specific dietary polyphenol intakes and 6-year anthropometric changes in a middleaged general population cohort. *Int J Obes (Lond)*. 2018;42:310-317.
- Vernarelli JA, Lambert JD. Flavonoid intake is inversely associated with obesity and C-reactive protein, a marker for inflammation, in US adults. Nutr Diabetes. 2017;7:e276. doi:10.1038/nutd.2017.22
- Guo X, Tresserra-Rimbau A, Estruch R, et al. Polyphenol levels are inversely correlated with body weight and obesity in an elderly population after 5 years of follow up (the randomised PREDIMED study). *Nutrients*. 2017;9:452. doi:10.3390/nu9050452
- Grosso G, Stepaniak U, Micek A, Stefler D, Bobak M, Pajak A. Dietary polyphenols are inversely associated with metabolic syndrome in Polish adults of the HAPIEE study. *Eur J Nutr.* 2017; 56:1409-1420.
- 19. Riboli E, Hunt KJ, Slimani N, et al. European Prospective Investigation into Cancer and Nutrition (EPIC): study populations and data collection. *Public Health Nutr*. 2002;5:1113-1124.
- May AM, Romaguera D, Travier N, et al. Combined impact of lifestyle factors on prospective change in body weight and waist circumference in participants of the EPIC-PANACEA study. *PLoS One*. 2012;7: e50712. doi:10.1371/journal.pone.0050712
- 21. Spencer EA, Appleby PN, Davey GK, Key TJ. Validity of self-reported height and weight in 4808 EPIC–Oxford participants. *Public Health Nutr.* 2002;5:561-565.
- 22. Slimani N, Deharveng G, Unwin I, et al. The EPIC nutrient database project (ENDB): a first attempt to standardize nutrient databases across the 10 European countries participating in the EPIC study. *Eur J Clin Nutr.* 2007;61:1037-1056.
- Knaze V, Rothwell R, Zamora-Ros R, et al. A new food-composition database for 437 polyphenols in 19,899 raw and prepared foods used to estimate polyphenol intakes in adults from 10 European countries. *Am J Clin Nutr.* 2018;108:517-524.

- 24. Orsini N, Greenland S. A procedure to tabulate and plot results after flexible modeling of a quantitative covariate. *Stata J.* 2011; 11:1-29.
- 25. Tomova GD, Arnold KF, Gilthorpe MS, Tennant PWG. Adjustment for energy intake in nutritional research: a causal inference perspective. *Am J Clin Nutr.* 2022;115:189-198.
- 26. Larsen SC, Mikkelsen ML, Frederiksen P, Heitmann BL. Habitual coffee consumption and changes in measures of adiposity: a comprehensive study of longitudinal associations. *Int J Obes (Lond)*. 2018;42: 880-886.
- 27. Bertoia ML, Rimm EB, Mukamal KJ, Hu FB, Willett WC, Cassidy A. Dietary flavonoid intake and weight maintenance: three prospective cohorts of 124 086 US men and women followed for up to 24 years. *BMJ*. 2016;352:i17. doi:10.1136/bmj.i17
- 28. Zhang W, Du J, Wang S, Ma H. Association of weight change patterns across adulthood with incident asthma: a retrospective cohort study. *Sci Rep.* 2022;12:9756. doi:10.1038/s41598-022-13555-w
- 29. Cui M, Chen B, Xu K, et al. Activation of specific bitter taste receptors by olive oil phenolics and secoiridoids. *Sci Rep.* 2021;11:22340. doi:10.1038/s41598-021-01752-y
- Hughes LA, Arts IC, Ambergen T, et al. Higher dietary flavone, flavonol, and catechin intakes are associated with less of an increase in BMI over time in women: a longitudinal analysis from the Netherlands Cohort Study. *Am J Clin Nutr.* 2008;88:1341-1352.
- Marranzano M, Ray S, Godos J, Galvano F. Association between dietary flavonoids intake and obesity in a cohort of adults living in the Mediterranean area. *Int J Food Sci Nutr.* 2018;69:1020-1029.
- Cercato C, Fonseca FA. Cardiovascular risk and obesity. Diabetol Metab Syndr. 2019;11:74. doi:10.1186/s13098-019-0468-0
- Farias-Pereira R, Park CS, Park Y. Mechanisms of action of coffee bioactive components on lipid metabolism. *Food Sci Biotechnol*. 2019; 28:1287-1296.
- Alam MA, Subhan N, Hossain H, et al. Hydroxycinnamic acid derivatives: a potential class of natural compounds for the management of lipid metabolism and obesity. *Nutr Metab (Lond)*. 2016;13:27. doi:10. 1186/s12986-016-0080-3
- Rosengren A, Dotevall A, Wilhelmsen L, Thelle D, Johansson S. Coffee and incidence of diabetes in Swedish women: a prospective 18-year follow-up study. J Intern Med. 2004;255:89-95.
- Kim JH, Park YS. Light coffee consumption is protective against sarcopenia, but frequent coffee consumption is associated with obesity in Korean adults. *Nutr Res.* 2017;41:97-102.
- Lee J, Kim HY, Kim J. Coffee consumption and the risk of obesity in Korean women. *Nutrients*. 2017;9:1340. doi:10.3390/nu9121340

- Kotyczka C, Boettler U, Lang R, et al. Dark roast coffee is more effective than light roast coffee in reducing body weight, and in restoring red blood cell vitamin E and glutathione concentrations in healthy volunteers. *Mol Nutr Food Res.* 2011;55:1582-1586. doi:10.1002/ mnfr.201100248
- 39. Greenberg JA, Owen DR, Geliebter A. Decaffeinated coffee and glucose metabolism in young men. *Diabetes Care*. 2010;33:278-280.
- Benbouguerra N, Hornedo-Ortega R, Garcia F, El Khawand T, Saucier C, Richard T. Stilbenes in grape berries and wine and their potential role as anti-obesity agents: a review. *Trends Food Sci Technol.* 2021;112:362-381.
- 41. Karković Marković A, Torić J, Barbarić M, Jakobušić BC. Hydroxytyrosol, Tyrosol and derivatives and their potential effects on human health. *Molecules*. 2019;24:2001. doi:10.3390/molecules24102001
- 42. Romaguera D, Norat T, Vergnaud AC, et al. Mediterranean dietary patterns and prospective weight change in participants of the EPIC-PANACEA project. *Am J Clin Nutr.* 2010;92:912-921.
- 43. Chopra S, Sharma KA, Ranjan P, Malhotra A, Vikram NK, Kumari A. Weight management module for perimenopausal women: a practical guide for gynecologists. *J Midlife Health*. 2019;10:165-172.
- Sharifi-Rad M, Anil Kumar NV, Zucca P, et al. Lifestyle, oxidative stress, and antioxidants: back and forth in the pathophysiology of chronic diseases. *Front Physiol.* 2020;11:694. doi:10.3389/fphys. 2020.00694
- 45. Freisling H, Van Bakel MM, Biessy C, et al. Dietary reporting errors on 24 h recalls and dietary questionnaires are associated with BMI across six European countries as evaluated with recovery biomarkers for protein and potassium intake. *Br J Nutr.* 2012;107:910-920.

#### SUPPORTING INFORMATION

Additional supporting information can be found online in the Supporting Information section at the end of this article.

How to cite this article: Castañeda J, Gil-Lespinard M, Almanza-Aguilera E, et al. Association between classes and subclasses of polyphenol intake and 5-year body weight changes in the EPIC-PANACEA study. *Obesity (Silver Spring)*. 2023;1-13. doi:10.1002/oby.23689