Paraoxonase 1 Q192R (PON1-192) polymorphism is associated with reduced lipid peroxidation in healthy young men on a low-carotenoid diet supplemented with tomato juice

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The HDL-bound enzyme paraoxonase (PON) protects LDL from oxidation and may therefore attenuate the development of atherosclerosis. We examined the effect of tomato and carrot juice consumption on PON1 activity and lipid peroxidation in healthy young volunteers with different PON1-192 genotypes (Q/R substitution at position 192). In this randomized cross-over study twenty-two healthy, non-smoking men on a low-carotenoid diet received 330 ml/d tomato juice (37.0 mg lycopene, 1.6 mg β-carotene) or carrot juice (27.1 mg β-carotene, 13.1 mg α-carotene) for 2 weeks. Intervention periods were preceded by 2-week low-carotenoid intake. We determined the PON1-192 genotype by restriction fragment length polymorphism–polymerase chain reaction (RFLP-PCR) and measured LDL oxidation (lag time), plasma malondialdehyde and PON1 activity at the beginning and end of each intervention period. At baseline, lag time was higher (P<0.05) in QQ (111 (SD 9) min) than in QR/RR subjects (101 (SD 8) min). Neither tomato nor carrot juice consumption had significant effects on PON1 activity. However, tomato juice consumption reduced (P<0.05) plasma malondialdehyde in QR/RR (Δ: ~0.073 (SD 0.11) μmol/l) as compared to QQ subjects (Δ: +0.047 (SD 0.13) μmol/l). Carrot juice had no significant effect on malondialdehyde irrespective of the PON1-192 genotype. Male volunteers with the QR/RR genotype showed an increased lipid peroxidation at baseline. Although tomato and carrot juice fail to affect PON1 activity, tomato juice intake reduced lipid peroxidation in healthy volunteers carrying the R-allele of the PON1-192 genotype and could thus contribute to CVD risk reduction in these individuals.

Tomato juice: Antioxidants: Lipid peroxidation: Paraoxonase polymorphism

Diet plays an important role among the risk factors for atherosclerosis and CVD since it has an impact on, for example, plasma lipids, blood pressure and antioxidant mechanisms. Large prospective cohort studies revealed that a high intake of fruit and vegetables (Liu et al. 2000, 2001; Joshipura et al. 2001; Bazzano et al. 2002) and of dietary carotenoids from fruit and vegetables (Osganian et al. 2003) reduce CVD risk. In contrast, supplemental carotenoids from a non-food source have no beneficial effects on CVD risk and supplementation should not be further recommended (Vivekananthan et al. 2003).

Oxidation of LDL and the subsequent generation of lipid hydroperoxides and other lipid oxidation products are hypothesized to play a crucial role in the development of atherosclerosis and CVD (Diaz et al. 1997). Dietary antioxidants such as ascorbic acid, vitamin E and β-carotene can inhibit LDL oxidation in vitro (Frei et al. 1996; Jialal & Devaraj, 1996) and may therefore protect from CVD although clinical trials testing antioxidant supplementation revealed conflicting results (Salonen, 2002; Upston et al. 2003). LDL oxidation is also reduced by paraoxonase (PON), an enzyme bound to HDL (Mackness et al. 1991, 2000a; Aviram et al. 1998), and may therefore attenuate the development of atherosclerosis (Mackness et al. 1998a; Aviram, 1999). Recent studies proposed that paraoxonase activity predicts CVD risk (Jarvik et al. 2000, 2003; Mackness et al. 2003). So far, it is not yet clear whether the activity polymorphism for PON1-192, with a Q/R substitution at position 192 (Adkins et al. 1993; Humbert et al. 1993), has an impact on atherosclerosis (Mackness et al. 2003). The PON1 QQ-genotype exhibits low activity towards hydrolysis of paraoxon, while the RR-genotype shows high activity towards paraoxon. However, this isoenzyme corresponding to the RR-genotype exhibits a low hydrolysing activity towards lipid hydroperoxides. The phenylacetate substrate activity is not affected by the PON1-192 genotype (Mackness et al. 1998a).

PON1 activity can be modulated by dietary factors such as alcohol (vander Gaag et al. 1999), pomegranate juice (Aviram et al. 2000) and dietary fat (Sutherland et al. 1999; Kuchchodkar et al. 2000; Mackness et al. 2000b; Tomás et al. 2001) and may thus have an impact on CVD risk. A high vegetable intake also lowers PON1 activity and this effect was most prominent in women with the PON1-192 QR/RR-genotype (Rantala et al. 2002). In a similar study, Kleemola et al. (2002) showed that PON1 activity is reduced during high vegetable intake and that it is negatively correlated with the intake of β-carotene. Data on other carotenoids, e.g. lycopene, were not presented in that study.

Abbreviations: PON1, paraoxonase 1; RFLP, restriction fragment length polymorphism; TBARS, thiobarbituric acid reactive substances; MDA, malondialdehyde.

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Recently, we demonstrated that consumption of a lycopene-rich tomato juice for 2 months reduced lipid peroxidation in QR/RR elderly subjects, but had no effect on PON1 activity (Bub et al. 2002). Here we investigated whether tomato or carrot juice consumption modulates PON1 activity and parameters of lipid peroxidation in healthy young volunteers on a low-carotenoid diet. Furthermore, we determined the PON1-192 genotype in order to see whether possible intervention effects are related to the PON1-192 polymorphism.

Material and methods

Subjects

Twenty-two healthy, non-smoking men participated in this study. All subjects were in good health as was determined by a screening history and medical examination. Subjects on medication or taking supplements were excluded from the study. The study was approved by the Medical Ethics Committee of the Landesa¨rztekammer Baden-W¨urttemberg and all participants gave their consent in writing.

Study design

This study was designed as a randomized cross-over study of two intervention periods (2 weeks each) with tomato juice or carrot juice (330 ml/d; Schoenenberger, Magstadt, Germany). Study participants were allocated to two groups (A: tomato juice/carrot juice; B: carrot juice/tomato juice). The tomato juice supplement (330 ml/d) provided 37·0 mg lycopene and 1·6 mg ß-carotene. Carrot juice supplementation (330 ml/d) supplied 27·1 mg ß-carotene and 13·1 mg α-carotene. Intervention periods were preceded by 2-week low-carotenoid periods and a 2-week washout period between the two interventions. After the second intervention period a third 2-week low-carotenoid period followed resulting in a total study period of 10 weeks. From a previous study we knew that a 2-week depletion or supplementation period is long enough to induce significant changes in plasma carotenoid concentrations and lipid peroxidation (Bub et al. 2000). Subjects were told to consume the juices with their main meals and were not restricted in their daily diet, except that they had to abstain from fruit and vegetables high in carotenoids throughout the whole study period. A list of the fruit and vegetables the subjects were not allowed to eat was provided (Müller et al. 1999).

Collection and preparation of blood samples

Fasting blood samples were taken at the beginning of the study and at the end of each experimental week in the morning between 7:00 and 9:00 hours. Blood was drawn from an antecubital vein into prechilled tubes containing EDTA (1·6 g/l; Monovette-Sarstedt, Nümbrecht, Germany) and immediately placed on ice in the dark. Plasma was collected after centrifugation at 1500 g for 10 min at 4°C. For the lipid peroxidation assay sucrose (15 g/l) was added to the plasma to prevent LDL aggregation and stored at −80°C until analysis. Tubes without anticoagulant ( Serum-Monovette-Sarstedt) were used for serum collections. Blood was allowed to clot at room temperature for 30 min, then centrifuged at 1500 g for 10 min at room temperature and the serum was stored at −80°C until analysis.

Analytical methods

The ex vivo oxidation of isolated LDL was performed by using a modified method of Esterbauer et al. (1999) which has been described previously (Bub et al. 2000). Intraset and inter-assay variability for LDL oxidation was <5% and <8%, respectively. Plasma malondialdehyde (MDA) was measured as thiobarbituric acid reactive substances (TBARS) by HPLC with fluorescence detection (Burkart et al. 2000) with minor modifications as described previously (Briviba et al. 2004). Intra- and inter-assay variabilities were <5%. Serum triacylglycerol, cholesterol and HDL-cholesterol were determined by enzymatic kits (Boehringer, Mannheim, Germany). LDL-cholesterol was calculated using the ‘Friedewald’ formula.

PON activity

Arylesterase activity of PON was determined spectrophotometrically using phenylacetate as substrate as described by Gan et al. (1991) with minor modifications. The assay mixture contained 5·0 mmol/l of phenylacetate and 0·9 mmol/l CaCl₂ in 20 mmol/l Tris-HCl, pH 8·0, at 25°C. The reaction was recorded at 270 nm. Non-enzymatic hydrolysis of phenylacetate was subtracted from the total rate of hydrolysis. Results are expressed as U/ml. The E₅₇₀ for the reaction is 1310 mol/l per cm and 1 U of arylesterase activity is equal to 1 μmol of phenylacetate hydrolysed/min per ml. The activity of PON towards paraoxon (paraoxonase activity) was measured as described by Mackness et al. (1998b). Serum was added to Tris-HCl buffer (100 mmol/l, pH 8·0) containing 2 mmol/l CaCl₂ and 5·5 mmol/l paraoxon (diethyl-p-nitrophenylphosphate; Sigma-Aldrich, Steinheim, Germany). The rate of p-nitrophenol generation during incubation at 25°C was monitored at 405 nm. One U of paraoxonase activity is equal to 1 nmol of p-nitrophenol formed/min per ml. Intra- and inter-assay variabilities were <5%.

Determination of the paraoxonase 192 genotype

The PON1-192 genotype was determined by restriction fragment length polymorphism–polymerase chain reaction (RFLP-PCR) using a previously published protocol (Humbert et al. 1993) with some minor modifications as described earlier (Bub et al. 2002). Each genotype was read by two independent observers.

Statistical analysis

Since genotyping revealed only two subjects to be homozygous for the R-allele, we merged the QR and the RR subjects to a new group QR/RR for further analysis. The cross-over trial was analysed using a standard method for the AB/BA design (Altman, 1991). As sequence effects were not significant, data from both intervention groups (group A: tomato/carrot juice, group B: carrot/tomato juice) were merged and analysed for PON genotype and vegetable juice consumption, respectively. Possible differences at baseline among treatment groups or among PON genotype groups were compared by unpaired t tests. Treatment effects were calculated as the difference between post- and pre-experimental values; within group treatment effects for tomato and carrot juice were tested by one-sample t test; between group treatment effects for tomato v. carrot juice were tested by unpaired t test. Comparison of PON genotypes within...
the tomato or carrot juice group were performed using ANOVA with ‘vegetable juice’ and ‘genotype’ as factor and the Tukey–Kramer post hoc test. All statistical calculations were performed with the StatView program (SAS Institute, Cary, NC, USA); significance level was set at P ≤ 0.05. Results are given as means with their standard deviations.

Results

All subjects complied with the study protocol and completed both dietary treatments. None of the volunteers reported any illness during the study. PON1-192 genotyping revealed seven subjects carrying the QQ-genotype, thirteen the QR-genotype and two the RR-genotype. The QR and the RR group have been merged and results are given for the QR/RR group (Table 1). At baseline, paraoxonase activity of PON was 123 (SD 29) and 217 (SD 44) U/l in QQ and QR/RR subjects, respectively (P < 0.001). Arylesterase activity was not significantly different in QQ and QR/RR subjects (156 (SD 26) and 144 (SD 19) U/ml). Tomato and carrot juice consumption had no statistically significant effects on arylesterase and paraoxonase activity of PON (Table 2). Additionally, tomato and carrot juice consumption had no effect on PON activity in either QQ or QR/RR subjects. We also observed no effects on total cholesterol, LDL-cholesterol or HDL-cholesterol in serum (data not shown).

Previous results from this study showed that tomato and carrot juice consumption had only marginal effects on lipid peroxidation in plasma (Briviba et al. 2004). Since PON can prevent lipid peroxidation depending on the PON genotype, we reanalysed our lipid peroxidation data with focus on the PON genotype. At baseline, lag time was significantly higher during LDL oxidation in QQ than in QR/RR subjects (111 (SD 9) v. 101 (SD 8) min; P < 0.05), while MDA in plasma was not significantly different in QQ and QR/RR subjects (0.44 (SD 0.12) v. 0.49 (SD 0.18) μM). Tomato and carrot juice consumption had no significant effect on lag time in QQ or QR/RR subjects (Fig. 1A). However, tomato juice consumption significantly reduced plasma MDA in QR/RR subjects (P < 0.05; Fig. 1B), while carrot juice had no significant effect on MDA irrespective of the PON1-192 genotype.

In plasma we also analysed lycopene and β-carotene, the major carotenoids in tomato juice and carrot juice, respectively. Results are given in Table 3. Tomato juice consumption significantly increased plasma lycopene concentrations (P < 0.001), while β-carotene in plasma did not change significantly. Carrot juice consumption increased β-carotene in plasma significantly (P < 0.001), but had no effect on lycopene plasma concentrations. Before carrot juice consumption plasma lycopene concentrations were lower in QR/RR subjects compared with QQ subjects (P = 0.04). However, after carrot juice consumption plasma lycopene significantly decreased in QQ subjects but not in the QR/RR group (P = 0.016). Although β-carotene plasma concentrations decreased during tomato juice consumption, these effects were not significant.

We also compared the data of washout phases at the beginning, during and after intervention (Table 4). There were no significant differences among the washout phases for lag time, plasma MDA, PON activities and β-carotene plasma concentrations. However, plasma lycopene concentrations were significantly lower (P < 0.05) at the end of the last washout phase.

Discussion

Supplementation of a low-carotenoid diet with either tomato or carrot juice for 2 weeks had no effect on the arylesterase and paraoxonase activity of PON in healthy young men. This is in line with a previous report which showed that tomato juice consumption over a period of 8 weeks had no effect on arylesterase activity in healthy elderly people (Bub et al. 2002). Other groups, however, have found that arylesterase activity increased after consumption of pomegranate juice (Aviram et al. 2000) and alcoholic beverages (van der Gaag & Biasi 1999) but decreased after used cooking fat (Sutherland et al. 1999).

The PON1-192 genotype predicts paraoxonase activity in human serum (Humbert et al. 1995). Our data on differences in basal paraoxonase activity are in line with this observation: low paraoxonase activity in QQ subjects, high activity in RR subjects and intermediate activity in QR heterozygous volunteers. We found no effect of tomato or carrot juice consumption on paraoxonase activity of PON in men regardless of PON1-192 genotype. Recently, it has been shown that a 5-week diet high in vegetables reduced paraoxonase activity in women, especially in those with the QR and RR genotype (Rantala et al. 2002). The authors discussed the possibility that changes in dietary fatty acid intake and changes in HDL-cholesterol may partly account for the observed effects. In our study HDL-cholesterol was unaffected, which may explain why PON activity was unchanged. Sex differences may also contribute to the observed discrepancies between our study

Table 1. Baseline characteristics of study participants based on treatment groups and on PON1 polymorphism†

<table>
<thead>
<tr>
<th></th>
<th>Total (n 22)</th>
<th>Group A† (n 11)</th>
<th>Group B (n 11)</th>
<th>QQ§ (n 7)</th>
<th>QR/RR (n 15)</th>
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<tr>
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<td>Mean</td>
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<td>Mean</td>
<td>SD</td>
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<tr>
<td>Age (years)</td>
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<td>30</td>
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<td>BMI (kg/m²)</td>
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<td>2</td>
<td>24</td>
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<td>Cholesterol (mmol/l)</td>
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<td>4.1</td>
<td>0.8</td>
<td>4.6</td>
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<tr>
<td>LDL-cholesterol (mmol/l)</td>
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<tr>
<td>HDL-cholesterol (mmol/l)</td>
<td>1.3</td>
<td>0.3</td>
<td>1.2</td>
<td>0.3</td>
<td>1.3</td>
</tr>
</tbody>
</table>

† There were no significant differences at baseline among groups A and B or QQ and QR/RR, respectively (unpaired t-test).
‡ Group A denotes the subgroup of subjects after randomisation with the intervention sequence tomato juice/carrot juice; group B consumed carrot juice followed by tomato juice.
§ QQ indicates homozygous subjects with glutamine at position 192 of the PON1 gene; QR/RR are heterozygous and homozygous, respectively, with arginine substituted for glutamine.
The differences in lag time before intervention may be partly attributable to differences in plasma carotenoid concentrations. Carotenoids are potent antioxidants and can inhibit lipid peroxidation (Krinsky 1998). High concentrations of carotenoids in plasma at baseline could protect LDL against oxidation. Lycopene and β-carotene in plasma tended to be lower in the QR/RR group than in QQ subjects, even before intervention, and this may possibly contribute to the lower lag time in the QR/RR group, plasma MDA was not significantly lower in the QR/RR group, although R-allele carriers show higher paraoxonase activity of PON1, lipid peroxide degradation is lower than in the QQ genotype (Mackness et al. 1998b). Although R-allele carriers show higher paraoxonase activity of PON1, lipid peroxide formation (Mackness et al. 1998b), which might explain why lag time was shorter in the QR/RR subjects, MDA, an end product of lipid peroxidation, circulates in plasma. One would expect an inverse association between lag time and plasma MDA. However, despite lower lag time in the QR/RR group, plasma MDA was not significantly higher in this group before intervention (QR/RR: 0.49 (SD 0.18); QQ: 0.44 (SD 0.12); P=0.51). It is therefore doubtful whether, under the given study conditions, there is any link between plasma MDA and LDL oxidation in healthy volunteers.

**Table 2. Effect of tomato and carrot juice consumption for 2 weeks on arylesterase and paraoxonase activities of human serum paraoxonase in healthy men with respect to PON1 genotype†**

(Mean values and standard deviations)

<table>
<thead>
<tr>
<th></th>
<th>Before intervention</th>
<th>After intervention</th>
</tr>
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<tbody>
<tr>
<td></td>
<td>Total (n 22)</td>
<td>QQ (n 7)</td>
</tr>
<tr>
<td></td>
<td>QQ§ (n 7)</td>
<td>QR/RR (n 15)</td>
</tr>
<tr>
<td></td>
<td>Mean</td>
<td>SD</td>
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<tr>
<td>Tomato juice</td>
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<td></td>
</tr>
<tr>
<td>Arylesterase (U/ml)</td>
<td>147</td>
<td>25</td>
</tr>
<tr>
<td>Paraoxonase (U/l)</td>
<td>185</td>
<td>60</td>
</tr>
<tr>
<td>Carrot juice</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Arylesterase (U/ml)</td>
<td>146</td>
<td>23</td>
</tr>
<tr>
<td>Paraoxonase (U/l)</td>
<td>190</td>
<td>62</td>
</tr>
</tbody>
</table>

† P¿0.001. Subjects carrying the R-allele (QR and RR) have a significantly higher paraoxonase activity as compared to the QQ wild-type (unpaired t test) before and after intervention, respectively.

‡ There were no significant differences at baseline among the tomato juice group and the carrot juice group for arylesterase and paraoxonase activity, respectively (unpaired t test). Intervention with either juice had no significant effect on enzyme activities. Intervention effects were calculated by subtracting pre- from post-experimental values and tested by one-sample t test within each juice intervention group. Unpaired t test was applied to compare treatment effects induced by tomato juice and carrot juice.

§ PON1 activity has been determined by using phenylacetate (arylesterase) and paraoxon (paraoxonase) as substrates, respectively. One U of arylesterase activity is equal to 1 μmol phenylacetate hydrolysed/min per ml. One U of paraoxonase activity is equal to 1 nmol paraoxon hydrolysed/min per ml.

†† QQ indicates homozygous subjects with glutamine at position 192 of the PON1 gene; QR/RR are heterozygous and homozygous, respectively, with arginine substituted for glutamine.

with men and the study by Rantala with women. Paraoxonase activity is higher in women than in men on a habitual diet (Kleemola et al. 2002). The authors found a significant negative correlation between paraoxonase activity and vegetable, fruit and berry intake in women only and they discussed the hormonal status which they thought might be responsible for the observed sex differences. Unfortunately, they did not determine the PON1-192 genotype which is a predictor of paraoxonase activity (Humbert et al. 1993), and could also account for the observed gender-related differences.

Consumption of tomato products reduces lipid peroxidation in healthy volunteers (Agarwal & Rao, 1998; Bub et al. 2000; Visioli et al. 2003). However, in the present study tomato and carrot juice consumption had only marginal effects on lipid peroxidation in plasma (Briviba et al. 2004). We therefore reanalysed our data focusing on the respective PON genotypes. At baseline we found that lag time was shorter in QR/RR subjects than in QQ subjects, thus LDL oxidation in the QR/RR group was higher. HDL-associated PON1 protects LDL from oxidation as seen by reduced lipid peroxide formation (Mackness et al. 1998b). Although R-allele carriers show higher paraoxonase activity of PON1, lipid peroxide degradation is lower than in the QQ genotype (Mackness et al. 2000b), which might explain why lag time was shorter in the QR/RR subjects. MDA, an end product of lipid peroxidation, circulates in plasma. One would expect an inverse association between lag time and plasma MDA. However, despite lower lag time in the QR/RR group, plasma MDA was not significantly higher in this group before intervention (QR/RR: 0.49 (SD 0.18); QQ: 0.44 (SD 0.12); P=0.51). It is therefore doubtful whether, under the given study conditions, there is any link between plasma MDA and LDL oxidation in healthy volunteers.

Fig. 1. Changes in lag time (A) as a measure of LDL oxidation, and plasma malondialdehyde (MDA) (B) in men after drinking 330 ml tomato or carrot juice for 2 weeks, respectively. Lag time reflects the antioxidant status of LDL and is determined by monitoring conjugated diene formation during copper-induced LDL oxidation. It is defined as the time from reaction start to the beginning of the rapid increase in diene production (propagation phase). Values are means with their standard deviations represented by vertical bars, n 7 for the QQ subjects (III) and n 15 for the QR/RR group (III). A significant reduction in plasma MDA was detected after tomato juice consumption (ANOVA, Tukey-Kramer test, P¿0.05). QQ indicates homozygous subjects with glutamine at position 192 of the PON1 gene. QR/RR are heterozygous and homozygous, respectively, with arginine substituted for glutamine.
RR group. However, we cannot rule out the possibility that other factors, e.g. vitamin E, may be responsible for the lower lag time in R-allele carriers. Vitamin E plays a crucial role in the protection of LDL from oxidation and the pathogenesis and progression of atherosclerosis (Salonen, 2002; Upston et al. 2003). Unfortunately, we did not determine vitamin E plasma concentrations in our volunteers. However, in a previous study supplementation with identical juice preparations had no significant effect on plasma vitamin E of healthy men (Müller et al. 2003). We cannot rule out the possibility that other factors, e.g. vitamin E, may be responsible for the lower lag time in R-allele carriers. Vitamin E plays a crucial role in the protection of LDL from oxidation and the pathogenesis and progression of atherosclerosis (Salonen, 2002; Upston et al. 2003). Unfortunately, we did not determine vitamin E plasma concentrations in our volunteers. However, in a previous study supplementation with identical juice preparations had no significant effect on plasma vitamin E of healthy men (Müller et al. 2003). Therefore, it is possible that the difference in lag time between QQ and RR group is due to other factors than PON1 polymorphism, such as age and gender possibly contributing to the observed genotype-related differences.

So far little is known about the impact of dietary intervention on lipid peroxidation in healthy volunteers when PON polymorphisms are considered. In a study with elderly subjects (Bub et al. 2002), recommending a low-carotenoid diet is essential to lower plasma carotenoid concentrations, with effects on plasma MDA in either QQ or QR/RR subjects, despite the fact that it led to plasma lycopene concentrations following tomato juice consumption. Carrot juice intake and subsequent high β-carotene plasma concentrations per se may not be sufficient to protect LDL from oxidation in healthy men. Additionally, the copper-induced ex vivo oxidation of LDL used in this study may not be an adequate measure of LDL oxidizability since it does not reflect physiological in vivo conditions and may mask possible protective effects attributable to carrot juice consumption. During the entire study the volunteers were kept on a low-carotenoid diet which has been shown to lower plasma carotenoid concentrations after vitamin E plasma concentrations were no significant differences in lag time between the two groups. The relatively short intervention period (2 weeks rather than 2 months in the study with elderly subjects) and the small number of volunteers (twenty-two rather than twenty-nine) may explain the absence of an effect on lag time in this study. Carrot juice consumption had no effect on lag time or on plasma MDA in either QQ or QR/RR subjects, despite the fact that it led to plasma β-carotene concentrations which were twice as high as plasma lycopene concentrations following tomato juice consumption. Carrot juice intake and subsequent high β-carotene plasma concentrations per se may not be sufficient to protect LDL from oxidation in healthy men. Additionally, the copper-induced ex vivo oxidation of LDL used in this study may not be an adequate measure of LDL oxidizability since it does not reflect physiological in vivo conditions and may mask possible protective effects attributable to carrot juice consumption. During the entire study the volunteers were kept on a low-carotenoid diet which has been shown to lower plasma carotenoid concentrations after 10 weeks (Watzl et al. 2003). Depletion of carotenoids (and probably other nutrients from fruit and vegetables) may induce ’stress’ which might not be completely
alleviated by supplementation with carrot and tomato juice. This could explain the absence of effects on lag time and MDA after carrot juice consumption. However, despite a 10-week-long low-carotenoid diet we found no increase in lipid peroxidation measurements and no negative effect on immune functions.

In order to minimize seasonal effects, we used a crossover study design with washout phases of 2-week duration, before, between and after the intervention periods. Comparing the three washout phases we found no differences in lag phase, MDA or PON1 activities. This shows that our study design was effective in achieving similar conditions for each intervention and that the low-carotenoid diet had no effect on these measurements. However, lycopene plasma concentrations were reduced after the last washout phase, probably due to the low-carotenoid diet and the fact that there are few sources of lycopene in food. Plasma β-carotene concentrations did not significantly decrease during the study. The widespread occurrence of β-carotene in food means that even on a low-carotenoid diet a constant intake of small amounts of β-carotene is probably unavoidable, which could explain why the decrease in β-carotene after the last washout phase was non-significant.

In conclusion, male volunteers with the QR/RR genotype showed increased lipid peroxidation under baseline conditions. Although tomato and carrot juice failed to affect PON1 activity, tomato juice did reduce lipid peroxidation in volunteers with the PON1-192 R-allele, and could therefore play a useful role in explaining both activities.

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