

Empirically-derived food patterns and the risk of total mortality and cardiovascular events in the PREDIMED study

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1 **ABSTRACT**

2 **Background and aims:** There is little evidence on *post hoc*- derived dietary patterns (DP)
3 and all-cause mortality in Southern-European populations. Furthermore, the potential effect
4 modification of a DP by a nutritional intervention has not been sufficiently assessed. We
5 assessed the association between *a posteriori* defined baseline major DP and total mortality or
6 cardiovascular events within each of the three arms of a large primary prevention trial
7 (PREDIMED) where participants were randomized to two active interventions with
8 Mediterranean-type diets or to a control group (allocated to a low-fat diet).

9 **Design:** We followed-up 7216 participants for a median of 4.3 years. A validated 137-item
10 food-frequency questionnaire was administered. Baseline DP were ascertained through factor
11 analysis based on 34 predefined groups. Cox regression models were used to estimate
12 multivariable-adjusted hazard ratios (HR) for cardiovascular disease (CVD) or mortality
13 across quartiles of DP within each of the three arms of the trial.

14 **Results:** We identified two major baseline DP: the first DP was rich in red and processed
15 meats, alcohol, refined grains and whole dairy products and was labeled Western Dietary
16 Pattern (WDP). The second DP corresponded to a “Mediterranean-type” dietary pattern
17 (MDP). During follow-up, 328 participants died. After controlling for potential confounders,
18 higher baseline adherence to the MDP was associated with lower risk of CVD (adjusted HR
19 for fourth vs. first quartile: 0.52; 95% CI (Confidence Interval): 0.36, 0.74; p-trend <0.001)
20 and all-cause mortality (adjusted HR: 0.53; 95% CI: 0.38, 0.75; p-trend <0.001), regardless of
21 the allocated arm of the trial. An increasing mortality rate was found across increasing
22 quartiles of the WDP in the control group (allocated to a low-fat diet), though the linear trend
23 was not statistically significant (p=0.098).

24 **CONCLUSIONS:** Higher adherence to an empirically-derived MDP at baseline was
25 associated with a reduced risk of CVD and mortality in the PREDIMED trial regardless of the

* CVD: Cardiovascular Disease, DP:Dietary Pattern, EVOO: Extra Virgin Olive Oil, FFQ: Food Frequency Questionnaire, IPW: Inverse Probability Weighting, MeDiet:Mediterranean Diet, MDP: Mediterranean Dietary Pattern, WDP: Western Dietary Pattern.

26 allocated arm. The WDP was not associated with higher risk of mortality or cardiovascular
27 events.

28 **INTRODUCTION**

29 The main causes of death in developed countries are cancer and cardiovascular disease (CVD)*,
30 chronic conditions in which prevention by lifestyle, particularly a healthy diet, plays an
31 important role.

32 In the context of overall dietary patterns (DP), many epidemiological studies have examined the
33 health benefits of an *a priori* defined Mediterranean diet (MeDiet) and found consistent evidence
34 that individuals who better adhere to this DP have a healthier ageing and a longer life span [1,
35 2].

36 In the last decade there has been growing interest in assessing the relationships between diet and
37 disease through the study of whole DP instead of focusing on single nutrients or foods [3]. The
38 approach consisting in collecting food data and using them afterwards to identify the DP actually
39 followed by the study subjects is known as the *a posteriori* approach (post hoc). It detects DP
40 empirically derived from available data using principal component analysis or factor analysis [4-
41 6]. Though several studies have assessed the relationship between *a posteriori* DP and various
42 health outcomes [7], including all-cause mortality [7-10], there is little evidence on *post hoc* DP
43 and mortality in Southern European populations [11]. An interesting issue that has not been
44 explored is whether the adverse effects of a DP can be modified by a nutritional intervention.
45 We evaluated the association between *a posteriori* defined major DP and CVD incidence or
46 total mortality in each of the three arms of a large randomized trial testing an intervention
47 with a Mediterranean-type diets for the primary prevention of CVD (the PREDIMED study).
48 Participants in the PREDIMED trial were randomized to two active interventions with
49 Mediterranean-type diets or to a control group (allocated to a low-fat diet). We assessed

50 whether this intervention was able to modify the association between baseline DP and two
51 outcomes: all-cause mortality and the risk of cardiovascular events.

52 **METHODS**

53 **Subjects**

54 The PREDIMED (PREvención con DIeta MEDiterránea) trial (ISRCTN35739639) is a
55 multicenter, randomized, controlled, parallel group, primary prevention trial conducted in
56 Spain to assess the effects of MeDiet on major cardiovascular events. The study protocol has
57 been published elsewhere [12, 13] and full details are available at www.predimed.es. Between
58 October 2003 and June 2009 we recruited 7,447 high-risk participants and randomly assigned
59 them to one of three dietary interventions: two MeDiets supplemented with either extra virgin
60 olive oil (EVOO) or mixed nuts and a control group (low-fat diet). Participants in the two
61 MeDiet groups received either extra-virgin olive oil (to consume 50 g/d) or mixed nuts per
62 day (15 g/d walnuts, 7.5 g/d hazelnuts, and 7.5 g/d almonds) at no cost. They received
63 instructions directed to upscale the traditional MeDiet 14-item score, including 1) the use of
64 olive oil for cooking and dressing; 2) increased consumption of vegetables, nuts, and fish
65 products; 3) consumption of white meat instead of red or processed meat; 4) preparation of
66 home-made sauce by simmering tomato, garlic, onion and aromatic herbs with olive oil to
67 dress vegetables, pasta, rice and other dishes; and 5) for alcohol drinkers, to follow a
68 moderate pattern of red wine consumption. Participants allocated to the control group
69 received small nonfood gifts and were advised to reduce all types of fat and were given
70 written recommendations according to American Heart Association guidelines. No total
71 calorie restriction was advised, nor was physical activity promoted in any of the three groups.
72 The primary endpoint for the main trial was a composite of cardiovascular events. Total
73 mortality was also used as a secondary outcome. All participants provided written informed
74 consent to a protocol approved by the institutional review boards of the recruiting centers.

75 Eligible participants were community-dwelling men aged 55-80 years and women aged 60-80
76 years without previous CVD who fulfilled at least one of the following criteria: type-2
77 diabetes or 3 or more cardiovascular risk factors, namely smoking, hypertension,
78 dyslipidemia, overweight (BMI \geq 25 kg/m²) or a family history of premature CVD. For the
79 present study we excluded 231 participants, 79 of them because their baseline food-frequency
80 questionnaires (FFQ) were missing and 152 who displayed out-of-range total energy intake
81 (<500 or >3500 kcal/d in women or <800 or >4000 kcal/d in men) [14]. Consequently, the
82 final sample size included 7,216 participants.

83 **Dietary assessment**

84 Data on dietary intake were collected at baseline with a semi-quantitative 137-item validated
85 FFQ [15]. For each item, a typical portion size was included, and consumption frequencies
86 were registered in 9 categories that ranged from “never or almost never” to “ \geq 6 times/day”.
87 Daily food consumption was estimated by multiplying the portion size of each food item by
88 its consumption frequency. Energy and nutrient intake were derived using a computer
89 program based on available information in Spanish food composition tables.

90 *Assessment of non- dietary variables*

91 Several questionnaires were used at baseline examination to collect sociodemographic data,
92 lifestyle variables, history of illnesses, and medication use. A validated questionnaire [16]
93 was used to collect information on physical activity.

94 *Ascertainment of mortality*

95 During follow-up dietitians delivered the nutritional intervention with quarterly individual
96 visits and quarterly group sessions [13]. The questionnaires and examinations carried out at
97 baseline were repeatedly administered every year to all participants. Besides, once a year a
98 team of medical doctors reviewed medical records to collect information on the main
99 outcomes, both in primary care centers and hospitals. Also, yearly inquiries were made of the

100 National Death Index. Considering all the sources of information, we are reasonably confident
101 that ascertainment of mortality outcomes was complete. When a death was identified using
102 any of these primary sources, medical records were requested where the patient had been
103 cared for and submitted to the end-point adjudication committee for assignment of the cause
104 of death. This committee was blinded with respect to the intervention group and the food
105 habits of participants.

106 *Statistical analyses*

107 The 137 food items included in the FFQ were grouped into 34 predefined food categories. A
108 principal component analysis was applied to these 34 categories in order to identify a reduced
109 number of factors that could explain the maximum proportion of the variance from the
110 original groups. Food groups with absolute loading >0.29 were considered relevant
111 components of the identified patterns (**Table 1**).

112 The score for each participant was calculated by summing up the consumption of each food
113 group weighted by each factor score obtained in the factor analysis. The resulting quantitative
114 scores were categorized into quartiles. The Scree plot and the criteria of eigenvalues >2 were
115 used to select factors. We selected 2 factors that accounted for 13% of the total variance.

116 Baseline characteristics and nutritional habits of participants according to their quartiles of
117 adherence to both selected factors (dietary patterns) were analyzed (tables 2 and 3) and p
118 values for linear trend tests were calculated for each variable. Nutritional variables were log-
119 transformed to calculate the p values for linear trend.

120 We used Cox regression models with length of follow-up since randomization for the trial as
121 the primary time variable. The exposure time was calculated as the time elapsed between
122 recruitment and the date of death for deceased participants, the last study visit, or the last
123 recorded clinical event of participants still alive. Hazard ratios (HR) with 95% confidence
124 intervals (CI) for the three upper quartiles compared to the lowest quartile of each DP were

125 calculated. In multivariable model 1, potential confounders included as covariates were sex,
126 age (continuous), intervention group and recruitment center. We constructed a second model
127 (multivariable model 2) that also included as covariates smoking status (never, former or
128 current smoker), baseline BMI (kg/m^2 , continuous), physical activity during leisure time
129 (METs min/day, continuous), baseline self-reported hypertension, hypercholesterolemia,
130 diabetes, history of previous depression and educational level (three categories). We
131 conducted analyses stratified by each of the three arms of the trial. We repeated the main
132 analyses adjusting also for total energy intake (kcal/day, continuous). Tests of linear trend
133 across successive quartiles of adherence to each of the two food patterns were calculated.
134 In order to assess the effects of the intervention across the quartiles of baseline adherence to
135 each DP, a new variable was created combining the joint exposure to the quartile of adherence
136 to the studied DP and the intervention. This cross-classification was conducted with the aim
137 of assessing whether the intervention was able to modify the association between baseline
138 adherence to DP and the risk of death or cardiovascular events. We used inverse probability
139 weighting (IPW) not only to control for baseline confounding but also to estimate the absolute
140 risks according to the joint classification of participants in 12 groups by both quartile of the
141 dietary pattern and intervention arm allocated in the trial [17]. To calculate the weights we
142 used as predictors baseline values of age, sex, smoking habit, BMI, physical activity,
143 hypertension, hypercholesterolemia, diabetes, educational level and history of depression.
144 This method allows analyzing observational studies (the effects of baseline adherence to DP
145 in our case) in a way similar to a randomized trial, under the assumption that all relevant
146 confounders are included in the computation of weights. We estimated the absolute risks of
147 mortality (or cardiovascular events) for each combined category of the DP (quartiles) and the
148 intervention group (3 categories), using the weighted pseudo-population obtained with the
149 IPW procedure. Relative risks were also calculated for upper quartiles of each DP using as

150 reference the first quartile of the DP, within each intervention group and the p for linear trend
151 was calculated.

152 We used STATA version 12.0 (StataCorp) for all analyses.

153 **RESULTS**

154 The mean (\pm SD) age of the 7,216 participants was 67.0 (6.2) y. Participants were followed-
155 up for an average of 4.3 y. During this period 328 deaths were recorded; the mean age at
156 death was 73.8 (6.9).

157 Factor analysis revealed two major DP. Absolute factor loadings >0.29 for each DP are
158 presented in **Table 1**. The first factor was characterized by a high consumption of high fat
159 processed meats and red meats, alcohol, refined grains, canned fish, whole-fat dairy products,
160 sauces, eggs, processed meals, commercial bakery and chocolates, whereas consumption of
161 low-fat dairy products was inversely loaded. We labeled it as “Western Dietary Pattern
162 (WDP)”.

163 The second factor was loaded with vegetables, EVOO, walnuts, oily fish and canned fish,
164 fruits, other nuts, whole-wheat bread, white fish and low fat dairy products. In addition, it was
165 defined by a low consumption (negative loadings) of refined grains and of other olive oils
166 different from EVOO. We labeled this second factor as Mediterranean-type dietary pattern
167 (MDP).

168 The baseline characteristics of participants by quartiles of the WDP and the MDP are shown
169 in **Tables 2** and **3** respectively. Subjects with a higher adherence to the WDP were more
170 likely to be men, current smokers and were more physically active. They also showed higher
171 total energy intake and significantly higher intakes of fat, except for monounsaturated fatty
172 acids, and also greater consumption of most food groups with the exception of vegetables and

Table 1. Factor loadings for the two major dietary patterns in the PREDIMED study

	Western Dietary Pattern	Mediterranean dietary pattern
High fat processed meat	0.55	
Alcohol	0.45	
Red meat	0.45	
Refined grains	0.40	-0.30
Canned fish/Seafood	0.38	0.38
Sauces	0.33	
Processed meal	0.32	
Whole dairy products	0.31	
Eggs	0.30	
Commercial bakery	0.29	
Chocolates	0.29	
Low fat dairy products	-0.33	0.29
Olive oil (not extra-virgin)		-0.39
White fish		0.33
Whole grain bread		0.35
Nuts (not walnut)		0.36
Fruit		0.36
Oily fish		0.39
Walnuts		0.40
Extra-virgin olive oil		0.47
Vegetables		0.51

Table 2. Baseline characteristics of the 7216 participants according to quartiles of adherence to Western Dietary Pattern (WDP)

	WDP1 (n=1804)	WDP2 (n=1804)	WDP3 (n=1804)	WDP4 (n=1804)	p for trend
Men (%)	23.0	33.0	47.5	67.0	<0.001
Smoking status					
Former smokers (%)	16.4	20.4	27.0	34.5	<0.001
Current smokers (%)	7.1	10.7	15.0	22.7	<0.001
Diabetes (%)	55.7	49.0	45.2	44.5	<0.001
Hypertension (%)	84.0	84.1	82.9	79.9	<0.001
Age (y)	68.2±6.0	67.4±6.1	66.7±6.1	65.8±6.3	<0.001
Weight (kg)	74.3±11.2	75.4±12.0	77.4±12.0	80.0±11.8	0.006
BMI (kg/m ²)	30.2±4.0	29.9±3.9	29.9±3.8	29.8±3.6	<0.001
Physical activity during leisure time (METs-min/day)	210.2±223.5	210.4±205.6	242.9±253.7	260.8±264.2	<0.001
Total energy intake (kcal/d)	1839.3±444.5	2053.8±394.2	2294.2±389.2	2757.4±462.4	<0.001
Carbohydrate intake (% total energy)	43.1±7.2	42.4±7.0	41.3±6.9	40.3±7.0	<0.001
Protein intake (% total energy)	17.8±3.1	16.9±2.6	16.3±2.5	15.4±2.4	<0.001
Fat intake (% total energy)	38.3±7.5	39.0±6.7	39.7±6.5	39.8±6.3	<0.001
Monounsaturated fatty acid intake (% total energy)	19.5±5.2	19.5±4.6	19.6±4.3	19.4±4.0	0.339
Polyunsaturated fatty acid intake (% total energy)	5.9±2.1	6.2±2.0	6.4±2.1	6.4±2.0	<0.001
Saturated fatty acid intake (% total energy)	9.3±2.1	9.7±2.1	10.2±2.1	10.7±2.3	<0.001
Alcohol intake (g/d)	2.2±5.2	4.6±7.7	8.8±11.9	17.7±20.6	<0.001
Vegetables (g/d)	336.3±157.3	335.0±145.1	329.1±142.5	335.8±143.3	0.927
Fruits (g/d)	365.0±202.9	369.7±200.9	371.1±199.4	367.7±202.7	0.355
Whole-fat dairy products (g/d)	42.3±76.6	78.0±126.4	110.8±152.5	163.8±192.2	<0.001
Low-fat dairy products (g/d)	390.5±239.6	300.0±221.0	243.8±206.5	189.9±202.1	<0.001

Eggs (g/d)	15.9±9.9	18.8±9.8	21.1±9.7	24.3±12.9	<0.001
Red meat (g/d)	30.8±24.4	43.3±28.9	55.2±33.3	72.2±45.2	<0.001
White fish (g/d)	46.4±33.3	44.6±28.9	43.1±26.4	42.0±31.0	<0.001
Oily fish (g/d)	22.3±21.5	24.5±20.9	24.8±20.2	27.0±22.0	<0.001
Other fish and seafood (g/d)	18.4±16.1	25.8±18.9	31.3±21.7	41.1±30.7	<0.001
High-fat processed meat (g/d)	1.8±3.4	4.5±5.9	7.5±8.7	16.2±16.8	<0.001
Processed meals (g/d)	0.5±1.2	0.9±1.8	1.9±2.3	2.5±4.1	<0.001
Refined grains (g/day)	62.7±60.4	104.0±76.1	126.4±86.3	160.5±103.7	<0.001
Whole bread (g/d)	55.3±77.2	24.5±52.9	18.7±46.2	12.5±38.6	<0.001
Chocolates (g/d)	2.1±4.6	3.9±7.6	5.3±8.1	9.4±15.1	<0.001
Commercial bakery (g/d)	3.1±7.1	5.9±10.5	9.0±14.6	14.1±20.9	<0.001
Sauces (g/d)	0.9±1.5	1.4±2.0	2.0±2.7	3.2±3.9	<0.001
Extra-virgin olive oil (g/d)	16.8±21.1	19.7±22.6	22.2±23.4	25.7±24.5	<0.001
Other olive oils (g/d)	19.7±20.3	17.9±19.1	17.7±20.3	16.5±20.6	0.224
Nuts (g/d)	7.1±11.3	5.4±8.3	11.1±14.2	13.3±16.1	<0.001

Table 3. Baseline characteristics of the 7216 participants according to quartiles of adherence to Mediterranean Dietary Pattern (MDP)

	MDP1 (n=1804)	MDP2 (n=1804)	MDP3 (n=1804)	MDP4 (n=1804)	p for trend
Men (%)	46.0	42.0	41.0	40.5	<0.001
Smoking status					
Former smokers (%)	22.8	24.5	24.8	26.3	0.242
Current smokers (%)	17.0	15.5	11.6	11.5	<0.001
Diabetes (%)	50.0	47.1	48.8	48.7	0.728
Hypertension (%)	83.1	83.4	81.3	83.0	0.540
Age (y)	67.6±6.4	67.4±6.1	66.9±6.0	66.3±6.2	<0.001
Weight (kg)	76.8±12.1	76.8±12.1	76.3±11.7	76.5±12.2	0.007
BMI (kg/m ²)	30.2±3.7	30.0±3.8	29.9±3.9	29.8±4.0	0.001
Physical activity during leisure time (METs-min/week)	211.2±219.4	224.7±247.9	235.0±225.2	253.2±258.9	<0.001
Total energy intake (kcal/d)	2156.1±575.9	2122.2±5239	2189.7±487.5	2476.6±510.9	<0.001
Carbohydrate intake (% total energy)	43.2±7.5	42.2±7.0	41.2±6.7	40.5±6.9	<0.001
Protein intake (% total energy)	15.4±2.6	16.6±2.7	17.1±2.8	17.3±2.8	<0.001
Fat intake (% total energy)	38.7±7.1	38.6±6.6	39.4±6.6	40.2±6.7	<0.001
Monounsaturated fatty acid intake (% total energy)	19.3±4.8	19.2±4.4	19.6±4.5	19.9±4.5	<0.001
Polyunsaturated fatty acid intake (% total energy)	5.9±2.2	5.9±1.8	6.2±1.9	6.8±2.1	<0.001
Saturated fatty acid intake (% total energy)	10.3±2.4	10.0±2.2	9.9±2.1	9.6±2.1	<0.001
Alcohol intake (g/d)	9.2±16.1	8.3±13.7	8.1±13.8	7.7±12.4	0.001
Vegetables (g/d)	254.1±101.2	300.4±107.9	345.5±121.2	436.6±178.9	<0.001
Fruits (g/d)	285.2±156.5	340.0±171.1	385.8±197.8	462.6±229.3	<0.001
Whole-fat dairy products (g/d)	159.9±199.6	94.4±141.6	72.5±112.1	68.2±110.2	<0.001
Low-fat dairy products (g/d)	186.9±196.0	270.6±215.5	312.1±231.2	354.5±241.1	<0.001

Eggs (g/d)	19.6±11.7	19.5±10.8	19.8±10.1	21.1±11.6	<0.001
Red meat (g/d)	53.2±40.3	50.5±34.6	49.2±35.7	48.6±36.7	<0.001
White fish (g/d)	33.0±24.7	41.0±25.8	46.0±26.3	56.1±37.0	<0.001
Oily fish (g/d)	16.1±14.1	21.5±18.0	25.9±20.3	35.1±26.0	<0.001
Other fish and seafood (g/d)	19.2±16.5	25.4±19.6	30.4±21.9	41.6±30.0	<0.001
High-fat processed meat (g/d)	8.4±13.3	8.0±11.1	7.0±10.3	6.7±10.7	<0.001
Processed meals (g/d)	1.6±3.4	1.3±2.6	1.1±2.3	1.1±2.2	0.010
Refined grains (g/day)	153.2±102.4	121.1±89.4	97.2±75.7	82.2±74.6	<0.001
Whole bread (g/d)	6.3±22.7	17.4±41.6	29.9±55.2	57.2±82.1	<0.001
Chocolates (g/d)	4.3±8.3	4.2±7.8	5.0±10.2	7.2±12.8	<0.001
Commercial bakery (g/d)	10.0±18.7	7.7±14.1	7.5±13.6	6.9±12.0	<0.001
Sauces (g/d)	1.9±2.55	1.9±2.7	1.8±2.8	1.9±3.2	0.213
Extra-virgin olive oil (g/d)	6.5±13.6	16.3±20.2	25.3±22.7	36.3±23.6	<0.001
Other olive oils (g/d)	30.0±20.5	20.0±19.4	13.6±18.2	8.2±15.1	<0.001
Nuts (g/d)	4.1±6.3	7.1±8.9	10.3±11.7	18.9±19.1	<0.001

173 fruit. They had a statistically significant lower consumption of white fish, whole bread, and
174 low-fat dairy products. Participants with a higher score for the MDP were more likely to be
175 women, non-smokers and more physically active. They presented a reduced intake of
176 carbohydrate, and higher intake of protein and fat, specifically mono- and poly-unsaturated
177 fat.

178 **Table 4** shows the hazard ratios (HR) for all-cause mortality according to baseline adherence
179 to both DP in the whole PREDIMED cohort. After adjustment for sex, age, intervention group
180 and recruitment center (model 1), there was no significant association between the upper
181 quartile of WDP and death (HR 1.07; 95% CI: 0.77,1.50). Model 2, including additional
182 confounders, showed similar results (HR 1.04; 95% CI: 0.74,1.47).

183 However, adherence to the empirically-derived MDP did show an inverse association that was
184 statistically significant for the linear trend ($p < 0.001$) and suggested strong risk reductions in
185 the two upper quartiles, with HR 0.74 (95% CI: 0.54,0.99) for the third versus the first
186 quartile and HR 0.53 (95% CI: 0.38, 0.75) for the fourth versus the first quartile (all in model
187 2).

188 When we assessed the association between empirically derived DP and cardiovascular events
189 (myocardial infarction, stroke or cardiovascular death, i.e. the primary end-point of the main
190 trial) no significant association was found between the WDP and the risk of cardiovascular
191 events (HR for the upper versus the lowest quartile: 1.00; 95% CI: 0.70,1.43; adjusted for sex,
192 age, intervention group and recruitment center) (**table 5**). Model 2, including additional
193 confounders, showed similar results (HR 1.05; 95% CI: 0.73,1.51).

194 In contrast, a decreased risk of cardiovascular events was observed among subjects classified
195 in the third and fourth baseline quartiles of the empirically-derived MDP (HR 0.60; 95% CI:
196 0.42, 0.81 and HR 0.52; 95%CI: 0.36, 0.74; respectively in model 1 and 2). Results were
197 similar

Table 4. Hazard ratios for total mortality according to quartiles of baseline adherence to the Western dietary pattern and Mediterranean dietary pattern in the PREDIMED study with the three intervention groups considered together (n= 7216)

Western dietary pattern	Q1	Q2	Q3	Q4	
	n= 1804	n= 1804	n= 1804	n=1804	p for trend
Overall mortality (n)	71	69	86	102	
Person-years	7635	7605	7805	8034	
Men (%)	23.0	33.0	47.5	67.0	<0.001
Age (y)	68.2±6.0	67.4±6.1	66.7±6.1	65.8±6.3	<0.001
Multivariable model ¹	1(ref)	0.94 (0.67-1.32)	1.09 (0.78-1.51)	1.07 (0.77-1.50)	0.53
Multivariable model ²	1(ref)	0.93 (0.66-1.30)	1.05 (0.75-1.46)	1.04 (0.74-1.47)	0.65
Mediterranean dietary	Q1	Q2	Q3	Q4	
Pattern	n= 1804	n= 1804	n= 1804	n= 1804	p for trend
Overall mortality (n)	115	85	73	55	
Person-years	7882	7613	7619	7963	
Men (%)	46.0	42.0	41.0	40.5	<0.001
Age (y)	67.6±6.4	67.4±6.1	66.9±6.0	66.3±6.2	<0.001
Multivariable model ¹	1(ref)	0.80 (0.60-1.06)	0.70 (0.52-0.95)**	0.51 (0.36-0.71)**	<0.001
Multivariable model ²	1(ref)	0.82 (0.62-1.10)	0.74 (0.54-0.99)**	0.53 (0.38-0.75)**	<0.001

¹Adjusted for sex, age, recruitment center and interventional group.

²Adjusted for sex, age, recruitment center and interventional group, smoking status (never smoker, former smoker and current smoker), baseline body mass index, physical activity during leisure time, self-reported hypertension, self-reported depression, self-reported diabetes, self-reported hypercholesterolemia and education level (three categories)

*p<0.05 ** p<0.001

Table 5. Hazard Ratios for primary cardiovascular event according to quartiles of baseline adherence to the Western dietary pattern and Mediterranean dietary pattern in the PREDIMED study with the three intervention groups considered together (n= 7216)

Western dietary pattern	Q1	Q2	Q3	Q4	
	n= 1804	n= 1804	n= 1804	n= 1804	p for trend
Primary cardiovascular event (n)	67	61	65	84	
Person-years	7627	7590	7799	8024	
Men (%)	23.0	33.0	47.5	67.0	<0.001
Age (y)	68.2±6.0	67.4±6.1	66.7±6.1	65.8±6.3	<0.001
Multivariable model ¹	1 (ref)	0.95 (0.67-1.35)	0.94 (0.66-1.35)	1.00 (0.70-1.43)	0.93
Multivariable model ²	1 (ref)	0.96 (0.68-1.38)	0.97 (0.68-1.39)	1.05 (0.73-1.51)	0.73
Mediterranean dietary	Q1	Q2	Q3	Q4	
Pattern	n= 1804	n= 1804	n= 1804	n= 1804	p for trend
Primary cardiovascular event (n)	104	70	54	49	
Person-years	7881	7599	7610	7950	
Men (%)	46.0	42.0	41.0	40.5	<0.001
Age (y)	67.6±6.4	67.4±6.1	66.9±6.0	66.3±6.2	<0.001
Multivariable model ¹	1(ref)	0.75 (0.55-1.02)	0.58 (0.41-0.81)**	0.50 (0.35-0.71)**	<0.001
Multivariable model ²	1(ref)	0.76 (0.56-1.04)	0.60 (0.42-0.83)**	0.52 (0.36-0.74)**	<0.001

¹Adjusted for sex, age, recruitment center and interventional group.

²Adjusted for sex, age, recruitment center and interventional group, smoking status (never smoker, former smoker and current smoker), baseline ²Adjusted for sex, age, recruitment center and interventional group, smoking status (never smoker, former smoker and current smoker), baseline body mass index, physical activity during leisure time, self-reported hypertension, self-reported depression, self-reported diabetes, self-reported hypercholesterolemia and education level (three categories)

*p<0.05 ** p<0.001

198 in both multivariable models (P for trend <0.001), suggesting an inverse association between
199 baseline adherence to the MDP and the risk of major cardiovascular events.

200 To further analyze the association between empirically derived DP and cardiovascular events
201 we repeated the analysis using each time as outcome each of the three components of the
202 primary end-point of the trial, namely acute myocardial infarction, stroke and cardiovascular
203 death (Table 6). For each of these analyses participants with the other 2 end-points were
204 excluded. We observed that the MDP was inversely and strongly associated with the risk of
205 myocardial infarction and also with the risk of cardiovascular death (fourth quartile vs first
206 quartile: HR 0.52 ; 95% CI: 0.36, 0.74 and HR 0.37 ; 95%CI: 0.18, 0.76; respectively for each
207 outcome in both adjusted models 2). Moreover, we found that higher baseline WDP
208 adherence was significantly associated with a higher risk of cardiovascular death (p for linear
209 trend = 0.033 in the multiple-adjusted model).

Table 6. a) Hazard Ratios for acute myocardial infarction according to quartiles of baseline adherence to the Western dietary pattern and Mediterranean dietary pattern in the PREDIMED study with the three intervention groups considered together

Western dietary pattern	Q1	Q2	Q3	Q4	p for trend
Acute Myocardial Infarction (n)	24	23	26	30	
Person-years	7508	7482	7662	7846	
Multivariable model ¹	1 (ref)	0.96 (0.54-1.72)	0.94 (0.52-1.68)	0.83 (0.46-1.50)	0.50
Multivariable model ²	1 (ref)	0.97 (0.54-1.74)	0.96 (0.53-1.72)	0.85 (0.47-1.56)	0.58
Mediterranean dietary Pattern	Q1	Q2	Q3	Q4	p for trend
Acute Myocardial Infarction (n)	42	22	22	17	
Person-years	7675	7450	7517	7857	
Multivariable model ¹	1(ref)	0.57 (0.34-0.96)*	0.55 (0.32-0.93)*	0.38 (0.21-0.68)*	0.001
Multivariable model ²	1(ref)	0.59 (0.35-1.00)	0.58 (0.34-0.97)*	0.41 (0.23-0.75)*	0.003

b) Hazard Ratios for stroke according to quartiles of baseline adherence to the Western dietary pattern and Mediterranean dietary pattern in the PREDIMED study with the three intervention groups considered together

Western dietary pattern	Q1	Q2	Q3	Q4	p for trend
Stroke (n)	37	33	31	34	
Person-years	7551	7520	7676	7856	
Multivariable model ¹	1 (ref)	0.99 (0.61-1.59)	0.91 (0.55-1.50)	0.94 (0.57-1.57)	0.78
Multivariable model ²	1 (ref)	1.03 (0.63-1.66)	0.98 (0.59-1.63)	1.05 (0.62-1.77)	0.89
Mediterranean dietary Pattern	Q1	Q2	Q3	Q4	p for trend
Stroke (n)	48	34	27	26	
Person-years	7688	7492	7526	7897	
Multivariable model ¹	1(ref)	0.81 (0.52-1.26)	0.67 (0.41-1.08)	0.66 (0.40-1.10)	0.073
Multivariable model ²	1(ref)	0.81 (0.52-1.26)	0.66 (0.41-1.08)	0.68 (0.41-1.13)	0.086

c) Hazard Ratios for cardiovascular death according to quartiles of baseline adherence to the Western dietary pattern and Mediterranean dietary pattern in the PREDIMED study with the three intervention groups considered together

Western dietary pattern	Q1	Q2	Q3	Q4	p for trend
Cardiovascular death (n)	14	14	20	33	
Person-years	7493	7467	7649	7872	
Multivariable model ¹	1 (ref)	1.03 (0.48-2.19)	1.41 (0.69-2.85)	1.90 (0.95-3.77)	0.031
Multivariable model ²	1 (ref)	1.03 (0.48-2.19)	1.37 (0.67-2.81)	1.91 (0.95-3.86)	0.033
Mediterranean dietary Pattern	Q1	Q2	Q3	Q4	p for trend
Cardiovascular death (n)	32	23	15	11	
Persons-years	7670	7472	7482	7857	
Multivariable model ¹	1(ref)	0.80 (0.47-1.38)	0.53 (0.28-0.99)*	0.36 (0.18-0.73)*	0.002
Multivariable model ²	1(ref)	0.84 (0.48-1.45)	0.56 (0.30-1.05)	0.37 (0.18-0.75)*	0.003

¹Adjusted for sex, age, recruitment center and interventional group.

²Adjusted for sex, age, recruitment center and interventional group, smoking status (never smoker, former smoker and current smoker), baseline ²Adjusted for sex, age, recruitment center and interventional group, smoking status (never smoker, former smoker and current smoker), baseline body mass index, physical activity during leisure time, self-reported hypertension, self-reported depression, self-reported diabetes, self-reported hypercholesterolemia and education level (three categories)

*p<0.05 ** p<0.001

210 In order to assess the potential effect modification by the nutritional intervention we obtained
211 the absolute risks of death or cardiovascular events for each of the 12 groups formed by the
212 joint combination according to the nutritional intervention (3 levels) and the baseline
213 adherence to DP (quartiles). We used inverse probability weighting in this analysis to avoid
214 confounding and obtain exchangeable groups.

215 **Figure 1a** displays the absolute overall risk of all-cause death (rate/1000 person-year) for
216 each category made by the cross-classification according to quartiles of baseline adherence to
217 the WDP and to the 3 randomized arms of the trial (the 2 intervention groups and the control
218 group). Although statistically non-significant (p for trend=0.098), a suggestion for a trend of
219 higher total mortality with higher quartiles of the WDP was present only in the control group,
220 but not in the two active intervention groups. The highest absolute risk of death was observed
221 in participants who were in the highest quartile of baseline adherence to the WDP and did not
222 receive the intervention with MeDiet (i.e. they were allocated to the control group).

223 Likewise, **figure 1b** shows that regardless of the intervention group, better baseline adherence
224 to the MDP seemed to be associated with lower absolute risks for overall mortality in the
225 three groups, being the inverse dose-response trend statistically significant in the control and
226 in the PREDIMED active intervention group supplemented with EVOO (p for trend=0.012
227 and 0.027 respectively).

228

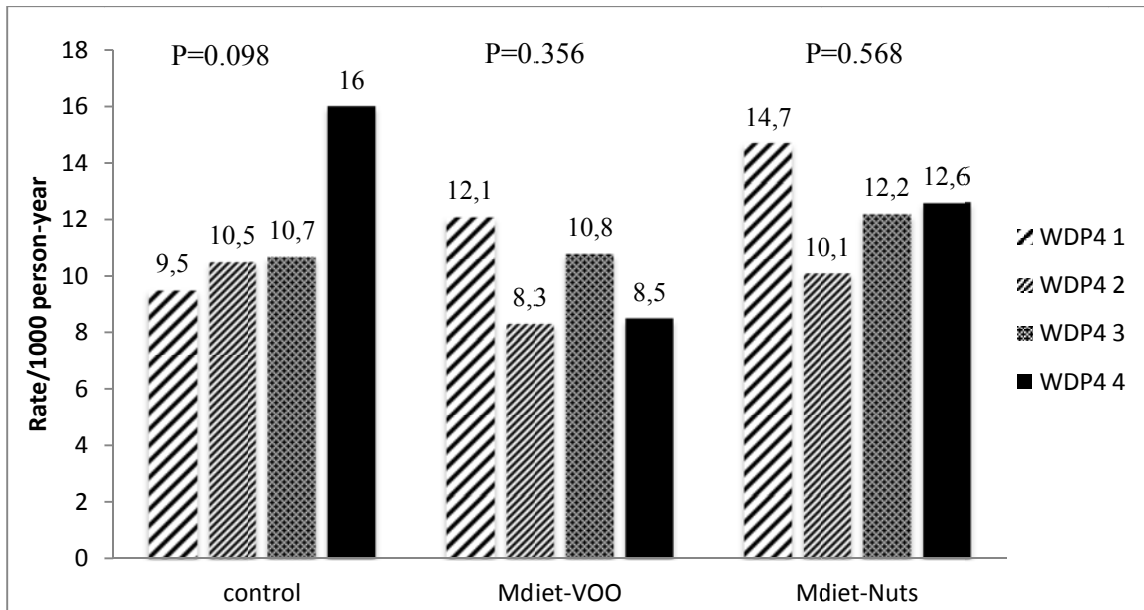


Figure 1a)

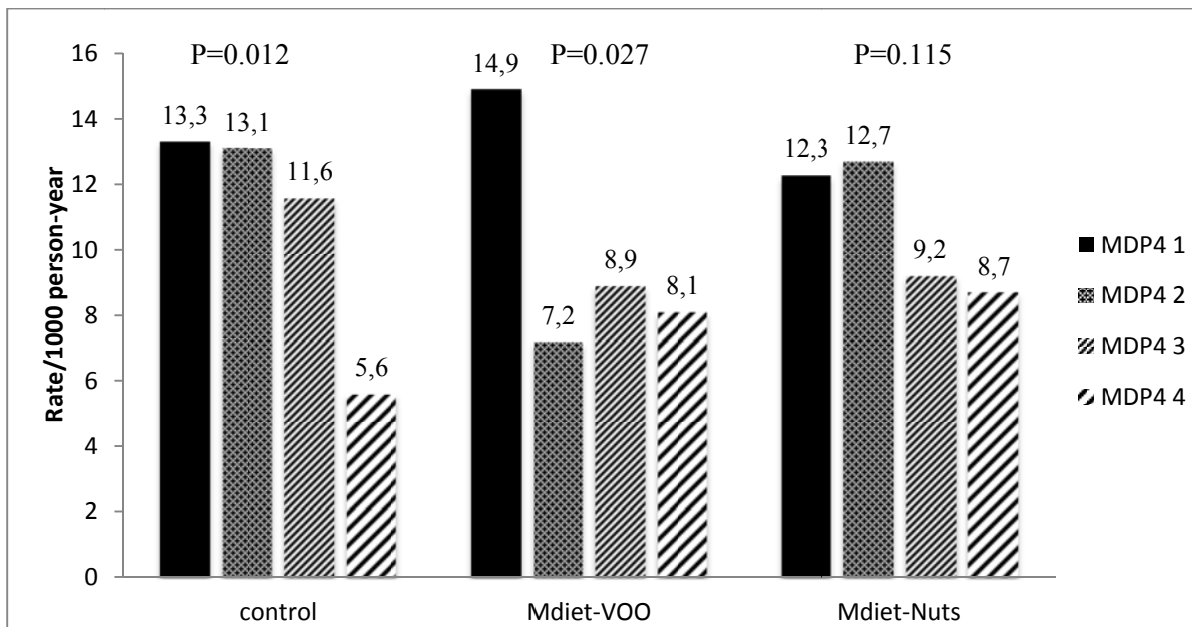


Figure 1b)

203 **Figure 2a** shows the absolute all-cause mortality risk (rate/1000 person-year) for each
204 category built according to the joint classification by quartiles of WDP and allocated
205 intervention arm in the trial. No statistically significant interactions were found.
206 Finally, **figure 2b** represents the cardiovascular risk according to the joint classification by
207 both baseline adherence to the empirically-derived MDP and the allocated intervention arm.
208 These absolute rates might suggest that, even in absence of the PREDIMED nutritional
209 intervention (i.e., in the control group), the inverse association between baseline adherence to
210 MDP and cardiovascular events was clearly apparent. Interestingly, the lowest absolute rates
211 of cardiovascular events were observed in both intervention groups for the 2 upper quartiles
212 of adherence to the baseline MDP (in the group supplemented with nuts) and for the fourth
213 quartile (in the group supplemented with EVOO). The inverse dose-response trend between
214 baseline adherence to the MDP and cardiovascular events remained significant within both
215 intervention groups (p for trend=0.013 for EVOO and 0.007 for nuts).

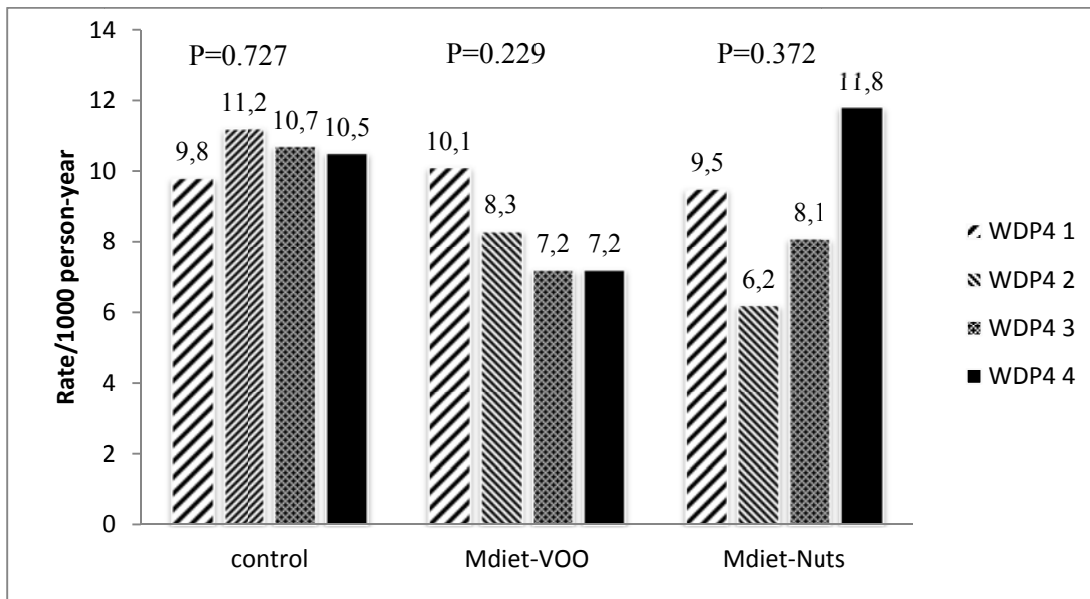


Figure 2 a)

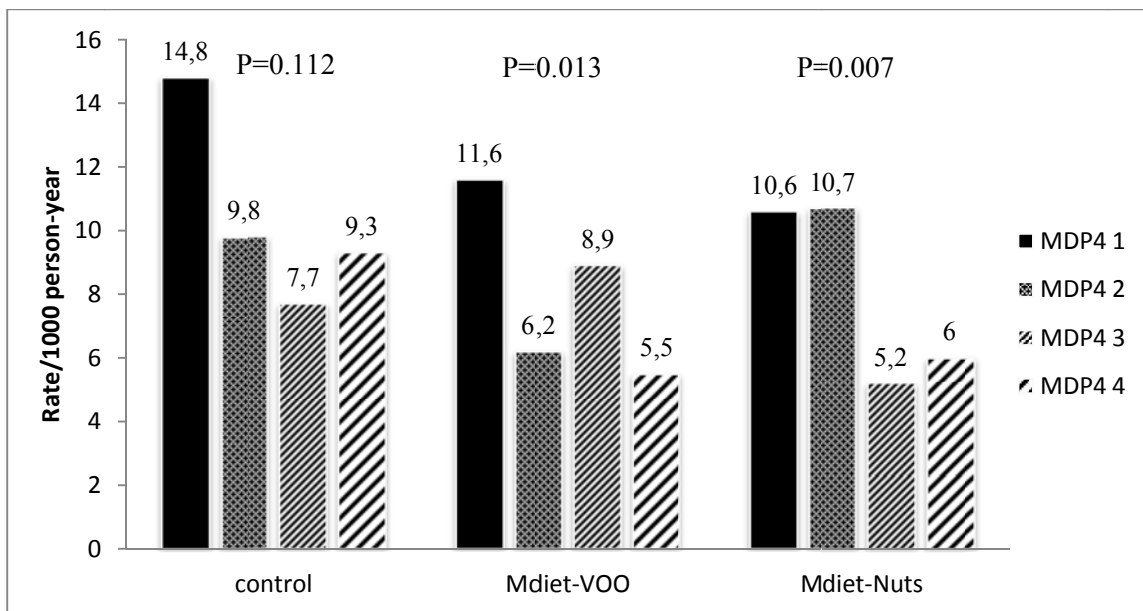


Figure 2 b)

218 To deepen into the combined effects of baseline DP and intervention on overall mortality or
219 cardiovascular events, we assessed the potential interactions between both factors. When the
220 two intervention groups with MeDiet were merged together, we found a p-value=0.054 for the
221 interaction between the highest quartile of WDP and the MeDiet intervention in the analysis
222 of all-cause death. However, the likelihood ratio test assessing the overall effect modification
223 was not statistically significant (3 degrees of freedom, p=0.27)

224 **DISCUSSION**

225 Using factor analysis, we investigated the association between baseline adherence to
226 empirically-derived DP (i.e. post hoc patterns) and all-cause mortality or major cardiovascular
227 events in a cohort of older Spanish subjects at high risk of CVD who underwent a nutritional
228 intervention in the PREDIMED trial [18, 19]. Two DP were identified: WDP and MDP (table
229 1). The results showed that a closer baseline adherence to the MDP was associated with a
230 statistically significant 47% reduction in all-cause mortality along the follow-up period. An
231 inverse association of similar magnitude was also observed between the baseline adherence to
232 the MDP and cardiovascular events. These inverse associations between baseline adherence to
233 the MDP and overall mortality or cardiovascular events were always present, in a low or high
234 magnitude, regardless of the allocated intervention group. However, the intervention with the
235 MeDiet, supplemented with EVOO or nuts, in conjunction with a good baseline conformity
236 with a MDP led to the lowest absolute cardiovascular risk. On the other hand, although no
237 significant results were found, a (non-significant) suggestion for a detrimental association of
238 the WDP with a higher risk of all-cause death was present in the control group. Interestingly,
239 when specific cardiovascular events were separately analyzed, an association of the WDP
240 with a higher risk of cardiovascular death was found. Both findings suggest a detrimental
241 effect of WDP leading to a higher mortality risk.

242 An interesting point of our study is that we were able to assess the combined effects on hard
243 clinical end-points of both the baseline adherence to empirically-derived DP and of a dietary
244 intervention. We observed that in the control group the differences between the highest and
245 lowest quartiles of both the WDP and the MDP appeared to be more apparent, especially
246 when mortality was analyzed. Control group subjects in the highest quartile of baseline
247 adherence to MDP exhibited a 42% relative reduction in their mortality risk compared to their
248 counterparts in the lowest quartile.

249 The present findings assume that the baseline DP might be a good proxy for lifetime dietary
250 exposures. In this line of thought, lifetime exposure can be more important to prevent
251 premature mortality than to follow a supplemented MeDiet or a low-fat diet for only 4 to 5
252 years. However, we were not able to demonstrate a potential interaction between a high
253 baseline adherence to WDP and the intervention with MeDiet (both active intervention groups
254 merged together versus control), though the p value approached the limit of statistical
255 significance (p=0.054).

256 Few studies have examined the association between a healthy DP using factor analysis and
257 the risk of death in high-risk elderly populations [8, 19]. Our results agree with previous
258 reports from the Mediterranean area [8] and non-Mediterranean countries [19], where a MDP
259 was not always identified. In the same context, there are few studies analyzing empirically
260 derived DP and CVD [8, 20], and they also found that in Mediterranean areas, better
261 adherence to a MDP was associated with lower CVD risk.

262 Our results are also consistent with previous studies that have reported inverse associations
263 between an *a priori* defined MDP (or a DP similar to this traditional diet) and total mortality
264 [19, 21, 22]. In relation to CVD, there are several studies analyzing the effects of a priori
265 defined MDP, that found a cardio-protective effect of this DP [23, 24]. However, different

266 country-specific DP have been described in Mediterranean populations and these results
267 should be interpreted with caution.

268 Regarding the Spanish context, a recent study in a large cohort found that higher adherence to
269 an empirically derived MDP in adults was associated with a reduction in the risk of all-cause
270 mortality [6]. Several explanations can account for the inverse association observed between
271 better baseline adherence to a MDP and mortality. First, the MDP has been shown to have a
272 beneficial effect on the incidence and prevalence of several diseases [25]. Second, plant-based
273 foods are protective and plant-based DP may decrease disease risk, whereas diets high in
274 animal foods may be more likely to increase the risk of mortality [26]. In fact, participants in
275 the upper baseline quartiles of MDP followed a diet rich in plant based foods and poor in
276 animal foods, and had the lowest risk of mortality. Third, the available evidence about olive
277 oil, suggests that it plays a role in the prevention of coronary heart disease, and cancer, and
278 may influence survival [1, 2, 21, 22, 25] . Besides, olive oil and particularly EVOO improves
279 the lipid profile and has potent antioxidant and anti- inflammatory properties [27, 28] .

280 Though we found a WDP associated with higher cardiovascular mortality, the absence of
281 association between WDP and all-cause mortality was unexpected. Several mechanisms
282 might also be proposed to explain this absence of association. A suggested explanation is that
283 the “WDP” described in the U.S.[7] and the “WDP” in our study are not entirely equivalent
284 and may not produce the same potential adverse effects on health and longevity. Thus, the
285 consumption of foods known to be associated with lower mortality, such as fish and seafood
286 and alcohol in moderation [29], was included in the so-called WDP in our cohort. Besides, it
287 is possible that residual confounding may have affected our results.

288 Our study has several strengths, including the opportunity to assess the combined effects of
289 the baseline diet and the dietary intervention, the large sample size, the Mediterranean setting,
290 the prolonged follow-up, the sub-studies conducted to validate the questionnaire [30], and the

291 objective, blinded and comprehensive ascertainment of events and close follow-up of
292 participants.

293 There are also limitations to our study. First, the results cannot be generalized to younger
294 and/or healthier individuals from other geographical locations. Second, there is an inherent
295 difficulty to change dietary habits in elderly subjects. Third, even though we adjusted the data
296 for the main known risk factors for mortality or CVD, residual confounding cannot be
297 completely excluded. Fourth, there is a potential for measurement error in the FFQ, which
298 provides only subjective information in comparison with the use of objective markers of food
299 intake. Fifth, the number of observed deaths was small. Despite this last limitation, which is
300 associated with lower statistical power, we found a significant inverse association between the
301 MDP and total mortality. Finally, the method used to define DP (factor analysis) involves
302 several questionable decisions that must be taken into account (e.g. the definition and
303 categorization of predefined food groups). Nevertheless, our results are in line with those of
304 other studies using similar factor analyses to define DP.

305 In conclusion, in a population of Spanish Mediterranean individuals at high cardiovascular
306 risk participating in a nutritional intervention trial, a greater baseline adherence to a MDP was
307 associated with a substantial reduction in CVD and overall mortality after follow-up for ≈ 5 y.
308 Further research is required to confirm the present findings in other Mediterranean and non-
309 Mediterranean settings, especially to better observe if the MeDiet could be able to reduce the
310 detrimental effect of a baseline WDP on cardiovascular or overall mortality risk.

311

312

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317 **Authors Contribution**

318 Conception and Design: MA Martinez-Gonzalez, I Zazpe, C Razquin.

319 Conducted research: A Sánchez-Tainta, I Zazpe and C Razquin.

320 Writing of the first draft: MA Martínez-González, C Razquin, I Zazpe, A Sánchez-Tainta.

321 Analysis and Interpretation of the Data: MA Martínez-González, C Razquin, I Zazpe.

322 Critical revision of the article for important intellectual content and final approval of the
323 article: all authors.

324 Statistical expertise: M.A. Martinez-Gonzalez and E.Toledo.

325 All authors have read and approved the final manuscript.

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368

References

1. Martinez-Gonzalez MA, Bes-Rastrollo M. Dietary patterns, Mediterranean diet, and cardiovascular disease. *Curr Opin Lipidol.* 2014; 25: 20-26.
2. Sofi F, Macchi C, Abbate R, Gensini GF, Casini A. Mediterranean diet and health status: an updated meta-analysis and a proposal for a literature-based adherence score. *Public Health Nutr.* 2013; 1-14.
3. Hu FB. Dietary pattern analysis: a new direction in nutritional epidemiology. *Curr Opin Lipidol.* 2002; 13: 3-9.
4. Martinez-Gonzalez MA, Garcia-Lopez M, Bes-Rastrollo M, Toledo E, Martinez-Lapiscina EH, Delgado-Rodriguez M, Vazquez Z, Benito S, Beunza JJ. Mediterranean diet and the incidence of cardiovascular disease: a Spanish cohort. *Nutr Metab Cardiovasc Dis.* 2011; 21: 237-244.
5. Kant AK, Leitzmann MF, Park Y, Hollenbeck A, Schatzkin A. Patterns of recommended dietary behaviors predict subsequent risk of mortality in a large cohort of men and women in the United States. *J Nutr.* 2009; 139: 1374-1380.
6. Zazpe I, Sanchez-Tainta A, Toledo E, Sanchez-Villegas A, Martinez-Gonzalez MA. Dietary patterns and total mortality in a Mediterranean cohort: the SUN project. *J Acad Nutr Diet.* 2014; 114: 37-47.
7. Heidemann C, Schulze MB, Franco OH, van Dam RM, Mantzoros CS, Hu FB. Dietary patterns and risk of mortality from cardiovascular disease, cancer, and all causes in a prospective cohort of women. *Circulation.* 2008; 118: 230-237.
8. Guallar-Castillon P, Rodriguez-Artalejo F, Tormo MJ, Sanchez MJ, Rodriguez L, Quiros JR, Navarro C, Molina E, Martinez C, Marin P, Lopez-Garcia E, Larranaga N, Huerta JM, Dorronsoro M, Chirlaque MD, Buckland G, Barricarte A, Banegas JR, Arriola L, Ardanaz E, Gonzalez CA, Moreno-Iribas C. Major dietary patterns and risk of coronary heart disease in middle-aged persons from a Mediterranean country: the EPIC-Spain cohort study. *Nutr Metab Cardiovasc Dis.* 2012; 22: 192-199.

9. Anderson AL, Harris TB, Tyllavsky FA, Perry SE, Houston DK, Hue TF, Strotmeyer ES, Sahyoun NR, Health ABC Study. Dietary patterns and survival of older adults. *J Am Diet Assoc.* 2011; 111: 84-91.
10. Hamer M, McNaughton SA, Bates CJ, Mishra GD. Dietary patterns, assessed from a weighed food record, and survival among elderly participants from the United Kingdom. *Eur J Clin Nutr.* 2010; 64: 853-861.
11. Masala G, Ceroti M, Pala V, Krogh V, Vineis P, Sacerdote C, Saieva C, Salvini S, Sieri S, Berrino F, Panico S, Mattiello A, Tumino R, Giurdanella MC, Bamia C, Trichopoulou A, Riboli E, Palli D. A dietary pattern rich in olive oil and raw vegetables is associated with lower mortality in Italian elderly subjects. *Br J Nutr.* 2007; 98: 406-415.
12. Martinez-Gonzalez MA, Corella D, Salas-Salvado J, Ros E, Covas MI, Fiol M, Warnberg J, Aros F, Ruiz-Gutierrez V, Lamuela-Raventos RM, Lapetra J, Munoz MA, Martinez JA, Saez G, Serra-Majem L, Pinto X, Mitjavila MT, Tur JA, Portillo MP, Estruch R, PREDIMED Study Investigators. Cohort profile: design and methods of the PREDIMED study. *Int J Epidemiol.* 2012; 41: 377-385.
13. Zazpe I, Sanchez-Tainta A, Estruch R, Lamuela-Raventos RM, Schroder H, Salas-Salvado J, Corella D, Fiol M, Gomez-Gracia E, Aros F, Ros E, Ruiz-Gutierrez V, Iglesias P, Conde-Herrera M, Martinez-Gonzalez MA. A large randomized individual and group intervention conducted by registered dietitians increased adherence to Mediterranean-type diets: the PREDIMED study. *J Am Diet Assoc.* 2008; 108: 1134-44; discussion 1145.
14. Willet W, Stampfer S. Implications of total energy intake for epidemiologic analyses. In: Willet W, ed. *Nutritional Epidemiology*. New York: Oxford University Press; 1998: 273.
15. Fernandez-Ballart JD, Pinol JL, Zazpe I, Corella D, Carrasco P, Toledo E, Perez-Bauer M, Martinez-Gonzalez MA, Salas-Salvado J, Martin-Moreno JM. Relative validity of a semi-quantitative food-frequency questionnaire in an elderly Mediterranean population of Spain. *Br J Nutr.* 2010; 103: 1808-1816.
16. Elosua R, Marrugat J, Molina L, Pons S, Pujol E. Validation of the Minnesota Leisure Time Physical Activity Questionnaire in Spanish men. The MARATHOM Investigators. *Am J Epidemiol.* 1994; 139: 1197-1209.

17. Robins JM, Hernan MA, Brumback B. Marginal structural models and causal inference in epidemiology. *Epidemiology*. 2000; 11: 550-560.
18. Estruch R, Ros E, Martinez-Gonzalez MA. Mediterranean diet for primary prevention of cardiovascular disease. *N Engl J Med*. 2013; 369: 676-677.
19. Kant AK, Graubard BI, Schatzkin A. Dietary patterns predict mortality in a national cohort: the National Health Interview Surveys, 1987 and 1992. *J Nutr*. 2004; 134: 1793-1799.
20. Panagiotakos D, Pitsavos C, Chrysohoou C, Palliou K, Lentzas I, Skoumas I, Stefanadis C. Dietary patterns and 5-year incidence of cardiovascular disease: a multivariate analysis of the ATTICA study. *Nutr Metab Cardiovasc Dis*. 2009; 19: 253-263.
21. Tognon G, Rothenberg E, Eiben G, Sundh V, Winkvist A, Lissner L. Does the Mediterranean diet predict longevity in the elderly? A Swedish perspective. *Age (Dordr)*. 2011; 33: 439-450.
22. Trichopoulou A, Kouris-Blazos A, Wahlqvist ML, Gnardellis C, Lagiou P, Polychronopoulos E, Vassilakou T, Lipworth L, Trichopoulos D. Diet and overall survival in elderly people. *BMJ*. 1995; 311: 1457-1460.
23. Kastorini CM, Millionis HJ, Kantas D, Bika E, Nikolaou V, Vemmos KN, Goudevenos JA, Panagiotakos DB. Adherence to the mediterranean diet in relation to ischemic stroke nonfatal events in nonhypercholesterolemic and hypercholesterolemic participants: results of a case/case-control study. *Angiology*. 2012; 63: 509-515.
24. Agnoli C, Krogh V, Gioni S, Sieri S, Palli D, Masala G, Sacerdote C, Vineis P, Tumino R, Frasca G, Pala V, Berrino F, Chiodini P, Mattiello A, Panico S. A priori-defined dietary patterns are associated with reduced risk of stroke in a large Italian cohort. *J Nutr*. 2011; 141: 1552-1558.
25. Sofi F, Abbate R, Gensini GF, Casini A. Accruing evidence on benefits of adherence to the Mediterranean diet on health: an updated systematic review and meta-analysis. *Am J Clin Nutr*. 2010; 92: 1189-1196.

26. Orlich MJ, Singh PN, Sabate J, Jaceldo-Siegl K, Fan J, Knutsen S, Beeson WL, Fraser GE. Vegetarian dietary patterns and mortality in Adventist Health Study 2. *JAMA Intern Med.* 2013; 173: 1230-1238.
27. Schwingshackl L, Hoffmann G. Mediterranean dietary pattern, inflammation and endothelial function: A systematic review and meta-analysis of intervention trials. *Nutr Metab Cardiovasc Dis.* 2014; .
28. Cicerale S, Lucas LJ, Keast RS. Antimicrobial, antioxidant and anti-inflammatory phenolic activities in extra virgin olive oil. *Curr Opin Biotechnol.* 2012; 23: 129-135.
29. Gea A, Bes-Rastrollo M, Toledo E, Garcia-Lopez M, Beunza JJ, Estruch R, Martinez-Gonzalez MA. Mediterranean alcohol-drinking pattern and mortality in the SUN (Seguimiento Universidad de Navarra) Project: a prospective cohort study. *Br J Nutr.* 2014; 111: 1871-1880.
30. Schroder H, Fito M, Estruch R, Martinez-Gonzalez MA, Corella D, Salas-Salvado J, Lamuela-Raventos R, Ros E, Salaverria I, Fiol M, Lapetra J, Vinyoles E, Gomez-Gracia E, Lahoz C, Serra-Majem L, Pinto X, Ruiz-Gutierrez V, Covas MI. A short screener is valid for assessing Mediterranean diet adherence among older Spanish men and women. *J Nutr.* 2011; 141: 1140-1145.

Figure and table legends

Table 1. Factor loadings for the two major dietary patterns in the PREDIMED study

Table 2. Baseline characteristics of the 7216 participants according to quartiles of adherence to Western Dietary Pattern (WDP)

Table 3. Baseline characteristics of the 7216 participants according to quartiles of adherence to Mediterranean Dietary Pattern (MDP)

Table 4. Hazard ratios for total mortality according to quartiles of baseline adherence to the Western dietary pattern and Mediterranean dietary pattern in the PREDIMED study with the three intervention groups considered together (n= 7216)

Table 5. Hazard Ratios for primary cardiovascular event according to quartiles of baseline adherence to the Western dietary pattern and Mediterranean dietary pattern in the PREDIMED study with the three intervention groups considered together (n= 7216)

Table 6. a) Hazard Ratios for acute myocardial infarction according to quartiles of baseline adherence to the Western dietary pattern and Mediterranean dietary pattern in the PREDIMED study with the three intervention groups considered together.

b) Hazard Ratios for stroke according to quartiles of baseline adherence to the Western dietary pattern and Mediterranean dietary pattern in the PREDIMED study with the three intervention groups considered together.

c) Hazard Ratios for cardiovascular death according to quartiles of baseline adherence to the Western dietary pattern and Mediterranean dietary pattern in the PREDIMED study with the three intervention groups considered together.

Figure 1. Absolute risks¹ for overall mortality according to the quartiles of baseline adherence to WDP (a) or MDP (b) and to the randomly allocated arm of the trial (intervention groups or control group).

¹ Absolute risks were adjusted for potential confounders using inverse probability weighting.

MDP: Mediterranean Dietary Pattern, WDP: Western Dietary Pattern.

Figure 2. Absolute risks² for cardiovascular events according to the quartiles of baseline adherence to WDP (a) or MDP (b) and to the randomly allocated arm of the trial (intervention groups or control group)..

MDP: Mediterranean Dietary Pattern, WDP: Western Dietary Pattern.

² Absolute risks were adjusted for potential confounders using inverse probability weighting.

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