In comparison with palm oil, dietary nut supplementation delays the progression of atherosclerotic lesions in female apoE-deficient mice

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Abstract
Epidemiological studies have demonstrated the benefits of nut consumption on cardiovascular risk factors and CHD, attributed to their fatty acid profile, rich in unsaturated fatty acids, and also to other nutrients. The effect of nuts on atherosclerotic lesions was studied in female and male apoE-knockout mice fed a diet supplemented with 3% (w/w) mixed nuts (mix: almonds, hazelnuts and walnuts in a proportion of 0.25:0.25:0.50, respectively), and compared with mice receiving an isoenergetic diet of similar fat content provided as palm oil. After 12 weeks, plasma lipid parameters and aortic lesions were measured. Males receiving nuts had lower plasma cholesterol than the palm oil group, and both sex groups had lower plasma non-HDL-cholesterol and lower content of reactive oxygen species in LDL than mice receiving the palm oil diet, the latter decrease being more pronounced in females than in males. Females consuming the nut diet showed a smaller aortic lesion area than those consuming palm oil, whereas no differences were observed in males. In females, hepatic paraoxonase 2 (Pon2) mRNA increased, and no change was observed in prenylcysteine oxidase 1 (Pcyox1) expression after the consumption of the nut-containing diet. In addition, aortic atherosclerotic lesions correlated directly with total plasma cholesterol and inversely with hepatic Pon2 expression. The results suggest that the beneficial effect of nut intake in female apoE-deficient mice may be attributed to reduced non-HDL-cholesterol levels and enhanced PON2 antioxidant activity.

Key words: Nuts; ApoE-deficient mice; Atherosclerosis; Lipoproteins

CVD, as a result of an atherosclerotic process, are the leading cause of morbidity and mortality in developed countries, in which the diet is one of the most important environmental factors associated with the atherothrombotic process. However, epidemiological studies have shown a lower incidence of CHD in populations of Mediterranean countries (1,2), a circumstance that could be attributed to the beneficial health effects of the so-called Mediterranean diet (3), which is characterised by a high intake of vegetables, legumes, fruits and nuts, together with a widespread use of olive oil (4). Nowadays, it is recognised that many of the nutrients in these foods have some protective effects against the atherogenic process. In the Mediterranean region, three nuts (walnuts, almonds and hazelnuts) are used to constitute part of the dietary source of energy, but have not been included in many dietary recommendations due to their high fat content (5–7). Nevertheless, recent clinical evidence suggests that consumption of nuts promotes a healthy lipid profile associated with a lower risk of CVD (8–16). Several studies have attributed this quality to their nutrients, mainly their MUFA and PUFA. Apart from that, nuts are complex foods that are sources of other nutrients: vegetable protein, fibre, vitamins, minerals and many bioactive constituents such as antioxidants, phytosterols and other phytochemicals (6,17). To provide an experimental understanding of the results of epidemiological studies in human subjects and to tackle the possible beneficial mechanisms of

Abbreviations: ITGA4, integrin α4; ITGAM, integrin αM; Pcyox1, prenylcysteine oxidase 1; Pon2, Paraoxonase 2.
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nut intake in atherosclerotic lesions, apoE-deficient mice were used in the present experimental study. The animals in this model develop spontaneous atherosclerosis, even with low-fat and low-cholesterol diets, and the disease features are similar to those observed in humans and other species\(^{18}\). Therefore, the aim of the present study was to compare the effect of supplementation with a mixture of three nuts on the development of atherosclerotic lesions with that of an isoenergetic diet containing palm oil in apoE-deficient mice.

Materials and methods

Animals

Female and male apoE-deficient mice (\(n = 23\) each group), aged 2.7 (SEM 0.03) months from a C57BL/6J × Ola129 genetic background, were fasted for 6 h and anaesthetised with isoflurane (Forane). Their blood was drawn from the retro-orbital plexus. Then, plasma cholesterol was determined. For each sex, two groups with similar plasma cholesterol concentrations were established. The animals had free access to food and water during the 12-week experimental period, and they were housed in a temperature-controlled facility and handled observing the criteria from the European Union for care and use of laboratory animals in research. The protocol was approved by the Ethics Committee for Animal Research of the University of Zaragoza.

Diets

In the present study, we established two study groups, one of which received a chow diet (Teklad Mouse/Rat Diet no. 2014, Harlan Teklad; Harlan Ibérica) supplemented with 3% (w/w) edible whole nuts (nut group) in a mix composed of 50% walnuts, 25% almonds and 25% hazelnuts (nut 11 for both female and male mice). This represents a daily dose of 3 g nuts/kg body weight per mouse which, considering the higher metabolic rate of mice\(^{19}\), would translate into the dose of 30 g/d or 0.4 g nuts/kg used in human interventions and the same proportion of nuts, so as to reproduce in mice the effects observed in human subjects\(^{20–22}\). The other group received the same chow diet but was supplemented with 2% (w/w) palm oil. Both diets were isoenergetic and provided the same amount of fat and MUFA (Table 1). Diets were prepared by milling the chow diet, adding the corresponding supplement and 3.0% (w/w) of wheat starch as agglutinant to facilitate pellet formation, and then drying at 60°C for 48 h. All diets were prepared weekly and stored in an N\(_2\) atmosphere at −20°C. Fresh food was provided daily. Preparation of extractable and non-extractable phenolic compounds from the diets was carried out according to Arranz et al.\(^{23}\) and quantified using the method of Singleton et al.\(^{24}\). The contents of extractable phenolic compounds and hydrolysable polyphenols were higher (21 and 26%, respectively) in the nut diet than in the palm oil diet (Table 1). The animals were fed the experimental diets for 12 weeks; both were well tolerated.

Experimental procedures

At 1 week before killing and after fasting, blood samples were obtained for integrin αM (ITGAM) and integrin α4 (ITGA4) expressions and paraoxonase activity. At killing, the animals were euthanised by suffocation with CO\(_2\) and exsanguinated by cardiac puncture. The liver was split; one piece was stored in neutral buffered formaldehyde for histological analysis and the remainder immediately frozen in liquid N\(_2\). The heart and the arterial tree were perfused with 5 ml of cold PBS (pH 7.4) under physiological pressure. After that, the heart was removed, embedded in OCT\({\text{T}}M\) (Tissue-Tek), frozen in dry ice-cold isopentane (Panreac) and stored at −80°C until analysis.

Plasma analysis

Total plasma cholesterol and TAG concentrations were measured in a microtitre assay, using Infinity\textsuperscript{TM} commercial kits (Thermo Scientific). Plasma HDL-cholesterol was quantified in the supernatant by a fluorometric enzyme assay (Amplex Red; Molecular Probes) after precipitation of apoB particles with phosphotungstic acid–MnCl\(_2\) (Roche). Paraoxonase was assayed as arylesterase activity determined by the rate of phenylacetate hydrolysis, as described previously\(^{25}\). ApoA1 and apoA4 were quantified by ELISA using specific polyclonal antibodies (Biodesign and Santa Cruz Biotechnology), as described previously\(^{26}\). Plasma lipoprotein profile was determined in 100 μl of pooled plasma samples from each group by fast protein liquid chromatography gel filtration\(^{27}\) using a Superose 6B column (Amersham Pharmacia), and the cholesterol in each fraction was measured with the fluorescent method described above.

Table 1. Characteristics of the experimental diets

<table>
<thead>
<tr>
<th></th>
<th>Nut</th>
<th>Palm oil</th>
</tr>
</thead>
<tbody>
<tr>
<td>Metabolizable energy (MJ/kg)</td>
<td>13.2</td>
<td>13.4</td>
</tr>
<tr>
<td>Total fat (% feed)</td>
<td>5.7</td>
<td>5.8</td>
</tr>
<tr>
<td>Fatty acid content (% fat)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>SFA</td>
<td>15.0</td>
<td>27.7</td>
</tr>
<tr>
<td>MUFA</td>
<td>26.9</td>
<td>27.5</td>
</tr>
<tr>
<td>PUFA</td>
<td>58.1</td>
<td>44.9</td>
</tr>
<tr>
<td>P:S ratio</td>
<td>3.9</td>
<td>1.6</td>
</tr>
<tr>
<td>Phenolic compounds (μmol caffeic acid equivalents/g)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Extractable</td>
<td>17</td>
<td>14</td>
</tr>
<tr>
<td>Non-extractable proanthocyanidins</td>
<td>1300</td>
<td>1300</td>
</tr>
<tr>
<td>Hydrolysable polyphenols</td>
<td>335</td>
<td>266</td>
</tr>
</tbody>
</table>

P:S, polyunsaturated:saturated ratio.
Blood cell analysis

Expression of ITGAM and ITGA4 proteins in blood cells from samples taken 1 week before killing was analysed using a fluorescence-activated cell sorter (Becton-Dickinson), as described previously(28). The results are expressed as the proportion (%) of the marker-positive cells recovered in the region corresponding to monocytes.

LDL oxidation susceptibility

LDL oxidisability was assessed using modified procedures of Navab et al.(29) to determine the presence of reactive oxygen species by measuring the conversion of 2',7'-dichlorofluorescein diacetate into fluorescent dichlorofluorescein. Briefly, LDL fractions (5 μg cholesterol) separated by fast protein liquid chromatography were incubated, at 37°C with 2 μg dichlorofluorescein, in 25 μl of 0·1% sodium azide and 100 μl PBS, up to a total volume of 150 μl. Fluorescence was measured after 3 h of incubation at an excitation wavelength of 485 nm and an emission wavelength of 535 nm(30).

Evaluation of atherosclerotic lesions

Sections of the proximal aorta from the frozen heart were sliced on a cryostat (Microm HM505E; Thermo Scientific). From each heart, four cryosections (6 μm) were fixed with neutral buffered formaldehyde, stained for lipids with Sudan IV and counterstained with haematoxylin and eosin (Harris solution; Sigma Chemical Company). Images were captured by a digital camera fitted to a Nikon microscope. Aortic lesion areas were blindly quantified using Scion Image software (Scion Corporation).

Hepatic histological analysis

Liver samples stored in formaldehyde were embedded in paraffin. Then, sections (4 μm) were stained with haematoxylin and eosin and images were captured. Hepatic fat content was evaluated by quantifying the lipid droplets in each liver section with Adobe Photoshop CS2 (Adobe Systems Incorporated) and expressed as a percentage of the entire liver section(31).

RNA isolation

RNA was isolated using TRI Reagent Solution (Ambion; Applied Biosystems). Contaminant DNA was removed by TURBO DNA-free™ DNase treatment and removal reagents from Ambion. RNA was quantified by absorbance at A_{260/280} (the A_{260/280} ratio was greater than 1·75). The integrity of 28S and 18S ribosomal RNA was verified by agarose gel electrophoresis followed by ethidium bromide staining, and the 28S:18S ratio was greater than 2.

Quantification of mRNA

Equal amounts of DNA-free RNA from each sample of all animals were used in quantitative RT-PCR analyses. First-strand complementary DNA synthesis and PCR were performed using the First-Strand complementary DNA Synthesis Kit (Fermentas) and Power SYBR® Green PCR Master Mix (Applied Biosystems) according to the manufacturers’ instructions and as described previously(32). Primers were designed by Primer Express® (Applied Biosystems) and checked by BLAST analysis (NCBI) to verify specificity and selective amplification of the target gene, as well as to ensure amplification of complementary DNA and not of genomic DNA. The sequences were as follows: for Pla2g7 – sense, 5'-GGC TTT GTA CTG AGC TCA AGA-3', antisense, 5'-TGC AGT TGT CAG AGA ACCA-3'; for prenylcysteine oxidase 1 (Pcxyox1) – sense, 5'-GGT TCA GTC ATT CAC CCC CTA A-3', antisense, 5'-TAC AAA CCA GCT GCT CTC CTC A-3'; for paraoxonase 2 (Pon2) – sense, 5'-GCA CGC TGG TGG ACA ATT TATC-3', antisense, 5'-TGT CAC TGAT GCC TTC TCG GAT-3'. Real-time PCR were performed in an ABI PRISM 7700 Sequence Detector (Applied Biosystems) following the standard procedure. The specificity of the PCR was confirmed by observing a single dissociation curve. The relative amount of all mRNA was calculated using the comparative 2^{-DeltaCt} method. Ppib mRNA was used as the reference gene(33).

Statistical analyses

Data were analysed by the Kolmogorov–Smirnov test to check for the normal distribution of variables and by Bartlett’s test to assess the homogeneity of variance. Student’s t test was used when both tests satisfied the corresponding hypothesis, and when either of these tests did not, differences between pairs were tested using the Mann–Whitney U test. Unless otherwise stated, results are presented as means with their standard errors. Correlations between variables were tested by calculating Spearman’s correlation coefficient using SPSS software, version 15.0 (SPSS, Inc.). Differences were considered significant when \( P<0·05 \).

Results

Body and liver weight

Somatic variables in the animals are shown in Table 2. At killing, males and females had significant differences in body weight (\( P<0·001 \)) and body-weight changes (\( P=0·01 \)). However, we found no significant differences between the two diets in terms of body-weight change, liver weight, feed intake or energy intake.

Plasma lipid parameters

The effect of the different diets on plasma lipid parameters after the 12-week experimental period is shown in Table 3. The same-sex groups had similar initial cholesterol levels, which were significantly higher in males than in females, and this difference remained after the experimental period. In females, no significant differences were observed between the two diets in terms of total cholesterol. However, male mice consuming the nut diet had significantly lower plasma cholesterol than those in the palm oil group. There were no
significant differences in plasma HDL-cholesterol in the two dietary groups. VLDL + LDL-cholesterol were significantly lower in mice of both sexes receiving the nut diet. Plasma TAG levels were higher ($P < 0.001$) in males than in females, but neither sex showed significant diet-related differences. ApoA1, the principal protein in HDL particles, and in agreement with the results for HDL-cholesterol, was not modified by the dietary intervention. ApoA4 was significantly decreased in males on the nut diet. A significant increase was observed in arylesterase activity in female mice receiving the nut diet.

**Monocyte activation**

ITGAM and ITGA4 protein expressions of circulating monocytes are also shown in Table 3. The percentages of monocytes expressing ITGAM and ITGA4 showed no significant differences between the diets in either sex.

**Quantification of atherosclerotic lesion area**

Fig. 1 shows the atherosclerotic lesion area in apoE-knockout mice consuming the different diets. After consumption of a diet supplemented with nuts or palm oil for 12 weeks, the aortic atherosclerotic lesion areas tended to show higher values in males than in females, although the difference did not reach statistical significance (94.1 (SEM 12.0) vs. 70.8 (SEM 7.3) $\times 10^3$ $\mu$m$^2$, $P < 0.11$). A sex-related difference in the pattern of response to the different diets was noted; while female mice fed the nut diet showed significantly lower values than those on the palm oil diet, this effect was not observed in males receiving the same diets. Moreover, males consuming the nut diet showed two types of response: one group of mice ($n = 6$) presented a marked reduction in lesion area (34.5 (SEM 8.4) $\times 10^3$ $\mu$m$^2$), while the other ($n = 5$) showed a considerable increase (158.5 (SEM 9.6) $\times 10^3$ $\mu$m$^2$), suggesting an individual genetic response to nut supplementation.

**Oxidative stress variables**

As shown in Fig. 2(a), LDL prepared from the nut groups showed significantly lower levels of reactive oxygen species than LDL from palm oil-fed animals in both sexes. The decrease was particularly important in females.

### Table 3. Effect of dietary intervention on plasma parameters in apoE-knockout mice according to sex (Mean values with their standard errors)

<table>
<thead>
<tr>
<th></th>
<th>Female</th>
<th></th>
<th>Male</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Nut† ($n = 12$)</td>
<td>Palm oil ($n = 11$)</td>
<td>Nut† ($n = 11$)</td>
<td>Palm oil ($n = 12$)</td>
</tr>
<tr>
<td></td>
<td>Mean</td>
<td>SEM</td>
<td>Mean</td>
<td>SEM</td>
</tr>
<tr>
<td>Initial cholesterol (mmol/l)</td>
<td>12.9</td>
<td>0.6</td>
<td>12.1</td>
<td>0.7</td>
</tr>
<tr>
<td>Total cholesterol (mg/dl)</td>
<td>9.1</td>
<td>0.8</td>
<td>10.8</td>
<td>0.9</td>
</tr>
<tr>
<td>HDL-cholesterol (mmol/l)</td>
<td>0.11</td>
<td>0.04</td>
<td>0.14</td>
<td>0.03</td>
</tr>
<tr>
<td>VLDL- + LDL-cholesterol (mmol/l)</td>
<td>8.5</td>
<td>0.7</td>
<td>10.6</td>
<td>0.9$^*$</td>
</tr>
<tr>
<td>TAG (mmol/l)</td>
<td>2.0</td>
<td>0.2</td>
<td>1.9</td>
<td>0.1</td>
</tr>
<tr>
<td>ApoA1 (AU)</td>
<td>110</td>
<td>6</td>
<td>112</td>
<td>6</td>
</tr>
<tr>
<td>ApoA4 (AU)</td>
<td>41</td>
<td>3</td>
<td>37</td>
<td>4</td>
</tr>
<tr>
<td>Arylesterase activity (x 10$^3$ µmol/min per l)</td>
<td>12.2</td>
<td>0.5</td>
<td>10.3</td>
<td>0.8$^*$</td>
</tr>
<tr>
<td>ITGAM† (% monocytes)</td>
<td>24.4</td>
<td>1.5</td>
<td>25.7</td>
<td>1.7</td>
</tr>
<tr>
<td>ITGA4† (% monocytes)</td>
<td>78.5</td>
<td>2.5</td>
<td>81.1</td>
<td>1.9</td>
</tr>
</tbody>
</table>

AU, arbitrary units; ITGAM, integrin $\alpha$M; ITGA4, integrin $\alpha$4.

† Nut, diet supplemented with 3% nuts, composed of 50% walnuts, 25% almonds and 25% hazelnuts. Statistical analysis was carried out using Student’s $t$ test.

‡ ITGAM and ITGA4 protein levels are shown.
Since PON2 is a ubiquitous antioxidant enzyme (34), we investigated whether the test diets induced any differential hepatic mRNA expression. According to the results depicted in Fig. 2(b), female mice consuming the nut diet showed a significant increase in the hepatic expression of this message. On the other hand, the hepatic mRNA expression of Pcyox1 (a pro-oxidant enzyme of LDL (35)) was also tested (Fig. 2(c)), and no change was observed in females, while it increased in males consuming the nut diet.

**Association among parameters**

In an association analysis including all animals, atherosclerotic lesion areas showed a positive relationship with plasma cholesterol concentrations ($r = 0.49$, $P = 0.001$). When the two sexes were analysed separately, this association was stronger in females ($r = 0.71$, $P = 0.00$; Fig. 3(a)). Likewise, a significant inverse association was observed between atherosclerotic lesion areas and hepatic Pon2 mRNA expression ($r = -0.55$, $P = 0.02$; Fig. 3(b)).

**Discussion**

Atherosclerotic lesions in female and male apoE-deficient mice showed differential responses to a diet supplemented with 3% mixed nuts compared with a diet with a similar contribution of fat provided by palm oil (saturated fat). In both sexes, the addition of a small amount of nuts decreased non-HDL-cholesterol and the content of reactive oxygen species in LDL. However, an increase in serum paraoxonase activity and in the hepatic expression of Pon2 was observed only in females consuming the nut diet. The decrease in atherosclerotic lesion areas was significant and correlated positively with plasma cholesterol and negatively with Pon2 expression. These results indicate that female mice benefit from nut intake, as it lowers their LDL and endows them with a higher antioxidant defence through PON2.

The decreased non-HDL-cholesterol in males and females on the nut diet compared with the saturated fat-enriched diet, palm oil in this case, agrees with numerous human clinical studies that have consistently shown a favourable plasma lipid response to diets including almonds, hazelnuts or walnuts (8,9,11–16). The beneficial effects of nuts on the lipid profile are mainly attributed to their favourable fatty acid composition – low in saturated and high in unsaturated fatty acids. In the present experiment, the nut supplement contributed to the increase in the PUFA content of this diet, leading to a higher and more favourable polyunsaturated:saturated ratio.
ratio than palm oil (3·9 v. 1·6, respectively; Table 1). The decrease in plasma cholesterol produced by a high PUFA in comparison with a saturated fat diet has also been reported in male apoE-deficient mice (27), in both sexes in LDL receptor- and apoE-deficient mice (36), in male rabbits (37) and in male hamsters (38). Therefore, using this experimental design, non-HDL-cholesterol in apoE-deficient mice fed nuts follows the human pattern and could be attributed to the fatty acid composition of the diets.

With respect to the finding of a lack of effect on HDL-cholesterol, or on its main apoA1, human studies involving diets enriched in almonds, hazelnuts or walnuts have also shown contradictory results, reporting increases (8,11,14) or absence of changes (13,39). Furthermore, there is no agreement on the effect of PUFA-containing diets on plasma HDL-cholesterol in animal studies: PUFA tend to decrease HDL-cholesterol in rabbits receiving a diet enriched with maize oil compared with olive or avocado oil (37), but tend to increase it in comparison with coconut oil. Meanwhile in male hamsters, the substitution of maize oil as a source of polyunsaturated fat for palm oil decreased HDL-cholesterol (38). Moreover, a different HDL response to diets has been reported in LDL receptor- and apoE-deficient mice, two models of atherosclerosis (36). While apoE-deficient mice consuming PUFA diets compared with diets containing SFA showed decreased HDL-cholesterol in females as well as in males, no differences were observed between these diets in terms of plasma HDL in LDL receptor-deficient mice of either sex. Overall, these results point to an interaction among dietary composition, sex and genetic background that makes dietary response complex.

In the present study, there were sex-related differences in the response of atherosclerotic lesions to dietary nut supplementation, a circumstance that does not seem to be the case in human studies dealing with risk factors (16,40) or in an intervention analysing carotid atherosclerosis (44). Despite the fact that we used the same proportion of nuts as the latter study, we should point out the differences between the two reports, including the analysis of different arteries and the particular lipid metabolism in apoE-deficient mice (18), which, moreover, exhibit clear differences between sexes (27,42). In females, the nut-supplemented diet delayed the progression of aortic lesions compared with palm oil, a finding that may be attributed to a synergistic effect on plasma non-HDL-cholesterol levels, on the one hand, and on increased antioxidant defence of LDL, on the other hand. The latter could be executed through an increase in Pon2 expression as an ubiquitous antioxidant enzyme (54) and without any change in Pcyox1, the pro-oxidant agent of these particles (55). In fact, PON2 has been shown to protect against atherosclerosis development (45–48). In addition, its expression could also be modified by sex as occurs with Pon1, another member of the three-gene paraoxonase family (47).

Favourable effects of nuts may be attributed to their fatty acid profiles or the presence of other minor components. One limitation of our experimental design using palm oil for comparison is the inability to discriminate between the two components, but a positive aspect is that it proposes a practical and reasonable combination to be recommended for populations. Indirectly, the changes observed in hepatic Pon2 gene expression cannot be explained by the different fatty acid profile. In macrophages, this enzyme has been induced by polyphenols from pomegranate (48), and some of these substances are also present in walnuts, but not in hazelnuts or almonds (49). In fact, our nut diet provides a higher amount of hydrolysable polyphenols. Therefore, the present observation would represent an initial contribution to defining a role for these compounds in the regulation of PON2 via its mRNA expression as an ubiquitous antioxidant enzyme (54), instead, ellagic acid and polyphenol-rich extracts from walnuts were effective inhibitors of in vitro plasma LDL oxidation (51,52), and flavonoids from the almond peel protected LDL from oxidation in hamsters (53). However, other authors have found no differences in plasma antioxidant status following walnut consumption, despite an increase in thiols (49), nor have Bullo et al. (54) observed differences between walnuts and walnut-skin extracts. In this respect, despite the slightly high hydrolysable polyphenol content of the nut diet, the different response observed in females may also be attributed to possible differences in nut polyphenol species with respect to palm oil, and...
to combinations and interactions among them and with the matrix. Phenolic compounds may exert both anti- and pro-oxidant effects. In this respect, Acín et al. reported that the oral administration of hydroxytyrosol to apoE-deficient mice increased the atherosclerotic lesion area, suggesting that phenolic compounds, outside the original matrix, could not only fail to be useful, but could even be harmful. Another source of variation is introduced by sex, which, in this species, modifies the response to exogenous factors and genetic background. The clarification of these aspects will require further studies.

In conclusion, the present results in apoE-deficient mice show that, compared with consumption of a palm oil-enriched diet, consumption of a mixed nut diet is associated with decreased aortic atherosclerosis in females but not in males, an effect that appears to be related to both a reduction in non-HDL-cholesterol and an enhanced expression of Pon2 with protection of LDL from oxidation.

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