

Comparison of an Energy-Reduced Mediterranean Diet and Physical Activity Versus an Ad Libitum Mediterranean Diet in the Prevention of Type 2 Diabetes

A Secondary Analysis of a Randomized Controlled Trial

Miguel Ruiz-Canela, PharmD, PhD*; Dolores Corella, PharmD, PhD*; Miguel Ángel Martínez-González, MD, PhD; Nancy Babio, RD, PhD; J. Alfredo Martínez, MD, PhD; Luis Forga, MD, PhD; Ángel M. Alonso-Gómez, MD, PhD; Julia Wärnberg, PhD; Jesús Vioque, MD, PhD; Dora Romaguera, PharmD, PhD; José López-Miranda, MD, PhD; Ramón Estruch, MD, PhD; José Manuel Santos-Lozano, MD, PhD; Luis Serra-Majem, MD, PhD; Aurora Bueno-Cavanillas, MD, PhD; Josep A. Tur, PharmD, PhD; Vicente Martín-Sánchez, MD, PhD; Antoni Riera-Mestre, MD, PhD; Miguel Delgado-Rodríguez, MD, PhD; Pilar Matía-Martin, MD, PhD; Josep Vidal, MD, PhD; Clotilde Vázquez, MD, PhD; Lidia Daimiel, PhD; Pilar Buil-Cosiales, MD, PhD; Sangeetha Shyam, PhD; Jose V. Sorlí, MD, PhD; Olga Castañer, PhD; Antonio García-Rios, MD, PhD; Laura Torres-Collado, MPH, PhD; Enrique Gómez-Gracia, MD, PhD; M. Ángeles Zulet, PharmD, PhD; Jadwiga Konieczna, PhD; Rosa Casas, PhD; Naomi Cano-Ibáñez, PhD; Lucas Tojal-Sierra, MD, PhD; Rosa M. Bernal-López, MD, PhD; Estefanía Toledo, MD, PhD; Jesús García-Gavilán, RD, PhD; Rebeca Fernández-Carrión, PhD; Albert Goday, MD, PhD; Antonio P. Arenas-Larriva, MD, PhD; Sandra González-Palacios, PhD; Helmut Schröder, PhD; Emilio Ros, MD, PhD; Montserrat Fitó, MD, PhD; Frank B. Hu, MD, PhD; Francisco J. Tinahones, MD, PhD†; and Jordi Salas-Salvadó, MD, PhD†

Background: Limited research has been done to evaluate the combined effect of energy reduction, Mediterranean diet (MedDiet), and physical activity on type 2 diabetes incidence.

Objective: To evaluate whether an energy-reduced MedDiet (erMedDiet) plus physical activity reduces diabetes incidence compared with a standard MedDiet.

Design: Prespecified secondary outcome analysis in the PREDIMED (Prevención con Dieta Mediterránea)-Plus randomized, single-blinded, controlled trial. (ISRCTN Registry: ISRCTN89898870)

Setting: 23 centers across Spain.

Participants: 4746 adults aged 55 to 75 years with metabolic syndrome and overweight or obesity, without prior cardiovascular disease or diabetes.

Intervention: Participants were randomly assigned 1:1 to an intervention group receiving an erMedDiet (planned reduction of 600 kcal per day), increased physical activity, and behavioral strategies for reducing weight, or a control group receiving ad libitum MedDiet advice.

Measurements: Diabetes incidence was based on the American Diabetes Association criteria. Anthropometric measurements were obtained annually. Cox regression models were used to assess the intervention effect.

Results: The 6-year absolute risk was 12.0% (95% CI, 11.9% to 12.1%) in the control group (349 cases) and 9.5% (CI, 9.4% to 9.5%) in the intervention group (280 cases). Over a median 6-year follow-up, diabetes incidence was 31% (CI, 18% to 41%) relatively lower in the intervention group compared with the control group, with an absolute risk reduction of −2.6 cases (CI, −2.7 to −2.4) per 1000 person-years. The intervention group attained better adherence to the erMedDiet, higher physical activity levels, and greater reductions in body weight and waist circumference.

Limitation: Secondary outcome, single-blinded design, and self-reported dietary adherence.

Conclusion: An intensive intervention with the MedDiet adding caloric reduction, physical activity, and modest weight loss was more effective than only an ad libitum MedDiet in reducing diabetes incidence in overweight/obese persons with metabolic syndrome.

Primary Funding Source: Instituto de Salud Carlos III.

Ann Intern Med. doi:10.7326/ANNALS-25-00388

For author, article, and disclosure information, see end of text.

This article was published at Annals.org on 26 August 2025.

* Drs. Ruiz-Canela and Corella are co-first authors.

† Drs. Tinahones and Salas-Salvadó are co-senior authors.

The rising prevalence, incidence, and burden of diabetes (1–3), driven by the obesity epidemic (2), underscore the need for simple, sustainable, preventive strategies (4).

Robust evidence shows that diabetes is preventable through lifestyle modifications aiming at weight loss (5–8). A meta-analysis including 19 randomized

See also:

Editorial comment
Summary for Patients

Web-Only
Supplement

clinical trials confirmed the long-term beneficial effect of lifestyle interventions, even with modest weight loss (9). Most of these interventions targeted weight loss through an energy-reduced, healthy *low-fat* diet combined with increased physical activity (5). However, no previous trial has assessed the effect of energy reduction in the context of a relatively high-fat diet, such as the Mediterranean diet (MedDiet).

The PREDIMED (Prevención con Dieta Mediterránea) trial demonstrated that an ad libitum MedDiet supplemented with either extra-virgin olive oil or mixed nuts reduced diabetes incidence by 30%, compared with a low-fat diet (10, 11). However, this reduction occurred with only marginally reduced body weight (12). Building on this finding, the PREDIMED-Plus trial was designed to test whether adding caloric restriction and physical activity to the MedDiet would provide additional benefit beyond the effect observed only with the MedDiet. In a 1-year interim analysis, an energy-reduced MedDiet (erMedDiet) combined with increased physical activity significantly reduced body weight, waist circumference, and other cardiometabolic risk markers compared with an ad libitum MedDiet (13–15). The present analysis, conducted with data from all PREDIMED-Plus participants free of diabetes at baseline, tests whether an intensive lifestyle intervention aiming at weight loss reduces diabetes risk beyond the effects of an ad libitum MedDiet.

METHODS

Study Design

This is a secondary outcome analysis in the PREDIMED-Plus trial, whose methods were previously reported (13, 16). Briefly, the PREDIMED-Plus is a multicenter, randomized, parallel-group, single-blind, lifestyle intervention trial aiming to assess the effects of a weight-loss intervention with an erMedDiet and increased physical activity (intervention group) versus a control MedDiet group without energy reduction for primary cardiovascular prevention. Diabetes incidence was a predefined secondary end point.

The study protocol was approved by the research ethics committees of all recruitment centers, and all participants provided written informed consent. The ethics approval and study protocol are available at [Annals.org](https://www.annals.org) and on the PREDIMED-Plus website.

Population and Randomization

Eligible participants were community-dwelling men aged 55 to 75 years and women aged 60 to 75 years, with overweight or obesity (body mass index [BMI] ≥ 27 and < 40 kg/m²), without documented cardiovascular disease or diabetes (for the present analysis), and with at least 3 metabolic syndrome components: waist circumference 80 cm or more in women and 94 cm or more in men, elevated triglycerides (≥ 1.69 mmol/L [≥ 150 mg/dL]), a reduced high-density lipoprotein cholesterol level (< 1.04 mmol/L [< 40 mg/dL]), elevated

blood pressure (systolic ≥ 150 and diastolic ≥ 85 mm Hg), and an elevated fasting glucose level (≥ 5.55 mmol/L [≥ 100 mg/dL]) (17). Primary care physicians from 23 Spanish recruitment centers assessed eligibility. Participants were randomly allocated 1:1 to one of the study arms using a centralized computer-based system with stratification by center, sex, and age (< 65 , 65 to 70, > 70 years) in blocks of 6. Couples sharing a household were randomly assigned as a cluster. Randomization was masked to center staff and investigators. Recruitment lasted from 5 September 2013 to 23 December 2016.

This study reports the intervention's effects on diabetes incidence in all participants without diabetes at baseline ($n = 4746$, from 6874 total participants in the trial). Prevalent diabetes cases at baseline were initially self-reported and subsequently confirmed through a review of their medical records. In addition, clinical records of potential new incident diabetes cases were reviewed to confirm that these diagnoses occurred after participants were enrolled in the study.

Intervention

The interventions were scheduled to last for an average of 6 years. Given the study design, neither dietitians nor participants were blinded to the randomized arm. Descriptions of the intervention and its effectiveness after 1 year were previously reported (13, 16, 18). Observed 1-year improvements in adiposity, glucose level, and lipids for the intervention versus control were reported in previous publications (13–15).

Participants in the intervention group were instructed to follow an erMedDiet with physical activity recommendations and behavioral support for weight loss (**Supplement Table 1**, available at [Annals.org](https://www.annals.org)). The erMedDiet was defined as a traditional MedDiet with a planned reduction of 600 kcal per day, providing 35% to 40% of calories from fat, 40% to 45% from carbohydrates, and approximately 20% from protein. Participants were encouraged to consume more fruits, vegetables, whole cereals, legumes, nuts, and extra-virgin olive oil, while limiting meat, sugar-sweetened beverages, foods with added sugars, and cream. A validated 17-item questionnaire was used to assess adherence to the erMedDiet annually and to deliver recommendations (**Supplement Table 2**) (19). Physical activity recommendations included: a) progressively increased aerobic physical activity, mainly brisk walking, aiming for 45 minutes per day (or the equivalent) 6 days per week; b) strength exercises (≥ 2 days per week); and c) flexibility and balance exercises (≥ 3 days per week) (18).

The control group received educational sessions on the traditional MedDiet with ad libitum caloric intake following the PREDIMED-1 trial recommendations (20). Adherence to the MedDiet was assessed using a validated 14-item MedDiet adherence screener (21). No advice on physical activity or weight loss was provided to the control group.

The participants in the intervention group were contacted by dietitians 3 times a month (1 group session, 1 individual session, and 1 phone call) during the first year, then received monthly group sessions, individual sessions every 3 months, and 2 phone calls every 3 months. Control group participants had 1 individual visit, 1 phone call, and 1 group session every 6 months. Both groups received 1 L per month of extra-virgin olive oil to support adherence and retention. During the COVID-19 pandemic, the intervention was remotely delivered (22).

Variables

Self-reported dietary measurements were conducted at baseline, after 6 months, and then annually during the 6 years of the intervention and 2 additional years of follow-up, using a validated 143-item food-frequency questionnaire (23) and 2 MedDiet adherence questionnaires—one with 17 items to assess adherence to the erMedDiet (19) and one with 14 items to assess adherence to the traditional MedDiet (21). Physical activity was measured with the REGICOR short physical activity questionnaire (24) and the Physical Activity Readiness and Rapid Assessment of Physical Activity questionnaires (25).

Body weight, height, and waist circumference were measured by trained personnel, not blinded to group assignment, and medication use was recorded. Fasting blood glucose level, lipid profile, and hemoglobin A_{1c} (HbA_{1c}) level were determined by standard methods at baseline, after year 1, and every other year thereafter. Baseline insulin was centrally measured by an electrochemiluminescence immunoassay using an Elecsys immunoanalyzer (Roche Diagnostics). Insulin resistance was estimated using the Homeostasis Model Assessment of Insulin Resistance (HOMA-IR) index (26).

The definition of prediabetes was based on fasting plasma glucose (FPG) levels from 5.55 to 6.94 mmol/L (100 to 125 mg/dL) or HbA_{1c} levels between 5.7% and 6.4% (27, 28).

Outcome

Diabetes incidence was assessed yearly for 7 years of follow-up (a 6-year intervention period and 1 additional year of observation) (13). The American Diabetes Association criteria were used to ascertain incident diabetes (29): a) HbA_{1c} levels of 6.5% or more, performed in a laboratory using a method that is National Glycohemoglobin Standardization Program certified and standardized to the Diabetes Control and Complications Trial (DCCT) assay; b) a fasting plasma glucose level of 7.0 mmol/L or more (≥ 126 mg/dL); c) a 2-hour plasma glucose level of 11.1 mmol/L or more (≥ 200 mg/dL) during an oral glucose tolerance test using a glucose load containing the equivalent to 75 g of anhydrous glucose dissolved in water; or d) a random plasma glucose level of 11.1 mmol/L or more (≥ 200 mg/dL) in patients with classic symptoms of

hyperglycemia or a hyperglycemic event. In the absence of unequivocal hyperglycemia, results should be confirmed by repeated testing.

Physicians, blinded to group assignment, reviewed participants' medical records annually. New-onset diabetes cases were identified based on medical diagnoses reported in medical charts or repeated glucose tests from routine analyses. Potential cases were reviewed by the Clinical Events Ascertainment Committee, blinded to treatment allocation. Only confirmed new-onset diabetes cases occurring between randomization and 31 December 2022 were included.

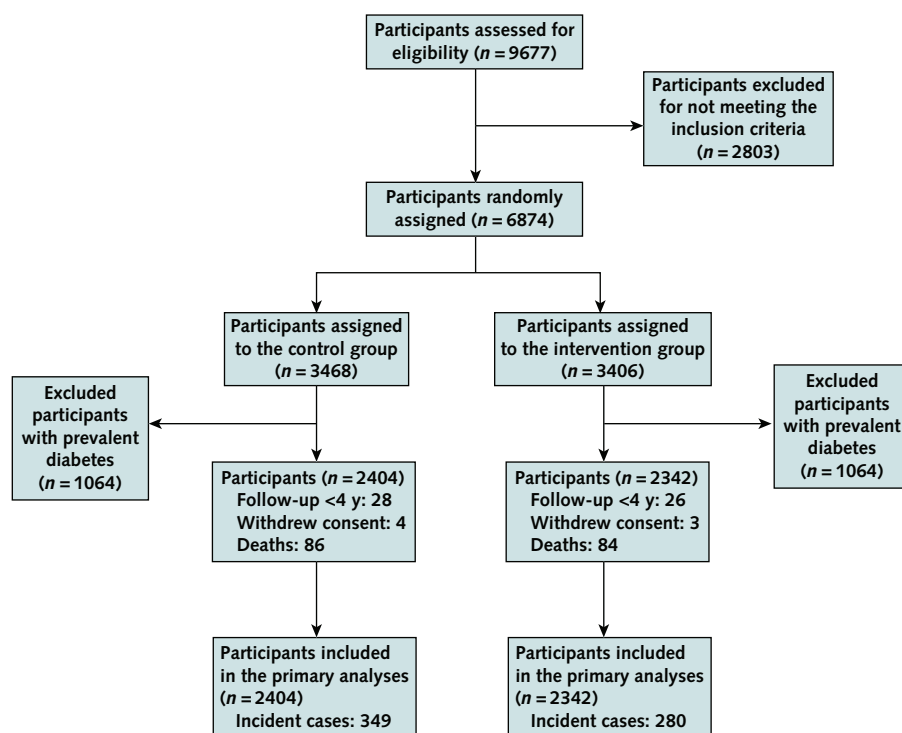
Statistical Analysis

The original trial was designed to recruit approximately 6000 participants primarily to detect a 30% relative reduction in the incidence of cardiovascular disease in the intervention versus the control group (16). For incident diabetes, we anticipated sufficient statistical power based on projections from the Diabetes Prevention Impact Toolkit (29).

Primary between-group comparisons followed the intention-to-treat principle. Time-to-first event was defined as the time from randomization to diabetes diagnosis or the last follow-up date for those without this diagnosis (last visit or last date in the medical record, whichever occurred later, or date of death) (30). Cox models assessed the intervention's effect, with results expressed as hazard ratios and 95% CIs (31, 32). Robust SEs were used to account for intracluster correlations among participants from the same households (569 pairs out of the 4746 participants). Absolute risk differences at 6 years were estimated using Cox regression models, with robust SEs and the same covariate adjustments as described in the next paragraph for hazard ratios, to estimate standardized differences in incidence.

A multivariable model adjusted for sex, age, and baseline glucose level, and stratified according to recruitment center and educational level (5 categories), was fitted. To assess the robustness of the results, a second multivariable model included additional baseline covariates: BMI (continuous), leisure-time physical activity (metabolic equivalent of task minutes per week), smoking habit (never/current/former), hypercholesterolemia, hypertension, alcohol intake (grams per day, including a quadratic term), total energy intake (kilocalories per day), baseline erMedDiet adherence score (17-point score, continuous), and family history of diabetes.

Linear mixed models (with robust SEs) assessed changes in erMedDiet, physical activity, and anthropometric variables from baseline by intervention group during the first 6 years of active follow-up, with random intercepts at recruitment site and family cluster level, and time categorically modeled (visit number) to capture potential nonlinear trends (a sample of the STATA code is available in **Supplement 1**, available at Annals.org).

Figure 1. Flow chart of participants in the PREDIMED-Plus study.

PREDIMED = Prevención con Dieta Mediterránea.

A simple imputation method was applied for continuous variables with missing values included in multivariable models using the command *impute* from Stata: baseline fasting plasma glucose ($n = 31$), total energy intake ($n = 29$), alcohol intake ($n = 29$), erMedDiet adherence ($n = 7$), and body weight ($n = 2$). The impute command uses regression-based methods, assuming that missing values are missing at random or missing completely at random. The variables included in the imputation model were age, sex, waist-to-height ratio, waist circumference, BMI, blood pressure, alcohol intake, dietary fiber, and smoking status. Missing values from categorical variables were imputed assuming a negative response to questions about family history of diabetes ($n = 178$), hypercholesterolemia ($n = 46$), hypertension ($n = 35$), level of education (lowest, $n = 7$), and smoking ($n = 2$).

Subgroup analyses were conducted according to the following variables: age categories (≤ 65 vs. > 65 years), sex, educational level (primary or less vs. secondary or higher), BMI (< 30 vs. ≥ 30 kg/m²), waist-to-height ratio (< 0.5 vs. ≥ 0.5), and baseline glucose level (< 5.55 vs. ≥ 5.55 mmol/L [< 100 vs. ≥ 100 mg/dL]). An interaction term between intervention group and each one of these variables was included in the Cox model, and it was assessed using the likelihood ratio test.

Several ancillary analyses were conducted including a per protocol-like approach (Supplement Figure, available at [Annals.org](https://annals.org)). In addition, a composite

variable strategy was implemented to account for deaths occurring before the diagnosis of diabetes, and both absolute and relative risks were recalculated.

In all statistical analyses (performed with STATA v.16.1), the PREDIMED-Plus database version 21-11-2023 was used. All tests were 2-sided with a significance threshold of $P < 0.05$.

Role of the Funding Source

The funding organizations had no role in the design and conduct of the study; collection, management, analysis, and interpretation of the data; preparation, review, or approval of the manuscript; and decision to submit the manuscript for publication.

RESULTS

Baseline Characteristics and Follow-up

Of the 6874 randomly assigned participants (Figure 1), 2128 with diabetes at baseline were excluded, leaving 4746 participants without diabetes (mean age, 64.9 years [SD, 4.9]; 50% women). Baseline characteristics were similar between included participants and those excluded due to prevalent diabetes (Supplement Table 3). Retained participants' baseline characteristics were balanced between both groups (Table 1). Baseline medication use was similar between groups (Supplement Table 4).

The number of participants with less than 6 years of follow-up was 241 (10.3%) in the intervention group

Table 1. Baseline Characteristic of Participants Without Type 2 Diabetes at Baseline in the PREDIMED-Plus Trial*

Characteristic	Control Group (n = 2404)	Intervention Group (n = 2342)
Age, y	64.9 (4.9)	64.8 (5.0)
Sex, n (%)		
Female	1189 (49.5)	1170 (50.0)
Male	1215 (50.5)	1172 (50.0)
Body weight, kg	86.0 (13.0)	86.3 (13.1)
BMI, kg/m ²	32.5 (3.5)	32.5 (3.4)
Waist circumference, cm	106.9 (9.7)	106.8 (9.7)
Smokers, n (%)		
Former	1060 (44.1)	959 (40.9)
Current	269 (11.2)	315 (13.5)
Never	1075 (44.7)	1068 (45.6)
Physical activity, MET-min/wk	2582 (2352)	2383 (2186)
Primary education or less, n (%)	1165 (48.5)	1095 (46.8)
erMedDiet score	8.5 (2.7)	8.3 (2.7)
Alcohol intake, g/d	11.9 (15.8)	11.0 (15.0)
Total energy intake, kcal/d	2457 (678)	2425 (620)
Systolic blood pressure, mm Hg	138.7 (16.4)	139.4 (17.1)
Diastolic blood pressure, mm Hg	81.0 (9.8)	81.4 (9.9)
Baseline hypertension, n (%)	1946 (81.0)	1945 (83.1)
Baseline dyslipidemia, n (%)	1598 (66.5)	1583 (67.6)
Baseline prediabetes,† n (%)	585 (24.3)	595 (25.4)
Baseline prediabetes,‡ n (%)	1744 (72.6)	1670 (71.3)
Fasting plasma glucose level		
mmol/L	5.67 (0.72)	5.66 (0.75)
mg/dL	102.1 (13.0)	102.0 (13.6)
Fasting plasma insulin, mIU/L	18.5 (10.1)	18.2 (10.0)
HOMA-IR	4.8 (2.9)	4.8 (3.2)
HbA _{1c} , %	5.8 (0.4)	5.7 (0.4)
Total cholesterol level		
mmol/L	5.29 (0.94)	5.26 (0.93)
mg/dL	204.3 (36.3)	203.2 (35.9)
HDL cholesterol level		
mmol/L	1.27 (0.30)	1.27 (0.31)
mg/dL	49.2 (11.6)	49.0 (12.1)
LDL cholesterol level		
mmol/L	3.26 (0.81)	3.25 (0.81)
mg/dL	126.0 (31.1)	125.3 (31.2)
Triglycerides		
mmol/L	1.69 (0.85)	1.67 (0.81)
mg/dL	149.5 (75.2)	148.0 (72.1)
Family history of diabetes, n (%)	828 (34.4)	818 (34.9)

erMedDiet = energy-reduced Mediterranean diet (score, 0 to 17 points); HbA_{1c} = hemoglobin A_{1c}; HDL = high-density lipoprotein; HOMA-IR = Homeostasis Model Assessment for Insulin Resistance; LDL = low-density lipoprotein; MET-min/wk = metabolic equivalent task minutes per week; PREDIMED = Prevención con Dieta Mediterránea.

* Data are mean (SD) unless otherwise specified.

† According to World Health Organization (WHO) criteria.

‡ According to American Diabetes Association (ADA) criteria.

and 242 (10.1%) in the control group. No statistically significant differences were observed in terms of sex, age, weight, BMI, or adherence to the erMedDiet between participants with less than 6 years versus 6 or more years of follow-up.

Effect of the Intervention on Diabetes Risk

In the intention-to-treat analysis, after a median follow-up of 6.6 years (IQR, 6.0 to 7.1 years), 629 incident diabetes cases occurred. The incidence curves progressively diverged between the intervention and the control group starting at 6 months (Figure 2).

Table 2 shows the number of new cases, rates, absolute risks, and relative risks by groups. After 6 years, the adjusted absolute risk difference for incident diabetes between the intervention and control groups was −2.4% (95% CI, −3.1 to −1.8), corresponding to a 31% (CI, 18% to 41%) relative risk reduction in the intervention group as compared with the control group.

In prespecified subgroup analyses, no statistically significant interactions were found for age, educational level, BMI, waist-to-height ratio, or baseline glucose level, except for sex, with greater diabetes risk reduction in men than in women (*P* for interaction = 0.035) (Supplement Table 5).

Ancillary analyses yielded similar results (Supplement Figure). During follow-up, 148 participants died without a diagnosis of diabetes (77 in the control group and 71 in the intervention group). The intervention effect remained largely unchanged when either diabetes incidence or death occurring before diabetes diagnosis was used as a composite outcome (Supplement Table 6).

Both groups showed similar increased use of anti-hypertensive and hypolipidemic medication (Supplement Table 4). Participants in the control group had greater antidiabetic drug use than those in the intervention group during the follow-up (11.7% vs. 9.5% at year 6; *P* = 0.018).

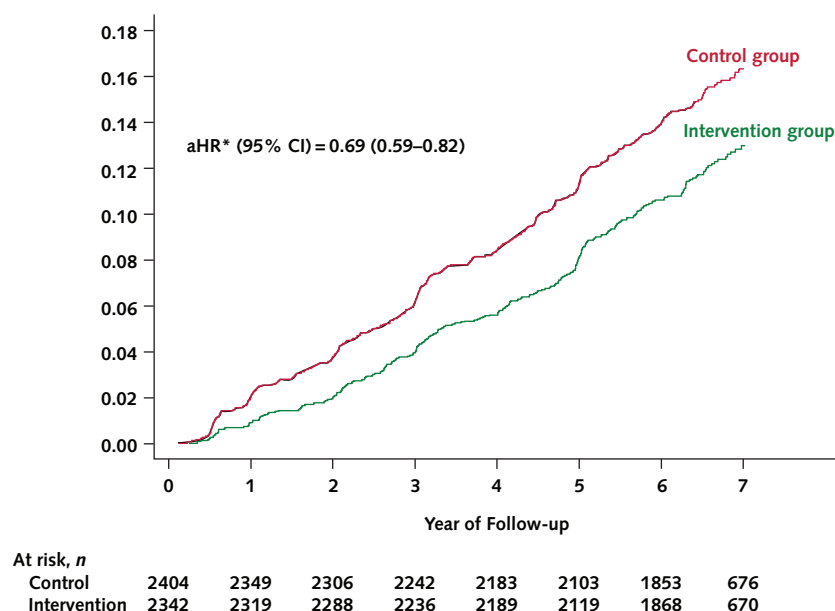
Changes in erMedDiet Adherence and Physical Activity

From year 1 on, both the intervention and the control groups improved their average adherence to the erMedDiet, although the former showed significantly greater improvement (Figure 3). The mean between-group difference in the erMedDiet score during the 6 years of intervention was 2.2 (CI, 2.0 to 2.3; *P* < 0.001). Detailed results for each of the 17 items of the erMedDiet score at baseline, and at years 1, 3, and 6 of follow-up, are shown in Supplement Table 7. An increase in the percentage of participants reaching the target for 16 of the 17 individual food items, excluding alcohol intake, was observed in both groups at years 3 and 6. However, this improvement was greater in the intervention than in the control group, with larger increments in the consumption of vegetables, fruits, legumes, fish, and whole grains, and reduced consumption of red and processed meat, refined grain, pasta, and white rice in the intervention than in the control group.

Compared with the control group, greater increases in physical activity were observed in the intervention group (Figure 3). During the first 6 years of intervention, the between-group mean difference was 800 (CI, 680 to 921) metabolic equivalent of task minutes per week (*P* < 0.001).

Effect on Adiposity Measures

Greater reductions in body weight and waist circumference occurred in the intervention group than in

Figure 2. Cumulative incidence of type 2 diabetes by randomized arm of the PREDIMED-Plus trial.

aHR = adjusted hazard ratio; PREDIMED = Prevención con Dieta Mediterránea.

* Adjusted for age, sex, and fasting glucose level (≤ 5.55 ; > 5.55 to < 6.11 ; 6.11 to < 6.99 ; and ≥ 6.99 mmol/L [≤ 100 ; > 100 to < 110 ; 110 to < 126 ; and ≥ 126 mg/dL]), body mass index (kg/m^2), smoking status (never, former, current smoker), baseline prevalence of dyslipidemia (yes/no) and hypertension (yes/no), family history of diabetes, leisure-time physical activity level (metabolic equivalent of task minutes per day), adherence to energy-reduced MedDiet, and alcohol intake (grams per day, adding quadratic term), and stratified by center and educational level.

the control group (Figure 3 and Supplement Table 8). In the analysis of completers, the 6-year mean body weight reduction was -3.3 kg (CI, -3.4 to -3.1 kg) in the intervention group and -0.6 kg (CI, -0.7 to -0.4 kg) in the control group, representing weight reductions of 3.7% in intervention and 0.6% in control. During this period, mean waist circumference was reduced by -3.6 cm (CI, -3.8 to -3.3 cm) in the intervention group and -0.3 cm (CI, -0.4 to -0.1 cm) in the control group.

DISCUSSION

In the PREDIMED-Plus randomized trial including adults aged 55 to 75 years with overweight or obesity and metabolic syndrome, an intervention consisting of advice on a calorie reduction using a traditional MedDiet pattern combined with increased physical activity and modest weight loss resulted in a 31% lower relative risk for incident diabetes compared with an ad libitum MedDiet replicating the PREDIMED trial intervention (20) after a median follow-up of almost 7 years. Moreover, analyses of adjusted risk differences revealed an absolute 2.4% reduction in the cumulative incidence of diabetes at 6 years, further reinforcing the clinical impact of the intervention. The reduction in diabetes incidence was consistent across subgroups of age, education, and baseline metabolic status, with a greater effect observed in men than women. The

consistency of results across multiple ancillary analyses underscores the robustness of the PREDIMED-Plus intervention as a strategy to lower diabetes risk beyond the effects of a strong comparator, which was the conventional MedDiet.

Some postulated mechanisms could explain our results. The MedDiet, rich in fiber, whole grains, antioxidants, and anti-inflammatory compounds, is known to reduce diabetes risk, even in the absence of weight loss (12). The erMedDiet fosters low glycemic index foods as well as more fiber and whole grains, which are known to improve insulin sensitivity (33). In addition, moderate physical exercise may protect mitochondrial function and protect against obesity-induced insulin resistance (34). Synergistically, reductions in weight and visceral fat, achieved through caloric reduction and the increased physical activity implemented in our intervention, could significantly lower insulin resistance and inflammation (12, 35, 36). The MedDiet and physical activity also trigger the secretion of incretin hormones such as glucagon-like peptide-1, which plays a crucial role in glucose metabolism (37). Finally, the intervention effect could be modulated by favorable changes in gut microbiota and metabolome (38). However, in our study, we cannot disentangle whether the greater reduction in diabetes incidence resulted from weight loss, enhanced adherence to the MedDiet, physical activity promotion, or any combination of their effects.

Table 2. Absolute and Relative Risk Estimates for Incident Type 2 Diabetes After 6 Years of Intervention

Estimate	Control Group (n = 2404)	Intervention Group (n = 2342)
Cases, n	349	280
Person-years, n	15 022	14 989
Absolute risk (95% CI)	12.0 (11.9 to 12.1)	9.5 (9.4 to 9.5)
Crude RD (95% CI)	0 (reference)	−2.6 (−2.7 to −2.4)
Model 1,† RD (95% CI)	0 (reference)	−2.4 (−3.1 to −1.8)
Model 2,‡ RD (95% CI)	0 (reference)	−2.4 (−3.1 to −1.8)
Incidence rate (95% CI) per 1000 person-years	23.2 (20.9 to 25.8)	18.7 (16.6 to 21.0)
Crude HR (95% CI)	1 (reference)	0.80 (0.68 to 0.94)
Model 1,† HR (95% CI)	1 (reference)	0.71 (0.60 to 0.84)
Model 2,‡ HR (95% CI)	1 (reference)	0.69 (0.59 to 0.82)

HR = hazard ratio; RD = risk difference.

* Absolute risk and RDs were obtained from the predicted 6-year cumulative incidence derived from Cox models, using the baseline survival and average linear predictors for each group (see also Supplement Table 9).

† Adjusted for age (years), sex and fasting plasma glucose level at baseline (≤ 5.55 ; >5.55 to <6.11 ; 6.11 to <6.99 ; and ≥ 6.99 mmol/L [≤ 100 ; >100 to <110 ; 110 to <126 ; and ≥ 126 mg/dL]) and stratified by center and educational level (primary education vs. secondary or higher). Robust SEs to account for intracluster correlations (clusters are family members of the same household).

‡ Additionally adjusted for body mass index (kg/m²), smoking status (never, former, current smoker), baseline prevalence of dyslipidemia (yes/no) and hypertension (yes/no), family history of diabetes, leisure-time physical activity level (metabolic equivalent of task minutes per day), adherence to the energy-reduced Mediterranean diet, and alcohol intake (grams per day, adding a quadratic term), and stratified by center and educational level (primary education vs. secondary or higher).

Previous landmark trials supported the efficacy of lifestyle interventions for diabetes prevention. The Diabetes Prevention Program (DPP) reported that an intensive lifestyle intervention, including a low-fat diet and enhanced physical activity, led to a 58% reduction in diabetes incidence compared with placebo over a mean follow-up of 2.8 years among persons with impaired glucose tolerance (8). Similarly, the Diabetes Prevention Study (DPS) reported comparable results after 3.2 years, also including patients with impaired glucose tolerance (7). More recently, 15-year follow-up of the DPP reported a 27% (CI, 17% to 35%) relative reduction in diabetes risk after lifestyle sessions were extended to all participants (39).

Our study differs from these previous trials in several important ways. First, we used a considerably stronger comparator, an ad libitum MedDiet supplemented with free extra-virgin olive oil, which has been previously shown to reduce diabetes risk by 30% (11). It is plausible that if the control intervention from the DPP trial had been used in PREDIMED-Plus, the observed risk reduction might have approached the 58% reduction reported in the DPP trial. The MedDiet is inherently high in fat (35% to 40% of daily energy), in contrast to the low-fat diets used in previous trials (40). Second, our participants were older persons with a mean age of 64.9 years versus 50.6 years in the DPP trial (8) and 53.0 years in the DPS trial (7), and they were selected on metabolic syndrome rather than impaired glucose tolerance. Third, we observed substantially lower incidence rates in both intervention and control groups

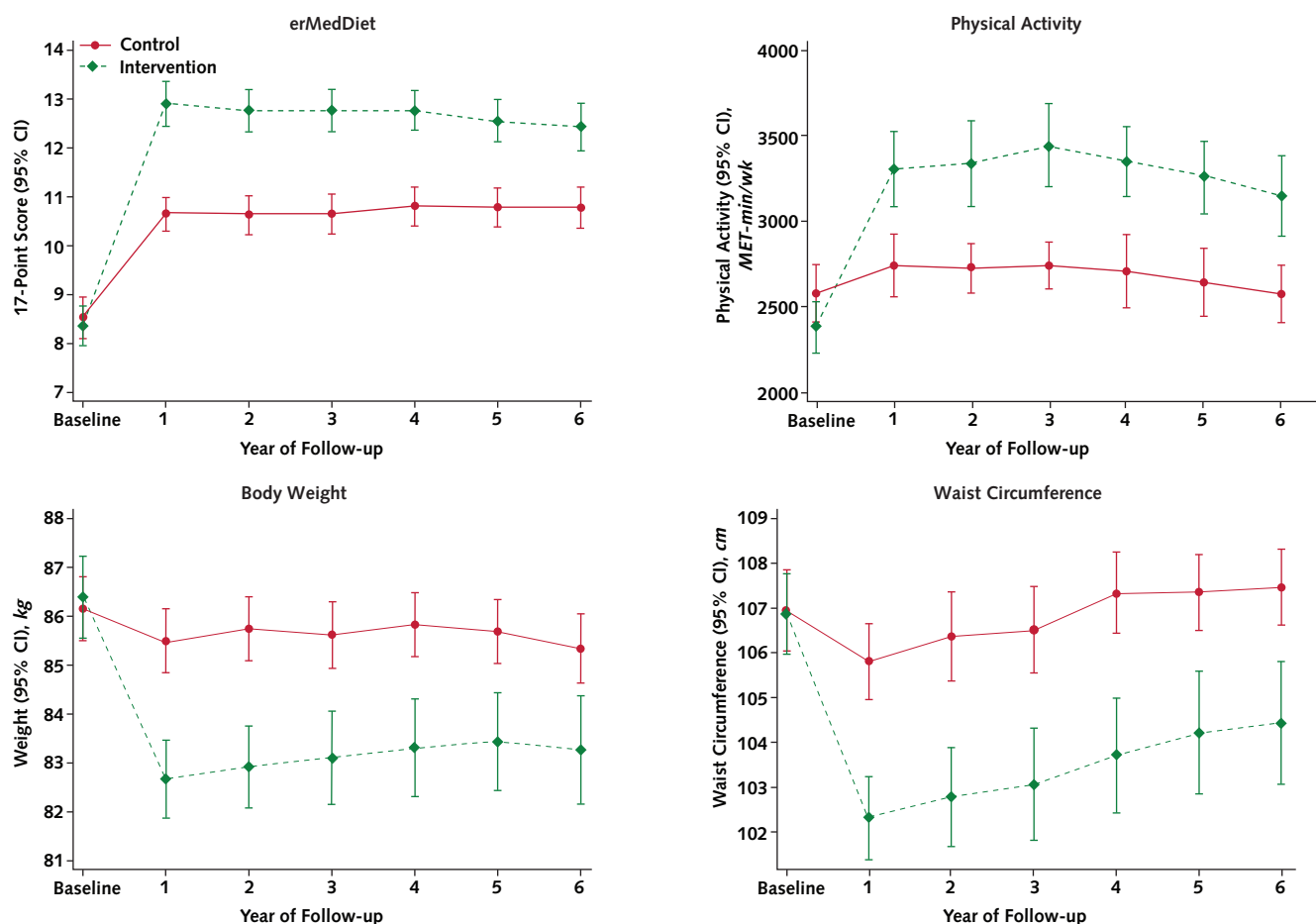
(1.87 and 2.32 cases per 100 person-years, respectively) compared with those reported in the DPP trial (4.8 in the intervention and 11.0 cases per 100 person-years in the placebo group) (8). Therefore, it is challenging to directly compare the effect size of our interventions with those from these studies.

Nevertheless, in general, weight loss is crucial for reducing diabetes risk. The DPP trial achieved an average weight loss in the lifestyle intervention group of 5.6 kg (a 7% reduction of initial body weight) (8), whereas the DPS trial saw 5% weight loss (7). In PREDIMED-Plus, 3.7% body weight loss over a period of 6 years was associated with long-term diabetes risk reduction. Therefore, even modest weight losses can lead to reductions in diabetes risk, consistent with previous studies (41–43). The potential effect modification by sex may be explained by greater weight losses in men (3.4 kg) than in women (2.3 kg), which was also observed in the DPP trial (44). However, these subgroup findings should be interpreted cautiously considering the number of comparisons performed (45).

Our findings build on the previous PREDIMED-1 trial (11), showing that an ad libitum MedDiet versus a low-fat diet reduced diabetes incidence by 30%. The intensity of the intervention in the control group of PREDIMED-Plus was comparable to that of the active intervention group with MedDiet in the previous PREDIMED-1 trial. However, we acknowledge that the higher intensity and frequency in the intervention than in the control group of PREDIMED-Plus may partially explain the improved adherence and outcomes. Undoubtedly, weight loss, increased physical activity, and reduction in caloric intake represent greater challenges than only changing the dietary pattern. That was the reason why higher intensity and frequency were required in the protocol of PREDIMED-Plus for the intervention than for the control group.

Some limitations are acknowledged. Diabetes was a prespecified secondary outcome, but randomization was effective in balancing baseline characteristics in all participants without diabetes. Dietary adherence was self-reported; however, we strengthened assessments through periodic in-person administration of a validated tool specific for the trial (19). Staff performing anthropometric measures—though trained to follow a standardized protocol—were not blinded to group assignment. In addition, the multifaceted intervention makes it difficult to disentangle the proportion of benefits due to the changes in each of the 3 components (erMedDiet, physical activity, or weight loss) of the intervention or a potential synergistic effect among them. Lastly, our results derive from participants aged 55 to 75 years with metabolic syndrome, living in a European Mediterranean country, potentially limiting their generalizability to younger populations or those with different ethnic backgrounds or diverse dietary traditions.

Figure 3. Mean values (95% CIs) of adherence to the erMedDiet (17-point score), physical activity, body weight, and waist circumference during follow-up by intervention group.



erMedDiet = energy-reduced Mediterranean diet; MET-min/wk = metabolic equivalent of task minutes per week. Mixed linear model with random intercepts at recruitment site, participant, and cluster family level. Conducted with completers only. $P = 0.027$ (between-group differences at baseline); $P < 0.001$ (between-group differences during the follow-up period).

Future research should address several important questions, including the identification of which components of the intervention are most critical for diabetes prevention and which subgroups derive the greatest benefit. Implementation studies exploring how to effectively translate this approach into diverse clinical and community settings are essential. Longer-term follow-up studies would also help determine whether the observed benefits persist beyond the intervention period. Finally, the greater improvements in both MedDiet adherence scores and physical activity levels in the intervention group demonstrate that more frequent participant contact likely contributed to better adherence to lifestyle recommendations. This highlights the potential importance of contact intensity in lifestyle interventions, which should be considered when interpreting our results and designing future diabetes-prevention strategies.

In conclusion, among adults with overweight or obesity with metabolic syndrome, a MedDiet intervention

including caloric reduction, physical activity, and modest weight loss reduced diabetes incidence beyond that achieved with only a traditional ad libitum MedDiet intervention, whose protective effects against diabetes were previously demonstrated. Although there is no "one-size-fits-all" dietary strategy plan for diabetes prevention (46), the MedDiet's higher palatability and cultural acceptance (47) could make it a highly sustainable, long-term, weight loss option when combined with moderately reduced energy intake. Clinicians should consider recommending this approach for patients with overweight or obesity, particularly when conventional Mediterranean dietary advice alone has proven insufficient. This multicomponent lifestyle modification represents a practical and sustainable strategy that could be incorporated into routine clinical practice for diabetes prevention.

From CIBER Fisiopatología de la Obesidad y Nutrición (CIBERObn), Instituto de Salud Carlos III, Madrid, Spain; and University of Navarra, Department of Preventive Medicine and

Public Health, Instituto de Investigación Sanitario de Navarra (IdiSNA), Pamplona, Spain (M.R.-C.); CIBER Fisiopatología de la Obesidad y Nutrición (CIBERObn), Instituto de Salud Carlos III, Madrid, Spain; and Department of Preventive Medicine, University of Valencia, Valencia, Spain (D.C., J.V.S., R.F.-C.); CIBER Fisiopatología de la Obesidad y Nutrición (CIBERObn), Instituto de Salud Carlos III, Madrid, Spain; Department of Preventive Medicine and Public Health, Instituto de Investigación Sanitario de Navarra (IdiSNA), University of Navarra, Pamplona, Spain; and Department of Nutrition, Harvard T.H. Chan School of Public Health, Boston, Massachusetts (M.Á.M.-G.); Universitat Rovira i Virgili, Departament de Bioquímica i Biotecnologia, Alimentació, Nutrició, Desenvolupament i Salut Mental (ANUT-DSM), Unitat de Nutrició Humana, Reus, Spain; and CIBER Fisiopatología de la Obesidad y Nutrición (CIBERObn), Instituto de Salud Carlos III, Madrid, Spain (N.B.); CIBER Fisiopatología de la Obesidad y Nutrición (CIBERObn), Instituto de Salud Carlos III, Madrid, Spain; Department of Nutrition, Food Sciences, and Physiology, Center for Nutrition Research, University of Navarra, Pamplona, Spain; Precision Nutrition and Cardiometabolic Health Program. IMDEA Food, CEI UAM + CSIC, Madrid, Spain; and Center of Endocrinology and Nutrition Research, Universidad de Valladolid, Valladolid, Spain (J.A.M.); Department of Endocrinology and Nutrition, Hospital Universitario de Navarra, Instituto de investigación Sanitaria de Navarra (IdiSNA), Pamplona, Spain (L.F.); CIBER Fisiopatología de la Obesidad y Nutrición (CIBERObn), Instituto de Salud Carlos III, Madrid, Spain; Bioaraba Health Research Institute, Cardiovascular, Vitoria-Gasteiz, Spain; Osakidetza Basque Health Service, Araba University Hospital, Vitoria-Gasteiz, Spain; and University of the Basque EHU, Vitoria-Gasteiz, Spain/EHU, Vitoria-Gasteiz, Spain (Á.M.A.-G.); CIBER Fisiopatología de la Obesidad y Nutrición (CIBERObn), Instituto de Salud Carlos III, Madrid, Spain; and Department of Nursing, Institute of Biomedical Research in Malaga (IBIMA), University of Málaga, Málaga, Spain (J.W.); CIBER de Epidemiología y Salud Pública (CIBERESP), Instituto de Salud Carlos III, Madrid, Spain; and Instituto de Investigación Sanitaria y Biomédica de Alicante. Universidad Miguel Hernández (ISABIAL-UMH). Alicante, Spain (J.V., S.G.-P.); CIBER Fisiopatología de la Obesidad y Nutrición (CIBERObn), Instituto de Salud Carlos III, Madrid, Spain; and Research Group on Nutritional Epidemiology & Cardiovascular Physiopathology (NUTRECOR), Health Research Institute of the Balearic Islands (IdISBa), Palma de Mallorca, Spain (D.R., J.K.); CIBER Fisiopatología de la Obesidad y Nutrición (CIBERObn), Instituto de Salud Carlos III, Madrid, Spain; and Department of Internal Medicine, Maimonides Biomedical Research Institute of Cordoba (IMIBIC), Reina Sofía University Hospital, University of Cordoba, Cordoba, Spain (J.L.-M., A.G.-R., A.P.A.-L.); CIBER Fisiopatología de la Obesidad y Nutrición (CIBERObn), Instituto de Salud Carlos III, Madrid, Spain; and Department of Internal Medicine, Institut d'Investigacions Biomèdiques August Pi Sunyer (IDIBAPS), Hospital Clinic, INSA-UB Nutrition and Food Safety Research Institute, University of Barcelona, Barcelona, Spain (R.E., R.C.); CIBER Fisiopatología de la Obesidad y Nutrición (CIBERObn), Instituto de Salud Carlos III, Madrid, Spain; and Department of Medicine, University of Sevilla, Sevilla, Spain (J.M.S.-L.); CIBER Fisiopatología de la Obesidad y Nutrición (CIBERObn), Instituto

de Salud Carlos III, Madrid, Spain; and Research Institute of Biomedical and Health Sciences (IUIBS), University of Las Palmas de Gran Canaria & Centro Hospitalario Universitario Insular Materno Infantil (CHUIMI), Canarian Health Service, Las Palmas de Gran Canaria, Spain (L.S.-M.); CIBER de Epidemiología y Salud Pública (CIBERESP), Instituto de Salud Carlos III, Madrid, Spain; Department of Preventive Medicine and Public Health, University of Granada, Granada, Spain; and Instituto de Investigación Biosanitaria ibs.GRANADA, Granada, Spain (A.B.-C., N.C.-I.); CIBER Fisiopatología de la Obesidad y Nutrición (CIBERObn), Instituto de Salud Carlos III, Madrid, Spain; and Research Group on Community Nutrition & Oxidative Stress, University of Balearic Islands, Palma de Mallorca, Spain (J.A.T.); CIBER de Epidemiología y Salud Pública (CIBERESP), Instituto de Salud Carlos III, Madrid, Spain; and Institute of Biomedicine (IBIOMED), University of León, León, Spain (V.M.-S.); CIBER Fisiopatología de la Obesidad y Nutrición (CIBERObn), Instituto de Salud Carlos III, Madrid, Spain; and Department of Internal Medicine, Hospital Universitari de Bellvitge-Bellvitge Biomedical Research Institute (IDIBELL), L'Hospitalet de Llobregat, Faculty of Medicine and Health Sciences, Universitat de Barcelona, Barcelona, Spain (A.R.-M.); CIBER de Epidemiología y Salud Pública (CIBERESP), Instituto de Salud Carlos III, Madrid, Spain; and Division of Preventive Medicine, Faculty of Medicine, University of Jaén, Jaén, Spain (M.D.-R.); Department of Endocrinology and Nutrition, Instituto de Investigación Sanitaria Hospital Clínico San Carlos (IdISSC), Madrid, Spain; and CIBER Diabetes y Enfermedades Metabólicas (CIBERDEM), Instituto de Salud Carlos III (ISCIII), Madrid, Spain (P.M.-M.); CIBER Diabetes y Enfermedades Metabólicas (CIBERDEM), Instituto de Salud Carlos III (ISCIII), Madrid, Spain; and Department of Endocrinology, Institut d'Investigacions Biomèdiques August Pi Sunyer (IDIBAPS), Hospital Clinic, University of Barcelona, Barcelona, Spain (J.V.); CIBER Fisiopatología de la Obesidad y Nutrición (CIBERObn), Instituto de Salud Carlos III, Madrid, Spain; and Department of Endocrinology and Nutrition, Hospital Fundación Jiménez Díaz, Instituto de Investigaciones Biomédicas (IISFDJ), University Autonoma, Madrid, Spain (C.V.); CIBER Fisiopatología de la Obesidad y Nutrición (CIBERObn), Instituto de Salud Carlos III, Madrid, Spain; and Nutritional Control of the Epigenome Group, Precision Nutrition and Obesity Program, IMDEA Food, CEI UAM + CSIC, Madrid, Spain (L.D.); CIBER Fisiopatología de la Obesidad y Nutrición (CIBERObn), Instituto de Salud Carlos III, Madrid, Spain; Department of Preventive Medicine and Public Health, Instituto de Investigación Sanitario de Navarra (IdiSNA), University of Navarra, Pamplona, Spain; and Atención Primaria, Servicio Navarro de Salud, Pamplona, Spain (P.B.-C.); CIBER Fisiopatología de la Obesidad y Nutrición (CIBERObn), Instituto de Salud Carlos III, Madrid, Spain; Departament de Bioquímica i Biotecnologia, Alimentació, Nutrició, Desenvolupament i Salut Mental (ANUT-DSM), Unitat de Nutrició Humana, Universitat Rovira i Virgili, Reus, Spain; and Institut D'Investigació Sanitària Pere Virgili (IISPV), Reus, Spain (S.S., J.G.-G.); CIBER de Epidemiología y Salud Pública (CIBERESP), Instituto de Salud Carlos III, Madrid, Spain; and Unit of Cardiovascular Risk and Nutrition, Hospital del Mar Research Institute (IMIM), Barcelona, Spain (O.C., H.S.); CIBER de Epidemiología y Salud Pública (CIBERESP), Instituto de Salud

Carlos III, Madrid, Spain; and Instituto de Investigación Sanitaria y Biomédica de Alicante, Universidad Miguel Hernández (ISABIAL-UMH), Alicante, Spain (L.T.-C.); Department of Preventive Medicine. Institute of Biomedical Research in Malaga (IBIMA), University of Málaga, Málaga, Spain (E.G.-G.); CIBER Fisiopatología de la Obesidad y Nutrición (CIBERObn), Instituto de Salud Carlos III, Madrid, Spain; and Department of Nutrition, Food Sciences, and Physiology, Center for Nutrition Research, University of Navarra, Pamplona, Spain (M.Á.Z.); CIBER Fisiopatología de la Obesidad y Nutrición (CIBERObn), Instituto de Salud Carlos III, Madrid, Spain; Cardiovascular, Respiratory and Metabolic Area, Bioaraba Health Research Institute, Vitoria-Gasteiz, Spain; Osakidetza Basque Health Service, Araba University Hospital, Vitoria-Gasteiz, Spain; and University of the Basque Country UPV/EHU, Vitoria-Gasteiz, Spain (L.T.-S.); CIBER Fisiopatología de la Obesidad y Nutrición (CIBERObn), Instituto de Salud Carlos III, Madrid, Spain; and Regional University Hospital of Malaga, Department of Internal Medicine. Instituto de Investigación Biomédica de Málaga (IBIMA-Plataforma Bionand), University of Málaga, Málaga, Spain (R.M.B.-L.); CIBER Fisiopatología de la Obesidad y Nutrición (CIBERObn), Instituto de Salud Carlos III, Madrid, Spain; and Department of Preventive Medicine and Public Health, Instituto de Investigación Sanitario de Navarra (IdiSNA), University of Navarra, Pamplona, Spain (E.T.); Fisiopatología de la Obesidad y Nutrición (CIBERObn), Instituto de Salud Carlos III, Madrid, Spain; Unit of Cardiovascular Risk and Nutrition, Hospital del Mar Research Institute (IMIM), Barcelona, Spain; and Department of Medicine and Life Sciences, Universitat Pompeu Fabra, Barcelona, Spain (A.G.); CIBER Fisiopatología de la Obesidad y Nutrición (CIBERObn), Instituto de Salud Carlos III, Madrid, Spain; and IDIBAPS, Hospital Clínic, Barcelona, Spain (E.R.); CIBER Fisiopatología de la Obesidad y Nutrición (CIBERObn), Instituto de Salud Carlos III, Madrid, Spain; and Unit of Cardiovascular Risk and Nutrition, Hospital del Mar Research Institute (IMIM), Barcelona, Spain (M.F.); Department of Nutrition, Harvard T.H. Chan School of Public Health, Boston; Department of Biostatistics, Harvard T.H. Chan School of Public Health, Boston; and Channing Division of Network Medicine, Department of Medicine, Brigham and Women's Hospital and Harvard Medical School, Boston, Massachusetts (F.B.H.); CIBER Fisiopatología de la Obesidad y Nutrición (CIBERObn), Instituto de Salud Carlos III, Madrid, Spain; and Virgen de la Victoria Hospital, Department of Endocrinology, Instituto de Investigación Biomédica de Málaga (IBIMA). University of Málaga, Málaga, Spain (F.J.T.); and Universitat Rovira i Virgili, Departament de Bioquímica i Biotecnologia, Alimentació, Nutrició, Desenvolupament i Salut Mental (ANUT-DSM), Unitat de Nutrició Humana, Reus, Spain; CIBER Fisiopatología de la Obesidad y Nutrición (CIBERObn), Instituto de Salud Carlos III, Madrid, Spain; and Institut d'Investigació Sanitària Pere Virgili (IISPV), Reus, Spain (J.S.-S.).

Note: Drs. Ruiz-Canela and Salas-Salvadó had full access to all of the data in the study and take responsibility for the integrity of the data and the accuracy of the data analysis.

Acknowledgment: The authors especially thank the PREDIMED-Plus participants for the enthusiastic collaboration, the PREDIMED-Plus personnel for outstanding support, and the personnel

of all associated primary care centers for the exceptional effort. CIBEROBN, CIBERESP, and CIBERDEM are initiatives of Instituto de Salud Carlos III (ISCIII), Madrid, Spain. The authors also thank the PREDIMED-Plus Biobank Network as a part of the National Biobank Platform of the ISCIII for storing and managing the PREDIMED-Plus biological samples.

Members of the PREDIMED-Plus Steering, Executive, Dietary and Lifestyle Intervention, and Clinical Event Ascertainment Committees, as well as the Support groups, can be found in the Collaborators section of **Supplement 1** (available at [Annals.org](https://annals.org)).

Grant Support: This work was supported by the official Spanish Institutions for funding scientific biomedical research, CIBER Fisiopatología de la Obesidad y Nutrición (CIBEROBN) and Instituto de Salud Carlos III (ISCIII), through the Fondo de Investigación para la Salud (FIS), which is co-funded by the European Regional Development Fund (6 coordinated FIS projects led by Drs. Vidal and Salas-Salvadó, including the following projects: PI13/00673, PI13/00492, PI13/00272, PI13/01123, PI13/00462, PI13/00233, PI13/02184, PI13/00728, PI13/01090, PI13/01056, PI14/01722, PI14/00636, PI14/00618, PI14/00696, PI14/01206, PI14/01919, PI14/00853, PI14/01374, PI14/00972, PI14/00728, PI14/01471, PI16/00473, PI16/00662, PI16/01873, PI16/01094, PI16/00501, PI16/00533, PI16/00381, PI16/00366, PI16/01522, PI16/01120, PI17/00764, PI17/01183, PI17/00855, PI17/01347, PI17/00525, PI17/01827, PI17/00532, PI17/00215, PI17/01441, PI17/00508, PI17/01732, PI17/00926, PI19/00957, PI19/00386, PI19/00309, PI19/01032, PI19/00576, PI19/00017, PI19/01226, PI19/00781, PI19/01560, PI19/01332, PI20/01802, PI20/00138, PI20/01532, PI20/00456, PI20/00339, PI20/00557, PI20/00886, PI20/01158); by National Institutes of Health National Institute of Diabetes and Digestive and Kidney Diseases grant 1R01DK127601 (Drs. Ruiz-Canela, Hu, and Salas-Salvadó); by European Research Council advanced research grant 2014-2019, agreement 340918 (M.Á. Martínez-González); by the Especial Action Project entitled Implementación y evaluación de una intervención intensiva sobre la actividad física Cohorte, PREDIMED-Plus grant (Dr. Salas-Salvadó); by Recercaixa grant 2013ACUP00194 (Dr. Salas-Salvadó); by grants from the Consejería de Salud de la Junta de Andalucía (PI0458/2013, PS0358/2016, PI0137/2018); by the PROMETEO/2017/017 and PROMETEO/2021/021 grants from the Generalitat Valenciana; by the SEMERGEN grant; by INSA-Ma María de Maeztu Unit of Excellence grant CEX2021-001234-M (funded by MICIN/AEI/FEDER, UE); by de Gestió d'Ajuts Universitaris i de Recerca (2021 SGR 00144) to Dr. Fitó; by Miguel Servet grant CP24/00089 from Instituto de Salud Carlos III (ISCIII), co-funded by the European Union (Dr. Konieczna); and by MINECO grant CNS2022-135862 (Dr. Romaguera). Dr. Salas-Salvadó was partially supported by ICREA under the ICREA Academia program.

Disclosures: Disclosure forms are available with the article online.

Data Sharing Statement: A controlled data sharing model is followed in the PREDIMED-Plus study using pseudoanonymized (deidentified) study data only, for collaborating with

approved researchers. Requestors wishing to access the PREDIMED-Plus trial data used in this study for collaboration can request it from the PREDIMED-Plus Trial Steering Committee at predimed_plus_scommittee@googlegroups.com. Requests are considered by the PREDIMED-Plus Steering Committee composed of J. Salas-Salvadó (Chair), M.Á. Martínez-González, M. Fitó, E. Ros, F. Tinahones, D. Corella, and R. Estruch. Decisions on data access are based on the scientific legitimacy of the requester and of their institution, and on assurances on information security and governance; and with regard to the study's scientific reputation, the needs of funded study team research, the terms of participant consent, and regulatory requirements. All research using PREDIMED-Plus must be within the scope of the ethically approved PREDIMED-Plus Protocol. Studies where data and/or biological samples are requested are subject to ethics approval by the institutional review boards of the corresponding PREDIMED Plus recruitment centers.

Corresponding Author: Jordi Salas-Salvadó, MD, PhD, Universitat Rovira i Virgili, Department of Biochemistry and Biotechnology, Human Nutrition Unit. C/Sant Llorenç 21, 43201, Reus, Tarragona, Spain (e-mail, jordi.salas@urv.cat); and Montserrat Fitó, MD, PhD, Hospital del Mar Medical Research Institute (IMIM), Barcelona 08003 Spain (e-mail, mfito@researchmar.net).

Author contributions are available at Annals.org.

References

- Zheng Y, Ley SH, Hu FB. Global aetiology and epidemiology of type 2 diabetes mellitus and its complications. *Nat Rev Endocrinol*. 2018;14:88-98. [PMID: 29219149] doi:10.1038/nrendo.2017.151
- GBD 2021 Diabetes Collaborators. Global, regional, and national burden of diabetes from 1990 to 2021, with projections of prevalence to 2050: a systematic analysis for the Global Burden of Disease Study 2021. *Lancet*. 2023;402:203-234. [PMID: 37356446] doi:10.1016/S0140-6736(23)01301-6
- Emerging Risk Factors Collaboration. Life expectancy associated with different ages at diagnosis of type 2 diabetes in high-income countries: 23 million person-years of observation. *Lancet Diabetes Endocrinol*. 2023;11:731-742. [PMID: 37708900] doi:10.1016/S2213-8587(23)00223-1
- Diabetes and Nutrition Study Group (DNSG) of the European Association for the Study of Diabetes (EASD). Evidence-based European recommendations for the dietary management of diabetes. *Diabetologia*. 2023;66:965-985. [PMID: 37069434] doi:10.1007/s00125-023-05894-8
- Uusitupa M, Khan TA, Vigiouliou E, et al. Prevention of type 2 diabetes by lifestyle changes: a systematic review and meta-analysis. *Nutrients*. 2019;11:2611. [PMID: 31683759] doi:10.3390/nu11112611
- Pan XR, Li GW, Hu YH, et al. Effects of diet and exercise in preventing NIDDM in people with impaired glucose tolerance. The Da Qing IGT and Diabetes Study. *Diabetes Care*. 1997;20:537-544. [PMID: 9096977] doi:10.2337/diacare.20.4.537
- Tuomilehto J, Lindström J, Eriksson JG, et al; Finnish Diabetes Prevention Study Group. Prevention of type 2 diabetes mellitus by changes in lifestyle among subjects with impaired glucose tolerance. *N Engl J Med*. 2001;344:1343-1350. [PMID: 11333990] doi:10.1056/NEJM200105033441801
- Knowler WC, Barrett-Connor E, Fowler SE, et al; Diabetes Prevention Program Research Group. Reduction in the incidence of type 2 diabetes with lifestyle intervention or metformin. *N Engl J Med*. 2002;346:393-403. [PMID: 11832527] doi:10.1056/NEJMoa012512
- Haw JS, Galaviz KI, Straus AN, et al. Long-term sustainability of diabetes prevention approaches: a systematic review and meta-analysis of randomized clinical trials. *JAMA Intern Med*. 2017;177:1808-1817. [PMID: 29114778] doi:10.1001/jamainternmed.2017.6040
- Salas-Salvadó J, Bulló M, Babio N, et al; PREDIMED Study Investigators. Reduction in the incidence of type 2 diabetes with the Mediterranean diet: results of the PREDIMED-Reus nutrition intervention randomized trial. *Diabetes Care*. 2011;34:14-19. [PMID: 20929998] doi:10.2337/dc10-1288
- Salas-Salvadó J, Bulló M, Estruch R, et al. Prevention of diabetes with Mediterranean diets: a subgroup analysis of a randomized trial. *Ann Intern Med*. 2014;160:1-10. [PMID: 24573661] doi:10.7326/M13-1725
- Estruch R, Martínez-González MA, Corella D, et al; PREDIMED Study Investigators. Effect of a high-fat Mediterranean diet on body-weight and waist circumference: a prespecified secondary outcomes analysis of the PREDIMED randomised controlled trial. *Lancet Diabetes Endocrinol*. 2019;7:e6-e17. [PMID: 31003626] doi:10.1016/S2213-8587(19)30074-9
- Salas-Salvadó J, Díaz-López A, Ruiz-Canela M, et al; PREDIMED-Plus Investigators. Effect of a lifestyle intervention program with energy-restricted Mediterranean diet and exercise on weight loss and cardiovascular risk factors: one-year results of the PREDIMED-Plus trial. *Diabetes Care*. 2019;42:777-788. [PMID: 30389673] doi:10.2337/dc18-0836
- Sayón-Orea C, Razquin C, Bulló M, et al. Effect of a nutritional and behavioral intervention on energy-reduced Mediterranean diet adherence among patients with metabolic syndrome: interim analysis of the PREDIMED-Plus randomized clinical trial. *JAMA*. 2019;322:1486-1499. [PMID: 31613346] doi:10.1001/jama.2019.14630
- Konieczna J, Ruiz-Canela M, Galmes-Panades AM, et al. An energy-reduced Mediterranean diet, physical activity, and body composition: an interim subgroup analysis of the PREDIMED-Plus randomized clinical trial. *JAMA Netw Open*. 2023;6:e2337994. [PMID: 37851444] doi:10.1001/jamanetworkopen.2023.37994
- Martínez-González MA, Buil-Cosiales P, Corella D, et al; PREDIMED-Plus Study Investigators. Cohort profile: design and methods of the PREDIMED-Plus randomized trial. *Int J Epidemiol*. 2019;48:387-388. [PMID: 30476123] doi:10.1093/ije/dyy225
- Alberti KGMM, Eckel RH, Grundy SM, et al; International Association for the Study of Obesity. Harmonizing the metabolic syndrome: a joint interim statement of the International Diabetes Federation Task Force on Epidemiology and Prevention; National Heart, Lung, and Blood Institute; American Heart Association; World Heart Federation; International Atherosclerosis Society; and International Association for the Study of Obesity. *Circulation*. 2009;120:1640-1645. [PMID: 19805654] doi:10.1161/CIRCULATIONAHA.109.192644
- Schröder H, Cárdenas-Fuentes G, Martínez-González MA, et al; PREDIMED-Plus Investigators. Effectiveness of the physical activity intervention program in the PREDIMED-Plus study: a randomized controlled trial. *Int J Behav Nutr Phys Act*. 2018;15:110. [PMID: 30424822] doi:10.1186/s12966-018-0741-x
- Schröder H, Zomeño MD, Martínez-González MA, et al; PREDIMED-Plus Investigators. Validity of the energy-restricted Mediterranean Diet Adherence Screener. *Clin Nutr*. 2021;40:4971-4979. [PMID: 34364236] doi:10.1016/j.clnu.2021.06.030
- Estruch R, Ros E, Salas-Salvadó J, et al; PREDIMED Study Investigators. Primary prevention of cardiovascular disease with a Mediterranean diet supplemented with extra-virgin olive oil or nuts. *N Engl J Med*. 2018;378:e34. [PMID: 29897866] doi:10.1056/NEJMoa1800389
- Schröder H, Fitó M, Estruch R, et al. A short screener is valid for assessing Mediterranean diet adherence among older Spanish men and women. *J Nutr*. 2011;141:1140-1145. [PMID: 21508208] doi:10.3945/jn.110.135566

22. Paz-Graniel I, Fitó M, Ros E, et al. Impact of COVID-19 pandemic on the PREDIMED-Plus randomized clinical trial: effects on the interventions, participants follow-up, and adiposity. *Front Nutr*. 2022;9:1098269. [PMID: 36712515] doi:10.3389/fnut.2022.1098269
23. Martin-Moreno JM, Boyle P, Gorgojo L, et al. Development and validation of a food frequency questionnaire in Spain. *Int J Epidemiol*. 1993;22:512-519. [PMID: 8359969] doi:10.1093/ije/22.3.512
24. Molina L, Sarmiento M, Peñafiel J, et al. Validation of the Regicor Short Physical Activity Questionnaire for the Adult Population. *PLoS One*. 2017;12:e0168148. [PMID: 28085886] doi:10.1371/journal.pone.0168148
25. Topolski TD, LoGerfo J, Patrick DL, et al. The Rapid Assessment of Physical Activity (RAPA) among older adults. *Prev Chronic Dis*. 2006;3:A118. [PMID: 16978493]
26. Matthews DR, Hosker JP, Rudenski AS, et al. Homeostasis model assessment: insulin resistance and beta-cell function from fasting plasma glucose and insulin concentrations in man. *Diabetologia*. 1985;28:412-419. [PMID: 3899825] doi:10.1007/BF00280883
27. World Health Organization. Definition and diagnosis of diabetes mellitus and intermediate hyperglycemia: report of a WHO/IDF consultation. World Health Organization; 2006.
28. American Diabetes Association. 2. Classification and Diagnosis of Diabetes: Standards of Medical Care in Diabetes-2020. *Diabetes Care*. 2020;43:S14-S31. [PMID: 31862745] doi:10.2337/dc20-S002
29. Lanza A, Soler R, Smith B, et al. The Diabetes Prevention Impact Tool Kit: an online tool kit to assess the cost-effectiveness of preventing type 2 diabetes. *J Public Health Manag Pract*. 2019;25:E1-E5. [PMID: 31348170] doi:10.1097/PHH.0000000000000961
30. American Diabetes Association. Standards of medical care in diabetes-2014. *Diabetes Care*. 2014;37 Suppl 1:S14-S80. [PMID: 24357209] doi:10.2337/dc14-S014
31. Kaplan EL, Meier P. Nonparametric estimation from incomplete observations. *J Am Stat Assoc*. 1958;53:457-481. doi:10.1080/01621459.1958.10501452
32. Cox DR. Regression models and life-tables. *J R Stat Soc Series B Stat Methodol*. 1972;34:187-220. doi:10.1111/j.2517-6161.1972.tb00899.x
33. Jenkins DJA, Willett WC, Yusuf S, et al; Clinical Nutrition & Risk Factor Modification Centre Collaborators. Association of glycaemic index and glycaemic load with type 2 diabetes, cardiovascular disease, cancer, and all-cause mortality: a meta-analysis of mega cohorts of more than 100000 participants. *Lancet Diabetes Endocrinol*. 2024;12:107-118. [PMID: 38272606] doi:10.1016/S2213-8587(23)00344-3
34. Heo J-W, No M-H, Cho J, et al. Moderate aerobic exercise training ameliorates impairment of mitochondrial function and dynamics in skeletal muscle of high-fat diet-induced obese mice. *FASEB J*. 2021;35:e21340. [PMID: 33455027] doi:10.1096/fj.202002394R
35. Konieczna J, Abete I, Galmés AM, et al; PREDIMED-Plus Investigators. Body adiposity indicators and cardiometabolic risk: cross-sectional analysis in participants from the PREDIMED-Plus trial. *Clin Nutr*. 2019;38:1883-1891. [PMID: 30031660] doi:10.1016/j.clnu.2018.07.005
36. Lee DH, Jin Q, Shi N, et al. The metabolic potential of inflammatory and insulinaemic dietary patterns and risk of type 2 diabetes. *Diabetologia*. 2024;67:88-101. [PMID: 37816982] doi:10.1007/s00125-023-06021-3
37. Fujiwara Y, Eguchi S, Murayama H, et al. Relationship between diet/exercise and pharmacotherapy to enhance the GLP-1 levels in type 2 diabetes. *Endocrinol Diabetes Metab*. 2019;2:e00068. [PMID: 31294084] doi:10.1002/edm2.68
38. García-Gavilán JF, Atzeni A, Babio N, et al. Effect of 1-year lifestyle intervention with energy-reduced Mediterranean diet and physical activity promotion on the gut metabolome and microbiota: a randomized clinical trial. *Am J Clin Nutr*. 2024;119:1143-1154. [PMID: 38428742] doi:10.1016/j.ajcnut.2024.02.021
39. Diabetes Prevention Program Research Group. Long-term effects of lifestyle intervention or metformin on diabetes development and microvascular complications over 15-year follow-up: the Diabetes Prevention Program Outcomes Study. *Lancet Diabetes Endocrinol*. 2015;3:866-875. [PMID: 26377054] doi:10.1016/S2213-8587(15)00291-0
40. Martínez-González MA, Gea A, Ruiz-Canela M. The Mediterranean diet and cardiovascular health. *Circ Res*. 2019;124:779-798. [PMID: 30817261] doi:10.1161/CIRCRESAHA.118.313348
41. Ramachandran A, Snehalatha C, Mary S, et al; Indian Diabetes Prevention Programme (IDPP). The Indian Diabetes Prevention Programme shows that lifestyle modification and metformin prevent type 2 diabetes in Asian Indian subjects with impaired glucose tolerance (IDPP-1). *Diabetologia*. 2006;49:289-297. [PMID: 16391903] doi:10.1007/s00125-005-0097-z
42. Saito T, Watanabe M, Nishida J, et al; Zensharen Study for Prevention of Lifestyle Diseases Group. Lifestyle modification and prevention of type 2 diabetes in overweight Japanese with impaired fasting glucose levels: a randomized controlled trial. *Arch Intern Med*. 2011;171:1352-1360. [PMID: 21824948] doi:10.1001/archinternmed.2011.275
43. Egúaras S, Bes-Rastrollo M, Ruiz-Canela M, et al. May the Mediterranean diet attenuate the risk of type 2 diabetes associated with obesity: the Seguimiento Universidad de Navarra (SUN) cohort. *Br J Nutr*. 2017;117:1478-1485. [PMID: 28625175] doi:10.1017/S0007114517001404
44. Perreault L, Ma Y, Dagogo-Jack S, et al; Diabetes Prevention Program. Sex differences in diabetes risk and the effect of intensive lifestyle modification in the Diabetes Prevention Program. *Diabetes Care*. 2008;31:1416-1421. [PMID: 18356403] doi:10.2337/dc07-2390
45. Wang R, Lagakos SW, Ware JH, et al. Statistics in medicine-reporting of subgroup analyses in clinical trials. *N Engl J Med*. 2007;357:2189-2194. [PMID: 18032770] doi:10.1056/NEJMSr077003
46. Evert AB, Dennison M, Gardner CD, et al. Nutrition therapy for adults with diabetes or prediabetes: a consensus report. *Diabetes Care*. 2019;42:731-754. [PMID: 31000505] doi:10.2337/dci19-0014
47. Martínez-González MA, Hershey MS, Zazpe I, et al. Transferability of the Mediterranean diet to non-Mediterranean countries. What is and what is not the Mediterranean diet. *Nutrients*. 2017;9:1226. [PMID: 29117146] doi:10.3390/nu9111226

Author Contributions: Conception and design: D. Corella, M.Á. Martínez-González, J.A. Martínez, J. Wärnberg, J. López-Miranda, R. Estruch, J.A. Tur, J. Vidal, L. Daimiel, E. Gómez-Gracia, E. Toledo, E. Ros, M. Fitó, F.B. Hu, F.J. Tinahones, J. Salas-Salvadó.

Analysis and interpretation of the data: M. Ruiz-Canela, D. Corella, M.Á. Martínez-González, N. Babio, J.A. Martínez, J. Vioque, J. López-Miranda, R. Estruch, J.M. Santos-Lozano, L. Serra-Majem, J.A. Tur, A. Riera-Mestre, M. Delgado-Rodríguez, L. Daimiel, S. Shyam, E. Gómez-Gracia, R. Casas, E. Toledo, A. Goday, H. Schröder, M. Fitó, F.B. Hu, F.J. Tinahones, J. Salas-Salvadó.

Drafting of the article: M. Ruiz-Canela, D. Corella, J.A. Martínez, J. López-Miranda, M. Delgado-Rodríguez, J.V. Sorlí, A. García-Ríos, E. Gómez-Gracia, J.F. García-Gavilán, A. Goday, M. Fitó, J. Salas-Salvadó.

Critical revision of the article for important intellectual content: D. Corella, M.Á. Martínez-González, N. Babio, J.A. Martínez, J. Wärnberg, J. Vioque, D. Romaguera, J. López-Miranda, R. Estruch, A. Bueno-Cavanillas, A. Riera-Mestre, P. Matía-Martín, J. Vidal, L. Daimiel, P. Buil-Cosiales, S. Shyam, O. Castañer, L. Torres-Collado, J. Konieczna, R. Casas, N. Cano-Ibañez, E. Toledo, J. García-Gavilán, A. Goday, S. González-Palacios, H. Schröder, E. Ros, M. Fitó, F.B. Hu, F.J. Tinahones, J. Salas-Salvadó.

Final approval of the article: M. Ruiz-Canela, D. Corella, M.Á. Martínez-González, N. Babio, J.A. Martínez, L. Forga, Á.M. Alonso-Gómez, J. Wärnberg, J. Vioque, D. Romaguera, J. López-Miranda, R. Estruch, J.M. Santos-Lozano, L. Serra-Majem, A. Bueno-Cavanillas, J.A. Tur, V. Martín-Sánchez, A. Riera-Mestre, M. Delgado-Rodríguez, P. Matía-Martín, J. Vidal, C. Vázquez, L. Daimiel, P. Buil-Cosiales, S. Shyam, J.V. Sorlí, O. Castañer, A. García-Ríos, L. Torres-Collado, E. Gómez-Gracia, M.Á. Zulet, J. Konieczna, R. Casas, N. Cano-Ibañez, L. Tojal-Sierra, R.M. Bernal-

López, E. Toledo, J. García-Gavilán, R. Fernández-Carrión, A. Goday, A.P. Arenas-Larriva, S. González-Palacios, H. Schröder, E. Ros, M. Fitó, F.B. Hu, F.J. Tinahones, J. Salas-Salvadó.

Provision of study materials or patients: D. Corella, M.Á. Martínez-González, N. Babio, J.A. Martínez, J. Wärnberg, J. Vioque, D. Romaguera, J. López-Miranda, J.M. Santos-Lozano, L. Serra-Majem, A. Bueno-Cavanillas, J.A. Tur, V. Martín-Sánchez, M. Delgado-Rodríguez, P. Matía-Martín, C. Vázquez, L. Daimiel, J.V. Sorlí, O. Castañer, M.Á. Zulet, R.M. Bernal-López, R. Fernández-Carrión, F.J. Tinahones, J. Salas-Salvadó.

Statistical expertise: M. Ruiz-Canela, D. Corella, M.Á. Martínez-González, J.A. Tur, L. Daimiel, S. Shyam, E. Gómez-Gracia, J. García-Gavilán, J. Salas-Salvadó.

Obtaining of funding: M. Ruiz-Canela, D. Corella, M.Á. Martínez-González, J.A. Martínez, Á.M. Alonso-Gómez, J. Wärnberg, J. Vioque, D. Romaguera, J. López-Miranda, R. Estruch, J.M. Santos-Lozano, L. Serra-Majem, A. Bueno-Cavanillas, J.A. Tur, V. Martín-Sánchez, M. Delgado-Rodríguez, J. Vidal, L. Daimiel, J.V. Sorlí, M.Á. Zulet, J. Konieczna, R.M. Bernal-López, A. Goday, M. Fitó, F.B. Hu, F.J. Tinahones, J. Salas-Salvadó.

Administrative, technical, or logistic support: M.Á. Martínez-González, L. Forga, R. Estruch, L. Serra-Majem, J.A. Tur, E. Gómez-Gracia, R. Casas, E. Toledo, F.B. Hu, J. Salas-Salvadó.

Collection and assembly of data: D. Corella, M.Á. Martínez-González, N. Babio, J.A. Martínez, L. Forga, Á.M. Alonso-Gómez, J. Wärnberg, J. Vioque, D. Romaguera, J. López-Miranda, L. Serra-Majem, A. Bueno-Cavanillas, J.A. Tur, V. Martín-Sánchez, A. Riera-Mestre, M. Delgado-Rodríguez, J. Vidal, C. Vázquez, L. Daimiel, P. Buil-Cosiales, J.V. Sorlí, O. Castañer, A. García-Ríos, J. Konieczna, L. Tojal-Sierra, R.M. Bernal-López, E. Toledo, R. Fernández-Carrión, A. Goday, M. Fitó, F.J. Tinahones, J. Salas-Salvadó.

Supplementary Material*

Ruiz-Canela M, Corella D, Martínez-González MÁ, et al. Comparison of an energy-reduced Mediterranean diet and physical activity versus an ad libitum Mediterranean diet in the prevention of type 2 diabetes. A secondary analysis of a randomized controlled trial. *Ann Intern Med*. 26 August 2025. [Epub ahead of print]. doi:10.7326/ANNALS-25-00388

I. Collaborators	2
II. Supplement Methods	7
III. Supplement Tables and Figures.....	8
Supplement Table 1. Intensive intervention program (intervention group)	8
Supplement Table 2. 17-item questionnaire of adherence to an energy-restricted Mediterranean diet.....	9
Supplement Table 3. Baseline characteristic of participants in the PREDIMED-Plus trial.....	10
Supplement Table 4. Use of Drugs (%) during the follow-up by intervention group	11
Supplement Table 5. Hazard ratios for incident type 2 diabetes in the PREDIMED-Plus trial: Subgroup analyses.....	12
Supplement Table 6. Absolute and relative risk estimates for the composite outcome of incident type 2 diabetes and death before diabetes diagnosis after 6 years of intervention.....	14
Supplement Table 7. Participants with positive answers to each of the 17 Items of the Mediterranean diet score by intervention group during Follow-up.....	15
Supplement Table 8. Body weight and waist circumference reduction throughout the trial by intervention group	17
Supplement Table 9. Absolute risk and rates difference estimates for incident type 2 diabetes throughout the entire 7-year follow-up.....	18
Supplement Figure. Ancillary analyses: Hazard ratios and confidence intervals (CI) for intervention versus control group under different assumptions and/or exclusions.	19

* This supplementary material was provided by the authors to give readers further details on their article. The material was not copyedited.

I. Collaborators: They have contributed to the participants recruitment and follow-up or provided technical support for the trial.

Rovira i Virgili University, Department of Biochemistry and Biotechnology, Human Nutrition Unit, University Hospital of Sant Joan de Reus, Pere Virgili Institute for Health Research, Reus, Spain: R. Pedret Llaberia, R. Gonzalez, R. Sagarra Álamo, F. París Palleja, J. Balsells, J.M. Roca, T. Basora Gallisa, J. Vizcaino, P. Llobet Alpizarte, C. Anguera Perpiñá, M. Llauredó Vernet, C. Caballero, M. Garcia Barco, M.D. Morán Martínez, J. García Rosselló, A. Del Pozo, C. Poblet Calaf, P. Arcelin Zabal, X. Floresví, M. Ciutat Benet, A. Palau Galindo, J.J. Cabré Vila, F. Dolz Andrés, M. Soler, M. Gracia Vidal, J. Vilalta J. Boj Casajuana, M. Ricard, F. Saiz, A. Isach, M. Sanchez Marin Martinez, E. Granado Font, C. Lucena Luque, C. Mestres Sola, J. Basora, F. Martín-Lujan, N. Babio, I. Abellán Cano, JF García-Gavilán, I. Paz-Graniel, C. Gomez-Martinez, L. Sánchez Niembro, A. Atzeni, C. Valle, M. Fernández de la Puente, J. Ni, S. De las Heras, C. Munné, N. Khoury, S Segura, A. Hernandez-Cacho, H.J. Margara-Escudero, H. Vázquez, I. Valverde, M.A Martínez-González, S. Shyam, S. Nishi, A. Pérez-Acosta, A. Salas-Huetos, M. Murphy.

Department of Preventive Medicine and Public Health, University of Navarra-Navarra Institute for Health Research (IdiSNA), Pamplona, Spain: Z. Vázquez, C. Razquin, M. Bes-Rastrollo, A. Gea, A. Sanchez Tainta, B. SanJulian Aranguren, E. Goñi, L. Goñi, M.J. Cobo, A. Rico-Campa, F.J. Basterra Gortari, A. Garcia Arellano, J. Diez-Espino, O. Lecea-Juarez, J. Carlos Cenoz-Osinaga, I. Alvarez-Alvarez, M.C. Sayon-Orea, C.I. Fernandez-Lázaro, L. Ruiz-Estigarribia, J. Bartolome-Resano, A. Sola-Larraz (†), E. Lozano-Oloriz, B. Cano-Valles, S. Eguaras, E. Pascual Roquet-Jalmar, I. Galilea-Zabalza, H. Lancova, R. Ramallal, M.L. Garcia-Perez, V. Estremera-Urabayen, M.J. Ariz-Arnedo, C. Hijos-Larraz, C. Fernandez-Alfaro, B. Iñigo-Martinez, R. Villanueva Moreno, S. Martin-Almendros, L. Barandiaran-Bengoetxea, C. Fuertes-Goñi, A. Lezaun-Indurain, M.J. Guruchaga-Arcelus, O. Olmedo-Cruz, L. Escriche-Erviti, R. Ansorena-Ros, R. Sanmartin-Zabaleta, J. Apalategi-Lasa, J. Villanueva-Telleria, M.M. Hernández-Espinosa, L. Herrera-Valdez, L. Dorronsoro-Dorronsoro, Lourdes Echeverria-Lizarraga (†), J.A. Cabeza-Beunza, P. Fernández-Urretavizcaya, P. Gascó-García, C. Royo-Jimenez, J. Moran-Pi, F. Salazar-Fernández, F.J. Chasco-Ros, F. Cortés-Ugalde, J.J. Jurio-Burgui, P. Pascual-Pascual, A.I. Rodríguez-Ezpeleta, M. Esparza-Cáceres, C. Arroyo-Azpa, M. Rodríguez-Sanz de Galdeano, T. Forcen-Alonso, M. Armendariz-Marcotegui, A. Brugos-Larumbe, A. Arillo, B. López-Aisa, R. Sanmartin-Zabaleta, M. Hermoso de Mendoza-Macua.

Department of Preventive Medicine, University of Valencia, University Jaume I, Conselleria de Sanitat de la Generalitat Valenciana, Valencia, Spain: J.I. González, O. Portolés, R. Fernández-Carrión, C. Ortega-Azorín, R. Barragán, E.M. Asensio, O. Coltell, I. Giménez-Alba, C. Sáiz, R. Osma, E. Ferriz, I. González-Monje, P. Guillém-Sáiz, F. Giménez-Fernández, E.C. Pascual, L. Quiles, P. Carrasco, A. Carratalá-Calvo, C. Valero-Barceló, C. Mir, S. Sánchez-Navarro, J. Navas, J.J. Tamarit, I. González-Gallego, L. Bort-Llorca, L. Pérez-Ollero, M. Giner-Valero, R. Monfort-Sáez, J. Nadal-Sayol, V. Pascual-Fuster, M. Martínez-Pérez, C. Riera, M.V. Belda, A. Medina, E. Miralles, M.J. Ramírez-Esplugues, M. Rojo-Furió, G. Mattingley, M.A. Delgado, M.A. Pages, Y. Riofrío, L. Abuomar, N. Blasco-Lafarga, R. Tosca, L. Lizán, A.M. Valcarce, M.D. Medina, S. de Valcárcel, N. Tormo, O. Felipe-Román, I.M. Romero-Martínez, F.A. Rodríguez-Lagos, J.V. Crespo, J.L. Llosa, L. González-García, R. Raga-Marí.

Cardiovascular Risk and Nutrition Research Group, Endocrinology Service, Epidemiology and Public health and Neurosciences Programmes, Clinical Research Unit at the Hospital del Mar Medical Research Institute (IMIM), Barcelona. Medicine Departament, Universitat Autònoma de Barcelona, Barcelona, Spain: M.A. Muñoz, M.D. Zomeño, A. Hernaéz, L.

Torres, M. Quifer, R. Llimona, G. Freixer, K.A. Pérez-Vega, M. Malcampo, J. Muñoz, M. Farràs, R. Elosua, J. Vila, I. Subirana, M. Cabañero, S. Pérez, M. Vila, V. Munoa, N. Herrera, A. Goday, J.J. Chillaron Jordan, J.A. Flores Lerroux, D. Benaiges Boix, G. Llauredó, M. Farré, E. Menoyo, A. Aldea-Perona, M. Pérez-Otero, D. Muñoz-Aguayo, S. Gaixas, G. Blanchart, A. Sanllorente, M. Soria, J. Valussi, A. Cuenca, L. Forcano, A. Pastor, A. Boronat, S. Tello, M. Cabañero, L. Franco, H. Schröder, R. De la Torre, J.L. del Val, C. Medrano, J. Bayó, M.T. García, V. Robledo, P. Babi, E. Canals, N. Soldevila, L. Carrés, C. Roca, M.S. Comas, G. Gasulla, X. Herraiz, A. Martínez, E. Vinyoles, J.M. Verdú, M. Masague Aguade, E. Baltasar Massip, M. López Grau, M. Mengual, V. Moldon, M. Vila Vergaz, R. Cabanes Gómez, Ciurana, M. Gili Riu, A. Palomeras Vidal, F. Peñas F, A. Raya, M.A. Sebastian, M. Valls, J. Guerrero, M. Marne, E. Minguella, M. Montenegro, A. Sala, M.R. Senan, N. Talens, N. Vera.

Unidad de Epidemiologías de la Nutrición. Universidad Miguel Hernández, Instituto de Investigación Sanitaria y Biomédica, (UMH-ISABIAL), CIBERESP. Alicante, Spain: M. García-de-la-Hera, S. González-Palacios, L. Torres-Collado, L. Compañ-Gabucio, A. Oncina-Cánovas, A.J. Signes-Pastor, J.M. Zazo-Menargues, I. Candela-García, M. Espina, I. Palazón, V. Martínez-Avilés, E.P. Cases-Pérez, C. Tercero-Maciá, L.A. Mira-Castejón, C. Pastor-Polo, A. Bernabé-Casanova, E. Puig-Agulló, S. Pertusa-Martínez, C. Barceló-Iglesias, M.V. Hernández-Marsán, M. García-Muñoz, R. Lloret-Macián, I. Vilanova-Martínez, A. Pastor-Morel, A. Asencio, M.A. Sempere-Pascual, C.M. López-García, P. Jiménez-Sellés, D. Vivancos-Aparicio, N. Gómez-Bellvert, F. Ortiz-Díaz, C. Sánchez-Botella.

Hospital Son Espases (HUSE) and Institute for Health Research Illes Balears (IdISBa), Palma de Mallorca, Spain: M. Fiol, M. Moñino, A. Colom, A. Chaplin, A. Curto, M. Morey, L. Prohens, A.M. Galmés-Panadés, E. Rayó, M. Nafria, J. Llobera, C. Fernández-Palomeque, E. Fortuny, M. Noris, L. López, C. Más, X. Rosselló, S. Munuera, F. Tomás, F. Fiol, A. Jover, J.M. Janer, C. Vallespir, I. Mattei, N. Feuerbach, M. del Mar Sureda, S. Vega, L. Quintana, A. Fiol, M. Amador, S. González, J. Coll, A. Moyá, T. Piqué Sistac, M.D. Sanmartín Fernández, M.C. Piña Valls, M.A. Llorente San Martín, J. Pou Bordoy.

Department of Nutrition, Food Sciences, and Physiology, Center for Nutrition Research, University of Navarra, Pamplona, Spain: I. Abete, J. Abad-Vicente, Ágreda-Peiró, A. Andueza-Azcárate, E. Arina-Vergara, J.I. Armendáriz-Artola, C. Arroniz, F. Artal-Moneva, F. Bárcena-Amigo, R. Bartolomé-Resano, I. Blanco-Platero, F. Calle-Irastoza, E. Cano-Cáceres, I. Cantero, B. Churio-Beraza, C. Cristobo, B. de Cuevillas, E. Echarte-Osacain, T. Elcarte-López, J. Escribano-Jarauta, A. Gimeno-Aznar, N. Goñi-Ruiz, M.V. Güeto-Rubio, I. Ibero-Baraibar, P. Iñigo-Cibrian, M.D. Lezáun-Burgui, E. Madoz-Zubillaga, J. Martínez-Jarauta, M.D. Martínez-Mazo, D. Martínez-Urbistondo, Y. Monzón-Martínez, C. Napal-Lecumberri, V. de la O, B. Ojeda-Bilbao, A. Parra-Osés, B. Pérez-Sanz, B. Rípodas-Echarte, A.L. Tobaruela, L. Ugalde-Sarasa, J. Ulibarri-del Portillo, E. Ursúa-Sesma, M. Zulet.

University of Málaga, Institute of Biomedical Research in Málaga (IBIMA), Málaga, Spain: J. Wärnberg, N. Pérez-Farinós, F.J. Barón-López, J.C. Fernández García, J.C. Benavente-Marín, A. Colom Fernández, J. Aguirre Bustamante, E. Crespo Oliva, V. Castillo Antúnez, O. Fernández Barceló, F.J. Carmona González, R. Carabaño Moral, S. Torres Moreno, M.V. Martín Ruíz, M. Alcalá Cornide.

Lipids and Atherosclerosis Unit, Department of Internal Medicine, Maimonides Biomedical Research Institute of Cordoba (IMIBIC), Reina Sofia University Hospital, University of Cordoba, Cordoba, Spain: J. Criado García, A.I. Jiménez Morales, A. Ortiz Morales, J.D. Torres Peña, F.J. Gómez Delgado, J.F. Alcalá, A. León Acuña, A.P. Arenas Larriva, J.L. Romero-

Cabrera, F. Rodríguez Cantalejo, J. Caballero Villaraso, I. Nieto Eugenio, P. Coronado Carvajal, M.C del Campo Molina, P.J. Peña Orihuela, I. Perez Corral, G. Quintana Navarro.

Department of Internal Medicine, Institut d'Investigacions Biomèdiques August Pi i Sunyer (IDIBAPS), Hospital Clínic, University of Barcelona, Barcelona, Spain: C. Viñas, S. Castro-Barquero, A.M. Ruiz-León, F. Casanovas-Garriga, L. Doblas, A. Jordán, M. Camafort, C. Sierra, E. Sacanella, J. Naval, E. Viñas-Esmel, O. Rodríguez, J. M. Cots, I. Sarroca, M. García, N. Bermúdez, A. Pérez, I. Duaso, A. de la Arada, R. Hernández, C. Simón, M.A. de la Poza, I. Gil, M. Vila, C. Iglesias, N. Assens, M. Amatller, LL. Rams, T. Benet, G. Fernández, J. Teruel, A. Azorin, M. Cubells, D. López, J.M. Llovet, M.L. Gómez, P. Climente, L. de Paula, J. Soto, C. Carbonell, C. Llor, X. Abat, A. Cama, M. Fortuny, C. Domingo, A. I. Liberal, T. Martínez, E. Yañez, M. J. Nieto, A. Pérez, E. Lloret, C. Carrazoni, A. M. Belles, C. Olmos, M. Ramentol, M. J. Capell, R. Casas, I. Giner, A. Muñoz, R. Martín, E. Moron, A. Bonillo, G. Sánchez, C. Calbó, J. Pous, M. Massip, Y. García, M.C. Massagué, R. Ibañez, J. Llaona, T. Vidal, N. Vizcay, E. Segura, C. Galindo, M. Moreno, M. Caubet, J. Altirriba, G. Fluxà, P. Toribio, E. Torrent, J. J. Anton, A. Viaplana, G. Vieytes, N. Duch, A. Pereira, M. A. Moreno, A. Pérez, E. Sant, J. Gené, H. Calvillo, F. Pont, M. Puig, M. Casasayas, A. Garrich, E. Senar, A. Martínez, I. Boix, E. Sequeira, V. Aragunde, S. Riera, M. Salgado, M. Fuentes, E. Martín, A. Ubieto, F. Pallarés, C. Sala, A. Abilla, S. Moreno, E. Mayor, T. Colom, A. Gaspar, A. Gómez, L. Palacios, R. Garrigosa.

Departament of Preventive Medicine and Public Health, University of Granada, Granada, Spain: L. García-Molina, B. Riquelme-Gallego, N. Cano-Ibañez, E.M. Garrido Garrido, A. Baena-Dominguez, M. Terrón-Navas, C. Morata-Cespedes, E. Thomas-Carazo, F. Padilla-Ruiz, J. Machado-Santiago, R. Rodríguez-Tapioles, F. Dorador-Atienza, D. Rueda-Lozano, M. López-Alcázar, MC Rosales Sierra, P. Aguacil-Cubero, M. Rivera-Izquierdo; A. López-Rodríguez, A. Bouzalmate-Hajjaj, S. Extremera-Galán; MA. Castillo-Hermoso, S Martínez-Diz, M. Lozano-Lorca, R Román-Gálvez.

Bioaraba Health Research Institute, Cardiovascular, Respiratory and Metabolic Area; Osakidetza Basque Health Service, Araba University Hospital; University of the Basque Country UPV/EHU, Vitoria-Gasteiz, Spain: L. Tojal Sierra, L. Goicolea Güemez, C. Sorto Sanchez, J. Vaquero Luna, I. Salaverria, A. Casi Casanellas, M.L. Arnal Otero, J. Ortueta Martínez De Arbulo, J. Vinagre Morgado, J. Romeo Ollora, J. Urraca, M.I. Sarriegui Carrera, F.J. Toribio, E. Magán, A. Rodríguez, S. Castro Madrid, M.T. Gómez Merino, M. Rodríguez Jiménez, M. Gutiérrez Jodra, B. López Alonso, J. Iturralde Iriso, C. Pascual Romero, A. Izquierdo De La Guerra.

Research Group on Community Nutrition & Oxidative Stress, University of Balearic Islands-IUNICS, Palma de Mallorca, Spain: E. Angullo-Martinez, E. Argelich, M.M. Bibiloni, P.A. Borràs, C. Bouzas, V. Cepeda, J.M. Gámez, S. García, C. Gómez, I. Llompарт, D. Mateos, M, Monserrat-Mesquida, A. Pons, M. Quetglas-Llabrés, T. Ripoll, T. Rodríguez, M. Ródenas, M. Santés, A. Sureda, S. Tejada, L. Ugarriza.

Virgen de la Victoria Hospital, University of Málaga, Málaga, Spain: M. Macías González, J. Ruiz Nava, J.C. Fernández-García, A. Muñoz Garach, A. Vilches Pérez, A. González Banderas, A.V. Alarcón-Martín, M. García Ruiz de Mier, M. Quero Moreno, J. Alcaide Torres, A. Vargas Candela, M. León Fernández, R. Hernández Robles, S. Santamaría Fernández, J.M. Marín, A. Gómez-Pérez, M. Damas-Fuentes.

Research Institute of Biomedical and Health Sciences, University of Las Palmas de Gran Canaria, Las Palmas, Spain: J. Álvarez-Pérez, E. Doreste Ayala, C. Montesdeoca-Mendoza A. Sánchez-Villegas, I. Bautista-Castaño, V. Dávila-Batista, C.D. García-Méndez, M.D. Arencibia-

Caballero, C. Simón-García, A.J. Santana-Santana, A. Rabaza-Castillo, Z. Hernández-Cubas, R. Padrón-Jiménez, C.B. Bethencour-Sarmiento, J.C. Pérez-Hayes, A. Sosa-Pérez B.V. Díaz-González, J.M. Castillo-Anzalas, R.E. Sosa-Also, J. Medina-Ponce, J. González-Vega, O Pérez-Luzardo.

Biomedicine Institute (IBIOMED); University of León, and Primary Health Care Management of León (Sacyl), León, Spain: Biomedicine Institute (IBIOMED); University of León, and Primary Health Care Management of León (Sacyl), León, Spain: S. Abajo Olea, L. Álvarez-Álvarez, A. Carbó Jordà, E. Carriedo Ule, N. Cubelos Fernández, M. Escobar Fernández, E. Fernández Mielgo, J. P. Fernández Vázquez, J. I. Ferradal García, M. García Fernández, A. Gayubo Serrenes, F. González Rivero, J. I. López Gil, A. Marcos Delgado, S. Puerta Pérez, S. Reguero Celada, M. Rodríguez Bul, M. Rubín-García, F. Vitelli-Storelli.

Department of Family Medicine, Distrito Sanitario Atención Primaria Sevilla, Sevilla, Spain: J Lapetra, L. Miró-Moriano, C. Domínguez-Espinaco, S. Vaquero-Díaz, F.J. García-Corte, A. Santos-Calonge, C. Toro-Cortés, N. Pelegrina-López, V. Urbano-Fernández, M. Ortega-Calvo, J. Lozano-Rodríguez, I. Rivera-Benítez, M. Caballero-Valderrama, P. Iglesias-Bonilla, P. Román-Torres, Y. Corchado-Albalat, L. Mellado-Martín.

Department of Endocrinology and Nutrition, Hospital Fundación Jimenez Díaz. Instituto de Investigaciones Biomédicas IISFJD. University Autonoma, Madrid, Spain: A.I. de Cos, S. Gutierrez, S. Artola, A. Galdon, I. Gonzalo.

Lipids and Vascular Risk Unit, Internal Medicine, University Hospital of Bellvitge- IDIBELL, Hospitalet de Llobregat, Barcelona, Spain: A. Galera, C. Puig-Muñoz, E. de la Cruz, M. Gimenez-Gracia, F. Trias, I. Sarasa, M. Fanlo-Maresma, V. Esteve-Luque, L. Hidalgo-Peña, H. Lafuente, M. Liceran, A. Rodriguez-Sanchez, S. Cardona-Piñeiro, I. Arjona, E. Corbella, A. Riera, X. Pintó.

Department of Endocrinology, IDIBAPS, Hospital Clinic, University of Barcelona, Barcelona, Spain: A. Altés, I. Vinagre, C. Mestre, J. Viaplana, M. Serra, J. Vera, T. Freitas, E. Ortega, I. Pla, R. Olbeyra.

Nutritional Control of the Epigenome Group. Precision Nutrition and Obesity Program, Institute IMDEA-Food, CEI UAM+CSIC, Madrid, Spain: J.M. Ordovás, V. Micó, L. Berninches, L. Díez, E. Cuadrado-Soto, C. Martínez-Pérez, J.A. Celada, J. Tapia, M.J. J. Muñoz, Y. de la Fuente, C. Albertos, P. Martín, C. Cuesta. B. Cáceres Sánchez, ME. Jiménez Caravera, MA. Aldavero Palacios, S. Conti Fernández, MC. Rodríguez Romero, PJ. Jiménez Pérez, V. Fernández Gutiérrez, R. Baños Morras, C. Gómez Almodóvar, P. González Escobar, AM. Ibarra Sánchez, A. Manzanares Briega M. Renata Muñoz Bieber, L. Santos Larregola, C. Cassinello Espinosa, MC. Molins Santos, JM. Rodríguez Buitrago, E. Sánchez Balsalobre, C. Albertos Carrion, L. González Torres, ME. Villahoz Loureiro, E. Arrebola Vivas, AF. Fernández Garcia, ML. Cornejo Alonso, JA. Romo Martin, L. Carabias Jaen, C. Barbero Macías, MA. Blanca De Miguel Oteo, E. Bartolomé Cobeña, MM. Adrián Sanz, MA. Angel Álvaro Sánchez, MD. Cano Pérez, MP. Lopez Morandeira, A. Montero Costa, E. Robles Fernandez, I. Alba Llacer, J. Aroca Palencia, R. Sanz Merino, MJ. Concejo Carranza, A. Garcia Romero, MC. Gómez Tabera, C. Lesmes Lora, J. Zarco Montejó, A. Campo Lopez, ME. Collado Correa, MS. Díaz Moreno, B. Doval Segura, RM. Gómez Quiroga, S. Hernando Gómez, MJ. Martínez Sanz, AM. Yunquera Alonso.

Division of Preventive Medicine, University of Jaén, Jaén, Spain: J.J. Gaforio, S. Moraleda, N. Liétor, J.I. Peis, T. Ureña, M. Rueda, M.I. Ballesta.

Department of Endocrinology and Nutrition, Hospital Clínico San Carlos, Instituto de Investigación Sanitaria del Hospital Clínico San Carlos (IdISSC), Madrid, Spain: C. Moreno Lopera, C. Aragoneses Isabel, M.A. Sirur Flores, M. Ceballos de Diego, T. Bescos Cáceres, Y. Peña Cereceda, M. Martínez Abad, R. Cabrera Vélez, M. González Cerrajero, M.A. Rubio Herrera, M. Torrego Ellacuría, A. Barabash Bustelo, M. Ortiz Ramos, A. Larrad Sáinz, A.L. Calle Pascual.

Clinical Event Ascertainment Committee: F. Arós; A.M. Alonso-Gómez; M. Villas-Roca; Jon B. Toledo; O. García-Regata; L. Forga; A. García-Layana; A. González-Pinto; I. Zorrilla; M. Martínez-Kareaga; P. Seoane; Fernando Arós; Angel M. Alonso-Gómez; Fernando Arós; Mònica Villas-Roca; Jon B. Toledo; Óscar García Regata; Lluís Forga; Alfredo García-Layana; Ana González Pinto; Iñaki Zorrilla; Mireia Martínez-Kareaga; Patricia Seoane.

Steering Committee: J. Salas-Salvadó, M.A. Martínez-González, M. Fitó, E. Ros, FJ. Tinahones, D. Corella and R. Estruch.

Executive Committee: J. Salas-Salvadó, M.A. Martínez-González, D. Corella, M. Fitó, J. Vioque, D. Romaguera, J.A. Martínez, J. Wärnberg, J. Lopez-Miranda, R. Estruch, A. Bueno-Cavanillas, Á.M. Alonso-Gómez, J.A. Tur, FJ. Tinahones, L. Serra-Majem, V. Martín, J. Lapetra, C. Vázquez, X. Pinto, J. Vidal, L. Daimiel, M. Delgado-Rodríguez, M.A. Rubio and E. Ros.

Dietary and Lifestyle Intervention Committee: J. Salas-Salvadó, M.A. Martínez-González, M. Fitó and R. Estruch; Dietary Intervention: J. Salas-Salvadó, N. Babio, E. Ros, A. Sánchez-Tainta; Physical Exercise: M. Fitó, H. Schröder, A. Marcos, D. Corella, J. Warnberg; Behavioural support: R. Estruch, F. Fernández-Aranda, C. Botella and J. Salas-Salvadó.

Support groups: C. Botella; F. Fernandez-Aranda; R. Lamuela; A. Marcos; M.P. Portillo; E. Ros; G. Sáez; F. Arós; E. Gómez-Gracia.

II. Supplement Methods

Stata code used to estimate changes in the Energy-restricted MedDiet (Figure 3.A)

Excerpt of Stata Code

```
mixed pl7_total_v0 interv#visita || nodo: || idcluster: || id: if visita < 7,  
vce(robust)  
  
margins visita#interv  
  
marginsplot, ///  
    title("") ///  
    ytitle("17-point score (95% CI)") ///  
    xtitle("Follow-up (years)") ///  
    ylabel(, angle(hor) grid) ///  
    legend(order(1 "Control" 2 "Intervention") size(small)) ///  
    ylab(7(1)14) ///  
    plotlopts(lcolor(cranberry) mcolor(cranberry)) cilopts(color(cranberry)) ///  
    plot2opts(lcolor(green) mcolor(green) lpattern(dash)) ci2opts(color(green))
```

III. Supplement Tables and Figures

Supplement Table 1. Intensive intervention program (intervention group)

Energy-restricted Mediterranean diet*

Dietary pattern

Traditional dietary patterns based on whole foods or minimally processed foods

Foods

High consumption of extra-virgin olive oil, nuts (especially walnuts), fruits and vegetables, salads, whole grains, fiber-rich foods and low-fat yogurts.

Avoid consumption of sugar-sweetened beverages, fast foods, refined grain products (especially white bread, which is widely consumed in Spain), white rice, pasta (except for whole-grain pasta), French fries, potatoes, trans fats (mainly present in commercial bakery products in Spain), sweets, cakes, pies, sugar, precooked meals, sausages or cold cuts of processed meats, and patés have been consistently associated with weight gain

Nutrients

- Fat intake should be 35-40 % of total calories: Olive oil and nuts must be the preferred sources of fat, and fat restriction involves fat from animal foods.
- Protein intake should be approximately 20% of total calories. Proteins should be derived first from plant and second from lean animal sources (like fish or poultry).
- Carbohydrates should be 40-45 % or more of total calories (of low glycemic index). Carbohydrates should be derived from solid, minimally processed and fiber-rich foods with a low glycemic index, such as vegetables, fruits and whole grains, all of which are good sources of vitamins, minerals, and fiber.
- Dietary fiber should be 30-35 grams/day.

Physical Activity**

- a) Aerobic physical activity. Gradually increase the level of moderate-to-vigorous physical activity to at least 45 minutes per day (6 days per week) after 6 months of intervention. The physical activity program includes aerobic activities, such as gentle walking or any equivalent activity of moderate intensity and resistance training. Recommendations are adapted to personal preferences and encourage participants to switch between activities with the same metabolic equivalence of tasks.
- b) Strength physical activity. To promote physical activities for developing strength in the main muscle groups at least 2 days per week, regardless of whether these activities are supervised or performed independently in a sports environment, outdoors, or at home. The duration of each strength training session were 30 to 40 minutes per day (2 days per week), and each exercise consisted of 2 to 4 sets with 8 to 12 repetitions of the movement.
- c) Flexibility and balance exercises. To promote physical activities for developing flexibility and balance 3 or more times per week. Flexibility exercises should be performed at the end of physical activity and/or exercise sessions, especially after strength exercises. Additionally, participants were advised that, if motivated and with access, they should engage in guided balance activities such as Tai Chi or Yoga.

Psycho-behavioral therapy

Participants are instructed on strategies and provided tools for solving problems associated with consuming high calorie foods and performing sedentary activities. They are encouraged to learn how to recognize lack of control on food intake under stressful or anxious situations and how to exercise self-control.

* A detailed description is available at the PREDIMED-Plus dietary intervention and physical activity handbooks. ** Recommendations were made based on the assessment according to the classification of physical status at baseline and every 3 months.

Supplement Table 2. 17-item questionnaire of adherence to an energy-restricted Mediterranean diet

Foods and frequency of consumption	Criteria for 1 point*
Do you use extra-virgin olive oil as the main culinary fat?	Yes
How many servings of vegetables do you consume per day? (Count garnish and side servings as ½ point; 1 serving =200g)	≥ 2
How many pieces of fruit or 100% natural fruit juice do you consume per day?	≥ 3
How many servings of red meat, hamburgers, or meat products (ham, sausages, etc.) do you consume per week? (1 serving = 100-150g)	≤ 1
How many servings of butter, margarine, or cream do you consume per week? (1 serving = 12g)	< 1
How many sugar-sweetened beverages (sodas, tonic waters, energy drinks, fruit juices with added sugar) do you consume per week?	< 1
How many servings of legumes do you consume per week? (1 serving = 150g)	≥ 3
How many servings of fish/shellfish do you consume per week? (1 serving = 100-150g, or 4 – 5 pieces of fish, or 200g of shellfish)	≥ 3
How many times per week do you consume pastries, such as cookies, sweets, or cakes?	< 3
How many times per week do you consume nuts [†] ? (1 serving = 30g)	≥ 3
Do you preferentially consume chicken, turkey, or rabbit meat instead of beef, pork, hamburgers, or sausage?	Yes
How many times per week do you consume vegetables, pasta, rice, or other dishes seasoned with <i>sofrito</i> (sauce made with tomato and onion, leek or garlic and simmered with olive oil)	≥ 2
Do you add sugar to beverages (coffee, tea)?	No/use of artificial sweeteners
How many servings of white bread do you consume per day? (1 serving = 75 g)	≤ 1
How many servings of whole grains (bread, rice, pasta) do you consume per week?	≥ 5
How many servings of refined bread, rice, and/or pasta do you consume per week?	< 3
Do you drink wine? How much do you consume per week? (One glass=100ml)	Men: 2-3 glasses/day Women:1-2 glasses /day

*0 points if the criteria are not met.

[†]including peanuts

Supplement Table 3. Baseline characteristic of participants in the PREDIMED-Plus trial

	All	With prevalent diabetes (excluded)	Without prevalent diabetes
N	6874	2128	4746
Age	65.0 (4.9)	65.2 (4.9)	64.9 (4.9)
Sex, No. (%)			
Female	3327 (48.4)	940 (44.2)	2387 (50.3)
Male	3547 (51.6)	1188 (55.8)	3363 (49.7)
Education, No. (%)			
Primary or less	3363 (48.9)	1103 (51.8)	2260 (47.5)
Weight, kg	86.6 (13.0)	87.5 (12.8)	86.2 (13.1)
BMI, kg/m ²	32.6 (3.5)	32.8 (3.5)	32.5 (3.4)
Waist circumference, cm	107.6 (9.7)	109.3 (9.4)	106.8 (9.7)
Tobacco smoking status, No. (%)			
Current	857 (12.5)	273 (12.8)	584 (12.3)
Former	2981 (43.4)	962 (45.2)	2109 (42.5)
Never	3036 (44.2)	893 (42.0)	2143 (45.2)
Physical activity, MET-min/wk	2456 (2271)	2394 (2264)	2484 (2273)
er-MedDiet score	8.5 (2.7)	8.7 (2.6)	8.4 (2.7)
Total energy intake, kcal/d	2416 (633)	2360 (588)	2441 (650)
Baseline hypertension, No. (%)	5711 (83.1)	1820 (85.5)	3891 (82.0)
Baseline dyslipidemia, No. (%)	4762 (69.3)	1581 (74.3)	3181 (67.0)

Abbreviations: er-MedDiet, Energy-reduced Mediterranean diet score (0-17 points); MET, Metabolic Equivalent Task

Supplement Table 4. Use of Drugs (%) during the follow-up by intervention group

	Baseline		Year 1		Year 3		Year 6	
	Control group	Intervention group	Control group	Intervention group	Control group	Intervention group	Control group	Intervention group
N*	2403	2341	2360	2315	2328	2271	1809	1726
Antihypertensive drugs	76.4	76.5	79.2	77.8	80.2	79.4	85.3	84.0
Diuretics	38.3	37.9	39.7	39.1	43.4	41.4	45.4	44.7
Beta-blockers	15.1	15.1	15.4	16.3	17.0	17.5	21.1	20.1
Lipid-lowering agents	44.9	46.1	45.6	45.4	47.8	47.2	54.7	52.5
Statins	41.2	41.7	42.1	41.2	44.0	43.5	50.9	48.8
Antidiabetic drugs	0	0	1.8	1.4	6.4	5.3	16.0	13.2
Insulin	0	0	0	0	0.1	0	0.9	1.2
Metformin	0	0	1.6	1.3	5.9	4.8	12.5	10.6
DPP-4 inhibitors	0	0	0.1	0.1	0.4	0.4	1.8	1.2
SGLT2 inhibitors	0	0	0.1	0	0.4	0.4	2.3	1.3
Other	0	0	0.1	0	0.3	0.2	2.5	2.4

Abbreviations: DPP-4 Dipeptidyl peptidase 4; SGLT2: Sodium glucose cotransporter 2.

*Completers only

Supplement Table 5. Hazard ratios for incident type 2 diabetes in the PREDIMED-Plus trial: Subgroup analyses

	Control group	Intervention group	P for interaction
Age			0.692
≤65 yrs			
Cases/Total	201/1290	171/1281	
HR ^a (95% CI)	1 (ref.)	0.74 (0.60-0.91)	
>65 yrs			
Cases/Total	148/1114	109/1061	
HR ^a (95% CI)	1 (ref.)	0.71 (0.55-0.91)	
Sex			0.035
Women			
Cases/Total	158/1215	142/1172	
HR ^a (95% CI)	1 (ref.)	0.87 (0.69-1.09)	
Men			
Cases/Total	191/1189	138/1170	
HR ^a (95% CI)	1 (ref.)	0.63 (0.51-0.79)	
BMI			0.429
<30 kg/m ²			
Cases/Total	69/660	49/641	
HR ^a (95% CI)	1 (ref.)	0.72 (0.49-1.04)	
≥30 kg/m ²			
Cases/Total	280/1744	231/1701	
HR ^a (95% CI)	1 (ref.)	0.74 (0.62-0.88)	
Waist-to-height ratio			0.344
< 0.5			
Cases/Total	109/860	75/807	
HR ^a (95% CI)	1 (ref.)	0.71 (0.53-0.95)	
≥0.5			
Cases/Total	240/1544	205/1535	
HR ^a (95% CI)	1 (ref.)	0.75 (0.62-0.91)	
Educational level			0.199
Primary or less			

Cases/Total	187/1164	136/1094	
HR ^a (95% CI)	1 (ref.)	0.72 (0.57-0.89)	
Secondary or higher			
Cases/Total	162/1240	144/1248	
HR ^a (95% CI)	1 (ref.)	0.76 (0.61-0.96)	
Baseline fasting plasma glucose			0.933
<100 mg/dl			
Cases/Total	45/1071	38/1055	
HR ^a (95% CI)	1 (ref.)	0.80 (0.52-1.24)	
≥100 mg/dl			
Cases/Total	304/1333	242/1287	
HR ^a (95% CI)	1 (ref.)	0.72 (0.61-0.86)	

* Robust standard errors to account for intracluster correlation (clusters are members of the same household)

^aAdjusted for age, sex and fasting glucose level (≤100; >100&<110; 110&<126; ≥126 mg/dl), body mass index (kg/m²), smoking status (never, former, current smoker), baseline prevalence of dyslipidemia (yes/no) and hypertension (yes/no), family history of diabetes, leisure-time physical activity level (Metabolic Equivalent of Tasks, METs-min/d), adherence to energy-reduced Mediterranean diet, and alcohol intake (g/d, adding quadratic term), and stratified by center and educational level

Supplement Table 6. Absolute and relative risk estimates for the composite outcome of incident type 2 diabetes and death before diabetes diagnosis after 6 years of intervention

	Control group	Intervention group
N	2,404	2,342
Cases	426	351
Person-years, n	15022	14989
Absolute risk (95% CI)	13.5 (13.3-13.6)	11.3 (11.1-11.4)
Crude RD (95% CI)	0 (ref.)	-2.2 (-2.4 to -2.0)
Model 1. ^a RD (95% CI)	0 (ref.)	-2.2 (-2.9 to -1.5)
Model 2. ^b RD (95% CI)	0 (ref.)	-2.2 (-2.9 to -1.4)
Incidence rate/1000 person-years (95% CI)	28.4 (25.8-31.2)	23.4 (21.1-26.0)
Crude HR (95% CI)	1 (ref)	0.82 (0.71-0.94)
Model 1. ^a HR (95% CI)	1 (ref.)	0.75 (0.65-0.86)
Model 2. ^b HR (95% CI)	1 (ref.)	0.73 (0.63-0.85)

Abbreviations: HR, Hazard Ratio; RD, Risk Difference

^a Adjusted for age (years), sex and fasting plasma glucose at baseline (≤ 100 ; >100 & <110 ; 110 & <126 ; ≥ 126 mg/dl), and stratified by center and educational level (primary education vs. secondary or higher). Robust standard errors to account for intra-cluster correlations (clusters are family members of the same household)

^b Additionally adjusted for body mass index (kg/m^2), smoking status (never, former, current smoker), baseline prevalence of dyslipidemia (yes/no) and hypertension (yes/no), family history of diabetes, leisure-time physical activity level (Metabolic Equivalent of Task, MET-min/d), adherence to energy-reduced Mediterranean diet, and alcohol intake (g/d, adding a quadratic term), and stratified by center and educational level (primary education vs. secondary or higher)

Supplement Table 7. Participants with positive answers to each of the 17 Items of the Mediterranean diet score by intervention group during Follow-up

	Baseline		Year 1		Year 6	
	Control group	Intervention group	Control group	Intervention group	Control group	Intervention group
N	2402	2337	2208	2104	1781	1665
Olive oil as main culinary fat	80.7%	80.2 %	95.2%	98.0 %	97.5%	98.3 %
≥ 2 servings/d vegetables	37.1%	35.9 %	58.0%	75.0 %	58.6%	70.7 %
≥ 3 servings/d fruits	47.7%	45.0 %	61.6%	77.4 %	62.7%	74.9 %
≤1 servings/wk red or processed meats						
	47.3%	46.5 %	57.4%	70.6 %	47.2%	64.5 %
<1 servings/wk butter or margarine or cream						
	80.7%	79.5 %	89.0%	94.1 %	86.9%	93.2 %
<1 servings/wk carbonated sugar-sweetened beverages						
	76.1%	72.4 %	81.7%	87.7 %	84.8%	90.7 %
≥ 3 servings/wk legumes						
	19.8%	18.1 %	34.2%	53.2 %	41.5%	52.9 %
≥ 3 servings/wk fish						
	47.4%	46.2 %	62.8%	76.7 %	62.7%	72.0 %
<3 servings/wk commercial bakery, cakes, biscuits, or pastries						
	58.6%	56.0 %	72.6%	83.2 %	69.8%	79.7 %
≥ 3 servings/wk nuts						
	41.0%	40.9 %	76.6%	89.5 %	70.1%	78.1 %
Preference of poultry (chicken, turkey, or rabbit) over red meats (beef, pork, lamb, hamburgers, or sausages)						
	73.2%	75.4 %	82.2%	90.7 %	82.4%	89.4 %
≥ 2 servings/wk olive oil sauce with tomato, garlic, onion, or leek (sofrito)						
	57.0%	56.7 %	68.4%	76.8 %	75.7%	80.9 %

Avoidance of any added sugar to beverages	55.2%	55.2 %	61.7%	76.6 %	71.6%	81.3 %
≤ 1 serving/d of white bread	45.0%	43.9 %	59.2%	82.5 %	70.6%	84.8 %
≥5 servings/wk of whole grain cereals	27.1%	28.3 %	40.2%	69.4 %	41.3%	58.1 %
< 3 servings/wk of refined grain, pasta or white rice	30.0%	29.3 %	36.5%	62.2 %	37.8%	58.3 %
2-3 wine glasses/d for men	24.8%	22.4 %	26.2%	26.1 %	19.8%	22.1 %
1-2 wine glasses/d for women						

One serving is equivalent to 200 g of vegetables, 10-150 g of meat and meat products, 12 g of butter, margarine or cream, 150 g of legumes, 100-150g, or 4 – 5 pieces of fish, or 200g of shellfish, 30 g of nuts, 75 g of bread. One glass of wine is equivalent to 100 ml.

Including peanuts

Supplement Table 8. Body weight and waist circumference reduction throughout the trial by intervention group

	Baseline		Year 1		Year 3		Year 6	
	Control group	Intervention group	Control group	Intervention group	Control group	Intervention group	Control group	Intervention group
N*	2403	2341	2292	2215	2226	2142	1866	1781
Weight, kg	86.0 (13.0)	86.3 (13.1)	85.3 (13.4)	82.4 (13.3)	85.3 (13.6)	82.7 (13.3)	84.9 (13.7)	82.8 (13.4)
Change from baseline	-	-	-0.7 (3.5)	-3.8 (4.3)	-0.5 (4.7)	-3.4 (4.9)	-0.8 (6.7)	-3.2 (5.6)
% of change from baseline			-0.8 (3.9)	-4.4 (4.8)	-0.6 (5.3)	-3.9 (5.5)	-0.9 (6.7)	-3.7 (6.4)
N	2403	2341	2197	2095	2077	1974	1676	1565
Waist circumference	106.9 (9.7)	106.8 (9.7))	105.7 (10.0)	102.0 (9.9)	106.2 (10.4)	102.7 (10.2)	107.2 (10.4)	103.9 (10.5)
Change from baseline	-	-	-1.1 (4.4)	-4.6 (5.5)	-0.4 (5.5)	-3.8 (6.2)	0.6 (6.4)	-2.4 (6.9)
% of change from baseline			-1.0 (4.2)	-4.2 (5.0)	-0.3 (5.2)	-3.5 (5.8)	0.7 (6.0)	-2.2 (6.5)
Mean (SD) unless otherwise stated								

*Completers only

Supplement Table 9. Absolute risk and rates difference estimates for incident type 2 diabetes throughout the entire 7-year follow-up.

	Control group	Intervention group
N	2,404	2,342
Cases of type 2 diabetes	349	280
Person-years, n	15022	14989
Cumulative incidence (95% CI)	14.5 (13.1-15.9)	12.0 (10.6-13.3)
Crude CI difference (%) (95% CI)	0 (ref.)	-2.6 (-4.5 to -0.6)
Model 1 [†] CI difference* (95% CI)	0 (ref.)	-2.9 (-4.7 to -1.1)
Model 2 [‡] CI difference * (95% CI)	0 (ref.)	-3.1 (-4.9 to -1.3)
Incidence rate (IR) per 1000 person-years (95% CI)	23.2 (20.9 to 25.8)	18.7 (16.6 to 21.0)
Crude IR difference	0 (ref)	-4.5 (-7.8 to -1.3)
Model 1 [†] IR difference*	0 (ref)	-7.0 (-10.7 to -3.3)
Model 2 [‡] IR difference*	0 (ref)	-7.5 (-11.2 to -3.7)

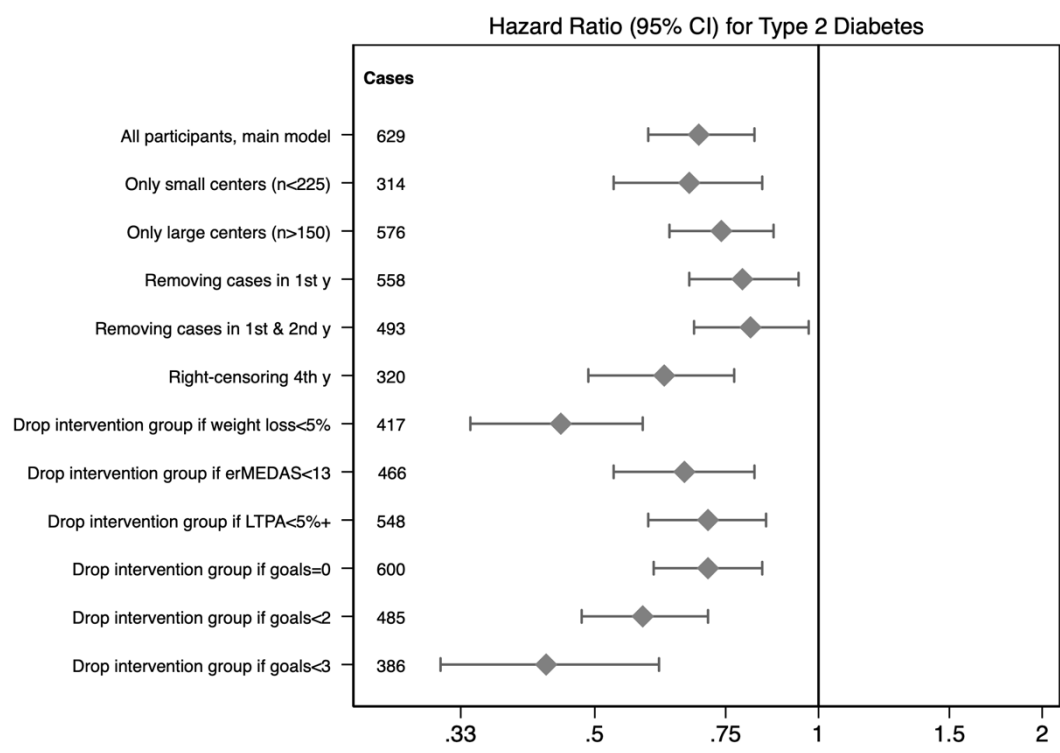
Abbreviations: IR, Incidence Rate; CI, Cumulative Incidence

This table provides complementary information to the risk difference estimates presented in Table 2 of the main manuscript. In Table 2, all risk differences were estimated from Cox models using risks predicted at the 6-year time point. In contrast, the present table includes all cases, including those that occurred after 6 years of follow-up

* Absolute risks (%) calculated with all cases diagnosed during all years of follow-up. Adjusted cumulative incidence differences were estimated from Poisson regression models with a log link and robust standard errors, using the margins command in Stata with the contrast option. Adjusted incidence rate differences were estimated from Poisson regression models with robust standard errors, using the margins command with the predict(ir) option.

[†] Adjusted for age (years), sex and fasting plasma glucose at baseline (≤ 100 ; >100 & <110 ; 110 & <126 ; ≥ 126 mg/dl), and stratified by center and educational level (primary education vs. secondary or higher). Robust standard errors to account for intra-cluster correlations (clusters are family members of the same household)

[‡] Additionally adjusted for body mass index (kg/m²), smoking status (never, former, current smoker), baseline prevalence of dyslipidemia (yes/no) and hypertension (yes/no), family history of diabetes, leisure-time physical activity level (Metabolic Equivalent of Task, MET-min/d), adherence to energy-reduced Mediterranean diet, and alcohol intake (g/d, adding a quadratic term), and stratified by center and educational level (primary education vs. secondary or higher)



Supplement Figure. Ancillary analyses: Hazard ratios and confidence intervals (CI) for intervention versus control group under different assumptions and/or exclusions.

LSM: Lifestyle modification; erMEDAS: energy-reduced Mediterranean Diet Adherence Score

Goals: (1) ≥5% mean weight reduction from baseline to year 6; (2) average erMEDAS>13 from baseline to year 6; and (3) ≥5% mean increase of leisure time physical activity (LTPA) from baseline to year 6

* Hazard Ratios adjusted for age, sex and fasting glucose level (≤100; >100&<110; 110&<126; ≥126 ≥126 mg/dl), BMI (kg/m²), smoking status (never, former, current smoker), baseline prevalence of dyslipidemia (yes/no) and hypertension (yes/no), family history of diabetes, leisure-time physical activity level (Metabolic Equivalent of Tasks, METs-min/d), adherence to energy-reduced Mediterranean diet, and alcohol intake (g/d, adding quadratic term), and stratified by center and education level

Reducing Diabetes Risk Through the Mediterranean Diet

Since the original PREDIMED (Prevención con Dieta Mediterránea) randomized controlled trial was published (1), scholars, clinicians, and health advocates have been fascinated with the potential for the Mediterranean diet (MedDiet) to reduce cardiovascular disease (CVD) events (for example, myocardial infarction, stroke, CVD mortality) and CVD risk factors (for example, hypertension, dyslipidemia, type 2 diabetes) substantially. The MedDiet focuses on high intake of plant-based foods; moderate consumption of fish, poultry, and dairy (along with optional red wine); and low intake of red meats, sweets, and sugar-sweetened beverages (2). Key components include extra-virgin olive oil, fruits, vegetables, legumes, nuts, and whole grains, which are known for their high-fiber, anti-inflammatory, and antioxidant properties, the mechanisms by which the MedDiet is presumed to decrease CVD risk (3).

Given the relatively high amount of dietary fat recommended in the MedDiet, the original PREDIMED study revealed that it only minimally reduced body weight though it reduced diabetes incidence by 30% (4). In 2013, the PREDIMED-Plus trial was launched to investigate the impact of an energy-reduced MedDiet (erMedDiet) plus physical activity and behavioral strategies delivered via dietician-led groups and 1:1 sessions on reducing body weight and CVD events in 6874 Spanish adults with overweight or obesity and metabolic syndrome (5). In their article, Ruiz-Canela and colleagues conducted a prespecified secondary analysis of the PREDIMED-Plus trial to examine the effect of the erMedDiet compared with an ad libitum MedDiet on diabetes incidence at the end of 6 years among the 4746 participants without diabetes at baseline (6).

After 6 years, the adjusted absolute risk difference for incident diabetes between the erMedDiet intervention and the ad libitum MedDiet control group was -2.4% (95% CI, -3.1% to -1.8%), corresponding to a 31% (CI, 18% to 41%) relative risk reduction in the intervention group compared with the control group (6). Of note, a greater effect was observed in men compared with women. Although the seminal Diabetes Prevention Program (DPP) reported greater reductions in absolute (-14.4%) and relative (58%) risks for incident diabetes from a low-fat diet with physical activity compared with placebo for 3 years (7), PREDIMED-Plus's longer duration of follow-up, use of a strong comparator (an ad libitum MedDiet), and older mean age of participants may in part explain the effect size differences. Weight loss was also greater in DPP than in PREDIMED-Plus, 7% versus 3.7%, respectively. This smaller reduction in weight, however, provides further evidence to clinicians and patients alike that even smaller reductions in body weight can lead to meaningful reductions in diabetes incidence.

Despite the promising findings in this study, the study design precludes the ability to disentangle key intervention components (that is, erMedDiet adherence, physical activity, number of dietetic contacts) and their individual versus synergistic effects on outcomes. It is also important to acknowledge that this study was conducted in Spain, and consumption of an erMedDiet may be particularly challenging in the United States where prevalence of CVD events, prediabetes, and type 2 diabetes remain critically high among adult populations, especially for those patients living in underresourced neighborhoods where healthy food options are limited (8). Participants in the study received extra-virgin olive oil to support adherence and retention; in the United States, prices of extra-virgin olive oil have nearly doubled since 2021 due to a combination of factors including climate change, rising production costs, supply chain disruptions, and now tariffs (9). Furthermore, the large number of dietician contacts during the study may prove difficult to scale broadly in the United States given challenges with health care access and reimbursement for prevention services. As the investigators acknowledge, more studies will be needed to replicate these results using a lower-intensity intervention to ensure adequate reach among the populations that need it most.

Because of the recent excitement around medications to treat obesity and diabetes and reduce CVD events (10), the tried-and-true effects of diet and physical activity have lost some of their popularity. The study by Ruiz-Canela and colleagues shows us that diet quality is still important to consider and adds to the large body of literature demonstrating the important role that dietary intake and lifestyle modification can play in chronic disease prevention (1, 6). Ultimately, this study contributes to the strong evidence base in support of the MedDiet as an optimal dietary pattern for long-term health (3).

Sharon J. Herring, MD, MPH

Program for Maternal Health Equity, Center for Health Justice and Bioethics, Department of Urban Health and Population Science, Lewis Katz School of Medicine at Temple University, Philadelphia; Department of Medicine, Lewis Katz School of Medicine at Temple University, Philadelphia; and Center for Obesity Research and Education, College of Public Health, Temple University, Philadelphia, Pennsylvania

Gina L. Tripicchio, PhD, MSc

Center for Obesity Research and Education, College of Public Health, Temple University, Philadelphia; and Department of Social and Behavioral Sciences, College of Public Health, Temple University, Philadelphia, Pennsylvania

Disclosures: Disclosure forms are available with the article online.

Corresponding Author: Sharon J. Herring, MD, MPH, Temple University, 3223 North Broad Street, Suite 175, Philadelphia, PA 19140; e-mail, herring01@temple.edu.

Ann Intern Med. doi:10.7326/ANNALS-25-02748

References

1. Estruch R, Ros E, Salas-Salvadó J, et al; PREDIMED Study Investigators. Primary prevention of cardiovascular disease with a Mediterranean diet supplemented with extra-virgin olive oil or nuts. *N Engl J Med.* 2018;378:e34. [PMID: 29897866] doi:10.1056/NEJMoa1800389
2. Davis C, Bryan J, Hodgson J, et al. Definition of the Mediterranean diet; a literature review. *Nutrients.* 2015;7:9139-9153. [PMID: 26556369] doi:10.3390/nu7115459
3. Schwingshackl L, Morze J, Hoffmann G. Mediterranean diet and health status: active ingredients and pharmacological mechanisms. *Br J Pharmacol.* 2020;177:1241-1257. [PMID: 31243760] doi:10.1111/bph.14778
4. Estruch R, Martínez-González MA, Corella D, et al; PREDIMED Study Investigators. Effect of a high-fat Mediterranean diet on body-weight and waist circumference: a prespecified secondary outcomes analysis of the PREDIMED randomised controlled trial. *Lancet Diabetes Endocrinol.* 2019;7:e6-e17. [PMID: 31003626] doi:10.1016/S2213-8587(19)30074-9
5. Martínez-González MA, Buil-Cosiales P, Corella D, et al; PREDIMED-Plus Study Investigators. Cohort profile: design and methods of the PREDIMED-Plus randomized trial. *Int J Epidemiol.* 2019;48:387-388o. [PMID: 30476123] doi:10.1093/ije/dyy225
6. Ruiz-Canela M, Corella D, Martínez-González MÁ, et al. Comparison of an energy-reduced Mediterranean diet and physical activity versus an ad libitum Mediterranean diet in the prevention of type 2 diabetes: a secondary analysis of a randomized controlled trial. *Ann Int Med.* 26 August 2025. [Epub ahead of print]. doi:10.7326/ANNALS-25-00388
7. Knowler WC, Barrett-Connor E, Fowler SE, et al; Diabetes Prevention Program Research Group. Reduction in the incidence of type 2 diabetes with lifestyle intervention or metformin. *N Engl J Med.* 2002;346:393-403. [PMID: 11832527] doi:10.1056/NEJMoa012512
8. Kris-Etherton PM, Petersen KS, Velarde G, et al. Barriers, opportunities, and challenges in addressing disparities in diet-related cardiovascular disease in the United States. *J Am Heart Assoc.* 2020;9:e014433. [PMID: 32200727] doi:10.1161/JAHA.119.014433
9. Moskin J. American Kitchens Face an Uncertain Mix: Olive Oil and Tariffs. *NY Times.* Accessed at www.nytimes.com/2024/12/18/dining/olive-oil-tariffs.html on 14 July 2025.
10. Anderer S. Obesity drugs considered first-line option for cardiovascular risk. *JAMA.* 11 July 2025. [Epub ahead of print]. [PMID: 40643905] doi:10.1001/jama.2025.10324