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Serum response to supplemental macular carotenoids in subjects with and without age-related macular degeneration

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Abstract

Macular pigment (MP) is composed of lutein (L), zeaxanthin (Z) and meso-zeaxanthin (MZ). The present study reports on serum response to three different MP supplements in normal subjects (n=27) and in subjects with age-related macular degeneration (AMD) (n=27). Subjects were randomly assigned to: Group 1 (20 mg L and 2 mg Z), Group 2 (10 mg L, 2 mg Z and 10 mg MZ) or Group 3 (3 mg L, 2 mg Z and 17 mg MZ). Serum carotenoids were quantified at baseline, and at 4 and 8 weeks using HPLC. Response data for normal and AMD subjects were comparable and therefore combined for analysis. We report response as the average of the 4- and 8-week concentrations (saturation plateau). Serum L increased significantly in Group 1 (0.036 μmol/l per mg (269%); P<0.001) and Group 2 (0.079 μmol/l per mg (340%); P<0.001), with no significant change in Group 3 (0.006 μmol/l per mg (7%); P=0.466). Serum Z increased significantly in Group 1 (0.037 μmol/l per mg (69%); P<0.001) and Group 2 (0.015 μmol/l per mg (75%); P<0.001), with no significant change in Group 3 (0.0002 μmol/l per mg (−6%); P=0.384). Serum MZ increased significantly in Group 1 (0.0094 μmol/l (absolute value); P=0.015), Group 2 (0.005 μmol/l per mg; P<0.001) and Group 3 (0.004 μmol/l per mg; P<0.001). The formulation containing all three macular carotenoids (Group 2 supplement) was the most efficacious in terms of achieving the highest combined concentration of the three MP constituent carotenoids in serum, thereby potentially optimising the bioavailability of these compounds for capture by the target tissue (retina).

Key words: Age-related macular degeneration: Macular carotenoids: Lutein: Zeaxanthin: meso-Zeaxanthin: HPLC separation

Carotenoids are a class of >700 tetraterpenoid compounds found in nature. These plant pigments contribute to a plethora of biological functions, due to their unique chemical features14, where they play important roles in both plants (e.g. the regulation of light in oxygenic photosynthesis15) and animals (e.g. precursors in the formation of vitamin A16). Carotenoids contain a conjugated system of double bonds, known as a polyene backbone, which is capped with two end groups. The backbone is responsible for their respective photochemical properties (short-wavelength light absorption) and chemical reactivity (antioxidant capacity), whereas the carotenoids are identified and defined primarily by their respective end groups. One or both end groups can undergo cyclisation as well as substitution with oxygen-containing groups (i.e. keto, hydroxy or epoxy groups), which is why there is such a large variety of carotenoids in nature. Oxygen-containing carotenoids are referred to as xanthophylls (e.g. lutein (L), zeaxanthin (Z) and meso-zeaxanthin (MZ)), while the true hydrocarbon carotenoids are referred to as carotenes (e.g. α- and β-carotene) (Fig. 1).

L, Z and MZ are the predominant xanthophyll carotenoids found in the macula17, the central part of the retina responsible for fine detail vision18. The macula accumulates these yellow carotenoids and excludes all the other thirty or so circulating carotenoids19, and are collectively known as macular pigments (MP) in this specialised tissue20.

Abbreviations: AMD, age-related macular degeneration; IS, internal standard; L, lutein; MP, macular pigment; MZ, meso-zeaxanthin; Z, zeaxanthin.

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There is a consensus that the macular carotenoids play an important role in protecting the macula from the damaging effects of reactive oxygen species. Reactive oxygen species, which can exist as either free radicals or non-radical subgroups, are produced as a result of oxygen metabolism, and their generation is increased when the tissue is exposed to short-wavelength (blue) light and to other environmental (pollution) and lifestyle (smoking) factors. Interestingly, the polyene chains of L, Z and MZ have the ability to quench reactive oxygen species and absorb short-wavelength (blue) light. Of interest, a study by Li et al. reported that a mixture of L, Z and MZ in a ratio of 1:1:1 (the ratio of these carotenoids typically seen in normal MP) can quench more singlet oxygen than the individual carotenoids at the same total concentration, and may explain the exquisite biological selectivity and spatial distribution of these pigments within this specialised retinal tissue.

The aforementioned properties of the macular carotenoids are believed to confer protection against age-related macular degeneration (AMD), the most common cause of blind registration in the developed world. Also, the optical (short-wavelength filtering) properties of the macular carotenoids suggest that they play a role in visual function, by reducing the effects of chromatic aberration (and therefore improving image quality) and light scatter (and therefore reducing the symptoms of glare).

L and Z cannot be synthesised de novo in mammals and must be obtained from the diet. L and Z are found in common foodstuffs such as fruits (e.g. kiwi) and vegetables (e.g. spinach and maize), whereas MZ has not been identified in these foods, although it is important to point out that there has been no published study on MZ concentrations in foods of a typical diet (including fruits and vegetables) as yet. However, Maoka et al. have shown MZ to be present in some unusual foods, such as fish skin and turtle fat. Interestingly, MZ accounts for one-third of MP at the macula, and simian experiments suggest that it is produced by isomerisation of L.

Many studies have reported serum responses to supplemental L and Z, but only three trials have reported on responses to supplemental MZ, as this carotenoid was only identified as being present at the macula in 1993. However, a recent clinical trial by Connolly et al. reported a rapid serum and MP response to a supplement containing all three macular carotenoids and, importantly, none of these trials reported any adverse effects associated with consumption of the macular carotenoids.

The present study was designed to investigate serum carotenoid responses to supplements containing at least two of the three macular carotenoids (i.e. a macular carotenoid supplement comparison study) in subjects with and without AMD.

Methods

Subjects

This was a randomised and double-blind study. All subjects signed an informed consent document and the experimental measures conformed to the Declaration of Helsinki. The study was reviewed and approved by the Research Ethics Committees, South East Region, Waterford Regional Hospital, and the Ethics Committee of the Waterford Institute of Technology, Waterford, Ireland.

We were interested in studying two different subject populations, those with and without AMD. These two populations were recruited under the following criteria. Normal subjects were those with no ocular pathology and in good health. Subjects suffering from AMD were defined as those with signs of early AMD, exhibiting drusen and pigmentedary changes. The AMD subjects were identified at a pre-project enrolment and screening visit, conducted by an ophthalmologist with a special interest in retinal disease and experienced in the classification of AMD for research purposes. Exclusion criteria comprised past or present use of supplemental macular carotenoids and/or pregnancy. Subject BMI was calculated (kg/m²). Height (m) was measured using a Leicester Height Measure and weight (kg) was measured using SECA weighing scales (SECA). Smoking status was identified as one of three positions: current smoker, ex-smoker and never smoker. A subject’s weekly intake of carotenoid-rich foods (eggs, broccoli, maize and dark leafy vegetables) were inputted into the ‘L/Z screener’ to give a carotenoid diet ‘score’. Values were weighted for frequency of intake of the food and for the bioavailability of L and Z within these foods, and a ranking score reflecting the relative intakes was generated. The range of scores from the L/Z screener is 0 to 75. After adding foods with known concentrations of L and Z into the screener, the following estimates were made. Low dietary carotenoid intake score is from 0 to 15 (i.e. ≤2 mg/d); medium dietary carotenoid intake score is from 16 to 30 (i.e. between 2 and 13 mg/d); and high dietary carotenoid intake score is from 31 to 75 (i.e. >13 mg/d).

Originally, seventy-two subjects were enrolled into the study. However, as the use of supplemental macular
The macular carotenoids was an exclusion criterion, and following analysis of baseline serum data (performed only after final study visit), we noted that eight subjects had substantial amounts of MZ in their serum. Given that these subjects suffered from AMD (and were based in the Republic of Ireland, where a supplement containing MZ is widely available for patients with AMD) we suspected that they had in fact been supplementing with the macular carotenoids, but had failed to disclose this fact at enrolment.

This was subsequently confirmed by a phone call to each of these volunteers, and data relating to these subjects were excluded from all analyses. Of the remaining sixty-four subjects, ten (one AMD, nine normal) did not attend all three study visits, and were therefore also excluded from the analysis, thus leaving twenty-one, twenty and thirteen subjects in the carotenoid intervention Groups 1, 2 and 3, respectively (see later).

Of these fifty-four subjects, twenty-seven had no ocular pathology (normal subjects) and twenty-seven had previously been diagnosed with AMD (AMD subjects). The fifty-four subjects were split into three different carotenoid intervention groups as follows: Group 1: (n 21; eleven normal, ten AMD) 20 mg of L and 2 mg of Z (Ultra Lutein™, provided by Nature's Plus, Natural Organics, Inc.); Group 2: (n 20; ten normal, ten AMD) 10 mg L, 2 mg Z and 10 mg MZ (Macushield™, provided by MacuVision Europe Limited); Group 3: (n 13; six normal, seven AMD) 3 mg L, 2 mg Z and 17 mg MZ (customised MZ formulation provided by Industrial Organica (not available commercially)). All supplements used in the present study consisted of oil-suspended, unesterified carotenoids provided in gelatine capsules. Each subject used in the present study consisted of oil-suspended, unesterified carotenoids provided in gelatine capsules. Each subject was required to consume one capsule daily, with the main meal, for the duration of the 8-week study period, with serum samples taken at baseline, and at 4 and 8 weeks. Significant efforts were made to ensure compliance to the study intervention. Compliance was monitored closely at the bi-weekly study visits. In addition, subjects were requested to return their supplement packs at their exit visit, and compliance was checked by tablet counting at this visit.

**Standards and solvents**

DSM Nutritional Products supplied the L and Z reference standards. The MZ standard was supplied by Industrial Organica as a soybean oil oleoresin. The internal standard (IS) α-tocopheryl acetate and all solvents (HPLC grade) used for extraction and HPLC analysis were supplied by Sigma-Aldrich.

**Serum carotenoid extraction**

Non-fasting blood samples were collected in 9 ml vacuette tubes containing a ‘Z Serum Sep Clot Activator’, usually in the morning when subjects arrived at the clinic. The blood samples were allowed to clot at room temperature for approximately 1 h and then centrifuged at 725 × g for 15 min in a Gruppe GC 12 centrifuge (Desaga Sarstedt) to separate the serum from the whole blood. The resulting serum samples were stored at −70°C until the time of extraction (maximum 12 months).

Serum (0.4 ml) was micropipetted into clear 1.5 ml Eppendorf tubes labelled according to subject and visit number. IS (0.2 ml), α-tocopheryl acetate (250 mg/l ethanol) and 0.3 ml of butylated hydroxytoluene (250 mg/l ethanol) were added and extracted into 0.5 ml of heptane using a Vortex Genie-2 (Scientific Industries) at the highest setting for 2 min, followed by centrifugation with an AccuSpin Micro 17 (Fisher Scientific Ireland) for 5 min at 4000 × g.

An aliquot of the upper heptane layer (0.4 ml) was removed to a light-resistant Eppendorf tube, and the heptane extraction was repeated once more, adding a further 0.5 ml of heptane to the original residue. The combined extracts were dried under N2 and stored at −70°C until analysis.

**HPLC analysis of serum L and total zeaxanthin (total Z) (Assay 1)**

The HPLC system used for the study was an Agilent 1200 Series (Agilent Technologies Limited) consisting of a quaternary pump, autosampler, thermostat column compartment and a photodiode array detector monitoring a wavelength of 450 nm for serum carotenoids and 292 nm for the IS. Sample analysis was carried out in order of subject number and time of visitation. In other words, subjects were batch assessed.

The dried samples were reconstituted in 0.2 ml of the isocratic mobile phase, vortexed at the lowest setting for 1 min and pipetted into 2.5 ml vials containing 0.35 ml glass inserts (Agilent Technologies Limited). The sample (0.1 ml) was injected via autosampler onto a Phenomenex Ultradic ODS 3μ C18 column, 250 × 4.6 mm (part number: 00G-025-E0) with a guard column (Phenomenex) and a 0.5 μm in-line filter (Upchurch; Sigma-Aldrich). The column was shown to separate the carotenoids of interest in previous studies.

The method used a premixed isocratic mobile phase consisting of 85% acetonitrile, 15% methanol and 0.1% triethylamine, with a stepwise dichloromethane gradient initiated at 15 min with 10% dichloromethane over 1 min and increased to 50% dichloromethane between 25 and 27 min. The initial flow rate was set at 1 ml/min, and then increased to 2 ml/min at 15 min and remained as such for the duration of the dichloromethane gradient. The system resumed initial settings at 34 min. The L and total Z (co-eluted Z and MZ) peaks eluted at approximately 9.9 and 10.5 min, respectively. The IS eluted at 17.8 min. The system temperature throughout was maintained at 15°C. The mixed Z fraction was collected manually by switching the waste tube to a collection tube a couple of seconds after the peak was observed to start on the monitor. The eluent was dried under N2 and stored for no more than a few days at −70°C for further analysis (Assay 2).

**HPLC analysis of serum meso-zeaxanthin (Assay 2)**

The enantiomers Z and MZ present in the total Z peak were separated using a 5 μm chiral column (Chiralpak™ AD column (250 × 4.6 mm)), a guard column (Apex Scientific Limited) and 2 μm filter. The total Z fractions were reconstituted by vortex in 0.1 ml of mobile phase (20% hexane–isopropanol, 90:10) and 0.05 ml was then injected using
normal-phase chromatography and a linear gradient during which the proportion of hexane increased to 95% over 30 min. MZ, Z and L eluted at approximately 15-8, 18 and 20 min, respectively.

The absolute concentrations of L and total Z were calculated directly from the peak areas obtained in Assay 1. The concentrations were quantified using their respective response factors determined by UV–VIS spectroscopy analysis of the individual L and Z standards in absolute ethanol. Individual Z and MZ concentrations were then quantified from Assay 2, by calculating the percentage proportion of Z and MZ and applying the resulting ratio to the corresponding total Z value (obtained in Assay 1). All chromatographic peaks of interest were manually integrated using the Agilent ChemStation software (Agilent Technologies Ltd).

Absolute carotenoid concentrations were calculated as μmol/L. For purposes of interpretation, we also report the responses of the individual carotenoids as the changes in concentrations per milligram of supplement carotenoid provided. That is, the response was calculated as the average of the 4- and 8-week concentrations, or saturation plateau minus baseline values. This allowed for direct comparison between the interventions (Groups 1, 2 and 3), in terms of individual and total serum macular carotenoid responses and controls for the amount of supplement provided.

**Capsule carotenoid analysis**

The carotenoid content of the three supplements used in the present study were analysed using the following protocol. A stock solution was made by dissolving the contents of one capsule in 250 ml of acetone. A 0.5 ml aliquot was taken from this stock solution and made up to 25 ml with acetone to give the working solution for the analysis. The working solution was analysed in triplicate.

A measure of 0.4 ml of the working solution was transferred to a glass tube and dried under N2. A measure of 0.1 ml of IS (ethanolic echinenone, 0.4 mg/500 ml) and 0.4 ml methanolic KOH (50 g/100 ml) was added to the sample. The samples were stoppered and allowed to saponify at 45°C for 1 h in a shaking incubator. The samples were removed and allowed to cool to room temperature. The remaining KOH was neutralised using 1.5 ml 1 M-HCl. Butylated hydroxytoluene in hexane (1 ml; 25 mg/100 ml) was added to the sample and mixed by vortex for 2 min. The sample layers were allowed to separate under gravity for 5 min and a 0.5 ml aliquot of the organic layer was removed to an evaporation tube. A measure of 1 ml of hexane was added to the remaining sample, which was vortexed and allowed to separate as earlier. A measure of 1 ml of the hexane layer was removed and combined with the initial extract, which was dried in a solvent concentrator and stored at −80°C until the time of analysis. These samples were analysed using the HPLC method described earlier in order to quantify L, Z and MZ concentrations in each capsule. Of note, tablet carotenoid assessment of the three supplements used in the study was also performed by Industrial Organica (supplier of Intervention 3) and the data are concordant.

**Statistical analysis**

Means and standard deviations are presented in the text and tables (SPSS version 17; SPSS, Inc., used for data analysis). SigmaPlot (version 8; SyStat Software) was used for graphical presentations. All data were tested using the non-parametric Kolmogorov–Smirnov test, and they exhibited normal distribution. Between-group differences for numeric data (age, BMI, diet score and serum carotenoid levels) were calculated using ANOVA. Between-group differences for categorical variables (sex, smoking habits, sex and ocular status (normal or AMD)) were calculated using the standard χ² test. Difference between baseline and saturation point (i.e. average of weeks 4 and 8) was investigated for L, Z and MZ using paired-sample t tests. Repeated measures analysis was used to test for differences in response of each carotenoid between normal subjects and subjects with AMD, by testing for a time/subject group (i.e. normal subjects v. AMD subjects) interaction effect (Greenhouse-Geisser significance values were used and presented in the results section). A 5% level of significance was implemented throughout the analysis.

**Results**

The demographic, lifestyle, ocular disease status (normal or AMD) and baseline serum carotenoid data for all three groups are presented in Table 1. There were no statistically significant differences between groups for baseline parameters, with the exception of significantly higher mean serum Z concentration for subjects in Group 3.

There was no significant difference between normal subjects and AMD subjects with respect to any of the known possible confounders for carotenoids (e.g. BMI, sex and smoking habits), with the exception of a significant difference between these populations for age, which was principally due to the older AMD subjects in Group 1 (P=0.024). Also, there were no significant differences between normal subjects and AMD subjects in terms of baseline serum concentrations of the macular carotenoids or in terms of the responses to any of the three carotenoids (with one exception, Group 3 response to MZ, discussed later). Therefore, data for normal and AMD subjects were combined for the main analyses reported here (Table 2). Furthermore, there were no differences in the serum carotenoid responses between weeks 4 and 8, and these data were therefore averaged to provide data on the serum response at saturation point.

**Supplement assessment**

Table 3 presents the findings of the capsule analysis for each intervention used in the trial. The chromatograms in Fig. 2 show the presence of MZ in the UltraLutein® (intervention Group 1) supplement.

**Serum lutein response**

There was no significant difference with respect to serum L response between normal subjects and subjects with AMD.
Serum response to the macular carotenoids

Table 1. Demographic, lifestyle, ocular status (normal or early age-related macular degeneration (AMD)) and baseline serum carotenoid data for the three intervention groups

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>All</th>
<th>Group 1 (n 21)*</th>
<th>Group 2 (n 20)†</th>
<th>Group 3 (n 13)‡</th>
<th>P§</th>
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</thead>
<tbody>
<tr>
<td></td>
<td>Mean</td>
<td>Mean</td>
<td>Mean</td>
<td>Mean</td>
<td></td>
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<tr>
<td>Number of subjects (n)</td>
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<td>20</td>
<td>13</td>
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<tr>
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<td>AMD</td>
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<td>10</td>
<td>10</td>
<td>7</td>
<td></td>
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<tr>
<td>Age (years)</td>
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<td>70**</td>
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<tr>
<td>BMI (kg/m²)</td>
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<td>22</td>
<td>8</td>
<td>1·113</td>
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<tr>
<td>Serum carotenoids</td>
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<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>L (µmol/l)</td>
<td>0·25</td>
<td>0·15</td>
<td>0·26</td>
<td>0·16</td>
<td>0·702</td>
</tr>
</tbody>
</table>
| Z (µmol/l)                   | 0·05      | 0·03            | 0·05            | 0·03            | 0·012†††
| MZ (µmol/l)                  | 0         | −                | 0               | −                | −    |

L, lutein; Z, zeaxanthin; MZ, meso-zeaxanthin.
* High L group (20 mg L/d and 2 mg Z/d).
† High MZ group (3 mg L/d, 2 mg Z/d and 17 mg MZ/d).
‡ High Z group (10 mg L/d, 2 mg Z/d and 17 mg MZ/d).
§ Significance difference between the three intervention groups.
‖ Significance difference between normal and AMD subjects in Group 1 (P=0·024).
¶ Significant difference with respect to age between normal and AMD subjects in Group 1 (P=0·005).
†† Group 3 baseline serum Z was significantly greater than Groups 1 and 2; BMI, sex, smoking habits and diet scores information did not differ between AMD and normal subjects and were therefore reported as combined values.

(Group 1: P=0·409; Group 2: P=0·843; and Group 3: P=0·571). Serum L concentrations (for normal and AMD subjects combined) increased significantly in Group 1 (0·036 µmol/l per mg (269 %) increase; P<0·001) and Group 2 (0·079 µmol/l per mg (340 %) increase; P<0·001), with no significant change seen in Group 3 (0·0006 µmol/l per mg (7 %) increase; P=0·466). Serum L concentrations at the three study visits and the saturation point responses are presented in Tables 2 and 4, respectively.

Serum zeaxanthin response

There was no significant difference with respect to serum Z response between normal subjects and subjects with AMD (Group 1: P=0·198; Group 2: P=0·626; and Group 3: P=0·404). Serum Z concentrations (for normal and AMD subjects combined) increased significantly in Group 1 (0·037 µmol/l per mg (69 %) increase; P=0·001) and Group 2 (0·015 µmol/l per mg (75 %) increase; P<0·001), with no significant change seen in Group 3 (−0·0002 µmol/l per mg (6 %) decrease; P=0·384). Serum Z concentrations at the three study visits and the saturation point responses are presented in Tables 2 and 4, respectively.

Serum meso-zeaxanthin response

There was no significant difference with respect to serum MZ response between normal subjects and subjects with AMD for either Group 1 (P=0·675) or for Group 2 (P=0·985). However, there was a significant difference for subjects with AMD demonstrating a significantly greater serum MZ response than normal subjects following supplementation with this carotenoid (AMD subjects: 0·094 µmol/l at saturation point (0·006 µmol/l per mg increase); normal subjects: 0·036 µmol/l at saturation point (0·002 µmol/l per mg increase); P=0·014) (Fig. 3).

Although there was no MZ declared in the Group 1 supplement, a small concentration of MZ appeared to be present in serum at weeks 4 and 8 (0·005 µmol/l at week 8; P=0·015). Serum MZ concentrations increased significantly in Group 2 (0·005 µmol/l per mg increase; P<0·001) and Group 3 (0·004 µmol/l per mg increase; P<0·001). As MZ was not included as a supplemental carotenoid in Group 1, the MZ response can only be presented as the absolute value of MZ in µmol/l (Table 2). Serum MZ concentrations at the three study visits and the saturation concentration are presented in Tables 2 and 4, respectively.

Total macular carotenoid serum response

There was no significant difference with respect to total serum carotenoid response between normal subjects and subjects with AMD (Group 1: P=0·393; Group 2: P=0·842; and Group 3: P=0·152). Total serum carotenoid concentrations (for normal and AMD subjects combined) increased significantly in Group 1
(0.036 μmol/l per mg (242%) increase; P<0.001), Group 2 (0.040 μmol/l per mg (321%) increase; P<0.001) and Group 3 (0.004 μmol/l per mg (24%) increase; P=0.030). Total serum macular carotenoid response over the three study visits and to saturation point is presented in Tables 2 and 4, respectively.

### Discussion

The present study was conducted to investigate serum responses in normal subjects and in those with early AMD to three different macular carotenoid interventions, and is the first study to do so. We believe detailed investigation into serum macular carotenoid response is required, given that serum is the transporter of these carotenoids to their target tissues, including the retina. Moreover, the question as to whether there are differences in response between normal subjects and subjects with AMD is important, and was uniquely answered in the present study.

To this point, we report that subjects with AMD are comparable with normal subjects in how they respond to macular carotenoid supplements in serum, with the exception of a difference between these subject populations in response to a particular carotenoid.

| Table 2. Concentrations of serum lutein (L), zeaxanthin (Z), meso-zeaxanthin (MZ) and total macular carotenoids for each of the three carotenoid intervention groups investigated* (Mean values and standard deviations)† |
|-----------------|-----------------|-----------------|-----------------|-----------------|
|                  | Baseline (μmol/l) | 4 weeks (μmol/l) | 8 weeks (μmol/l) | P‡             |
|                  | Mean  | SD    | Mean  | SD    | Mean  | SD    | Mean  | SD    |    |
| L                |       |       |       |       |       |       |       |       |    |
| Group 1§         | 0.264  | 0.162 | 1.085 | 0.766 | 0.866 | 0.617 | 0.000 |       |    |
| Group 2†         | 0.233  | 0.141 | 0.954 | 0.480 | 1.095 | 0.628 | 0.000 |       |    |
| Group 3¶         | 0.273  | 0.134 | 0.290 | 0.092 | 0.292 | 0.105 | 0.501 |       |    |
| Z                |       |       |       |       |       |       |       |       |    |
| Group 1§         | 0.046  | 0.027 | 0.081 | 0.049 | 0.075 | 0.044 | 0.000 |       |    |
| Group 2†         | 0.040  | 0.025 | 0.067 | 0.033 | 0.074 | 0.043 | 0.000 |       |    |
| Group 3¶         | 0.071**| 0.033 | 0.066 | 0.038 | 0.066 | 0.033 | 0.287 |       |    |
| MZ               |       |       |       |       |       |       |       |       |    |
| Group 1§         | –      | –     | 0.007 | 0.006 | 0.009 | 0.016 | 0.014 |       |    |
| Group 2†         | –      | –     | 0.046 | 0.032 | 0.063 | 0.049 | 0.000 |       |    |
| Group 3¶         | –      | –     | 0.061 | 0.044 | 0.073 | 0.056 | 0.000 |       |    |
| Total macular carotenoids†† |       |       |       |       |       |       |       |       |    |
| Group 1§         | 0.310  | 0.181 | 1.173 | 0.816 | 0.951 | 0.651 | 0.000 |       |    |
| Group 2†         | 0.273  | 0.160 | 1.068 | 0.535 | 1.233 | 0.714 | 0.000 |       |    |
| Group 3¶         | 0.344  | 0.156 | 0.417 | 0.150 | 0.431 | 0.180 | 0.042 |       |    |

* Repeated measures analysis found no differences in response of each carotenoid between normal subjects and subjects with AMD except in Group 3 (see text).
† Mean values were in response to each carotenoid component for each group.
‡ Significance (P) values represent paired-sample t tests significance for the increase between baseline and 8 weeks.
There were no differences in carotenoid concentrations between weeks 4 and 8.
§ Group 1 (n=21): high L group (20 mg L/d and 2 mg Z/d).
∥ Group 2 (n=20): combination group (10 mg L/d, 2 mg Z/d and 10 mg MZ/d).
¶ Group 3 (n=13): high MZ group (3 mg L/d, 2 mg Z/d and 17 mg MZ/d).
** There were no differences in baseline concentrations, except for Z, in Group 3 (see Table 1).
†† Mean values were in response to each carotenoid component for all three carotenoids in each supplement.

### Table 3. Declared and measured carotenoid content of the three study supplements

<table>
<thead>
<tr>
<th>Group</th>
<th>Supplement</th>
<th>Declared carotenoid content (mg/capsule)</th>
<th>Measured carotenoid content (mg/capsule)</th>
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</thead>
<tbody>
<tr>
<td></td>
<td>L</td>
<td>Z</td>
<td>MZ</td>
</tr>
<tr>
<td>1*</td>
<td>UltraLutein††</td>
<td>20</td>
<td>0.86</td>
</tr>
<tr>
<td>2†‡</td>
<td>Macushield‡‡§</td>
<td>10</td>
<td>2</td>
</tr>
<tr>
<td>3¶</td>
<td>Customised MZ¶</td>
<td>3</td>
<td>2</td>
</tr>
</tbody>
</table>

L, lutein; Z, zeaxanthin; MZ, meso-zeaxanthin.
* Group 1 (n=21: eleven normal and ten age-related macular degeneration (AMD)).
†† Provided by Nature’s Plus, Natural Organics, Inc., Melville, NY, USA.
‡ Group 2 (n=20: ten normal and ten AMD).
§§ Provided by MacuVision Europe Limited, Solihull, UK.
¶ Group 3 (n=13: six normal and seven AMD).
††† Provided by Industrial Organica, Monterrey, Mexico (not available commercially).
supplementation with the central macular carotenoid, MZ. Of note, we report no significant difference in serum L or serum Z response between these subject populations for any of the interventions tested. However, the Group 3 formulation (i.e. very high MZ group) yielded significantly different serum MZ responses between the AMD and normal populations, with the MZ response amongst AMD subjects significantly greater than that observed for normal subjects. This seemingly counterintuitive observation is difficult to explain, but may reflect enhanced absorption of this macular carotenoid in subjects who exhibit tissue deficiencies of MZ, reflected in the absence of a typical central peak in MP optical density spatial profile in association with risk factors for AMD(30). However, the main finding from the present study is that normal and AMD subjects responded comparably to the three different macular carotenoid interventions. This is an important finding, as it confirms that the known lack of MP seen in subjects afflicted with(31), or at high risk of developing (32), AMD is not due to an inability of such subjects to respond to carotenoid consumption, and is therefore due to either a defective capture of circulating carotenoids by, or stabilisation within, the central retina. Our data are consistent with a publication by Wang et al.(33) who studied subjects with and without AMD following dietary modification (increased consumption of spinach, 
maize, cabbage, circa 11 mg/d of L and Z (combined), in the high-carotenoid-fed group) and reported no difference between these group in terms of serum concentrations of L and Z.

Given that, by and large and with the exception of response to supplementation with high-dose MZ, there were no differences between subjects with and without AMD in terms of response to supplementation with MP’s constituent carotenoids, we elected to treat our dataset as a single and merged set of data.

We examined the literature to determine the carotenoid responses reported by other researchers, in order to allow comparison with our findings. Examination of these studies has shown that it has been a common practice for other researchers to report carotenoid response in terms of concentrations and/or percentage increases. Presenting the data as percentages and/or concentrations, however, makes it very difficult to interpret the data, as baseline concentrations can vary considerably between subjects. It was for this reason that we constructed and present Table 5. Table 5 presents data on serum L, Z and MZ responses of these other published studies, but we have converted their data to the unit reported in the present study, which is serum response per mg of supplemental carotenoid (controlling for the amount of carotenoid given and the baseline values). Interestingly, upon
examination of this table with respect to L response of the other published studies (and excluding our own data), we found that the mean response was 0.066 (SD 0.042) with a range of 0.01–0.17 mmol/l per mg. Also, for Z response (and excluding our own data), the mean response was lower at 0.050 (SD 0.035) with a range of 0.004–0.15 mmol/l per mg. Thus, the range of serum macular carotenoid response reported in the literature is very wide. Of note, the responses observed in the present study fall comfortably within the ranges reported in these other studies (see Table 5).

As seen in Table 5, studies investigating serum carotenoid response to MZ are few, although Connolly et al. (23) and Thurnham et al. (25) did study and report on MZ response in normal men and women to a supplement similar in composition to that used in Group 2 here. Although it is known that there is considerable inter-individual variation in terms of MP response to any dietary/supplement intervention(34), there are no published studies of reports designed to investigate serum response to differing macular carotenoid formulations.

**Serum lutein response**

L was present in each of the three carotenoid group interventions. However, Group 1 (the high L group) contained approximately double that of Group 2 and six times that of Group 3. As expected, Groups 1 and 2 demonstrated the greatest serum response to L (0.036 mmol/l per mg (269%)) and 0.079 mmol/l per mg (340%), respectively, with Group 3 demonstrating no response (0.006 mmol/l per mg (6%)).

The Group 1 response is consistent with a previous study by Johnson et al. (23), who used a similar formulation. Of interest, the Group 2 supplement, which contained only half the amount of L compared with that of Group 1, but also contained 10 mg of MZ and 2 mg of Z, achieved a significantly greater serum L response than that observed for Group 1 (0.079 v. 0.036 mmol/l per mg, respectively). The only other previous study that reported on serum response to a formulation similar to Group 2 was conducted by Thurnham et al. (25) in 2008. In that study, nineteen subjects were supplemented with 10 mg L, 1.2 mg of Z and 8 mg of MZ (LuteinPlus™), and a lower L response (0.056 mmol/l per mg) was reported than that observed in the present study (0.079 mmol/l per mg). In a different study, Thurnmann et al. (50) reported serum responses to two different dosages of almost pure L (41 and 20.5 mg/d). In that 42 d study, the
authors found that the rate of increase in serum L was greater for the low-dose L supplement when compared with the high-dose L supplement (0.093 and 0.064 μmol/l per mg, respectively). However, as expected, supplementation with high-dose L did result in a significantly higher absolute serum response to the macular carotenoids following supplementation with the macular carotenoids

Table 5. Serum carotenoid response per mg of supplemental carotenoid, following supplementation with the macular carotenoids

<table>
<thead>
<tr>
<th>Principal author</th>
<th>Journal</th>
<th>Year</th>
<th>Age (years)</th>
<th>L</th>
<th>Z</th>
<th>MZ</th>
<th>L response (μmol/l per mg)</th>
<th>Z response (μmol/l per mg)</th>
<th>MZ response (μmol/l per mg)</th>
<th>Duration (weeks)</th>
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<tr>
<td>Bone et al.</td>
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<td></td>
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<td>22</td>
<td>18–30</td>
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<td></td>
<td>0.053</td>
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* Data unavailable.
† Free (unesterified) carotenoid supplement.
‡ Includes MZ supplementation.

Interpretation of our data for serum L response suggests that uptake of L may be inhibited when MZ is present in the supplement formulation in very high amounts (e.g. Group 3 where MZ was 77%), but uptake of L may be unaffected or facilitated when the combined amounts of MZ and Z are present in more comparable amounts to L (i.e. as in Group 2). Whether high amounts of Z in a supplement has the same effect on uptake into serum, as seen here with MZ, has not yet been studied, but warrants attention. Thus, in the present study, our data demonstrate a greater serum response to supplemental L when the macular carotenoids are provided in a L:Z:MZ ratio of 10:2:10, possibly representing an interactively additive relationship that is dependent on a ratio approximating the Group 2 formulation.

**Serum zeaxanthin response**

Z was present in all three intervention groups as follows: Group 1 containing 2 mg Z, and Groups 2 and 3 each containing 2 mg Z in their respective formulations. Interestingly, the
increase in serum Z is comparable between Groups 1 and 2. Also, the Z response for all groups in the present study is lower, when compared with reports by Hartman et al. (25) and Thurnham et al. (25) (0.088 μmol/l per mg and 0.087 μmol/l per mg, respectively).

Surprisingly, the Group 3 intervention did not achieve any serum Z response, suggesting that very high amounts of MZ (i.e. 17 mg, as in Group 3) in a supplement inhibits uptake of Z and is also consistent with previously published data (29). Further, Bone et al. (29) and Schalch et al. (39) have each reported maximum serum Z response when this carotenoid is supplemented in the absence of other carotenoids, but this does not agree with the observations of Hartman et al. (25) and Thurnham et al. (25), who obtained high Z responses with very different supplement compositions. Nevertheless, several other studies report a reduced Z response in the presence of high amounts of L, as observed in the present study in Group 1 (see Table 5 for comparison) (29, 35, 38, 39).

Interpretation of our data for serum Z response suggests that uptake of Z is inhibited when supplemented with very high quantities of MZ (Group 3). It is possible that the inhibitions seen in Group 3 are due to the structural and chemical similarities between Z and MZ, which might cause them to compete for absorption. Differences between the present study and other studies with respect to absorption of Z and MZ was, indeed, present in the serum of these subjects. We then tested the Group 1 intervention formulation (Ultra Lutein® from Nature’s Plus®; L provided by FloraGLO® which is a registered trademark of Kemin Health, L.C.) and determined that this formulation did, in fact, contain MZ (0.3 mg per capsule), which we believe explains the observation that Group 1 subjects exhibited a trace peak with the spectrophotometric characteristics of MZ in serum at 4 and 8 weeks. We then went on to test the composition of the other interventions used in the present study and found that they were concordant with their respective label claims (see Table 3). These findings have implications on the ongoing research surrounding carotenoid supplementation. Indeed, any discrepancy between actual and alleged concentrations of the respective macular carotenoids in commercially available preparations is particularly important when such formulations are used for research. Consequently, the concentration of all three macular carotenoids in a wide array of commercially available formulations and foods, with particular attention directed towards MZ, will be the subject of further study by our laboratory.

Conclusion
The present study has yielded important and novel findings, such as the presence of MZ in the serum of normal and AMD subjects following supplementation with high doses of L, and the significantly greater serum MZ response amongst subjects with AMD v. normal subjects following supplementation in the high MZ group. A limitation of the present study rests on the absence of a placebo arm, and further research should therefore include a placebo control group. Moreover, a head-to-head trial of supplemental MZ v. supplemental Z (in comparable amounts) would also enhance our understanding of serum response to supplementation with the macular carotenoids.

We conclude that all three macular carotenoid interventions resulted in significant serum carotenoid response, albeit to varying extents. Group 2, an intervention containing 2 mg Z, 10 mg L and 10 mg MZ achieved the greatest composite serum response for these carotenoids (i.e. total macular carotenoid response). In other words, it appears that a formulation containing all three macular carotenoids was more efficacious in terms of achieving a higher concentration of circulating levels of total macular carotenoids, thereby potentially optimising the bioavailability of these compounds for capture by the target tissue (retina).
Acknowledgements

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