

TITLE

Mediterranean diet increases endothelial function in adults: A systematic review and meta-analysis of randomized controlled trials

AUTHORS

Oliver M Shannon¹, Inês Mendes¹, Christina Köchl¹, Mohsen Mazidi², Ammar W Ashor³, Sofia Rubele^{1,4}, Anne-Marie Minihane⁵, John C Mathers¹, Mario Siervo^{1,6}

AFFILIATIONS

¹Human Nutrition Research Centre, Institute of Cellular Medicine, Newcastle University, UK

² Department of Biology and Biological Engineering, Food and Nutrition Science, Chalmers University of Technology, SE-412 96 Gothenburg, Sweden

³ Department of Internal Medicine, College of Medicine, Mustansiriyah University, Baghdad, Iraq

⁴Faculty of Medicine & Surgery, University of Verona, Italy

⁵Norwich Medical School, Bob Champion Research & Education, University of East Anglia

⁶School of Life Sciences, The University of Nottingham Medical School, Queen's Medical Centre, Nottingham, NG7 2UH, UK

AUTHOR SURNAMES FOR PUBMED INDEXING

Shannon, Mendes, Köchl, Mazidi, Ashor, Rubele, Minihane, Mathers, Siervo

CONFLICT OF INTEREST STATEMENT AND FUNDING DISCLOSURE

Oliver M Shannon: No conflict of interest

Inês Mendes: No conflict of interest

Christina Köchl: No conflict of interest

Mohsen Mazidi: No conflict of interest

Ammar W Ashor: No conflict of interest

Sofia Rubele: No conflict of interest

Anne-Marie Minihane: No conflict of interest

John C Mathers: No conflict of interest

Mario Siervo: No conflict of interest

No specific funding was received for this work.

CORRESPONDING AUTHOR:

Name: Dr. Oliver M Shannon

Mailing address: Room 2.050, William Leech Building, Medical School, Newcastle University, Newcastle-upon-Tyne, NE2 4HH

Telephone number: 0191 208 1140

Email: Oliver.Shannon@Newcastle.ac.uk

WORD COUNT: 3479

NUMBER OF FIGURES: 4

NUMBER OF TABLES: 3

ONLINE SUPPORTING MATERIAL: 1 supplementary data file

RUNNING TITLE

Mediterranean diet and endothelial function

ABBREVIATIONS:

BP	Blood pressure
BMI	Body mass index
CRP	C-reactive protein
CVD	Cardiovascular disease
DASH	Dietary approach to stop hypertension
EVOO	Extra virgin olive oil
FBF	Forearm blood flow
FMD	Flow mediated dilation
IL-6	Interleukin-6
LDL	Low density lipoprotein
MedDiet	Mediterranean dietary pattern
MUFA	Monounsaturated fatty acids
NO	Nitric oxide
PREDIMED	Prevención con Dieta Mediterránea
RCT	Randomised controlled trial
sICAM-1	Soluble intercellular adhesion molecule-1
TNF- α	Tumour necrosis factor- α
VCAM-1	Vascular cell adhesion protein-1

1 ABSTRACT

2 Background: The endothelium plays a key role in the maintenance of vascular health, and
3 represents a potential physiological target for dietary and other lifestyle interventions designed
4 to reduce risk for cardiovascular diseases (CVD) including stroke or coronary heart disease.

5 Objective: To conduct a systematic review and meta-analysis of randomized controlled trials
6 investigating effects of the Mediterranean dietary pattern (MedDiet) on endothelial function.

7 Methods: Medline, Embase and Scopus databases were searched from inception until January
8 2019 for studies that met the following criteria: 1) RCTs including adult participants, 2)
9 interventions promoting a Mediterranean dietary pattern, 3) inclusion of a control group and 4)
10 measurements of endothelial function. A random-effects meta-analysis was conducted. Meta-
11 regression and sub-group analyses were performed to identify whether effects were modified
12 by health status (i.e., healthy participants vs participants with existing comorbidities), type of
13 intervention (i.e., MedDiet alone or with a co-intervention), study duration, study design (i.e.,
14 parallel or crossover), body mass index (BMI), and age of participants.

15 Results: Fourteen articles reporting data for 1930 participants were included in the meta-
16 analysis. Study duration ranged from 4 weeks to 2.3 years. We observed a beneficial effect of
17 the MedDiet on endothelial function (standardised mean difference (SMD): 0.35 95% CI: 0.17,
18 0.53, $P < 0.001$, $I^2 = 73.68\%$). MedDiet interventions improved flow mediated dilation (FMD)
19 - the reference method for non-invasive, clinical measurement of endothelial function - by
20 1.66% (absolute change; 95% CI: 1.15, 2.17, $P < 0.001$, $I^2 = 0\%$). Effects of the MedDiet on
21 endothelial function were not modified by health status, type of intervention, study duration,
22 study design, BMI, and age of participants ($P > 0.05$).

23 Conclusions: MedDiet interventions improve endothelial function in adults, which suggest that
24 the protective effects of the MedDiet are evident at early stages of the atherosclerotic process
25 with important implications for the early prevention of CVD.

26 **PROSPERO registration number:** CRD42018106188.

27 **KEY WORDS:** Mediterranean diet, endothelial function, flow mediated dilation,
28 cardiovascular disease, healthy ageing, dietary patterns

29

30

31

32

33

34

35

36

37

38

39

40

41

42

43

44

45

46

47

48

49

50

51

52

53

54

55

56 INTRODUCTION

57 The endothelium plays a key role in the maintenance of vascular health via the secretion of
58 multiple signalling molecules including nitric oxide (NO), endothelins, selectins and adhesion
59 molecules which, in concert, control vasomotor tone, and have anti-atherogenic and anti-
60 proliferative actions (1,2). Loss of functional and structural integrity of the endothelium is
61 thought to be an early pathogenic step in the development of atherosclerotic lesions and the
62 subsequent onset of cardiovascular diseases (CVD) (3). Therefore, the endothelium has been
63 identified as a tractable physiological target for therapeutic interventions designed to reduce
64 risk for CVD such as stroke, coronary heart disease or peripheral arterial disease (4,5).

65

66 The Mediterranean dietary pattern (MedDiet) is characterised by high consumption of olive
67 oil, fruits, vegetables, legumes, nuts and seeds, and unrefined grains, moderate-to-high
68 consumption of fish, and low consumption of red meat and sugar-sweetened products such as
69 sweets, cakes, and pastries and is considered as one of the healthiest dietary patterns (6,7).
70 Evidence from randomised controlled trials (RCTs) such as the Lyon Diet Heart Study in
71 France (8) and the Prevención con Dieta Mediterránea (PREDIMED) trial in Spain (9)
72 demonstrates that the MedDiet is effective in both primary and secondary prevention of CVD.
73 Improvements in endothelial function with the MedDiet may be one of the key mechanisms
74 underpinning these beneficial effects (10). Evidence from RCTs demonstrates that, in
75 isolation, a number of components of the MedDiet, particularly olive oil (11), nuts (12), and
76 oily fish (13,14), improve endothelial function. In addition, there is evidence that the
77 composite MedDiet enhances endothelial function in both healthy subjects and patients with
78 cardiovascular and metabolic diseases. For example, a 6 month MedDiet intervention in
79 healthy older individuals induced highly significant improvements in endothelial function
80 measured via flow mediated dilation (FMD), with the percentage FMD approximately double

81 baseline values (absolute increase of ~1.3%) (15). Similarly, in patients with pre-diabetes and
82 diabetes a 1.5 year MedDiet intervention increased FMD by 1.1% and 1.4%, respectively (16).

83 In addition, markers of endothelial structure such as carotid artery intima media thickness have
84 also been shown to be improved with MedDiet interventions (17)

85

86 A previous systematic review and meta-analysis in 2014 reported improved endothelial
87 function and decreased inflammation with MedDiet versus control interventions (18).

88 However, that review also included MedDiet-like interventions such as the Dietary Approach
89 to Stop Hypertension (DASH) in the same analysis, which may elicit differential effects on

90 endothelial function compared with the traditional MedDiet. Moreover, in that systematic
91 review, only studies with intervention period lasting ≥ 12 weeks were included, and the effects

92 of shorter MedDiet interventions were not evaluated. These limitations, and the emergence of
93 considerable new research in this area over the past 5 years (e.g. 14,15,17–19), provide the

94 rationale for an updated systematic review and meta-analysis of RCTs. Therefore, the aim of

95 this study was to undertake a systematic review and meta-analysis of published RCTs exploring
96 the effects of MedDiet interventions on structural (e.g., intima media thickness) and/or

97 functional (e.g., FMD) measures of endothelial function in humans. In an attempt to understand
98 potential differences in findings, we also investigated whether effects were modified by health

99 status, type of intervention, study duration, study design, body mass index (BMI), and age of
100 participants.

101

102 **METHODS**

103 The present systematic review was conducted according to the Preferred Reporting Items for
104 Systematic Review and Meta-analyses (PRISMA) guidelines (21).

105

106 **Literature search**

107 Three databases (Medline, Embase and Scopus) were used to search for articles from inception
108 until January 2019. In addition, a manual search of reference lists of relevant reviews and
109 articles included in the systematic review was performed. The search was conducted based on
110 the pre-defined search terms (Mediterranean AND diet*) AND (“endotheli*” [All Fields] OR
111 “endothelial function” [All Fields] OR “endothelial dysfunction” [All Fields] OR “vascular
112 function” [All Fields] OR “blood flow” [All Fields] OR “vascular reactivity” [All Fields] OR
113 “vasodilation” [All Fields]) and (Mediterranean AND diet*) AND (“Flow-mediated dilatation”
114 OR “Flow-mediated dilation” OR FMD OR “Venous occlusion plethysmography” OR
115 “Peripheral arterial tonometry” OR “Nitric oxide” OR “Endothelial function” OR “Endothelial
116 dysfunction” OR “Carotid Intima-Media Thickness” OR “Pulse Wave Velocity” OR
117 “Augmentation Index”). Further details of the search strategy are provided in **Supplemental**

118 **Methods 1.**

119

120 **Study selection**

121 The following criteria were applied to identify articles to be included in this systematic review
122 and meta-analysis: 1) RCTs (no further exclusion criteria were applied in relation to study
123 design or blinding); 2) studies involving adults aged ≥ 18 years and no exclusion criteria were
124 applied for health status or smoking history; 3) MedDiet (which was defined as a MedDiet by
125 the authors of each study) administered alone or with other clinical, pharmaceutical or lifestyle
126 interventions if a comparable and valid control group was included (for example, MedDiet plus
127 exercise compared to control group including exercise alone); 4) studies reporting changes in
128 endothelial function for intervention and control groups separately; 5) no language or time
129 restrictions were applied in searching the databases.

130

131 Two investigators (CH, IM) independently screened the titles and abstracts of the articles to
132 evaluate eligibility for inclusion. If consensus was reached, articles were either excluded or
133 moved to the next stage (full-text). If consensus was not reached the articles was moved to the
134 full-text stage. The full-texts of the selected articles were appraised critically to determine
135 eligibility for inclusion in the systematic review. Disagreements were resolved by discussion
136 between the reviewers (including MS) until consensus was reached.

137

138 **Data extraction and quality assessment**

139 Data extraction was completed by one investigator (CH) and data entries were checked for
140 accuracy by a second investigator (IM). The following information was extracted from the
141 eligible articles: author, year of publication, country, study design, inclusion and exclusion
142 criteria, study duration, run-in phase, intention to treat analysis, sample size, type of
143 intervention (control and MedDiet), age, sex, ethnicity, randomisation procedure, blinding of
144 exposure and outcome measurements, compliance with the interventions, BMI, dietary
145 intervention, weight loss during the study, baseline and post-intervention measurements of
146 systolic and diastolic blood pressure (BP), baseline and post-intervention measurements of
147 endothelial function. In addition, two independent reviewers (CH, IM) utilised the Cochrane
148 risk of bias tool to assess the risk of bias of the included studies which was classified as 1) high
149 risk, 2) low risk or 3) unknown risk of bias (22) and any discrepancy was resolved by consensus
150 with a third reviewer (MS)

151

152 **Statistical analysis**

153 Statistical analyses were performed by using Comprehensive Meta-Analysis Software Version
154 2 (Biostat, Englewood, NJ, USA). For this purpose, sample size, the mean and SD of the
155 endothelial function measurements before and after the intervention period (for both MedDiet

156 intervention and control) were extracted and used in the analyses. When no baseline
157 measurements were reported, the sample size, means and SDs after the intervention were used.
158 If the mean and SD were not given, the sample size and the *P* value of the difference between
159 MedDiet and control were used to calculate the effect size (Cohen's *d*). For studies that reported
160 changes in endothelial function at two or more time-points, the last endothelial function
161 measurement was used in the meta-analysis. The calculation of the effect sizes using different
162 sets of data is performed automatically by the software using integrated algorithms (23). Data
163 not provided in the main text or tables were extracted from the figures. Some trials used more
164 than one method to assess changes in endothelial function (Table 1) which may lead to a
165 reduced independence of the measurements and, consequently, to over-estimation of the effect
166 size derived from the meta-analysis. This potential confounding factor was taken into account
167 during analysis by estimating the mean of the standardised effect sizes derived from each
168 endothelial function measurement within each such study to provide a more conservative
169 estimate of the effect size.

170

171 Effect sizes and 95% confidence intervals for the MedDiet interventions were calculated using
172 a using a weighted DerSimonian-Laird random effects model (24). Forest plots were generated
173 to present graphically the cumulative effect of MedDiet on endothelial function. Analyses were
174 conducted on all endothelial function measurements but also stratified by type of endothelial
175 function measurement (structural and functional). Functional measurements include FMD
176 derived from ultrasound, forearm blood flow (FBF) derived from phlethysmography or
177 cutaneous microcirculation derived from laser Doppler. Structural measurements include
178 intima media thickness or vessel size measured both by ultrasound. In addition, we performed
179 a sensitivity analyses to test the effects of MedDiet on FMD only, on the basis that this is the

180 reference method for non-invasive, clinical measurement of endothelial function (25) and was
181 used in the majority of studies.

182

183 Subgroup analyses were undertaken to investigate variables which may have influenced the
184 effects of MedDiet on endothelial function. These factors included: health status (healthy
185 subjects *vs* patients with existing comorbidities), type of intervention (MedDiet alone or
186 administered with other clinical or pharmaceutical interventions), type of endothelial function
187 measurement (functional versus structural) and study design (parallel or crossover). Random-
188 effect meta-regression analyses were used to determine whether participant baseline
189 characteristics (age, BMI) and duration of the study influenced the effect of the MedDiet on
190 endothelial function. Funnel plots and Egger's regression tests were performed to evaluate the
191 risk of publication bias. Heterogeneity was assessed by using Cochrane Q statistic; $P < 0.1$
192 indicates significant heterogeneity. The I^2 test was also utilised to assess heterogeneity across
193 trials where a value $< 25\%$ indicates low risk, 25-75% indicates moderate risk, and $>75\%$
194 indicates a high risk (26). Sensitivity analyses were conducted to identify the source of
195 heterogeneity by conducting stratified analyses or selectively removing studies with larger
196 effect size. All of the data used in the meta-analysis can be found in **Supplemental Tables 1-**
197 **5.**

198

199 **RESULTS**

200 **Search results**

201 The process of screening and selection of the studies is summarised in **Figure 1**. The primary
202 search of the three databases produced 12857 articles, after removal of duplicates. After title
203 and abstract screening, 15 full-text papers were retrieved for further evaluation. A further four
204 studies were found by manual searching of references of relevant reviews and studies.

205 Examination of the full text of the 19 included articles yielded 14 studies which were eligible
206 to be included in this systematic review and meta-analysis. Some of these papers reported
207 results from independent studies testing the effects of MedDiet on endothelial function
208 generating a total of 20 sets of independent measures of endothelial function using different
209 methods that were included in the meta-analyses.

210

211 **Study characteristics**

212 The total number of participants from the 14 articles included in this systematic review was
213 1930 with a median of 131 (range 20 - 438) participants per study. The median participant age
214 was 55 (range 38 - 71) years. Eleven of the RCTs included in the meta-analysis were parallel
215 trials with a control group, while 3 were crossover studies (22). The paired nature of the cross-
216 over trials was taken into account in the meta-analysis to minimise unit-of-analysis errors and
217 underestimation of the effect size. The duration of the interventions ranged from 4 weeks to
218 121 weeks (**Table 1**). Five studies investigated the effect of MedDiet in healthy participants
219 (15,16,27–29), three in people with diabetes (16,17,19), two in patients with elevated risk of
220 CVD (30,31), two in overweight or obese participants (20,32), one in patients with metabolic-
221 syndrome (33), one in people with pre-diabetes (16), one in patients with acute coronary
222 syndrome (34), and one in hypercholesteraemic men (35). Various permutations of the
223 MedDiet were prescribed, including: a MedDiet (n=8), a MedDiet plus supplementary nuts
224 (n=2), a MedDiet plus supplementary extra virgin olive oil (n=5), a MedDiet plus mono-
225 unsaturated fatty acids (n=2), and a MedDiet plus exercise (n=1). Additionally, a variety of
226 different control groups were employed. These were: a low-fat diet (n=8), a typical Swedish
227 diet (n=1), the Atkins low-carbohydrate diet (n=1), the participants habitual diet (n=2), the
228 National Cholesterol Education Program Diet (n=1), a non-Mediterranean diet plus exercise
229 (n=1), a saturated fatty acid diet (n=1), and a prudent diet (n=1). Several methods were used

230 to assess endothelial function in the included trials. The most commonly used methods were
231 FMD and carotid intima-media thickness. Other methods included forearm blood flow,
232 cutaneous microvascular function, baseline vessel size, and calculation of an endothelial
233 function score (**Table 1**).

234

235 **Meta-analysis**

236 Meta-analysis of the 20 sets of independent results showed that, overall, MedDiet improved
237 endothelial function (SMD: 0.35, 95% CI: 0.17, 0.53, $P<0.001$; **Figure 2**). Heterogeneity
238 between studies was significant ($Q=64.60$, $I^2=73.68\%$, $P<0.001$). However, the removal of two
239 studies with wider effect estimates (27,33) (**Supplemental Fig. 1**) explained the heterogeneity
240 of the results ($Q=13.82$, $I^2=0\%$, $P=0.53$) while still confirming a significant effect of MedDiet
241 on endothelial function (SMD: 0.27, 95% CI: 0.18, 0.36, $P<0.001$). The subgroup analyses
242 showed that the effect was stronger on functional (SMD: 0.44, 95% CI: 0.19, 0.69, $P<0.001$;
243 $I^2=78.2\%$) compared with structural (SMD: 0.16, 95% CI: 0.02, 0.30, $P=0.01$; $I^2=0\%$)
244 measurements of endothelial function. MedDiet increased FMD by 1.66% (95% CI: 1.15, 2.17,
245 $P<0.001$; **Figure 3**). Subgroup analyses showed that MedDiet improved endothelial function
246 significantly in healthy participants (SMD: 0.29, 95% CI: 0.05, 0.53, $P=0.01$; $I^2=26.5\%$) and
247 in those with increased risk of CVD (SMD: 0.36, 95% CI: 0.15, 0.58, $P=0.001$; $I^2=79.1\%$). The
248 effects of the MedDiet on endothelial function were not modified significantly by the type of
249 study design (crossover or parallel) or type of intervention (MedDiet alone or combined; **Table**
250 **2**). Meta-regression analyses demonstrated no modification of the effect size by age, BMI or
251 study duration (**Table 3**). However, a significant association was found between study duration
252 (in weeks) with functional (slope: 0.006; SE: 0.003; $P=0.04$; **Figure 4A**) but not structural
253 measurements (slope: -0.001; SE: 0.001; $P=0.39$, **Figure 4B**) of endothelial function.

254

255 **Study quality and publication bias**

256 Overall, the quality of the trials was modest as the majority of the studies failed to report key
257 information to assess the presence of bias. Attrition bias was present in 50% of the studies (15–
258 17,20,31,33,34) and few studies reported selection bias (<10% of the studies) (20). Seven
259 studies described the randomisation method (15–17,19,28,33,34), and three studies stated the
260 methods of allocation concealment (17,29,33). Five studies reported and described participant
261 dropout (19,27–29,32), while three studies described selective reporting of the results
262 (20,29,34) (**Supplemental Fig. 2**). Visual inspection of the Funnel plot revealed two studies
263 with wider effect estimates but overall there was no evidence of publication bias, which was
264 confirmed by Egger’s Regression test ($P=0.71$; Supplemental Fig. 1).

265

266 **DISCUSSION**

267 Overall, the results of this meta-analysis demonstrate that a MedDiet improves endothelial
268 function. The beneficial effects are evident for both functional and structural measures of
269 endothelial function, although effects were stronger for functional measures. In addition, the
270 effects of a MedDiet were similar in both healthy participants and those at increased risk of
271 CVD and, overall, were not modified by the study design or duration, type of intervention, BMI
272 or age of participants.

273

274 In a previous systematic review and meta-analysis, Schwingshackl and Hoffmann (18) reported
275 improvements in endothelial function with MedDiet interventions. In particular, MedDiet
276 interventions increased FMD by 1.86%, which is similar to the 1.66% average improvement in
277 FMD observed in this analysis. Importantly, the pooled effect size reported by Schwingshackl
278 and Hoffmann (18) was based on the results of only two studies, where as our meta-analysis
279 included seven studies which examined the effects of the MedDiet on FMD, which adds greater

280 confidence to this result. To contextualise these findings, a meta-analysis by Inaba et al (36)
281 demonstrated a 13% decrease in the risk of cardiovascular events per 1% increase in FMD.
282 Thus, a 1.66% improvement in FMD with a MedDiet could potentially translate into a ~22%
283 reduction in cardiovascular events; however, these results require a cautious interpretation and
284 need corroboration in future, more robust studies. As the MedDiet may also reduce CVD risk
285 via a range of other mechanisms, some of which may be independent of effects on endothelial
286 function (e.g. reduced BP, decreased oxidative stress and inflammation, altered gut microbiome
287 (37)), the overall effects of this dietary pattern on CVD risk may be even greater. Indeed, in
288 the large-scale PREDIMED trial, CVD incidence was reduced by 31% and 28% with an
289 average 4.8 year MedDiet intervention supplemented with additional olive oil or nuts,
290 respectively (9). A novel finding of the present analysis was that MedDiet interventions also
291 improved structural measures of endothelial function (e.g. carotid intima-media thickness).
292 However, effects were less pronounced than for functional changes and meta-regression
293 revealed no relationship between study duration and effects of the MedDiet on structural
294 outcomes. By contrast, there was a positive association between study duration and
295 improvement in functional measures of endothelial function. This suggests that longer term
296 consumption of the MedDiet may maximise the effects of this dietary pattern on functional
297 measures of endothelial function, whilst structural changes appear to be relatively modest
298 irrespective of the duration of exposure to this dietary pattern.

299

300 There are several mechanisms through which the MedDiet could improve endothelial function,
301 which may account for the beneficial effects observed in this study. Firstly, the MedDiet
302 augments the bioavailability of nitric oxide (NO) (38), which is essential for healthy endothelial
303 function due to its vasodilatory, anti-atherogenic and anti-proliferative actions (5). The NO
304 'boosting' effects of the MedDiet may be due to antioxidant effects minimising superoxide

305 scavenging of NO (38,39), the provision of the NO precursors inorganic nitrate (green leafy
306 vegetables) and L-arginine (nuts, grains, legumes, and fish) (39), and/ or the upregulation of
307 endothelial NO synthase (oily fish) (40,41). In addition, the MedDiet may improve endothelial
308 function by reducing oxidation of low-density lipoprotein (LDL) (42), which plays a major role
309 in endothelial dysfunction and atherogenesis (43,44). The lower levels of oxidised LDL with
310 the MedDiet are likely mediated by both the antioxidant effects of this dietary pattern and the
311 increased provision of monounsaturated fatty acids which enhance the resilience of LDL to
312 oxidation (45). Finally, a strong link has been reported between inflammation and endothelial
313 dysfunction (46), and several studies have demonstrated beneficial effects of the MedDiet on
314 inflammatory markers including interleukin-6 (IL-6), c-reactive protein (CRP), tumour
315 necrosis factor- α (TNF- α), vascular cell adhesion protein-1 (VCAM-1), and soluble
316 intercellular adhesion molecule-1 (sICAM-1) (47–50). These effects are associated with
317 downregulation of the NF-kB pathway (51) and altered methylation of inflammation-related
318 genes (50), and may further contribute towards improvements in endothelial function with a
319 MedDiet.

320

321

322 **Limitations**

323 The overall quality of the studies included was modest. The majority of investigations did not
324 blind participants to the intervention arm, with only three studies reporting methods of
325 allocation concealment. This is a notable limitation, given the risk of expectation bias whereby
326 the expectation of beneficial effects could result in more favourable outcomes in the
327 intervention group (52). Nevertheless, it is acknowledged that blinding participants is very
328 difficult in dietary intervention studies, particularly those advocating dietary pattern changes,
329 and in many cases this may be unfeasible (53). Some studies had multiple assessments of

330 endothelial function over the duration of the trial and, for studies with longer duration, this may
331 result in a decline of the effects on endothelial function due to a gradual decrease of the
332 adherence to the interventions. However, we included the last measurement in the meta-
333 analysis to standardize the approach across studies and remove any bias related to the selection
334 of the intermediate measurements to be included in the analysis. A further limitation is that
335 most studies were conducted in older subjects. Although meta-regression revealed no influence
336 of age on the effects of the MedDiet, the lack of younger participants may limit our ability to
337 generalise our conclusions, and further research is warranted to determine if the results are
338 applicable to individuals at different life stages. Additionally, since there is no universal
339 definition of what constitutes a MedDiet, details of the dietary interventions differed between
340 studies, and it is possible that certain permutations of the MedDiet may be more effective than
341 others in improving endothelial function as was evident from the high heterogeneity in our
342 analysis. Likewise, the control group utilised was highly variable, such that there was no
343 uniform benchmark against which the MedDiet was compared, which may further contribute
344 towards the high heterogeneity in our analysis. Finally, there are also certain methodological
345 limitations of this review which warrant discussion. Notably, we decided to only include
346 studies where the authors identified their intervention as a MedDiet. This means that we may
347 have missed some studies which administered a Mediterranean-type diet but which was not
348 defined using this specific terminology. In addition, given the small number of studies included
349 in this review, our analysis may have been underpowered to detect differences in intervention
350 effectiveness based on health status, type of intervention, type of measurement, and study
351 design.

352

353

354

355 Conclusions

356 The present systematic review and meta-analysis demonstrates that the MedDiet improves both
357 functional and structural measures of endothelial function, which likely make a large
358 contribution towards the consistently observed beneficial effects of this dietary pattern on
359 cardiovascular health. However, the overall quality of the evidence was modest and more
360 robust and well-designed trials are need to corroborate the evidence highlighting positive
361 effects of the MedDiet on endothelial function.

362

363

364

**365 ACKNOWLEDGMENTS AND STATEMENT OF AUTHORS' CONTRIBUTIONS TO
366 MANUSCRIPT**

367 The systematic review was designed by OS and MS. IM, CK and MS searched, collected and
368 analysed the data. OS, IM, CK, MM, AW, SR, AMM, JCM and MS contributed to data
369 interpretation. OS and MS drafted the manuscript. IM, CK, MM, AWA, SR, AMM and JCM
370 further contributed towards the writing and critical revision of the paper. All authors have read
371 and approved the final manuscript. OS and MS had primary responsibility for the final content.

372 All authors have read and approved the final manuscript.

373

374

375

376

377

378

379

REFERENCES

1. Quyyumi AA. Endothelial function in health and disease: new insights into the genesis of cardiovascular disease. *Am J Med.* 1998;105:S32-S39.
2. Deanfield JE., Halcox JP., Rabelink TJ. Endothelial Function and Dysfunction. *Circulation.* 2007;115:1285–95.
3. Vanhoutte PM, Shimokawa H, Feletou M, Tang EHC. Endothelial dysfunction and vascular disease - a 30th anniversary update. *Acta Physiol.* 2017;219:22–96.
4. Versari D, Daghini E, Viridis A, Ghiadoni L, Taddei S. Endothelial dysfunction as a target for prevention of cardiovascular disease. *Diabetes Care.* 2009;32:S314–21.
5. Siervo M, Scialò F, Shannon OM, Stephan BCM, Ashor AW. Does dietary nitrate say NO to cardiovascular ageing? Current evidence and implications for research. *Proc Nutr Soc.* 2018;77:112–23.
6. Bach-Faig A, Berry EM, Lairon D, Reguant J, Trichopoulou A, Dernini S, Medina FX, Battino M, Belahsen R, Miranda G, et al. Mediterranean diet pyramid today. Science and cultural updates. *Public Health Nutr.* 2011;14:2274–84.
7. Shannon OM, Stephan BCM, Granic A, Lentjes M, Hayat S, Mulligan A, Brayne C, Khaw K-T, Bundy R, Aldred S, et al. Mediterranean diet adherence and cognitive function in older UK adults: the European Prospective Investigation into Cancer and Nutrition-Norfolk (EPIC-Norfolk) Study. *Am J Clin Nutr.* 2019; 110:938-948.
8. de Lorgeril M, Salen P, Martin JL, Monjaud I, Delaye J, Mamelle N. Mediterranean diet, traditional risk factors, and the rate of cardiovascular complications after myocardial infarction: Final report of the Lyon Diet Heart Study. *Circulation.* 1999;99:779–85.
9. Estruch R, Ros E, Salas-Salvadó J, Covas M-I, Corella D, Arós F, Gómez-Gracia E, Ruiz-Gutiérrez V, Fiol M, Lapetra J, et al. Primary prevention of cardiovascular disease with a

Mediterranean diet supplemented with extra-virgin olive oil or nuts. *N Engl J Med.* 2018; 378:e34.

10. Estruch R. Anti-inflammatory effects of the Mediterranean diet: The experience of the PREDIMED study. *Proc Nutr Soc.* 2010;69:333–40.

11. Schwingshackl L, Christoph M, Hoffmann G. Effects of olive oil on markers of inflammation and endothelial function - A systematic review and meta-analysis. *Nutrients.* 2015;7:7651–75.

12. Xiao Y, Huang W, Peng C, Zhang J, Wong C, Kim JH, Yeoh E-K, Su X. Effect of nut consumption on vascular endothelial function: A systematic review and meta-analysis of randomized controlled trials. *Clin Nutr.* 2018;37:831–9.

13. Kondo K, Morino K, Nishio Y, Kondo M, Nakao K, Nakagawa F, Ishikado A, Sekine O, Yoshizaki T, Kashiwagi A, et al. A fish-based diet intervention improves endothelial function in postmenopausal women with type 2 diabetes mellitus: A randomized crossover trial. *Metab Clin Exp.* 2014;63:930–40.

14. Xin W, Wei W, Li X. Effect of fish oil supplementation on fasting vascular endothelial function in humans: A meta-analysis of randomized controlled trials. *PLoS ONE.* 2012;7:e46028.

15. Davis CR, Hodgson JM, Woodman R, Bryan J, Wilson C, Murphy KJ. A Mediterranean diet lowers blood pressure and improves endothelial function: Results from the MedLey randomized intervention trial. *Am J Clin Nutr.* 2017;105:1305–13.

16. Torres-Peña JD, Garcia-Rios A, Delgado-Casado N, Gomez-Luna P, Alcala-Diaz JF, Yubero-Serrano EM, Gomez-Delgado F, Leon-Acuña A, Lopez-Moreno J, Camargo A, et al. Mediterranean diet improves endothelial function in patients with diabetes and prediabetes: A report from the CORDIOPREV study. *Atherosclerosis.* 2018;269:50–6.

17. Maiorino MI, Bellastella G, Petrizzo M, Gicchino M, Caputo M, Giugliano D, Esposito K. Effect of a Mediterranean diet on endothelial progenitor cells and carotid intima-media thickness in type 2 diabetes: Follow-up of a randomized trial. *Eur J Prev Cardiol.* 2017;24:399–408.
18. 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;24:929–39.
19. Ceriello A, Esposito K, La Sala L, Pujadas G, De Nigris V, Testa R, Bucciarelli L, Rondinelli M, Genovese S. The protective effect of the Mediterranean diet on endothelial resistance to GLP-1 in type 2 diabetes: A preliminary report. *Cardiovasc Diabetol.* 2014;13:140.
20. Jaacks LM, Sher S, Staercke CD, Porkert M, Alexander WR, Jones DP, Vaccarino V, Ziegler TR, Quyyumi AA. Pilot randomized controlled trial of a Mediterranean diet or diet supplemented with fish oil, walnuts, and grape juice in overweight or obese US adults. *BMC Nutr.* 2018;4:26.
21. Liberati A, Altman DG, Tetzlaff J, Mulrow C, Gøtzsche PC, Ioannidis JPA, Clarke M, Devereaux PJ, Kleijnen J, Moher D. The PRISMA statement for reporting systematic reviews and meta-analyses of studies that evaluate healthcare interventions: Explanation and elaboration. *BMJ.* 2009;339:b2700.
22. Higgins JPT, Green, S.E. *Cochrane Handbook for Systematic Reviews of Interventions Version 5.1.0.* London, UK: The Cochrane Collaboration; 2011.
23. Borenstein M, Hedges L., Higgins JP., Rothstein H. *Introduction to Meta-Analysis.* New Jersey, US: Wiley; 2019.
24. DerSimonian R, Laird N. Meta-analysis in clinical trials. *Control Clin Trials.* 1986;7:177–88.

25. Korkmaz H, Onalan O. Evaluation of endothelial dysfunction: Flow-mediated dilation. *Endothelium*. 2008;15:157–63.
26. Higgins JPT, Thompson SG, Deeks JJ, Altman DG. Measuring inconsistency in meta-analyses. *BMJ*. 2003;327:557–60.
27. Ambring A, Friberg P, Axelsen M, Laffrenzen M, Taskinen M-R, Basu S, Johansson M. Effects of a Mediterranean-inspired diet on blood lipids, vascular function and oxidative stress in healthy subjects. *Clin Sci*. 2004;106:519–25.
28. Klonizakis M, Alkhatib A, Middleton G, Smith MF. Mediterranean diet- and exercise-induced improvement in age-dependent vascular activity. *Clin Sci*. 2013;124:579–87.
29. Marin C, Ramirez R, Delgado-Lista J, Yubero-Serrano EM, Perez-Martinez P, Carracedo J, Garcia-Rios A, Rodriguez F, Gutierrez-Mariscal FM, Gomez P, et al. Mediterranean diet reduces endothelial damage and improves the regenerative capacity of endothelium. *Am J Clin Nutr*. 2011;93:267–74.
30. Murie-Fernandez M, Irimia P, Toledo E, Martínez-Vila E, Buil-Cosiales P, Serrano-Martínez M, Ruiz-Gutiérrez V, Ros E, Estruch R, Martínez-González MÁ, et al. Carotid intima-media thickness changes with Mediterranean diet: A randomized trial (PREDIMED-Navarra). *Atherosclerosis*. 2011;219:158–62.
31. Sala-Vila A, Romero-Mamani E-S, Gilabert R, Núñez I, de la Torre R, Corella D, Ruiz-Gutiérrez V, López-Sabater M-C, Pintó X, Rekondo J, et al. Changes in ultrasound-assessed carotid intima-media thickness and plaque with a Mediterranean diet: A sub-study of the PREDIMED trial. *Arterioscler Thromb Vasc Biol*. 2014;34:439–45.
32. Buscemi S, Verga S, Tranchina MR, Cottone S, Cerasola G. Effects of hypocaloric very-low-carbohydrate diet vs. Mediterranean diet on endothelial function in obese women. *Eur J Clin Invest*. 2009;39:339–47.

33. Esposito K, Marfella R, Ciotola M, Di Palo C, Giugliano F, Giugliano G, D'Armiento M, D'Andrea F, Giugliano D. Effect of a Mediterranean-style diet on endothelial dysfunction and markers of vascular inflammation in the metabolic syndrome: a randomized trial. *JAMA*. 2004;292:1440–6.
34. Thomazella MCD, Góes MFS, Andrade CR, Debbas V, Barbeiro DF, Correia RL, Marie SKN, Cardounel AJ, daLuz PL, Laurindo FRM. Effects of high adherence to Mediterranean or low-fat diets in medicated secondary prevention patients. *Am J Cardiol*. 2011;108:1523–9.
35. Fuentes F, López-Miranda J, Sánchez E, Sánchez F, Paez J, Paz-Rojas E, Marín C, Gómez P, Jimenez-Perepérez J, Ordovás JM, et al. Mediterranean and low-fat diets improve endothelial function in hypercholesterolemic men. *Ann Intern Med*. 2001;134:1115–9.
36. Inaba Y, Chen JA, Bergmann SR. Prediction of future cardiovascular outcomes by flow-mediated vasodilatation of brachial artery: a meta-analysis. *Int J Cardiovasc Imaging*. 2010;26:631–40.
37. Tosti V, Bertozzi B, Fontana L. Health benefits of the Mediterranean diet: Metabolic and molecular mechanisms. *J Gerontol A Biol Sci Med Sci*. 2018;73:318–26.
38. Medina-Remón A, Tresserra-Rimbau A, Pons A, Tur JA, Martorell M, Ros E, Buil-Cosiales P, Sacanella E, Covas MI, Corella D, et al. Effects of total dietary polyphenols on plasma nitric oxide and blood pressure in a high cardiovascular risk cohort. The PREDIMED randomized trial. *Nutr Metab Cardiovasc Dis*. 2015;25:60–7.
39. Shannon OM, Stephan BCM, Minihane A-M, Mathers JC, Siervo M. Nitric oxide boosting effects of the Mediterranean diet: A potential mechanism of action. *J Gerontol A Biol Sci Med Sci*. 2018; 73:902-904.
40. Armah CK, Jackson KG, Doman I, James L, Cheghani F, Minihane AM. Fish oil fatty acids improve postprandial vascular reactivity in healthy men. *Clin Sci*. 2008;114:679–86.

41. Balakumar P, Taneja G. Fish oil and vascular endothelial protection: Bench to bedside. *Free Radic Biol Med.* 2012;53:271–9.
42. Fitó M, Guxens M, Corella D, Sáez G, Estruch R, Torre R de la, Francés F, Cabezas C, López-Sabater M del C, Marrugat J, et al. Effect of a traditional Mediterranean diet on lipoprotein oxidation: A randomized controlled trial. *Arch Intern Med.* 2007;167:1195–203.
43. Ross R. Atherosclerosis - an inflammatory disease. *N Engl J Med.* 1999;340:115–26.
44. Valente AJ, Irimpen AM, Siebenlist U, Chandrasekar B. OxLDL induces endothelial dysfunction and death via TRAF3IP2: Inhibition by HDL3 and AMPK activators. *Free Radic Biol Med.* 2014;70:117–28.
45. Bonanome A, Pagnan A, Biffanti S, Opportuno A, Sorgato F, Dorella M, Maiorino M, Ursini F. Effect of dietary monounsaturated and polyunsaturated fatty acids on the susceptibility of plasma low density lipoproteins to oxidative modification. *Arterioscler Thromb.* 1992;12:529–33.
46. Zhang C. The role of inflammatory cytokines in endothelial dysfunction. *Basic Res Cardiol.* 2008;103:398–406.
47. Estruch R, Martínez-González MA, Corella D, Salas-Salvadó J, Ruiz-Gutiérrez V, Covas MI, Fiol M, Gómez-Gracia E, López-Sabater MC, Vinyoles E, et al. Effects of a Mediterranean-style diet on cardiovascular risk factors: a randomized trial. *Ann Intern Med.* 2006;145:1–11.
48. Mena M-P, Sacanella E, Vazquez-Agell M, Morales M, Fitó M, Escoda R, Serrano-Martínez M, Salas-Salvadó J, Benages N, Casas R, et al. Inhibition of circulating immune cell activation: A molecular antiinflammatory effect of the Mediterranean diet. *Am J Clin Nutr.* 2009;89:248–56.
49. Casas R, Sacanella E, Urpí-Sardà M, Chiva-Blanch G, Ros E, Martínez-González M-A, Covas M-I, Rosa Ma Lamuela-Raventos, Salas-Salvadó J, Fiol M, et al. The effects of the

Mediterranean diet on biomarkers of vascular wall inflammation and plaque vulnerability in subjects with high risk for cardiovascular disease. A randomized trial. PLoS ONE. 2014;9:e100084.

50. Arpón A, Riezu-Boj JI, Milagro FI, Martí A, Razquin C, Martínez-González MA, Corella D, Estruch R, Casas R, Fitó M, et al. Adherence to Mediterranean diet is associated with methylation changes in inflammation-related genes in peripheral blood cells. J Physiol Biochem. 2016;73:445–55.

51. Perez-Martinez P, Lopez-Miranda J, Blanco-Colio L, Bellido C, Jimenez Y, Moreno JA, Delgado-Lista J, Egido J, Perez-Jimenez F. The chronic intake of a Mediterranean diet enriched in virgin olive oil, decreases nuclear transcription factor kappaB activation in peripheral blood mononuclear cells from healthy men. Atherosclerosis. 2007;194:e141-146.

52. Staudacher HM, Irving PM, Lomer MCE, Whelan K. The challenges of control groups, placebos and blinding in clinical trials of dietary interventions. Proc Nutr Soc. 2017;76:203–12.

53. Schulze MB, Martínez-González MA, Fung TT, Lichtenstein AH, Forouhi NG. Food based dietary patterns and chronic disease prevention. BMJ. 2018;361:k2396.

Table 1: Summary of main characteristics of randomized clinical trials investigating the effects of Mediterranean Diet on endothelial function in adults

Author	Country	Study Design	Health Status	Outcome	Sample Size	Male (<i>n</i>)	Age (years)	BMI (kg/m ²)	SBP/DBP (mmHg)	Duration (Weeks)	Type of Intervention	Type of Control
Ambring et al. (27)	Sweden	Cross-over	Healthy	FBF	22	12	43	26.0	NR	12	MedDiet	Swedish Diet
Buscemi et al. (32)	Italy	Parallel	Obese	FMD	20	0	38	34.2	128/88	8	MedDiet	Atkins Low-carbohydrate
Ceriello et al. (19)	Spain	Parallel	DM2	FMD	24	17	NR	29.5	116/78	12	MedDiet + MUFA	Low Fat Diet
Davis et al. (15)	Australia	Parallel	Healthy	FMD	166	72	71	26.9	124/71	24	MedDiet	Habitual Diet
Esposito et al. (33)	Italy	Parallel	MetS	EFS	180	89	44	28.0	135/86	96	MedDiet	Prudent Diet
Fuentes et al. (35)	Spain	Cross-over	Hypercholesteremic	FMD, BVS	22	22	40	NR	NR	8	MedDiet + MUFA	NCEP-1 Diet
Jaacks et al. (20)	USA	Parallel	Overweight	FMD	30	8	51	31.5	NR	8	MedDiet	Habitual Diet
Klonizakis et al. (28)	UK	Parallel	Healthy	CM	22	7	55	30.5	127/79	8	MedDiet + Exercise	Non-MedDiet + Exercise
Maiorino et al. (17)	Italy	Parallel	DM2	CIMT	215	106	52	29.6	140/87	121	MedDiet	Low Fat Diet
Marin et al. (29)	Spain	Cross-over	Healthy	CM	20	10	67	31.9	NR	4	MedDiet	Saturated Fatty Acid Diet
Murie-Fernandez et al. (30)	Spain	Parallel	CVD risk	CIMT	187	91	67	29.4	NR	48	G1: MedDiet + EVOO G2: MedDiet + Nuts	Low Fat Diet
Sala-Vila et al. (31)	Spain	Parallel	CVD risk	ICA-IMT	175	42	66	29.6	150/81	115	G1: MedDiet + EVOO G2: MedDiet + Nuts	Low Fat Diet
Thomazella et al. (34)	Brazil	Parallel	ACS	FMD, BVS	42	42	55	26.4	136/84	12	MedDiet	Low Fat Diet
Torres-Peña et al. (16)	Spain	Parallel	DM2	FMD	438	NR	61	31.8	NR	72	MedDiet + EVOO	Low Fat Diet
Torres-Peña et al. (16)	Spain	Parallel	pDM2	FMD	289	NR	58	30.3	NR	72	MedDiet + EVOO	Low Fat Diet
Torres-Peña et al. (16)	Spain	Parallel	Healthy	FMD	78	NR	56	29.5	NR	72	MedDiet + EVOO	Low Fat Diet

ACS, Acute Coronary Syndromes; BVS, Baseline Vessel Size; CIMT, Carotid Intima-Media Thickness; CM, Cutaneous Microvascular Function; CVD risk, Risk of Cardiovascular Disease; DBP, Diastolic Blood Pressure; DM2, Type 2 Diabetes; EFS, Endothelial Function Score; EVOO, Extra Virgin Olive Oil; FBF, Forearm Blood Flow; FMD, Flow Mediated Dilatation; ICA-IMT, Internal Carotid Intima-Media Thickness; MedDiet, Mediterranean Diet; MetS, Metabolic Syndrome; MUFA, Monounsaturated Fatty Acids; *n* = number of subjects; NCEP-1, The National Cholesterol Education Program Diet; pDM2, prediabetes; SBP, Systolic Blood Pressure.

Table 2: Sensitivity analysis to evaluate the influence of health status, type of intervention, type of measurement, and study design on the effects of the Mediterranean dietary pattern on endothelial function (EF) in adults.

Category	No of EF measurements per subgroup	Effect size	95% CI	<i>P</i>	<i>P</i> between Groups	<i>I</i> ²
Health status					0.66	
• Healthy	7	0.29	0.05 - 0.53	0.01		26.5%
• Increased CVD Risk	13	0.36	0.15 - 0.58	0.001		79.1%
Type of Intervention					0.71	
• MedDiet	9	0.37	0.03 - 0.77	0.04		85.7%
• MedDiet + other	11	0.29	0.19 - 0.39	<0.001		0%
Type of Measurement					0.05	
• Functional	13	0.44	0.19 - 0.69	<0.001		78.2%
• Structural	7	0.16	0.02 - 0.30	0.01		0%
Study Design					0.55	
• Cross-over	4	0.25	-0.05 - 0.56	0.10		41.7%
• Parallel	16	0.36	0.16 - 0.56	<0.001		74.9%

CVD risk = Risk of Cardiovascular Disease. MedDiet = Mediterranean dietary pattern. *P* refers to the effect sizes of the subgroups in each category. *P* between Groups refers to the comparison of the effect sizes between sub-groups within each category.

Table 3: Meta-regression analysis to evaluate potential modifiers of the effects of Mediterranean dietary pattern on endothelial function in adults.

	Slope	SE	Q (df=1)	P
Age (years)	-0.002	0.008	0.12	0.72
Study Duration (weeks)	0.0008	0.002	0.17	0.67
Body mass index (kg/m ²)	-0.01	0.04	0.07	0.78

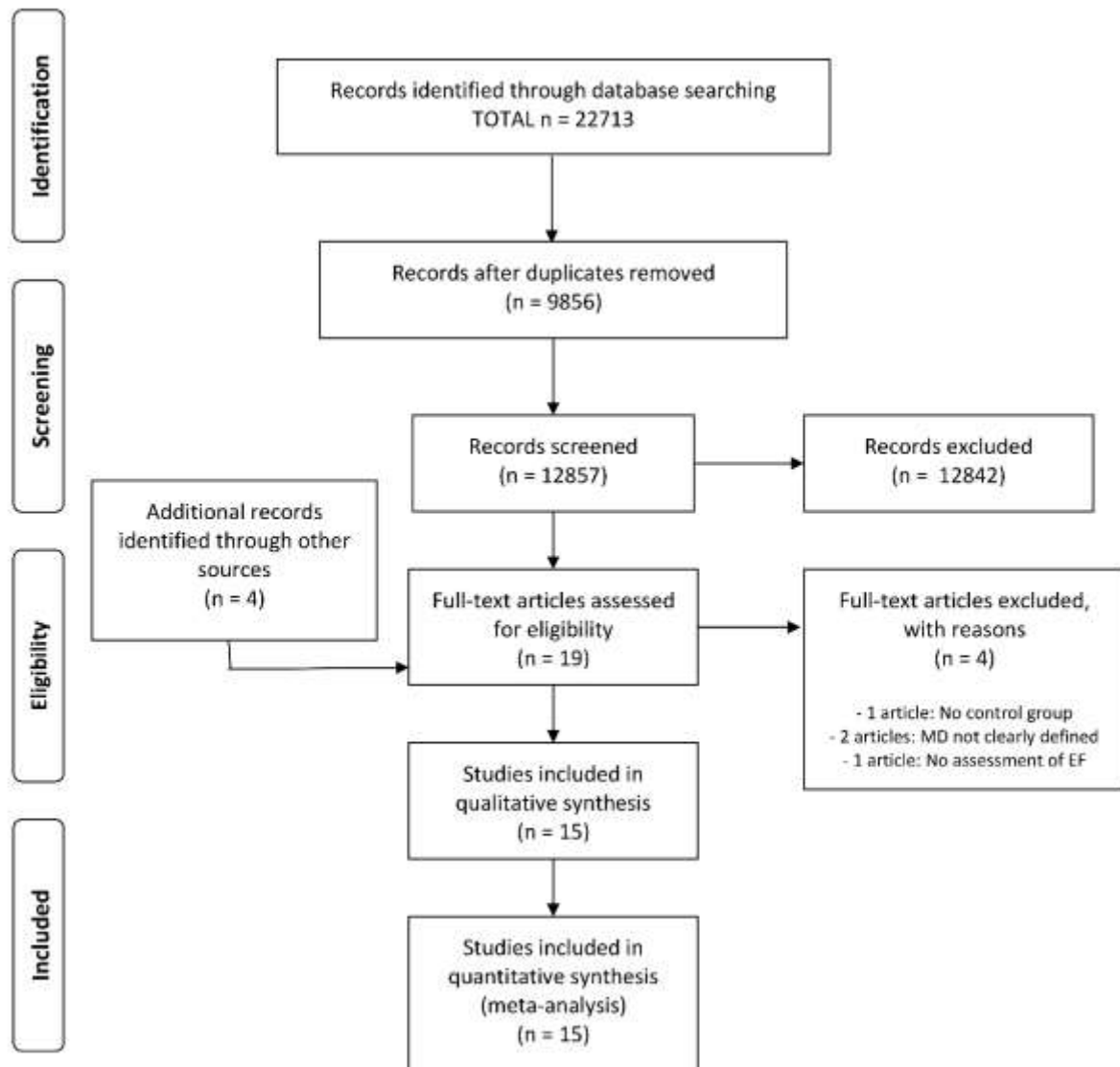


Figure 1: Flow diagram of the selection process of the randomized controlled trials included in the meta-analysis.

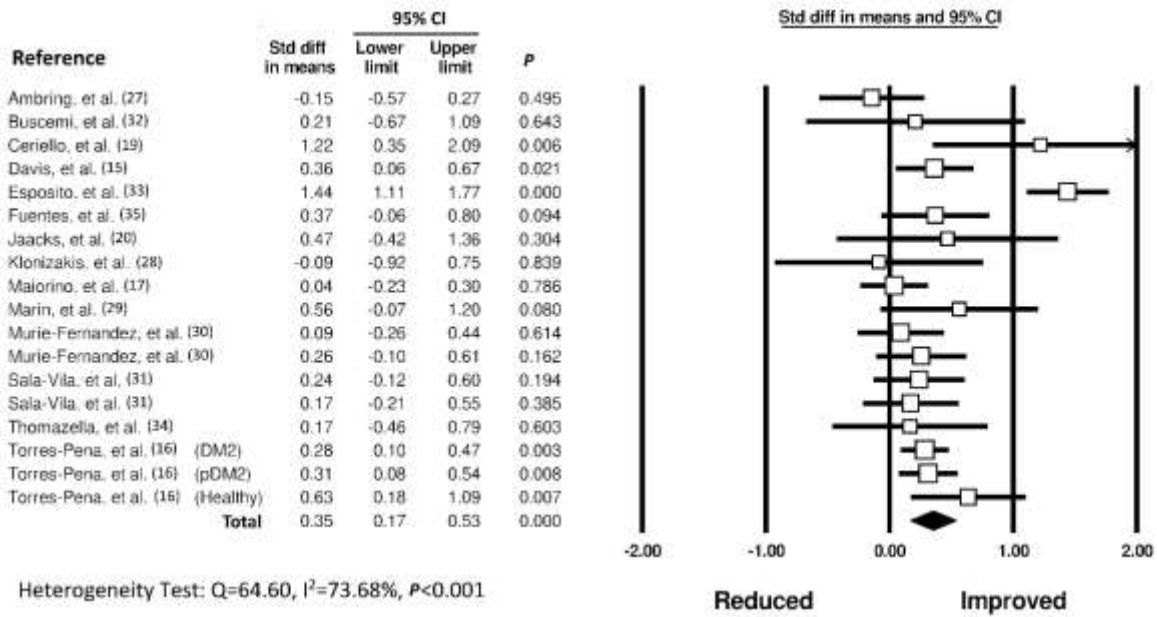


Figure 2: Forest plot showing the overall effect of the Mediterranean dietary pattern on endothelial function in adults. DM2 = type 2 diabetes; pDM2 = pre-diabetes; a = MedDiet + extra virgin olive oil; b = MedDiet + nuts. Data showed as standardised differences in means. Horizontal lines denote 95% confidence intervals (CI). The size of the boxes is proportionally scaled to the effect size for each study.

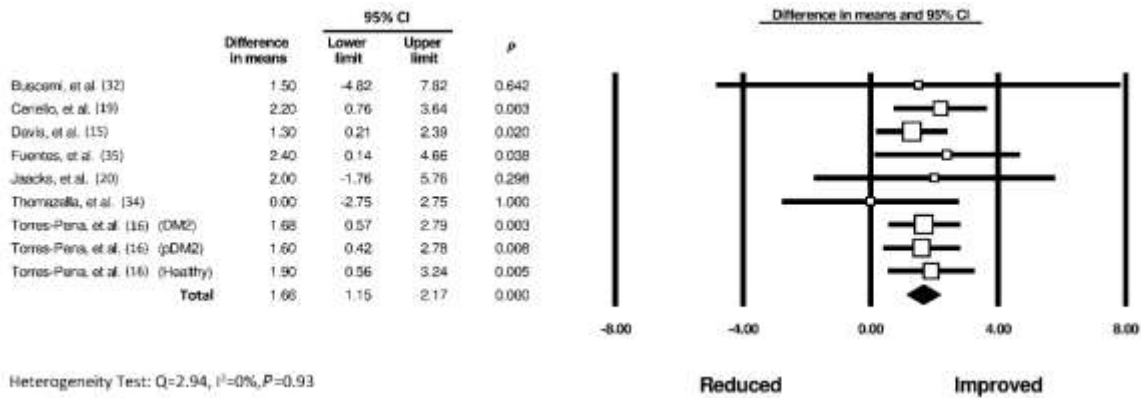


Figure 3: Forest plot showing the overall effect of the Mediterranean dietary pattern on flow mediated dilation in adults (expressed as percent change). DM2 = type 2 diabetes; pDM2 = pre-diabetes. Data showed as percent differences in means. Horizontal lines denote 95% confidence intervals (CI). The size of the boxes is proportionally scaled to the effect size for each study.

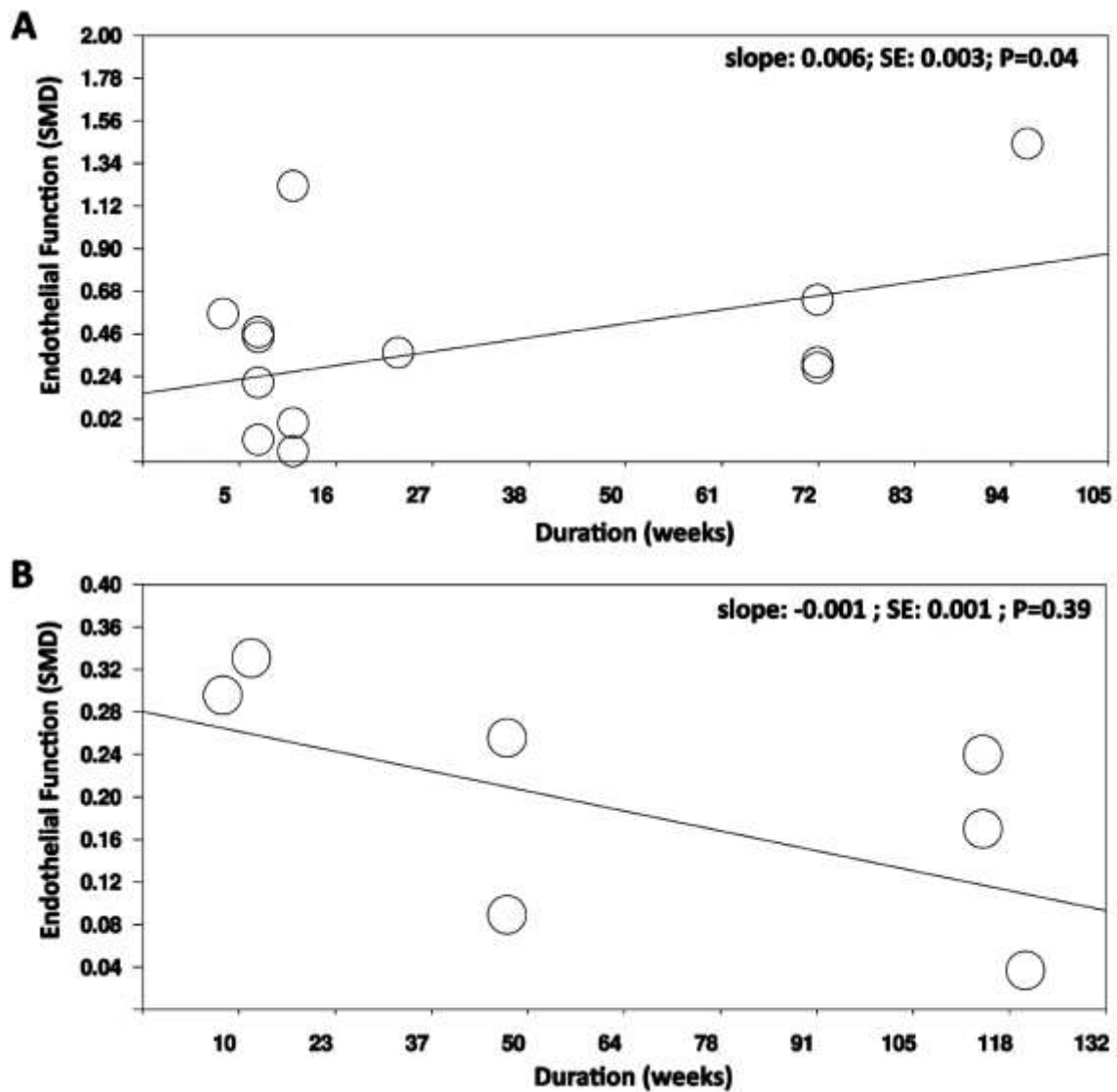


Figure 4: Meta-regression analysis of the association between study duration and the effect size (expressed as standardised mean difference [SMD]) of functional (A) and structural (B) alterations in endothelial function in adult subjects following consumption of a Mediterranean dietary pattern.