

Relative validity of a semi-quantitative food-frequency questionnaire in an elderly Mediterranean population of Spain

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The aim of the present study was to assess reproducibility and relative validity of a self-administered FFQ used in the PREDIMED Study, a clinical trial for primary prevention of CVD by Mediterranean diet in a population at high cardiovascular risk. The FFQ was administered twice (FFQ1 and FFQ2) to explore reproducibility at 1 year. Four 3 d dietary records (DR) were used as reference to explore validity; participants therefore recorded their food intake over 12 d in the course of 1 year. The degree of misclassification in the FFQ was also evaluated by a contingency table of quintiles comparing the information from the FFQ2 and the DR. A total of 158 men and women (aged 55–80 years) were asked not to modify their dietary habits during the study period. Reproducibility for food groups, energy and nutrient intake, explored by the Pearson correlation coefficient (r) ranged 0.50–0.82, and the intraclass correlation coefficient (ICC) ranged from 0.63 to 0.90. The FFQ2 tended to report higher energy and nutrient intake than the DR. The validity indices of the FFQ in relation to the DR for food groups and energy and nutrient intake ranged (r) from 0.24 to 0.72, while the range of the ICC was between 0.40 and 0.84. With regard to food groups, 68–83 % of individuals were in the same or adjacent quintile in both methods, a figure which decreased to 55–75 % for energy and nutrient intake. We concluded that FFQ measurements had good reproducibility and a relative validity similar to those of FFQ used in other prospective studies.

Validity: Reproducibility: FFQ: Dietary records: Mediterranean population: PREDIMED Study: Spain

Nutrition has been linked to diseases that cause the highest rates of mortality and morbidity in developed populations. There is convincing evidence of the relationship between dietary factors and the development of chronic diseases^(1–6). Several studies have suggested that individuals consuming a traditional Mediterranean diet have a reduced risk of CVD, cancer and neurodegenerative diseases^(7–12).

When we need to explore the nutritional status of a population and associate it to their health status, it is very important to accurately estimate food intake in that population. However, all the available methods for measuring food intake have some limitations that must be taken into account before choosing the one that is most appropriate for the study aims and design. Self-reporting of food intake can lead to a certain degree of measurement error that may dilute and/or bias the relationship between nutrient and/or

food intake and the disease of interest. Some suggested reasons for this measurement error are within-subject variability in food intake, difficulties in remembering which and how much food was consumed, and the ability or lack thereof to report food consumption.

FFQ are widely accepted as the most suitable assessment tool in large epidemiological studies which associate diet with chronic diseases because they provide dietary information over a long period of time without altering usual food habits and are relatively inexpensive when self-administered⁽¹³⁾.

The FFQ is currently the most frequent method used to assess food intake in large population-based studies. This method, with a single measurement, provides a convenient assessment of the habitual dietary intake of an individual. It is accepted that FFQ allow a reasonably accurate ranking

Abbreviations: DR, dietary record; ICC, intraclass correlation coefficient.

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of individuals; however, the ability of FFQ to quantify the absolute intake of foods and specific nutrients^(14,15) is limited.

Our aim was to assess the reproducibility and the relative validity of a self-administered FFQ designed to measure habitual food and nutrient intake and to be used in the PREDIMED Study^(7,16), a large multicentre randomised controlled clinical trial for the primary prevention of CVD by means of a traditional Mediterranean diet in a high-risk population.

Materials and methods

The design of the present validation study is shown in Fig. 1. The PREDIMED FFQ was adapted from a previously validated Spanish FFQ⁽¹⁷⁾. The original questionnaire had 118 items. After several panel discussions among the researchers of the PREDIMED network, another nineteen new items were added, including food items only recently introduced in Spain (kiwi, all-bran cereals) and fruits, vegetables and whole grains with special interest for a Mediterranean diet trial (for example, onion, garlic, aromatic herbs, mushrooms and some other varieties of fungi). In order to better assess the intake of flavonoids, phytosterols and polyphenols, several additional questions about varieties of olive oil, nuts, cocoa, tea and wines were also added. A complementary question about the frequency of consuming fruit as dessert was also added. Some portion sizes were updated with respect to the previous version to represent the serving size of that item now most frequently consumed currently in Spain. The FFQ was designed to be semi-quantitative and optically readable for data input. The FFQ included 137 food items, and the frequencies of consumption of the food items were reported on an incremental scale with nine levels (never or almost never, 1–3 times per month, once per week, 2–4 times per week, 5–6 times per week, once per d, 2–3 times per d, 4–6 times per d and more than six times per d). A full version of the FFQ can be downloaded (http://www.unav.es/departamento/preventiva/predi_educational). The reported frequencies of food consumption were converted to number of intakes per d and multiplied by the weight of the portion size indicated.

A total of 158 independent individuals (seventy-three men and eighty-five women) aged between 55 and 80 years old and from three different Spanish areas (Tarragona, Pamplona and Valencia) of the PREDIMED Study⁽⁷⁾ were evaluated. The participants were chosen from users of primary health centres and the compliance rate was 85%. Exclusion criteria were: dementia, severe terminal disease, enteral or parenteral nutrition, and medically prescribed changes in dietary habits. The participants were asked not to modify their dietary habits during the study's 1-year duration.

The present study was conducted according to the guidelines laid down in the Declaration of Helsinki and all procedures involving human subjects were approved by the Ethics Committees of the Fundació Jordi Gol, University of Valencia and University of Navarra. Written informed consent

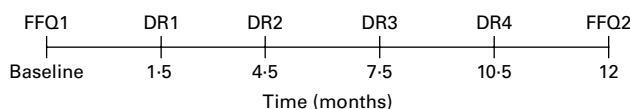


Fig. 1. Design of the PREDIMED's FFQ validation study. DR, dietary record.

was obtained from all subjects. After informed consent was obtained, the sociodemographic characteristics and health habits of the participants were recorded.

The dietary record (DR) was used as a reference to explore validity. The participants were asked to record their food intake for 12 d (3 d for each of the four times).

Food intake was grouped into the food groups (dairy products, eggs, meat or meat products, fish or seafood, vegetables, potatoes, fruits, total nuts, legumes, cereals, olive oil, pastries, cakes or sweets, and alcoholic drinks) that were used in the analysis.

To calculate energy and nutrient intake from both methods (FFQ and DR) we used the Spanish food composition tables⁽¹⁸⁾. The nutrient composition of each FFQ item was calculated as the mean of all explicitly mentioned foods.

Nobody reported extremes or implausible energy intakes according to DR (> 16 790 kJ/d (> 4013 kcal/d) and < 3360 kJ/d (< 803 kcal/d) for males and > 14 690 kJ/d (> 3511 kcal/d) and < 2100 kJ/d (< 502 kcal/d) for females⁽¹³⁾; therefore, nobody was excluded from the analysis of results on this basis.

Statistical methods

The mean daily intake of the twelve (four 3 d) dietary records was used as representative of DR. The days of the week recorded using DR for each individual were randomly selected, and the appropriate weight was given to weekends (3 or 4 d for each participant).

Intakes of food groups, energy and nutrients were transformed (\log_{10}) to optimise the normality of the distribution.

We calculated Pearson product-moment correlation coefficients and intraclass correlation coefficients (ICC) of log-transformed food and nutrient intakes and energy-adjusted nutrient intakes using the residual method⁽¹³⁾.

To correct for within-individual error in the measurement of the DR that tends to reduce correlation coefficients toward zero, the observed correlation coefficient was multiplied by the de-attenuation factor $(1 + (\sigma_w^2/\sigma_b^2)/n)^{0.5}$, where σ_w^2 is the within-individual variance, σ_b^2 is the between-individual

Table 1. Baseline characteristics of the 158 participants (Mean values and standard deviations or percentages)

Characteristics	Mean	SD
Age (years)	65.5	7.4
Weight (kg)	73.9	12.9
Height (cm)	164.2	9.4
BMI (kg/m ²)	27.3	4.1
Sex (%)		
Male	46.2	
Female	53.8	
Educational level (%)		
Primary	51.9	
Secondary	24.1	
University	24.0	
Marital status (%)		
Single	5.1	
Married	81	
Widowed	11.4	
Divorced	1.3	
Separated	1.3	
Current smokers (%)	19.2	
Physical activity (%)	62.6	

variance, and n represents the number of replicate measurements (here $n = 12$)⁽¹³⁾. The within- and between-individual variance components were determined by a random-effects model with the record intake as the dependent and subject identification number as the independent variable⁽¹⁹⁾.

Quintiles of intake were also calculated, and the degree of gross misclassification in FFQ with respect to the DR was evaluated using contingency tables of quintiles, which were constructed using the FFQ2 and the average DR.

To graphically check the agreement between the two methods in terms of absolute intake we used the analysis proposed by Bland & Altman⁽²⁰⁾, which plots the difference in intake between the two methods (FFQ2 – DR) against the mean intake of the two measures ((FFQ2 + DR)/2). The plots include lines for the mean difference and the so-called 'limits of agreement', defined as mean \pm 1.96 SD of the mean.

Results

The general characteristics of the 158 participants in the study are shown in Table 1. The majority of the participants were of

primary educational level, married, non-smokers and did some physical activity at least once a week (walking, yoga, swimming, gymnastics, bicycle, rowing, gardening, hunting, football, trekking, tennis, aerobics or jogging). Of the participants, 53.8, 37.7 and 12.0% stated that they had hypertension, dyslipidaemia and type 2 diabetes, respectively.

Daily intake of food groups, energy, macro- and micro-nutrients (as assessed by FFQ1, FFQ2 and DR) are shown in Table 2. To explore relative validity we chose the second FFQ instead of the first one because the former covers the same time period (i.e. previous year) as the four 3 d DR used as the reference. The FFQ2 tended to report higher energy and nutrient intake than the DR, which indicates that the former method may overestimate dietary intake to some degree. On the other hand, this may just be reflecting that the DR underestimates many food groups, with the notable exceptions of eggs, legumes and olive oil.

Table 3 show the reproducibility of the FFQ measurements (FFQ1 *v.* FFQ2) in terms of the Pearson correlation coefficient and ICC. For food groups the Pearson correlation coefficients ranged from 0.55 to 0.82 and were generally

Table 2. Daily consumption of food groups and nutrients estimated by two FFQ (FFQ1 and FFQ2) and four 3 d dietary records (DR)

(Mean values and standard deviations)

Item (units/d)	FFQ1		FFQ2		DR	
	Mean	SD	Mean	SD	Mean	SD
Food groups (g)						
Dairy products	382.7	227.8	374.5	216.7	279.8	129.7
Eggs	20.5	9.9	20.9	10.8	24.8	15.9
Meat or meat products	149.7	63.7	149.3	58.1	126.1	54.2
Fish or seafood	122.3	73.5	110.9	55.2	98.7	44.9
Vegetables	471.4	230.0	438.4	203.2	247.5	99.4
Potatoes	73.7	57.2	73.5	43.7	74.2	54.4
Fruits	308.4	188.6	311.1	207.1	318.6	157.8
Total nuts	8.1	15.8	6.8	11.1	4.2	7.4
Legumes	18.2	9.7	19.4	11.1	36.3	28.2
Cereals	159.8	100.9	153.2	90.0	158.0	75.2
Olive oil	29.0	16.8	33.9	20.1	43.7	15.9
Pastries, cakes or sweets	46.8	71.4	44.8	66.6	23.9	26.3
Alcoholic drinks	132.5	179.7	143.4	198.7	145.1	231.0
Energy and nutrients						
Energy (kJ)	10 015.2	3360.2	9910.6	3392.8	8932.0	2366.5
Energy (kcal)	2393.7	803.1	2368.7	810.9	2134.8	565.6
Protein (g)	103.1	27.8	99.7	26.2	93.4	23.5
Carbohydrates (g)	259.8	107.0	252.9	109.0	239.7	77.6
Fat (g)	96.1	38.1	97.6	36.3	89.3	25.7
SFA (g)	27.1	11.6	27.0	11.7	20.3	7.1
MUFA (g)	44.8	18.8	46.9	19.0	44.7	13.5
PUFA (g)	15.5	8.5	15.0	6.9	12.7	4.7
Cholesterol (mg)	419.3	150.9	409.0	137.4	210.3	87.6
Fibre (g)	28.2	10.7	26.6	10.2	22.0	8.4
Vitamin C (mg)	230.6	112.8	225.8	110.5	163.7	64.3
Thiamin (mg)	2.4	1.0	2.3	0.9	1.3	0.4
Riboflavin (mg)	2.1	0.7	2.1	0.7	1.5	0.3
Niacin (mg)	42.9	12.7	41.4	11.5	33.9	8.8
Pyridoxine (mg)	2.5	0.8	2.4	0.7	1.6	0.5
Cobalamin (μ g)	10.8	6.2	10.2	4.6	7.1	4.0
Vitamin A (IU)*	1678.2	925.0	1619.1	846.3	988.7	446.8
Vitamin D (μ g)	7.0	5.8	6.0	3.8	4.5	3.0
Vitamin E (mg)	11.2	4.9	10.6	4.1	6.7	2.7
Marine origin <i>n</i> -3 (g)	1.0	0.8	0.8	0.5	0.8	0.5
Non-marine origin <i>n</i> -3 (g)	1.5	1.0	1.4	0.8	1.1	0.4
Linolenic acid (g)	1.5	1.0	1.4	0.8	1.1	0.4

* 1 IU vitamin A = 0.3 μ g retinol or 0.6 μ g β -carotene.

Table 3. Reproducibility of FFQ: correlations between food groups, energy intake and nutrient intake in FFQ1 and FFQ2*

	Pearson correlation coefficient		Intraclass correlation coefficient	
	Unadjusted	Energy-adjusted	Unadjusted	Energy-adjusted
Food groups				
Dairy products	0.81	–	0.89	–
Eggs	0.62	–	0.76	–
Meat or meat products	0.70	–	0.82	–
Fish or seafood	0.49	–	0.65	–
Vegetables	0.80	–	0.89	–
Potatoes	0.61	–	0.75	–
Fruits	0.62	–	0.76	–
Total nuts	0.66	–	0.80	–
Legumes	0.47	–	0.63	–
Cereals	0.56	–	0.72	–
Olive oil	0.55	–	0.71	–
Pastries, cakes or sweets	0.73	–	0.84	–
Alcoholic drinks	0.82	–	0.90	–
Energy and nutrients				
Energy	0.66	–	0.79	–
Protein	0.55	0.50	0.71	0.67
Carbohydrates	0.71	0.67	0.83	0.80
Fat	0.61	0.58	0.75	0.73
SFA	0.67	0.67	0.81	0.80
MUFA	0.59	0.57	0.74	0.73
PUFA	0.63	0.58	0.77	0.73
Cholesterol	0.59	0.56	0.74	0.72
Fibre	0.76	0.78	0.86	0.88
Vitamin C	0.77	0.77	0.87	0.87
Thiamin	0.72	0.78	0.84	0.88
Riboflavin	0.70	0.75	0.83	0.86
Niacin	0.58	0.58	0.74	0.74
Pyridoxine	0.66	0.73	0.80	0.84
Cobalamin	0.54	0.53	0.70	0.69
Vitamin A	0.74	0.72	0.85	0.84
Vitamin D	0.53	0.51	0.69	0.68
Vitamin E	0.60	0.60	0.75	0.75
Marine origin <i>n</i> -3	0.52	0.52	0.68	0.68
Non-marine origin <i>n</i> -3	0.66	0.58	0.79	0.73
Linolenic acid	0.66	0.60	0.79	0.75

* Intakes of food groups, energy and nutrients were transformed (\log_{10}) to improve normality.

above 0.6 (mean 0.65). ICC ranged from 0.63 to 0.90 and were mostly above 0.75 (mean 0.78). For energy and nutrient intake, the Pearson correlation coefficient ranged from 0.52 to 0.76 (mean 0.64) and from 0.50 to 0.78 (mean 0.63) for energy-unadjusted and -adjusted data, respectively; ICC ranged from 0.68 to 0.87 (mean 0.78) and from 0.67 to 0.88 (mean 0.77) for energy-unadjusted and -adjusted data, respectively.

Correlation coefficients that explored the validity of FFQ in relation to DR are shown in Table 4. For food groups, the Pearson correlation coefficients ranged from 0.29 to 0.72 and were mostly above 0.4 (mean 0.53); ICC ranged from 0.40 to 0.84 and were mostly above 0.70 (mean 0.67). For energy and nutrient intake, the Pearson correlation coefficient ranged from 0.24 to 0.49 (mean 0.39) and from 0.23 to 0.68 (mean 0.44) for energy-unadjusted and -adjusted data, respectively; ICC ranged from 0.44 to 0.78 (mean 0.55) and from 0.37 to 0.80 (mean 0.59) for energy-unadjusted and -adjusted data, respectively. De-attenuated coefficients are not shown because their values were very similar, but are available upon request.

In Table 5, we present the percentage of gross misclassification (both over- and underestimation by the FFQ) and complete or adjacent agreement as indices of validity of FFQ in categorising individuals. For the food groups, between 68 and 83 % of individuals were in the same or adjacent quintile with both of the two methods (mean 72.3 %) and 55–75 % (mean 67.7 %) were in the same quintile for energy and nutrients. The overall extreme misclassification was always lower than 5 % (with the single exception of potatoes (5.1 %) among food groups, and the single exception of cholesterol (5.7 %) among nutrients).

The Bland–Altman analysis confirmed a general overestimation – as previously mentioned – of the consumption of food groups and of nutrient intake by the FFQ. Examples of the several patterns of relationship observed between the two methods for the consumption of food groups are plotted in Fig. 2 and for total energy and specific nutrient intakes are shown in Fig. 3. For most food groups, energy and macro- and micronutrients, the mean error in intake assessment by FFQ does not change with increased dietary intake (such as meat and meat products, fish or seafood, potatoes,

Table 4. Validity of FFQ: correlations between food groups, energy intake and nutrient intake in four 3 d dietary records and FFQ2*

	Pearson correlation coefficient		Intraclass correlation coefficient	
	Unadjusted	Energy-adjusted	Unadjusted	Energy-adjusted
Food groups				
Dairy products	0.72	–	0.84	–
Eggs	0.42	–	0.58	–
Meat or meat products	0.61	–	0.75	–
Fish or seafood	0.42	–	0.59	–
Vegetables	0.70	–	0.81	–
Potatoes	0.37	–	0.54	–
Fruits	0.57	–	0.72	–
Total nuts	0.38	–	0.55	–
Legumes	0.29	–	0.40	–
Cereals	0.58	–	0.71	–
Olive oil	0.60	–	0.71	–
Pastries, cakes or sweets	0.53	–	0.68	–
Alcoholic drinks	0.70	–	0.82	–
Energy and nutrients				
Energy	0.36	–	0.53	–
Protein	0.28	0.40	0.44	0.55
Carbohydrates	0.38	0.56	0.55	0.71
Fat	0.47	0.46	0.63	0.62
SFA	0.52	0.61	0.68	0.75
MUFA	0.51	0.44	0.67	0.61
PUFA	0.43	0.42	0.60	0.60
Cholesterol	0.37	0.23	0.54	0.37
Fibre	0.49	0.60	0.66	0.75
Vitamin C	0.65	0.68	0.78	0.80
Thiamin	0.32	0.38	0.45	0.46
Riboflavin	0.29	0.41	0.40	0.56
Niacin	0.35	0.50	0.51	0.66
Pyridoxine	0.36	0.56	0.52	0.71
Cobalamin	0.38	0.41	0.54	0.57
Vitamin A	0.44	0.42	0.60	0.59
Vitamin D	0.24	0.24	0.38	0.39
Vitamin E	0.35	0.38	0.51	0.55
Marine origin <i>n</i> -3	0.33	0.32	0.49	0.49
Non-marine origin <i>n</i> -3	0.36	0.37	0.50	0.52
Linolenic acid	0.37	0.39	0.52	0.55

* Intakes of food groups, energy and nutrients were transformed (\log_{10}) to improve normality.

fruits, total nuts, cereals, alcoholic drinks, energy, protein, carbohydrates, fat, SFA, PUFA, niacin, cobalamin, vitamin D, marine origin *n*-3 and linolenic acid), despite the wider scatter with increasing mean intake values that is sometimes observed. For other dietary variables a systematic variation in agreement between the two methods was observed with increased dietary intake; for dairy products, vegetables, pastries, cakes or sweets, MUFA, cholesterol, fibre, vitamin C, thiamin, riboflavin, pyridoxine, vitamin A, vitamin E and non-marine *n*-3, low intakes were underestimated whereas high intakes were overestimated by the FFQ; and for eggs, legumes and olive oil low intakes were overestimated and high intakes were underestimated by the FFQ.

Discussion

In the present study, we assessed the reproducibility and validity of a 137-item semi-quantitative FFQ used in a large multicentre randomised controlled clinical trial for the primary prevention of CVD with a traditional Mediterranean diet in a high-risk population (PREDIMED Trial). The present results show that this FFQ provides a highly reproducible assessment

of food and nutrient intake and reasonably good validity in relation to DR for many food groups and nutrients (correlation coefficients > 0.60). Agreement between these two methods was also reasonably acceptable and gross misclassification using the FFQ was found to be very unlikely.

Although there are good biomarkers for energy⁽²¹⁾ and some for nutrient intake^(22,23), there is no available biomarker that can provide a reliable measurement of food habits as a whole. Therefore, a dietary assessment method is frequently chosen as a reference when validating an FFQ⁽²⁴⁾, even for specific nutrients⁽¹⁴⁾. Since perfectly error-free reference methods for diet measurement cannot be obtained, it is understood that the reference method provides better data but of limited accuracy (i.e. it is not a true 'gold standard'). It has been suggested that the measurement errors of FFQ and reference methods may be independent⁽²⁵⁾, but there is some disagreement on this point^(26,27). In any case, because recovery biomarkers are not available for most foods or nutrients, usually there is no other choice but to use self-reported measures as reference methods, despite the bias associated with them. In this context, the DR has been widely used as a reference method in FFQ validation studies. In the present

Table 5. Gross misclassification (%) of food groups, energy and energy-adjusted nutrient distribution: classification in opposite quintiles or in the same/adjacent quintile in the second FFQ (FFQ2) v. classification according to the dietary records mean

Item	Lowest quintile in four 3 d dietary records and highest quintile in FFQ2	Highest quintile in four 3 d dietary records and lowest quintile in FFQ2	Classified in FFQ2 within one quintile in four 3 d dietary records
Food groups			
Dairy products	3.2	0.6	74.1
Eggs	1.3	0.0	75.3
Meat or meat products	0.0	1.3	68.4
Fish or seafood	1.9	2.5	69.0
Vegetables	1.3	0.0	76.6
Potatoes	3.2	1.9	62.7
Fruits	0.6	1.9	73.4
Total nuts	2.5	0.0	70.9
Legumes	0.0	3.8	69.6
Cereals	1.9	1.3	76.6
Olive oil	3.2	0.6	69.6
Pastries, cakes or sweets	0.0	1.9	75.3
Alcoholic drinks	0.0	0.6	82.9
Energy and nutrients			
Energy	1.3	2.5	68.4
Protein	0.6	2.5	68.4
Carbohydrates	0.6	1.3	72.2
Fat	1.3	0.6	68.4
SFA	0.0	0.6	74.8
MUFA	2.5	0.6	71.5
PUFA	0.6	1.3	64.6
Cholesterol	1.9	3.8	55.1
Fibre	0.0	1.3	71.5
Vitamin C	0.0	0.6	74.7
Thiamin	1.3	1.3	65.2
Riboflavin	0.6	2.5	68.4
Niacin	0.0	0.6	66.5
Pyridoxine	0.0	0.6	72.8
Cobalamin	1.3	1.9	69.6
Vitamin A	1.3	2.5	67.1
Vitamin D	0.6	3.2	58.9
Vitamin E	3.2	1.3	69.0
Marine origin <i>n</i> -3	0.6	3.8	65.8
Non-marine origin <i>n</i> -3	1.3	1.9	64.6
Linolenic acid	0.6	1.9	64.6

study, the DR was applied for 12 d (in four 3 d periods); although a longer recording period may have better reflected true intake, we chose to record no more than 12 d in order to prevent a possible lack of compliance and the subsequent detriment to the quality of the data obtained.

We found a general overestimation of food group and nutrient intake with the FFQ, though this result could be somewhat artificial given potential underestimations by the DR. FFQ usually reflect higher estimates for most food groups and nutrients than the reference method⁽²⁸⁾, particularly if the FFQ exceeds 100 items⁽²⁹⁾, as does ours. A recent quantitative review reported that FFQ with more items are better able to rank individuals according to their intake⁽³⁰⁾. This overestimation may be due to measurement errors introduced by differences in conceptualisation of portion sizes, misinterpretation of specific items, and frequency and serving size differences between FFQ and DR. This is especially the case when there are multiple foods in an FFQ item. Moreover, averaging amounts of intake over 1 year, as we have done, is prone to estimation misjudgement that may be different from estimation errors in recording very recent amounts of intake as in DR.

The test–retest interval between the two FFQ is an important factor which influences reproducibility. If the interval is too short, responses to the second FFQ may be influenced by the memory of the first FFQ and reproducibility will be overestimated. However, if the interval is too long, the dietary pattern may have actually changed, which could lead to an underestimation of reproducibility⁽³¹⁾. According to a comprehensive review, the time interval in published reports ranges from 2 h to 15 years⁽²⁴⁾. In the present study, we chose a time interval of 1 year to prevent the types of errors mentioned above, and asked participants not to change their dietary habits during the whole study period. We observed small differences in mean intakes (for food groups, energy and nutrients) between FFQ1 and FFQ2. Studies with the same interval time between the two FFQ have shown similar correlation coefficient ranges⁽³²⁾.

Validity of food group intake was better than that of energy and nutrients.

Correlation coefficients for food groups, energy and nutrient intake were comparable with other similar validation studies^(33–37).

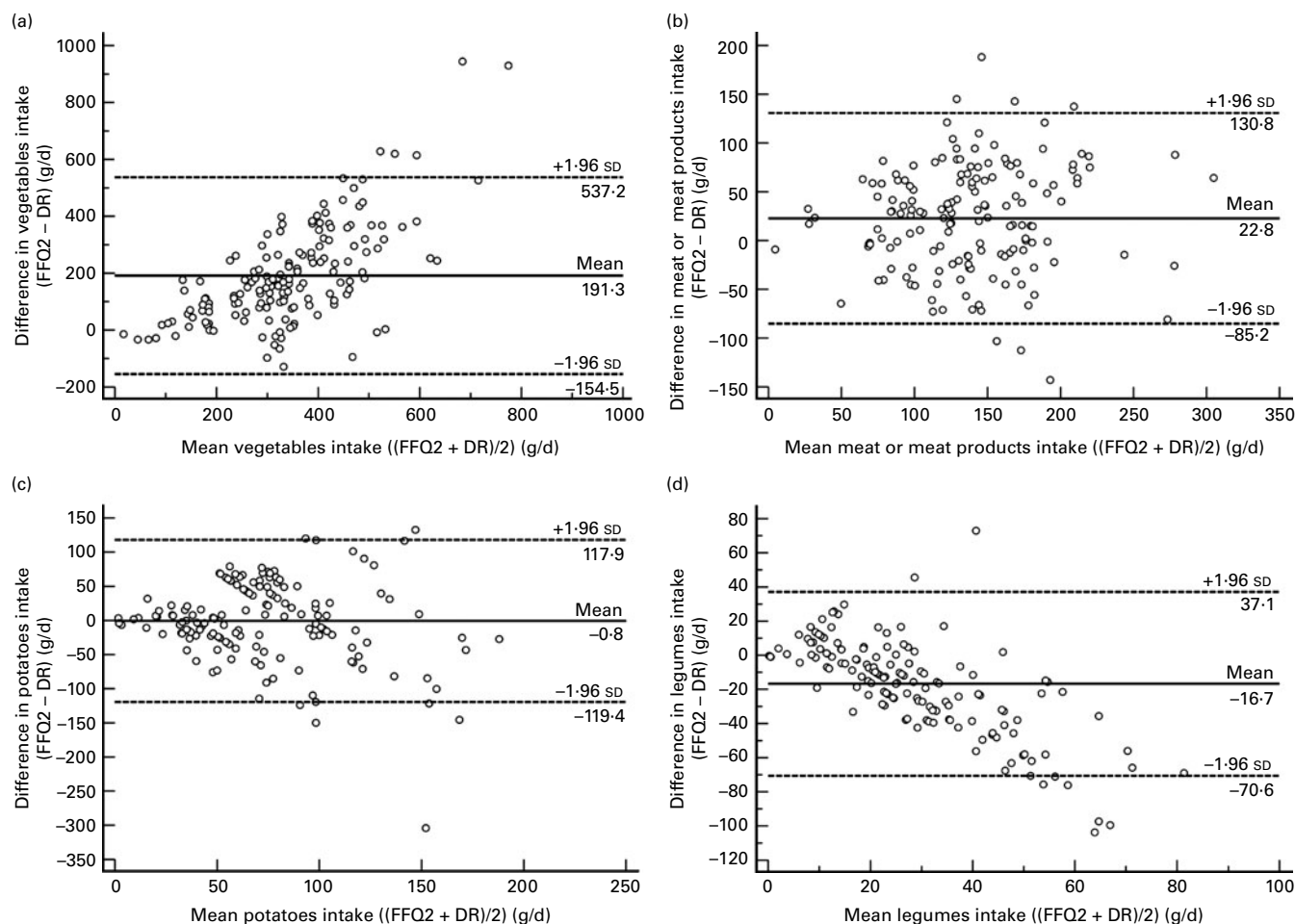


Fig. 2. Bland–Altman plots showing the relationship between difference in the daily intake of (a) vegetables, (b) meat or meat products, (c) potatoes and (d) legumes estimated with the second FFQ (FFQ2) and four 3 d dietary records (DR), and the corresponding mean daily intake estimated by the two methods. Lines are mean difference (—) and lower and upper 95% limits of agreement (---).

The validity study of an FFQ specifically designed to estimate PUFA intake using the weighted food record as its reference method found only slightly better results than those reported here⁽³⁸⁾.

When calculating nutrient intake from the information given in an FFQ, the assumption that each explicitly mentioned food item contributes equally to the intake of the nutrient may be wrong. Perhaps a more realistic approach would be to obtain – in the same population – the relative weight of each mentioned and unmentioned food and use this information to calculate the nutrient intake. Such information could also be used to determine portion size. These limitations may partially explain the relatively low level of disagreement between the two methods compared in the validation results.

We must take into account that our participants voluntarily agreed to take part in the present study, and they may tend to respond more accurately to FFQ and DR or have different dietary habits from non-participants.

The correction for random errors associated to within-individual variation almost always increases correlation values. Nevertheless, in the present results, de-attenuated correlations were similar to non-corrected estimates, perhaps because of a relatively low within-individual variation in relation to the

between-individual variation in our participants and/or because of the large number of days – 12 – recorded.

Given that FFQ are often used to categorise individual intake rather than to obtain an exact quantification of nutrient intake, it is appropriate to explore the degree of gross misclassification when quintiles are used to rank individuals according to their relative intake. In this aspect, we found a very low frequency of gross misclassification and a relatively high agreement, especially for the specific food groups and nutrients more strongly related to cardiovascular risk.

In the Bland–Altman plots, no apparent change in the magnitude of between-measurement (FFQ v. DR) differences was observed across mean values of both methods for most foods and nutrients. See, for example, the plot for total energy intake (Fig. 3(a)). The interpretation for these nutrients is that agreement of the FFQ with the DR was similar regardless of their average intake. However, in some other cases, a wider scatter was observed, suggesting worse agreement when mean intakes were higher (for example, meat and meat products, potatoes, protein or cobalamin in Fig. 2(b), 2(c), 3(b) and 3(d)). The interpretation here is that errors were somewhat proportional to mean intake.

In conclusion, this 137-item self-administered semi-quantitative FFQ appears to be a useful instrument for the

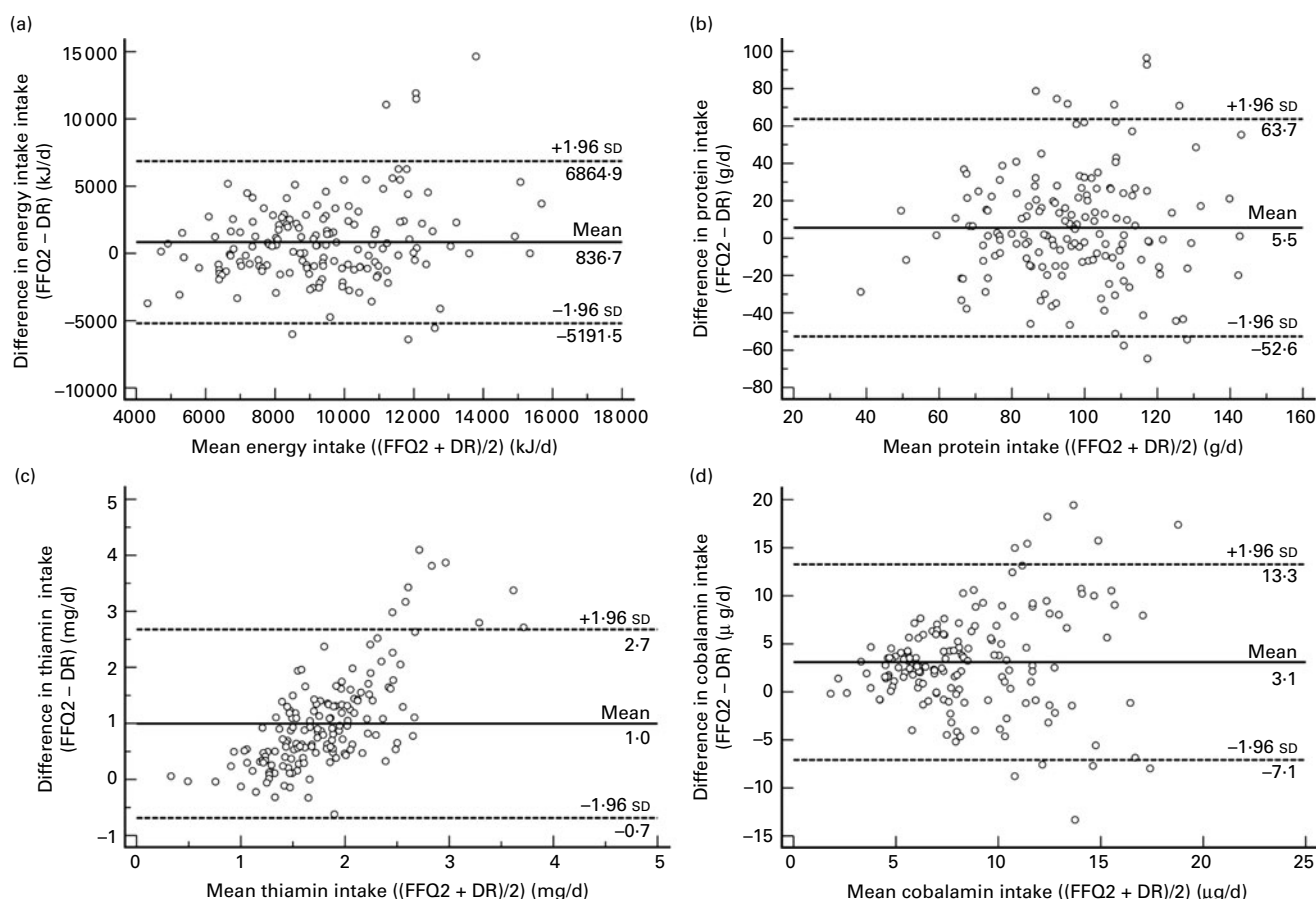


Fig. 3. Bland–Altman plots showing the relationship between difference in the daily intake of (a) energy, (b) protein, (c) thiamin and (d) cobalamin estimated with the second FFQ (FFQ2) and four 3 d dietary records (DR), and the corresponding mean daily intake estimated by the two methods. Lines are mean difference (—) and lower and upper 95% limits of agreement (---).

estimation of food habits and nutrient intake in the epidemiological research of diet as a risk factor for CVD and other chronic diseases and for dietary surveillance in Spain. It was found to have acceptable levels of reproducibility and validity.

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J. L. P. and M. P.-B. participated in design of the study, acquisition of data, analysis of data and interpretation of results, and revising the article for relevant intellectual content and contribution to drafting of the manuscript.

I. Z., P. C. and E. T. participated in acquisition of data and interpretation of results, and revising the article for relevant intellectual content and contribution to drafting of the manuscript.

D. C. participated in design of the study, interpretation of results and drafting of the manuscript with intellectual and scientific input.

M. A. M.-G. and J. S.-S. participated in design of the study, calculation of nutritional data, interpretation of results and contribution to drafting of the manuscript with intellectual and scientific input.

J. M. M.-M. participated in interpretation of results and revising the article for relevant intellectual content and contribution to drafting of the manuscript.

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