Plasma carotenoid concentrations in relation to acute respiratory infections in elderly people

Judith M. van der Horst-Graat, Frans J. Kok and Evert G. Schouten*

Division of Human Nutrition and Epidemiology, Wageningen University, Bomenweg 2, P.O. Box 8129, 6703 HD, Wageningen, The Netherlands

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A high plasma carotenoid concentration could improve the immune response and result in decreased risk of infectious diseases. However, data on the relationship of plasma carotenoid concentration with acute respiratory infections, which occur frequently in elderly people, are scarce. We investigated, therefore, the relationship of plasma concentrations of six major carotenoids (β-carotene, α-carotene, β-cryptoxanthin, lycopene, lutein and zeaxanthin) with the incidence and severity of acute respiratory infections. Baseline data from an intervention trial were used. Participants were 652 non-institutionalized elderly people (≥ 60 years old) enrolled via two community-based sampling strategies in the Wageningen area of The Netherlands in 1998–99. Plasma carotenoid concentrations were divided into quartiles, the lowest being the reference. Frequency and severity of episodes during the previous 1 year, i.e. staying in bed, medical consultation and episode-related medication, were self-reported by means of a questionnaire. On average 1·6 episodes per person were recorded. The incidence rate ratio of acute respiratory infections at high β-carotene status was 0·71 (95% CI 0·54–0·92) as compared with the low β-carotene concentration group. No association was observed between β-carotene and illness severity. α-Carotene, β-cryptoxanthin, lycopene, lutein and zeaxanthin were not related to incidence or severity of the infections. We conclude that elderly people with a high plasma β-carotene concentration may have a lower occurrence of acute respiratory infections.

Acute respiratory infection: Carotenoids: Elderly people

Elderly people are at high risk of morbidity from infections, especially from respiratory infections, which occur frequently (Miller, 1996). On average, community-dwelling elderly people suffer from one to two acute respiratory infections per year (Nicholson et al. 1997; Graat et al. 2002). β-Carotene, α-carotene, β-cryptoxanthin, zeaxanthin, lycopene and lutein comprise approximately 90% of the total plasma pool of carotenoids (Thurnham, 1994; Vogel et al. 1997). β-Carotene has been studied extensively in relation to immune response, whereas for other carotenoids such information is scarce (Jyonouchi et al. 1996; Hughes et al. 2000; Corridan et al. 2001). Some studies have shown an enhanced immune response at high plasma β-carotene status or after β-carotene supplementation (Watson et al. 1991; Fuller et al. 1992; van-Poppel et al. 1993; Murata et al. 1994; Santos et al. 1996; Hughes, 2001), whereas others have not (Ringer et al. 1991; Daudu et al. 1994; Murata et al. 1994; Santos et al. 1997; Corridan et al. 2001). Only one study has investigated acute respiratory infections, i.e. common cold incidence, in relation to both dietary β-carotene intake and supplementation in male smokers. High intake was associated with a slightly higher cold incidence, whereas supplementation had no effect on incidence (Hemila et al. 2002). The antioxidant properties and modulating effect on prostaglandin E_2 production of β-carotene might explain its possible beneficial effects (Khachik et al. 1995; Hughes, 2001).

Although clinical endpoints such as infectious diseases have much greater public health relevance than immune function, they have hardly been studied until now. The aim of our present study was to investigate the relationship between plasma carotenoid concentrations and acute respiratory infections during 1997–98 in elderly persons. We investigated the incidence of illness and severity in relation to β-carotene, α-carotene, β-cryptoxanthin, lycopene, lutein and zeaxanthin.

Methods

Subjects

Subjects were approached to participate in a 15-month double-blind intervention trial investigating the effect of daily micronutrients on acute respiratory infections. According to the 2 × 2 factorial design, participants were randomly assigned to the following groups using blocked randomization: 200 mg dl-α-tocopheryl acetate; a physiological dose of multivitamins–minerals; 200 mg dl-α-tocopheryl acetate plus a physiological dose of multivitamins–minerals;
placebo. The present study has been described in detail by Graat et al. (2002).

We used the baseline information of our present study population for this retrospective study concerning carotenoid status and respiratory infections in the previous 1 year. The study population included 652 participants: 325 men and 327 women. All persons were ≥60 years old, with an average age of 73 years. Only 2% of the subjects lived in homes for the aged. We therefore consider that our present study population were non-institutionalized. Participants had no history of cancer, liver disease or fat malabsorption in the 5 years before enrolment. A questionnaire was used to ask for nutritional supplementation during the previous year. Written informed consent was obtained from all participants before participation. The medical ethics committee of Wageningen University, The Netherlands, approved the research protocol.

Respiratory infections
The main outcomes were incidence and severity of acute respiratory infections during the previous year. At the time of blood collection, which was between 1 September 1998 and 23 March 1999, a detailed questionnaire about respiratory infections during the previous 12 months was filled out by all participants, with the help of a research assistant. Therefore, the recorded infections took place in 1997–98. Participants were asked about the frequency of common cold, influenza, pneumonia, sore throat and pain in facial sinuses. These different manifestations of infection were combined and simply referred to as ‘acute respiratory infection’ for two reasons: (1) it is difficult to distinguish between upper and lower respiratory infections, because the symptom patterns can be indistinguishable (Tannock et al. 1993; Nicholson et al. 1997; Carrat et al. 1999); (2) lower respiratory tract symptoms were reported to complicate 65% of upper respiratory tract infections (Nicholson et al. 1997). For each past episode of acute respiratory infection, information on illness severity, i.e. staying in bed (in bed for several hours because of infection-related symptoms at a time that one would not usually stay in bed), medical consultation (visit a general practitioner or other physician because of infection-related symptoms) and use of medication (any medication, including antibiotics, prescribed by a general practitioner or other physician), were also recorded in the questionnaire.

Analysis of carotenoids
Between 1 September 1998 and 23 March 1999 blood samples were drawn to determine the plasma concentrations of cis-β-carotene, trans-β-carotene, α-carotene, β-cryptoxanthin, lycopene, lutein and zeaxanthin. The β-carotene concentration was calculated from the sum of cis-β-carotene and trans-β-carotene values. Concentrations of retinol, α-tocopherol and ascorbic acid were also determined. Samples were collected between 08.30 and 11.00 hours. A light breakfast, without salads, fruits or fruit juices, was allowed before sampling. None of the subjects had been taking any nutritional supplementation in the 2 months before blood drawing. Plasma was immediately stored on ice in a closed box, and, within 6 h of blood collection, stored at −80°C. The reversed-phase HPLC method was used to analyse fat-soluble vitamin concentrations (Hess et al. 1991; Aebischer et al. 1999). Ascorbic acid concentration was obtained via standard procedures and assessed by fluorimetric assay (Vuilleumier & Keck, 1989).

Statistical analyses
Participants were divided into quartiles of plasma carotenoid concentrations. The two middle quartiles, i.e. 25–75th, were collapsed and referred to as ‘intermediate carotenoid status’. The low-carotenoid class was taken as the reference group. Frequencies including percentages were calculated for categorical data and were compared by χ² test or Fisher’s exact test. Continuous variables were compared by ANOVA for unbalanced data by the SAS procedure GLM (SAS version 8; SAS Institute, Cary, NC, USA) and were expressed as mean values and standard deviations.

To calculate incidence rate ratios of the infections, a Poisson regression model was used with the number of episodes as the dependent variable and the carotenoid status as the independent variable included in the model. Because some carotenoids possess pro-vitamin A activity, the relationship between vitamin A (retinol) and incidence of the infections was assessed.

Because we assumed that all carotenoids could have the same beneficial effect on the infections, all carotenoids were combined in a Poisson regression model to investigate whether the relative risk of such combination differed from the results of β-carotene only.

Logistic regression was used to calculate adjusted odds ratios and the corresponding 95% CI for the severity outcomes. Illness severity was assessed in participants who experienced at least one acute respiratory infection.

The variables age, BMI, self-rated health (score 1–10, 10 indicating highest self-rated health), retinol, α-tocopherol and ascorbic acid status, gender, chronic obstructive pulmonary disease, asthma, influenza vaccination, history of nutritional supplementation and smoking (current, former, never) were evaluated for confounding and effect modification. If variables were related to the carotenoid status and to the incidence of infection, they were entered into the model. Depending on the carotenoid investigated, gender, age, BMI, self-rated health and chronic obstructive pulmonary disease turned out to be confounders. Inclusion of age or gender into the models precluded adjustment for BMI, because of multicollinearity between gender or age and BMI. No effect modifiers turned out to influence the relationships.

Results
Population characteristics and plasma (pro-) vitamin concentrations of participants at high, intermediate and low β-carotene status are described in Table 1. Similar characteristics were observed for the α-carotene, β-cryptoxanthin, lycopene, lutein and zeaxanthin, and are therefore not shown. The correlation between β-carotene and α-carotene was 0.78 and between zeaxanthin and
Correlations between the remaining carotenoids ranged from 0.19 to 0.48.

Of the 652 participants, 5.2% chronically used multivitamin–mineral supplementation, 0.2% vitamin A and vitamin A plus D, 3.5% vitamin C and 1.2% vitamin E. The last 2 months of the observation period (2 months before blood-drawing) all participants refrained from any nutritional supplementation because of the upcoming intervention.

In total, 417 participants reported 754 events of common cold, 118 events of influenza, 108 events of sore throat, 43 events of pain in facial sinuses and 30 events of pneumonia. These events were combined to 1053 acute respiratory infections, which was the outcome in our present analysis.

In 26% (n = 235) of subjects there were no reports of respiratory infection during the previous 1 year. On average, 1.6 episodes were recorded per person per year.

A statistically significant inverse relationship was observed between β-carotene status and the incidence rate, but not severity, of the infections (Table 2). We observed a tendency for a similar relationship between the incidence rate of respiratory infections and α-carotene status and β-cryptoxanthin status. The incidence rate ratios at high and intermediate concentrations were 0.79 (95% CI 0.60–1.02) and 0.74 (95% CI 0.60–1.01) for α-carotene and 0.83 (95% CI 0.64–1.07) and 0.80 (95% CI 0.63–0.97) for β-cryptoxanthin. When combinations of β-carotene, α-carotene and β-cryptoxanthin were entered into one model, β-carotene turned out to have the greatest

### Table 1. Population characteristics and plasma concentrations of carotenoids and vitamins in 652 elderly Dutch subjects by β-carotene concentration*

<table>
<thead>
<tr>
<th>β-Carotene concentration</th>
<th>High (n = 162)</th>
<th>Intermediate (n = 325)</th>
<th>Low (n = 165)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Population characteristics</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Men (%)</td>
<td>37.0</td>
<td>47.1</td>
<td>47.9</td>
</tr>
<tr>
<td>Age (years)</td>
<td>74.0</td>
<td>73.6</td>
<td>72.2</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>25.9</td>
<td>27.5</td>
<td>28.4</td>
</tr>
<tr>
<td>Self-rated health (score 1 (lowest)–10)</td>
<td>7.6</td>
<td>7.5</td>
<td>7.2</td>
</tr>
<tr>
<td>Chronic obstructive pulmonary disease (%)</td>
<td>4.3</td>
<td>9.5</td>
<td>17.0</td>
</tr>
<tr>
<td>Asthma (%)</td>
<td>1.9</td>
<td>2.8</td>
<td>1.8</td>
</tr>
<tr>
<td>Allergy (%)†</td>
<td>17.3</td>
<td>20.0</td>
<td>18.2</td>
</tr>
<tr>
<td>Current smoker (%)</td>
<td>8.0</td>
<td>9.9</td>
<td>14.6</td>
</tr>
<tr>
<td>Influenza vaccination (%)</td>
<td>75.2</td>
<td>74.9</td>
<td>75.2</td>
</tr>
<tr>
<td>Plasma vitamins (μmol/l)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>β-Carotene</td>
<td>0.83</td>
<td>0.40</td>
<td>0.18</td>
</tr>
<tr>
<td>α-Carotene</td>
<td>0.15</td>
<td>0.07</td>
<td>0.04</td>
</tr>
<tr>
<td>β-Cryptoxanthin</td>
<td>0.43</td>
<td>0.28</td>
<td>0.19</td>
</tr>
<tr>
<td>Lycopene</td>
<td>0.43</td>
<td>0.25</td>
<td>0.21</td>
</tr>
<tr>
<td>Lutein</td>
<td>0.32</td>
<td>0.06</td>
<td>0.05</td>
</tr>
<tr>
<td>Zeaxanthin</td>
<td>0.07</td>
<td>0.23</td>
<td>0.21</td>
</tr>
<tr>
<td>Retinol</td>
<td>1.98</td>
<td>2.03</td>
<td>2.01</td>
</tr>
<tr>
<td>α-Tocopherol</td>
<td>31.4</td>
<td>29.4</td>
<td>26.9</td>
</tr>
<tr>
<td>Ascorbic acid</td>
<td>55.0</td>
<td>49.2</td>
<td>44.8</td>
</tr>
</tbody>
</table>

* For details of procedures, see p. 114.
† Allergy against pollen grains, domestic pets or house-dust.

### Table 2. Incidence and severity of acute respiratory infections according to plasma β-carotene concentration in elderly Dutch subjects from 1997 to 1998*

<table>
<thead>
<tr>
<th>Outcomes†</th>
<th>n</th>
<th>OR</th>
<th>95% CI</th>
<th>OR</th>
<th>95% CI</th>
<th>OR</th>
<th>P for trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>Incidence rate (per person per year)</td>
<td>652</td>
<td>1.66</td>
<td>1.73</td>
<td>2.34</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Incidence rate ratio</td>
<td>0.71</td>
<td>0.54</td>
<td>0.92</td>
<td>0.74</td>
<td>0.60</td>
<td>0.91</td>
<td></td>
</tr>
<tr>
<td>Staying in bed‡</td>
<td>417</td>
<td>0.74</td>
<td>0.38</td>
<td>1.44</td>
<td>1.20</td>
<td>0.71</td>
<td></td>
</tr>
<tr>
<td>Medical consultation‡</td>
<td>417</td>
<td>1.27</td>
<td>0.62</td>
<td>2.61</td>
<td>1.44</td>
<td>0.79</td>
<td></td>
</tr>
<tr>
<td>Episode-related medication‡</td>
<td>417</td>
<td>1.33</td>
<td>0.73</td>
<td>2.43</td>
<td>1.60</td>
<td>0.96</td>
<td></td>
</tr>
</tbody>
</table>

OR, Odds ratio.
† All outcomes were corrected for gender, age, self-rated health and chronic obstructive pulmonary disease.
‡ Adjusted OR (95% CI) of variables indicating infection severity, i.e. fever, staying in bed, medical consultation and use of medication were assessed in participants who experienced at least one respiratory infection (103 subjects in the high-, 206 in the intermediate- and 108 in the low-β-carotene group).
Discussion

In our large observational analysis in elderly people, a high plasma β-carotene status was associated with a lower incidence of acute respiratory infections, but not with a lower illness severity. No relationship was observed between plasma α-carotene, β-cryptoxanthin, lycopene, lutein and zeaxanthin status and the incidence and severity of the infections.

In the following discussion, we will address the possible threats to the internal validity in our present study. Error in the assessment of both exposure and outcome may lead to information bias. Participants and investigators were unaware of the carotenoid status at the time of outcome assessment, and the incidence rate ratio was 1.00 (0.77–1.30) for high v. low retinol status and 0.98 (95% CI 0.79–1.24) for intermediate v. low retinol status.

No significant results. Additional analysis to investigate the relationship between retinol (vitamin A) and the incidence of acute respiratory infections showed that the incidence rate ratio was 1.00 (0.77–1.30) for high v. low retinol status and 0.98 (95% CI 0.79–1.24) for intermediate v. low retinol status.

β-Carotene is a precursor of retinol. No clear favourable effects of vitamin A supplementation on infectious disease or immune response have been observed in elderly people in developed countries, which is in accordance with our present results (Murphy et al. 1992; Fortes et al. 1998). Beneficial effects of vitamin A supplementation on infectious diseases were predominantly observed in children of developing countries; these children were sometimes marginally deficient in vitamin A (Semba, 1994). Although we cannot exclude the possibility that β-carotene is inversely related to respiratory illness by influencing plasma retinol concentration, we suppose that this mechanism is less plausible.

We did not observe relationships between α-carotene, β-cryptoxanthin, lycopene, lutein and zeaxanthin, and acute respiratory infections. α-Carotene and β-cryptoxanthin have never been reported in relation to incidence or severity of acute respiratory infections, nor in relation to human immune response. Literature concerning lycopene, lutein and zeaxanthin, and immune function is scarce. No clear associations of lycopene, lutein and zeaxanthin were observed with blood monocytes and T-helper cell activities (Iynouchi et al. 1996; Hughes et al. 2000; Corridan et al. 2001). Correspondingly, no effect of tomato juice, with lycopene being the predominant carotenoid, on cell-mediated immunity was shown (Watzl et al. 2000). Those findings support our null-findings on respiratory infections. In contrast, Watzl et al. (1999) showed improved T-lymphocyte function by tomato juice consumption in subjects consuming a diet low in carotenoids.

One may question whether information on plasma carotenoid concentration can be extrapolated to the intake of certain foods. Some studies have shown a significant relationship between dietary intake of fruits and vegetables and plasma carotenoid concentration (Rook et al. 1997; Broekmans et al. 2000), whereas others did not
References


Santos MS, Leka LS, Ribaya-Mercado JD, Russell RM, Meydani M, Hennekens CH, Gaziano JM & Meydani SN...


