Olive oil and oxidative stress

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RESUMEN
Aceite de oliva y estrés oxidativo.

La composición del aceite de oliva virgen extra se caracteriza por su contenido en ácidos grasos, fundamentalmente monounsatuatedados (ácido oleico) beneficiosos para reducir el riesgo de enfermedad coronaria, y en componentes menores, particularmente polifenoles (p.e. hidroxitirosol y oleuropeína) responsables de su sabor y estabilidad. Diversos estudios demuestran el poder antioxidante de los compuestos fenólicos del aceite de oliva (virgen extra). Aunque la mayoría de ellos se han realizado in vitro, algunos in vivo parecen confirmar que los polifenoles se absorben dependiendo de la dosis y que retienen las actividades biológicas después de su ingestión. Estos resultados pueden explicar en parte la menor incidencia de enfermedad coronaria en los paises del área Mediterránea, donde el aceite de oliva (extra virgen) es la principal fuente de grasas.

PALABRAS-CLAVE: Aceite de oliva; Antioxidantes; Hidroxitirosol; Dieta Mediterránea; Polifenoles; Oleuropeína.

SUMMARY
Olive oil and oxidative stress.

In addition to the fatty acid profile of olive oil, which is high in the monounsaturated oleic acid and appears to be beneficial in reducing several risk factors for coronary heart disease and certain cancers, extra virgin olive oil contains a considerable amount of phenolic compounds, e.g. hydroxytyrosol and oleuropein, that are responsible for its peculiar taste and for its high stability. A body of evidence demonstrates that olive oil phenolics are powerful antioxidants. Although most of these studies have been carried out in vitro, some in vivo experiments confirm that olive oil phenolics are dose-dependently absorbed and that they retain their biological activities after ingestion. These data could in part explain the lower incidence of coronary heart disease in the Mediterranean area, where (extra virgin) olive oil is the principal source of fat.

KEY-WORDS: Olive oil; Antioxidants; Hydroxytyrosol; Mediterranean diet; Phenolics; Oleuropein

1. INTRODUCTION
Atherosclerosis begins as an injury to the endothelium of the blood vessel walls (Ross, 1999; Assanelli et al., 2004; Schulz et al., 2004). Many studies reported that hyperlipidemia might injure the cell wall as well as interfere with normal responses by tissues to injury, leading to an infiltration of blood platelets, the proliferation of smooth muscle cells, and accumulation of collagen and lipids at the site of damage. Atherosclerotic plaques evolve in the narrowing of the coronary arteries that supply blood to the heart.

There is evidence that classic risk factors for coronary heart disease (CHD), such as high serum cholesterol and blood pressure, are not much different between the populations of the Mediterranean area - where the incidence of CHD and certain cancers, e.g. breast and colon cancers, is lowest than those of other North European and Western countries. Further, there are several observations that do not completely link CHD incidence and fat intake and absorption (Mancini and Rubba, 2000). These data suggest that other, yet unexplored risk factors may be favorably affected by a healthful diet. Conversely, in order to lower the risk of onset and progression of CHD, nutritional recommendations have been primarily focused upon dietary regimes aimed at reducing the amount of circulating lipids as well as the formation of blood clots.

These observations, together with several studies indicating a key role played by oxidation of low-density lipoproteins (LDL) in the onset of atherosclerosis (Witztum and Steinberg, 2001), led to the formulation of an antioxidant/atherosclerosis hypothesis. Polyphenolic metabolites of plants, common in the Mediterranean area, are found in fruits, vegetables and olive oil (see this issue of Grasas y Aceites), and act as strong antioxidants in various systems. Their multiple biological actions
have been reviewed by many authors (Kris-Etherton et al., 2002). Moreover, many epidemiological studies have shown an inverse association between the consumption of fruit, vegetables, and their products and the incidence of cardiovascular diseases and cancer (Hu and Willett, 2002; Kromhout et al., 2002).

Clinical and epidemiological studies have shown that the dietary factor most closely correlated with high levels of blood cholesterol and CHD risk are saturated fatty acids (SFA). The healthful properties of olive oil have been, until recently, exclusively attributed to its high monounsaturated fatty acid (MUFA) content, mostly in the form of oleic acid (18:1\(\text{n}-9\)), which ranges from 56 to 84% of total fatty acids. However, several observations argue against this hypothesis. For example, the effects of MUFA on circulating lipids and lipoproteins are still equivocal. It is therefore unlikely that oleic acid is exclusively accountable for the healthful properties of olive oil. Finally, it is also noteworthy that several seed oils obtained through genetic selection, such as sunflower, soybean, and rapeseed oils are nowadays rich in MUFA, albeit devoid of phenolics (Owen et al., 2000), and are commercially available. Healthy effects of dietary MUFA, including lower endothelial activation (Massaro et al., 2002; Massaro and De Caterina, 2002) and susceptibility of LDL to oxidation (Berry et al., 1992; Bonanome et al., 1992), are indeed to be considered; but it is also remarkable to establish the amount and quality of phenolic compounds in extra virgin olive oils.

This issue focuses on the evidence that indicates how the phenolics of extra virgin olive oil may play a role in the protection from CHD and cancer observed in the Mediterranean area (see also issues 3 and 4). These effects go beyond those of oleic acid and we propose that consumption of good-quality (extra virgin) olive oil might have an important impact on our health.

2. OIL OIL PHENOLICS AND OXIDATIVE STRESS

Among the several minor compounds of virgin olive oil, there are vitamins such as \(\alpha\) and \(\gamma\)-tocopherols, \(\beta\)-carotene, phytosterols, squalene, pigments, terpenic acids, flavonoids such as luteolin and quercetin, and phenolics usually referred to as polyphenols (Boskou, 2000; Blekas et al., 2002).

As dietary antioxidants are likely capable of decreasing the incidence of CHD and certain cancers in the Mediterranean area (Trichopoulou et al., 1999), research is investigating the potentially protective activities of olive oil minor constituents, some of which (namely hydroxytyrosol and oleuropein) have been recently become commercially available.

As opposed to other vegetable oils, (extra virgin) olive oil possesses substantial amount of phenolic compounds. One of the current major limitations to natural antioxidant research is the current lack of appropriate methodologies to quantify the amount of phytochemicals, including phenolic molecules, in foods. This also applies to (extra virgin) olive oil: several methods to quantify and identify its phenolic components have been developed but they rarely agree. Currently, the most popular methods for evaluating the polyphenolic content of (extra virgin) olive oil are the Folin-Ciocalteau colorimetric assay (Visioli et al., 1995b) and the HPLC (Montedor et al., 1992). Although the colorimetric method is simple to perform and does not require expensive equipment, it has low specificity of the reagent toward phenolic compounds and it does not provide qualitative information of the composition of the phenolic fraction. Given the importance of catecholic compounds within the phenolic fraction, this information is actually of extreme importance. Conversely, HPLC is very sensitive and specific, it can identify relevant molecules, but it is time-consuming and does not provide information on phenolic molecules for which reference standards are unavailable. Another assay, sensitive, specific, and easy to perform has been proposed by Mosca et al. (2000). This assay is based on a substrate-recycling procedure that employs tyrosinase together with NADH as a reducing agent. Again, this method only provides quantitative information. Finally, a rapid and sensitive method to evaluate the phenolics of olive oil by Atmospheric Pressure Chemical Ionization-Mass Spectrometry (APCI-MS) has been described by Caruso et al. (2000). This method allows for quick analyses of crude methanolic extracts of olive oil, does not need extensive analytical workup, and makes it possible to quantify oleuropein aglycone. However, the apparatus is very expensive and requires trained personnel to operate.
In turn, the lack of reliable methodology to evaluate the quali/quantitative profile of olive oil phenolic fraction is one of the major obstacles that prevent dose-response, pharmacology-style trials in animals and humans.

3. OLIVE OIL PHENOLICS AS BIOACTIVE COMPOUNDS

3.1. In vitro studies

3.1.1. Antioxidant activities

In 1994, our laboratory initiated a series of experiments aimed at investigating the antioxidant activities of (extra virgin) olive oil phenolics on parameters that are considered relevant to human health. The first paper reported inhibition of LDL oxidation by oleuropein glycoside, which was, to our knowledge, the only phenol typical of olive oil that was commercially available (Extrasyntese, France).

Oleuropein was found to potently and dose-dependently inhibit LDL oxidation induced by copper sulphate (Visioli and Galli, 1994). This investigation was subsequently expanded to include hydroxytyrosol (which wasn’t commercially available but was isolated from olive oil and kindly provided by Professor G.F. Montedoro, University of Perugia) and other catechols (Visioli et al., 1995a). Metal-independent oxidation was also tested and revealed that hydroxytyrosol, in addition being a metal chelator, is also a scavenger of free radicals (Visioli et al., 1995a). Subsequent experiments showed that both oleuropein and hydroxytyrosol effectively scavenge superoxide anion generated by either human polymorphonuclear cells or by the xantine/xantine oxidase system (Visioli et al., 1998a); it is noteworthy that, in these experimental setups, both vitamin E and butylated hydroxytoluene were found to be inactive. A scavenging effect of oleuropein and hydroxytyrosol was also demonstrated with respect to hypochlorous acid (Visioli et al., 1998a), a potent and dangerous oxidant species produced in vivo by activated neutrophils at the site of inflammation (Aruoma and Halliwell, 1987). In addition to damaging nearby molecules such as α-antiprotease, hypochlorite is a major component of chlorine-based bleaches that can often be exposed to food during manufacturing. In terms of protection from atherosclerosis, the HOCI-scavenging property of hydroxytyrosol may bear important consequences: evidence is rapidly accumulating that the formation of chloramines via the myeloperoxidase-catalyzed formation of HOCI and subsequent chlorination of apo B-100 is responsible for LDL modification and peroxidation (Carr et al., 2000).

It is of interest that similar antioxidant activities were discovered in the major by-product of olive oil production (e.g. olive mill waste-water), which is very rich in phenolic molecules [actually, due to their partition coefficient, olive phenols are more abundant in waste-water than in olive oil (Rodis et al., 2002)] and might be employed in preservative chemistry (Visioli et al., 1995b; Visioli et al., 1999; Leger et al., 2000; Visioli and Galli, 2003). Indeed, the lack of appropriate disposal techniques for waste-waters (Demichel and Bontoux, 1996) and the increasing need for antioxidants of natural origin make olive mill waste-waters an appealing resource of bioactive molecules, which can be selectively recovered (Italian patent n° BO2001A0419) from this abundant source.

Following our initial observations, other laboratories joined in this line of research to further investigate the antioxidant properties of olive phenols. For example, hydroxytyrosol proved to be effective in a model of oxidative stress induced in intestinal epithelial cells (Manna et al., 1997). Conversely, tyrosol, which lacks the ortho-diphenolic structure, was found to be ineffective in this experimental model, as it was in the models of LDL oxidation described above (unpublished data). Manna et al. (1999) also described a protective effect of hydroxytyrosol toward hydrogen peroxide-induced damage to human erythrocyte. Vascular production of superoxide anion is associated with increased risk for endothelial dysfunction and cardiovascular disease (Schulz et al., 2004). As indicated above, oleuropein and hydroxytyrosol were found to be very effective scavengers of superoxide (Table 1).

Damage to DNA is associated with increased risk for cancer. The activities of hydroxytyrosol toward chemically-induced DNA and aminoacid modification have been investigated (Deiana et al., 1999) and it was found that low concentrations of hydroxytyrosol, e.g. 50 µM, are able to scavenge peroxynitrite and therefore to prevent ONOO⁻-dependent DNA damage and tyrosine nitration. Scavenging of peroxynitrite was further confirmed (de la Puerta et al., 2001).

A matter of concern in oxidant/antioxidant chemistry is the prooxidant potential of several molecules, including antioxidants with strong reducing capabilities. The prooxidant activities of hydroxytyrosol were investigated in a model of copper-induced DNA damage and were found to be 40-fold weaker than those of ascorbate. Further, these actions occurred only at very high, non-physiological concentrations (>500 µM) (Deiana et al., 1999).

The mechanism of action of olive oil phenolics is related to their o-diphenolic structure, which increase radical stability by forming an intra-molecular hydrogen bond between the free hydrogens of their hydroxyl group and their phenoxyl radicals (Visioli et al., 1998a). Although studies on structure-activity of
olive oil phenolics are yet to be carried out, similar investigations performed on phenolic molecules indicated that the degree of antioxidant activity is correlated with the number of hydroxyl substitutions (Rice-Evans et al., 1996). In particular, the o-diOH (catecholic) substitution confers a high antioxidant capacity, whereas single hydroxyl substitutions, as in the case of tyrosol, lack antioxidant activity.

3.1.2. Beyond antioxidants. Enzyme-modulating activities

One of the most interesting aspects of olive oil phenolics' biochemistry is the discovery of their enzyme-modulating properties, which, in some cases, are unrelated to their antioxidant potential. In fact, most olive oil phenolics are amphiphilic and they partition between the lipid (olive oil) and water (waste-water) phases. Therefore, they might be able to interact with enzymes relevant to human pathology, namely atherosclerosis.

For example, hydroxytyrosol was proven to inhibit chemically-induced in vitro platelet aggregation, the accumulation of the pro-aggregant agent thromboxane (TxA₂) in human serum, the production of the pro-inflammatory molecules leukotrienes by activated human leukocytes, and the arachidonate lipoxygenase (Petroni et al., 1995; Kohyama et al., 1997; de la Puerta et al., 1999; de la Puerta et al., 2000; Martinez-Dominguez et al., 2001). The mean inhibiting concentration (IC₅₀) were found to be in the 10⁻⁵ M range, indicating unpredicted biological activities of olive oil phenolics that go beyond their antioxidant properties.

Concerning the immuno-modulating properties of olive oil phenolics, in vitro studies on murine macrophages revealed that oleuropein increases the functional response of these immune-competent cells when they are stimulated with bacterial polysaccharide, as evaluated by a significant increase (+58.7 ± 4.6%, mean ± SD) in the production of nitric oxide (Visioli et al., 1998b). It is noteworthy that this increase is due to a direct tonic effect of oleuropein, which increases both the activity and the expression of the inducible form of the enzyme nitric oxide synthase (iNOS). As relevant to atherosclerosis, it should be noted that during acute sepsis and inflammation that take place in the arterial wall, macrophages react to the endotoxin challenge by increasing the production of nitric oxide, which inhibits platelet aggregation and adherence and maintains a proper end-organ perfusion rate through increased vaso-relaxation. In agreement with this finding, inhibition of nitric oxide synthesis during sepsis increases cellular damage and animal mortality (Lowenstein et al., 1996). Finally, macrophagic nitric oxide exerts antioxidant activities that play a protective role in preventing oxidative LDL modification that occur at the site of

Table 1

<table>
<thead>
<tr>
<th>Superoxide-producing system</th>
<th>Xantine-xantine oxidase</th>
<th>PMN + PMA</th>
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<tbody>
<tr>
<td>Hydroxytyrosol (IC₅₀)</td>
<td>9.1 µM</td>
<td>3.2 µM</td>
</tr>
<tr>
<td>Oleuropein (IC₅₀)</td>
<td>14.3 µM</td>
<td>29.3 µM</td>
</tr>
</tbody>
</table>

The mean inhibiting concentration (IC₅₀) was calculated by employing MacALLFIT as software. PMN, human polymorphonuclear neutrophils; PMA, phorbol-12-myristate-13 acetate. From Visioli et al. (1998a).

![Figure 1](image_url)

Hydroxytyrosol (HT) and oleuropein glycoside (OE) do not inhibit MMP-2 activity in bovine aortic endothelial cells (BAEC). BAEC were incubated for 24 hours at 37 °C with DMEM supplemented with 0.2% BSA in the absence (control) or in the presence of HT or OE 10 µM. Gelatinolytic activity was detected by SDS-PAGE zymography on 7.5% polyacrylamide gels containing 10% SDS and gelatin (1 mg/mL) under non-reductive conditions. After electrophoresis, SDS was removed from the gel by two washes with 2.5% Triton X-100. At the end of the incubation, gels were stained with a solution of NaCl (150 mmol/L), CaCl₂ (10 mmol/L), and ZnCl₂ (1 mol/L). Areas of lysis appeared after renaturing and staining of gels with Coomassie brilliant-blue. Gelatinolytic bands were quantified by densitometry (software: NIH Image 1.52). Statistical significance was investigated by Student’s t test with two–tailed distribution (software: SPSS 11.0 version). *p<0.01 compared to controls. Unpublished data.
inflammation during enhanced reactive oxygen species production (Jessup et al., 1999).

More recently, we studied the activities of olive phenols on other enzymes relevant to cardiovascular disease, e.g. metalloproteases (MMP). These enzymes can degrade the atherosclerotic plaque, leading to plaque instability and enhancing possibility to release thrombi. As shown in Figure 1, olive phenols are unable to down-regulate MMP's activity in bovine aortic endothelial cells, which was actually mildly enhanced by hydroxytyrosol supplementation. Further experiments will clarify whether this lack of effect applies to other MMP and if it is unique to oleuropein and hydroxytyrosol.

3.2. In vivo studies

Notwithstanding the hype that is currently surrounding natural antioxidants and their healthful effects, very limited number of controlled trials proves their efficacy in vivo. Experimental evidence that flavonoids and phenolic compounds are absorbed from the diet is accumulating (Bravo, 1998) and this includes olive oil phenolics. Studies with laboratory animals have demonstrated a higher resistance to oxidation of LDL obtained from animals fed virgin olive oil, as compared to LDL separated from animals that were only administered an equivalent amount of oleic acid as either triolein (Scaccini et al., 1992) or "plain" olive oil (Wiseman et al., 1996). Visioli et al. (2000b) demonstrated the dose-dependent absorption of olive oil phenolics in humans and that their urinary excretion as glucuronide conjugates; another interesting finding of that study was that increasing amounts of phenolics administered with olive oil stimulated the rate of conjugation with glucuronide. These findings were confirmed by other laboratories (Miro-Casas et al., 2001; Vissers et al., 2002; Miro-Casas et al., 2003a; Miro-Casas et al., 2003b), and recent data shed light on the human metabolism of hydroxytyrosol (Caruso et al., 2001). From studies carried out in human intestinal Caco-2 cells, it appears that hydroxytyrosol is absorbed from the gut by passive diffusion (Manna et al., 2000) and that it is extensively metabolized before being excreted with the urine. Whether circulating metabolites still exert biological actions is still under investigation, although data from other laboratories that study tea components, namely quercetin, suggest even higher antioxidant potential of the metabolites as compared with the parent molecule (Moon et al., 2001).

Interestingly, the bioavailability of hydroxytyrosol is higher when this molecule is administered as natural component of olive oil than when it is given after addition to olive oil or to yogurt, as approximation of a functional food (Visioli et al., 2003a). These data actually add to the notion that oligonutrients bioavailability is lower than such compounds isolated from their original matrix and administered as supplements (Visioli et al., 2003b).

In vivo evidence of the antioxidant activity of (extra virgin) olive oil phenolics, namely hydroxytyrosol, is also accumulating. In addition to the effects on LDL oxidation reported above, we have been recently able to demonstrate that hydroxytyrosol, administered to animals as the only bioactive component of an olive mill waste-water extract, is dose-dependently absorbed and is able to increase their plasma antioxidant capacity (Visioli et al., 2001). Further, low doses of hydroxytyrosol, e.g. 414 µg/animal, are able to inhibit passive smoking-induced oxidative stress in animals, as demonstrated by a reduced urinary excretion of the F₂-isoprostane 8-iso-PGF₂α (IPF₂α-III) (Visioli et al., 2000c). Finally, a dose-dependent inhibition of the rate of 8-iso-PGF₂α excretion was observed in human volunteers who ingested olive oils added with increasing amounts of phenolics (Visioli et al., 2000a). Interestingly, the urinary levels of 8-iso-PGF₂α inversely correlated with those of homovanillyl alcohol, e.g. a catechol-o-methyltransferase (COMT)-derived metabolite of hydroxytyrosol (Manna et al., 1999; Lamensdorf et al., 2000), suggesting that this phenol

<table>
<thead>
<tr>
<th>Concentration (mM)</th>
<th>Cell lines</th>
</tr>
</thead>
<tbody>
<tr>
<td>Test agent</td>
<td>GN61</td>
</tr>
<tr>
<td>oleuropein aglycone</td>
<td>0.66</td>
</tr>
<tr>
<td>oleuropein glycoside</td>
<td>1.0</td>
</tr>
<tr>
<td>caffeic acid</td>
<td>1.5</td>
</tr>
<tr>
<td>o-coumaric acid</td>
<td>6.0</td>
</tr>
<tr>
<td>cinnamic acid</td>
<td>6.75</td>
</tr>
<tr>
<td>tyrosol</td>
<td>&gt;10.0</td>
</tr>
<tr>
<td>syringic acid</td>
<td>9.0</td>
</tr>
<tr>
<td>protocatechuic acid</td>
<td>10.0</td>
</tr>
<tr>
<td>vanillic acid</td>
<td>10.0</td>
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Table 3
Antioxidant and additional biological actions of olive oil phenolics

<table>
<thead>
<tr>
<th>Action</th>
<th>Reference(s)</th>
</tr>
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<tbody>
<tr>
<td><strong>Antioxidant activities, in vitro</strong></td>
<td></td>
</tr>
<tr>
<td>Inhibition of LDL oxidation</td>
<td>(Scaccini et al., 1992; Grignaffini et al., 1994; Visioli and Galli, 1994;</td>
</tr>
<tr>
<td></td>
<td>Salami et al., 1995; Visioli et al., 1995a; Visioli et al., 1995b; Wiseman</td>
</tr>
<tr>
<td></td>
<td>et al., 1996; Aruoma et al., 1998; Leenen et al., 2002; Wiseman et al.,</td>
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<tr>
<td></td>
<td>2002; Benkhalti et al., 2003)</td>
</tr>
<tr>
<td>Inhibition of apolipoprotein modification</td>
<td>(Visioli et al., 1995a)</td>
</tr>
<tr>
<td>Scavenging of free radicals and other oxidants</td>
<td>(Le Ttour and Guedon, 1992; Aeschbach et al., 1994; Manna et al., 1997;</td>
</tr>
<tr>
<td></td>
<td>Saja et al., 1998; Speroni et al., 1998; Visioli et al., 1998a; Manna et</td>
</tr>
<tr>
<td></td>
<td>al., 1999; Fogliano et al., 1999; Leger et al., 2000; Benavente-Garcia et al.,</td>
</tr>
<tr>
<td></td>
<td>2000; de la Puerta et al., 2001; Casalino et al., 2002; Briante et al.,</td>
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<tr>
<td></td>
<td>2002; Quiles et al., 2002)</td>
</tr>
<tr>
<td>Inhibition of peroxynitrite-induced DNA damage and protein nitration</td>
<td>(Aruoma et al., 1998; Deiana et al., 1999)</td>
</tr>
<tr>
<td><strong>Modulation of enzymes, in vitro</strong></td>
<td></td>
</tr>
<tr>
<td>Inhibition of platelet aggregation</td>
<td>(Petroni et al., 1995)</td>
</tr>
<tr>
<td>Reduced cyclo- and lipoxygenase activation</td>
<td>(Petroni et al., 1995; Kohyama et al., 1997; Petroni et al., 1997; Visioli</td>
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<tr>
<td></td>
<td>et al., 1999; de la Puerta R. et al., 1999; Brzosko et al., 2002)</td>
</tr>
<tr>
<td>Increased nitric oxide production by LPS-challenged macrophages</td>
<td>(Visioli et al., 1998b)</td>
</tr>
<tr>
<td>Inhibition of neutrophil respiratory burst</td>
<td>(Visioli et al., 1998a)</td>
</tr>
<tr>
<td><strong>Cellular activities</strong></td>
<td></td>
</tr>
<tr>
<td>Inhibition of bacterial growth and activity</td>
<td>(Tranter et al., 1993; Capasso et al., 1995; Bisignano et al., 1999)</td>
</tr>
<tr>
<td>Decreased endothelial activation</td>
<td>(Carluccio et al., 2003)</td>
</tr>
<tr>
<td>Cytostasis</td>
<td>(Saenz et al., 1998)</td>
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<tr>
<td><strong>Ex vivo and in vivo activities</strong></td>
<td></td>
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<tr>
<td>Increased plasma antioxidant capacity and resistance of LDL to</td>
<td>(Ramirez-Tortosa et al., 1999; Visioli et al., 2001; Masella et al., 2001;</td>
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<tr>
<td>oxidation</td>
<td>Ochoa et al., 2002; Fito et al., 2002; Wiseman et al., 2002)</td>
</tr>
<tr>
<td>Antiinflammatory activity</td>
<td>(de la Puerta et al., 2000; Martinez-Dominguez et al., 2001)</td>
</tr>
<tr>
<td>Decreased isoprostane excretion in humans and in sidestream smoke-</td>
<td>(Visioli et al., 2000a; Visioli et al., 2000c)</td>
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<tr>
<td>exposed animals</td>
<td></td>
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<tr>
<td>Decreased TxB2 production by human serum</td>
<td>(Visioli and Galli, 2003)</td>
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<tr>
<td>Vasomodulation</td>
<td>(Panizzi et al., 1960; Benkhalti et al., 2003)</td>
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enters into cellular compartments where it exerts its antioxidant activity. As other laboratories failed to detect any biological effect of extra virgin olive oil administration (Vissers et al., 2001a,b), no clear conclusion can be drawn and further experiments are needed before any health claim can be made.

As far as toxicity of (extra virgin) olive oil phenolics is concerned, very few data are available that did not show toxic effects of hydroxytyrosol in animals (D’Angelo et al., 2001) and cell cultures (Table 2) (Babich and Visioli, 2003). Currently, there is a serious lack of methodology to allow for controlled human trials. In addition to the uncertainty that surrounds the determination of phytochemicals in foods, their detection in human fluids/tissues is even more problematic. Moreover, there is no reliable method to detect the effect of antioxidant supplementation on human oxidative status; until a reliable biomarker is validated (Halliwell, 2000; Halliwell et al., 2004), it is impossible to carry out controlled studies on the antioxidant activities of oligonutrients, including those of olive oil (Visioli, 2004).

4. CONCLUSIONS

It is now clear that the incidence of degenerative pathologies, including those in which the excessive free radical formation has been suggested, is very low in the Mediterranean area, where the diet is rich in antioxidant compounds (Keys, 1995; Willett et al., 1995; Trichopoulou et al., 2003). Among such antioxidants, the contribution of olive oil phenolics is under active investigation (Table 3). Even though researchers are far from being able to make claims, based on the current data it is advisable to suggest the use of extra virgin olive oil as the predominant source of fat of a healthful, Mediterranean diet. The intake of bioactive molecules (hydroxytyrosol, oleuropein) through olive oil might contribute to lower the risk of developing CHD and certain cancers.

ACKNOWLEDGMENTS

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