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Novel targeted approach to better understand how natural structural barriers govern carotenoid *in vitro* bioaccessibility in vegetable-based systems

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A B S T R A C T

An experimental approach, allowing us to understand the effect of natural structural barriers (cell walls, chromoplast substructures) on carotenoid bioaccessibility, was developed. Different fractions with different levels of carotenoid bio-encapsulation (carotenoid-enriched oil, chromoplasts, small cell clusters, and large cell clusters) were isolated from different types of carrots and tomatoes. An *in vitro* method was used to determine carotenoid bioaccessibility.

In the present work, a significant decrease in carotenoid *in vitro* bioaccessibility could be observed with an increasing level of bio-encapsulation. Differences in cell wall material and chromoplast substructure between matrices influenced carotenoid release and inclusion in micelles. For carrots, cell walls and chromoplast substructure were important barriers for carotenoid bioaccessibility while, in tomatoes, the chromoplast substructure represented the most important barrier governing bioaccessibility. The highest increase in carotenoid bioaccessibility, for all matrices, was obtained after transferring carotenoids into the oil phase, a system lacking cell walls and chromoplast substructures that could hamper carotenoid release.

Keywords:

Bioaccessibility
 Carotenoids
 Carrot
 Natural structural barriers
 Tomato

1. Introduction

Epidemiological studies often relate a healthy lifestyle, in which the consumption of adequate amounts of fruit and vegetables is important, to a reduced risk of chronic diseases such as cardiovascular diseases and cancers (Key, 2011; Ness & Powles, 1997; Steinmetz & Potter, 1996; Van't Veer et al., 2000). The health-related benefits that are associated with the consumption of fruit and vegetables, can be attributed to the presence of various bioactive compounds, of which micronutrients form an important class (Key, 2011; Van't Veer et al., 2000). Carotenoids, a group of natural food pigments, are major contributors to the nutritional value of several fruits and vegetables (Rao & Rao, 2007).

Carotenoids are isoprenoid compounds which typically have a tetraterpenoid structure, implying a long chain of conjugated double bonds. Due to their chemical structure, carotenoids are highly lipophilic molecules (Britton, 1995). In fruit and vegetable tissues,

specific structures are developed in chloroplasts and chromoplasts to sequester and store large amounts of carotenoids (Vishnevetsky, Ovadis, & Vainstein, 1999). Their lipophilic nature and their specific localisation in plant tissues (i.e. attached to cellular components and surrounded by organelle membranes, cell membrane and cell wall), hamper the absorption of carotenoids from fruits and vegetables in the human tract (Rich et al., 2003). This has been and still is an issue for food technologists and food processors. In this context, the concepts of carotenoid bioaccessibility and bioavailability are to be defined. Carotenoid bioaccessibility refers to the fraction of ingested carotenoids that is released from the food matrix and incorporated into micelles during digestion in the gastrointestinal tract, and thus becomes available for intestinal absorption. The amount of carotenoids that is bioavailable is always less than the amount that is bioaccessible, since carotenoid bioavailability additionally takes into account the fraction that is available for utilisation in normal physiological functions or for storage in the human body (Holst & Williamson, 2008; Parada & Aguilera, 2007).

In the literature, several studies describe the carotenoid bioaccessibility in fruit- and vegetable-based food products (e.g. Granado-Lorencio et al., 2007; O'Connell et al., 2007; O'Sullivan et al., 2010; Reboul et al., 2006; Ryan, O'Connell, O'Sullivan,

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Aherne, & O'Brien, 2008; Veda, Kamath, Platel, Begum, & Srinivasan, 2006; etc.). However, only a few authors have linked the observations for carotenoid bioaccessibility to structural characteristics of the food products (e.g. Lemmens, Van Buggenhout, Oey, Van Loey, & Hendrickx, 2009 (carotenoid bioaccessibility – pectin properties); Bengtsson, Brackmann, Enejder, Alminger, & Svanberg, 2010 (carotenoid bioaccessibility – microstructure); Colle, Van Buggenhout, Van Loey, & Hendrickx, 2010 (carotenoid bioaccessibility – strength of fibre network); Lemmens, Van Buggenhout, Van Loey, & Hendrickx, 2010 (carotenoid bioaccessibility – particle size); Tydeman et al., 2010 (carotenoid bioaccessibility – microstructure); Knockaert, Lemmens, Van Buggenhout, Hendrickx, & Van Loey, 2012 (carotenoid bioaccessibility – microstructure)). Reviews by Waldron, Parker, and Smith (2003), Parada and Aguilera (2007) and Van Buggenhout et al. (2010) have stressed the importance of the association between food structure and nutrient bioaccessibility. However, studies identifying and directly investigating the role of structural barriers against carotenoid bioaccessibility in fruit- and vegetable-based food products are lacking. In our opinion, such studies might give useful explanations and better insights of how crucial structural parameters can determine carotenoid bioaccessibility in fruit- and vegetable-based food products. Recently, some initiatives in this direction have been taken. Schweiggert, Mezger, Schimpf, Steingass, and Carle (2012) studied the relationship between chromoplast morphology and carotenoid bioaccessibility in different matrices, and found a strong correlation between the physical state of the chromoplast substructures and the efficiency of carotenoid release during digestion. Jeffery, Holzenburg, and King (2012) and Jeffery, Turner, and King (2012) studied different fruit and vegetable purees microscopically (cell wall thickness, cell size) (Jeffery, Holzenburg, et al., 2012) and they tried to link these observations with the carotenoid bioaccessibility results obtained for the same purees (Jeffery, Turner, et al., 2012). They concluded that the cell wall and the chromoplast substructures form the most important barriers against carotenoid release during digestion. Moreover, it turned out that a high amount of large plant cells, a low density of cell wall material and a high concentration of plastoglobuli (containing the carotenoids) are factors favouring carotenoid bioaccessibility (Jeffery, Turner, et al., 2012). However, for these experiments, it should be kept in mind that, during the mixing/blending process to obtain fruit and vegetable purees, a complex environment is created, which could lead to difficulties in the interpretation of the results. For example, it is possible that additional structural networks are being formed during mixing which entail new processed-induced barriers against carotenoid release during digestion, next to and/or replacing the natural barriers present in the fruit and vegetable matrix. Such examples have already been cited by Colle, Van Buggenhout, Lemmens, Van Loey, and Hendrickx (2012) and Anese, Mirolo, Beraldo, and Lippe (2013), respectively, in the framework of the effect of high pressure homogenisation and ultrasound treatments on tomato pulp microstructure and lycopene *in vitro* bioaccessibility.

Therefore, in this study, a specific experiment (including different matrices, different types of carotenoids, and different levels of bio-encapsulation) was designed to evaluate the role of natural barriers in carotenoid bioaccessibility. The fruit and vegetable matrices included in this study (orange carrots, red carrots, orange tomatoes, red tomatoes) were not used as such (e.g. as a puree), but specific fractions were isolated, each representing a different number of barriers encapsulating the carotenoids. The fractions included a carotenoid-enriched oil, a chromoplast, a small/single cell cluster and a large cell/multicellular cluster. In this way, a systematic and detailed understanding of the role of different structural barriers for the carotenoid bioaccessibility could be obtained. Carotenoids with different polarities (β -carotene, ζ -carotene and lycopene) in different matrices (e.g. carrot, tomato) were included.

2. Materials and methods

2.1. Materials

Red tomatoes (*Lycopersicon esculentum* cv. Patrona) were obtained from a Spanish supplier. Orange tomatoes (*Lycopersicon esculentum* cv. Bolzano) were purchased in an auction in Mechelen, Belgium. The two types of tomatoes were cut, frozen with liquid nitrogen, and stored at -40°C , until the start of the experiments.

Orange carrots (*Daucus carota* cv. Nerac) and red carrots (unknown Indian variety) were obtained fresh from local shops in Belgium and stored briefly at 4°C , prior to their use for the experiments.

2.2. Experimental set-up

In order to study the different physical barriers that determine the carotenoid *in vitro* bioaccessibility, various fractions from red tomatoes, red carrots, orange carrots and orange tomatoes were isolated. The fractions prepared include a carotenoid-enriched oil fraction, a chromoplast fraction, a small cell cluster fraction, and a large cell cluster fraction. The concomitant physical barriers surrounding the carotenoids were assumed to be the chromoplast structure/organisation, and (multiple) cell membranes and cell walls. An *in vitro* bioaccessibility assay was performed on each fraction. Light microscopy was used to visualise the chromoplast and small cell cluster fractions and to observe the structural characteristics of barriers involved. Fig. 1 represents a schematic overview of the experimental set-up.

2.3. Preparation of carotenoid-containing fractions with different barrier properties

2.3.1. Carotenoid-enriched oil fraction

Lycopene from red carrots and red tomatoes, β -carotene from orange carrots and ζ -carotene from orange tomatoes were extracted in olive oil.

To produce carotenoid-enriched oil fractions, carrots were peeled, cut into small pieces, mixed with deionized water (1:1) and blended (Waring Commercial, Torrington, CT, USA) for 1 min. Tomatoes were thawed, peeled, mixed three times (Büchi B-400 mixer, Flawil, Switzerland) for 5 s and then sieved to remove the seeds.

According to the method described by Colle et al. (2010), the obtained carrot and tomato purees were homogenised (Panda 2 K, Gea Niro Soavi, Parma, Italy) at 1000 bar for one cycle to decrease the particle size and facilitate the release of carotenoids from the matrix. Purees were then mixed with olive oil (ratio puree-oil 5:1) for 5 h while rotating, end-over-end. The carotenoid-enriched oil fraction was separated by centrifugation (J2 – HS centrifuge, Beckman, J2 – HS centrifuge, Palo Alto, CA, USA) at 18,900g, 4°C for 15 min. The carotenoid-enriched oil fraction was isolated and emulsified (5% carotenoid-enriched oil in water emulsion) with a 1% of $\text{l-}\alpha$ -phosphatidylcholine solution. The emulsion was stabilized by homogenisation (Gea Niro Soavi, Parma, Italy) at 1000 bar for one cycle.

2.3.2. Chromoplast fraction

The chromoplast fraction was obtained by following the procedure of Hansen and Chiu (2005) with some modifications. Both types of carrots and tomatoes (previously defrosted) were cut into pieces and mixed in a blender (Waring Commercial, Torrington, CT, USA) for 5 s with 0.05 M EDTA solution (1:1 ratio).

The obtained purees were filtrated using a cheesecloth. The filtrate was centrifuged (Beckman, J2-HS Centrifuge, Palo Alto, CA,

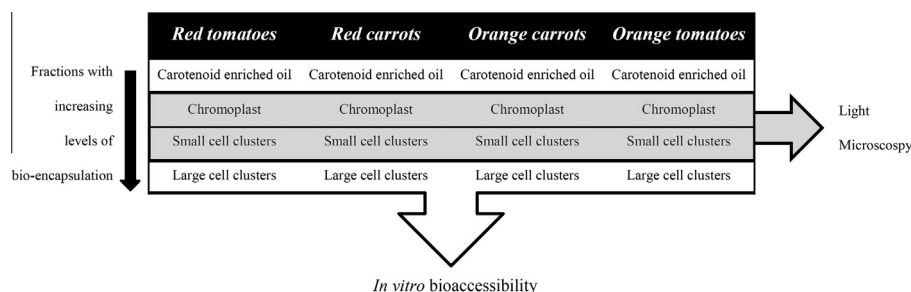


Fig. 1. Overview of the experimental set-up.

USA) at 27,200g, 4 °C for 30 min. The pellet, which corresponds to the chromoplast-enriched fraction, was re-dissolved in 100 ml of deionized water.

2.3.3. Cell cluster fractions

Cell cluster fractions were obtained by separating the particles of tomato and carrot purees into different sizes with a wet sieving equipment (Retsch AS200, Haan, Germany). The carrot and tomato purees were prepared as described in the section on the carotenoid-enriched oil fraction.

In the case of carrot purees, the size of the fraction considered as "small cell clusters" was 40–250 µm, whereas cell clusters with a size of 800–2000 µm were isolated and corresponded to the "large cell clusters". These ranges were determined according to the data reported by Lemmens et al. (2010) and confirmed by light microscopy observations, obtaining mean diameters of 62.9 ± 22.5 and 61.4 ± 15.2 µm for red carrot cells and orange carrot cells, respectively.

For the tomato purees, the collected fractions, ranging from 160 to 500 µm and from 1000 to 1400 µm corresponded to "small cell clusters" and "large cell clusters", respectively. These ranges were based on the data reported by Cheniclet et al. (2005), and on diameters of 446.4 ± 216.5 and 495.2 ± 272.5 µm, obtained by light microscopy, respectively, for red and orange tomato cells.

2.4. Light microscopy

Unstained carrot and tomato fractions were examined under an Olympus BX-41 light microscope (Olympus, Tokyo, Japan) equipped with an Olympus XC50 CCD camera (Olympus, Tokyo, Japan). Cell dimensions were measured from pictures obtained at ×10 and ×40 magnifications, using Cell* software (Olympus, Tokyo, Japan). The chromoplast fraction of the different matrices was analysed with ×100 oil objective lens.

2.5. Carotenoid concentration

Carotenoids (lycopene, β-carotene and ζ-carotene) were extracted from each fraction according to the method described by Sadler, Davis, and Dezman (1990) and Lemmens et al. (2010) with some modifications. Sodium chloride was added to a specific weight of the samples and then mixed for 20 min at 4 °C by continuous stirring with 50 ml of extraction solution, which consisted of hexane:acetone:ethanol (50:25:25 v/v/v) with 0.1% of butylated hydroxytoluene. The stirring was continued for 10 min at 4 °C after adding 15 ml of reagent grade water (18.2 MΩ cm).

The organic phase, containing the carotenoids, was separated from the polar phase in a separation funnel. The separation was performed under subdued light conditions, in order to prevent carotenoid isomerization and degradation. The organic phase was filtered (Chromafil PET filters, 0.2 µm pore size-25 mm diameter), and transferred into a dark vial for HPLC analysis.

The identification and quantification of carotenoids were achieved using an HPLC system equipped with a diode array detector (Agilent Technologies 1200 Series, Dinslaken, Belgium). A C₃₀-column (3 µm × 150 mm × 4.6 mm, YMC Europe, Dinslaken, Germany) was used to separate the carotenoids.

A linear gradient was applied to separate carotenoids. The starting conditions were 81% methanol, 15% methyl-*t*-butyl-ether and 4% reagent grade water. The end conditions for all-*trans*-lycopene determination were 16% methanol, 80% methyl-*t*-butyl-ether and 4% reagent grade water (18.2 MΩ cm) and, in the case of all-*trans*-β-carotene and all-*trans*-ζ-carotene, the end conditions corresponded to 41% methanol, 55% methyl-*t*-butyl-ether and 4% reagent grade water (18.2 MΩ cm). The gradient was built up in 38 min for all-*trans*-lycopene analysis and in 17 min for all-*trans*-β-carotene and all-*trans*-ζ-carotene analyses, with a constant flow rate of 1 ml/min. The column temperature was kept at 25 °C and the auto-sampler at 4 °C during the analyses. Identification and quantification were performed at 472 nm for all-*trans*-lycopene and 450 nm for all-*trans*-β-carotene and all-*trans*-ζ-carotene.

Standards for all-*trans*-lycopene, all-*trans*-β-carotene, and all-*trans*-ζ-carotene (CaroteNature, Lupsingen, Switzerland) were used to identify and quantify each carotenoid, using calibration curves.

The ζ-carotene in orange tomatoes, was identified by retention time and DAD spectra with the standard. However, DAD spectra clearly showed that the compound identified was not purely ζ-carotene. Other structurally similar carotenoid might be present as well, such as pro-lycopene, identified before in orange tomatoes by Mackinney and Jenkins (1949), having a maximal absorption at 400–430 nm. Nevertheless, in this study, our results on orange tomatoes, are expressed as equivalents of ζ-carotene as one of the major compounds.

2.6. Carotenoid in vitro bioaccessibility

The *in vitro* digestion procedure was based on the method described by Lemmens et al. (2010), where stomach and small intestinal digestion were simulated.

Gastric digestion consisted of the addition of 5 ml of NaCl / ascorbic acid solution (0.9% NaCl, 1% ascorbic acid in water) and 5 ml of electrolyte solution (0.3% NaCl, 0.11% KCl, 0.15% CaCl₂·2H₂O, 0.05% KH₂PO₄ and 0.07% MgCl₂·6H₂O) to a specific amount of each fraction. The pH was adjusted to 4 ± 0.05 and 5 ml of gastric pepsin solution (0.52% porcine pepsin in electrolyte solution) were added. The headspace of the tubes was flushed with N₂ and the samples were incubated at 37 °C while shaking, end-over-end, for 30 min. The pH was then adjusted to 2 ± 0.05 and the headspace was again flushed with N₂ and incubated for 30 min under the same conditions.

The small intestinal digestion was simulated by adjusting the pH to 6.9 ± 0.05 and adding 6 ml of pancreatin/bile solution (0.2% lipase from porcine pancreas, 0.4% porcine pancreatin, 2.5% bile extract, 0.5% pyrogallol and 1% tocopherol in water). The headspaces

of the tubes were flushed with N₂ and the samples were incubated for 2 h at 37 °C while shaking, end-over-end.

After digestion, samples were ultracentrifuged (Beckman, L7 Ultracentrifuge, Palo Alto, CA, USA) at 165,000g, 4 °C for 65 min. The soluble part, corresponding to the carotenoids incorporated into the micelles, was isolated. Carotenoids were extracted and quantified according to the procedure described above.

In the case of tomato samples, after phase separation in the funnel, the organic phase was concentrated, using a rotatory evaporator at 30 °C for 35 min. The concentrated extract was then redissolved in 400 µl of hexane:dichloromethane (4:1). A known amount of β-apo-8'-carotenal was used to calculate the concentration factor.

To determine the carotenoid bioaccessibility of the chromoplast and the small and large cell cluster fractions, 10 ml of an emulsion (5% olive oil in water, 1% L-α-phosphatidylcholine), were added at the beginning of the *in vitro* digestion assay. The emulsion was incorporated into the fractions that did not contain oil in order to obtain a carotenoid/oil ratio similar to that of the carotenoid-enriched oil fraction.

It should be kept in mind that phospholipids stabilizing the emulsions could decrease the transfer of carotenoids to the oil phase (Degrou, George, Renard, & Page, 2013). For the cell clusters and chromoplast fractions, the oil emulsion stabilized with L-α-phosphatidylcholine, added before the *in vitro* bioaccessibility procedure, could form an important barrier against carotenoids diffusing into oil. The same case will apply for the enriched oil phase fractions that were stabilized with the same emulsifier. Nevertheless, our results are comparable between each other, as the amount of phospholipids present was the same for the different fractions on the four matrices.

Measurements of the carotenoid concentration and carotenoid *in vitro* bioaccessibility of each fraction were performed in triplicate. Results are shown as the ratio of the bioaccessible carotenoids to the initial carotenoid concentration in the sample (B/C).

2.7. Data analysis

In order to identify statistically significant differences between the bioaccessibilities (B/C) of the different fractions in each matrix, a Tukey's standardized range test (SAS version 9.3, SAS Inst. Inc., Cary, NC, USA) was used for the analysis. The level of significance was set at $P < 0.05$. Significant differences among matrices in each fraction were analysed by the same approach.

3. Results and discussion

3.1. Carotenoid concentrations

The fractions isolated from each matrix (red carrots, red tomatoes, orange carrots and orange tomatoes) containing carotenoids with increasing levels of bio-encapsulation (carotenoid-enriched oil, chromoplast, small and large cell cluster fractions) were analysed for the initial carotenoid concentration.

Lycopene was the major carotenoid present in fractions obtained from red carrots and red tomatoes. β-Carotene was the main carotenoid in fractions obtained from orange carrots, whereas ζ-carotene and another unknown carotenoid were the major carotenoids present in orange tomato fractions. However, as already explained, results for orange tomato fractions were expressed as equivalents of ζ-carotene. Concentrations of the main carotenoids in the fractions obtained from each matrix are shown in Table 1.

3.2. Lycopene *in vitro* bioaccessibility in fractions of red tomatoes and red carrots

Fractions obtained from red tomato and red carrot fractions (carotenoid-enriched oil, chromoplast, small and large cell clusters) were prepared and the corresponding lycopene *in vitro* bioaccessibilities were measured.

The results for red carrot fractions (Fig. 2A) and red tomato fractions (Fig. 2B) show a clear decrease in lycopene *in vitro* bioaccessibility with increasing levels of lycopene bio-encapsulation.

It has been previously hypothesised that cell walls and chromoplast structures constitute important physical barriers against carotenoid release from the matrix and thus for subsequent incorporation into micelles (Jeffery, Holzenburg, et al., 2012; Lemmens et al., 2010; Tydeman et al., 2010; Xianquan, Shi, Kakuda, & Yue-ming, 2005). Cell wall polysaccharides may impede micelle formation by entrapping lipids and bile salts, and by preventing digestive enzymes reaching the active compounds inside the cells for further digestion and absorption (Palafox-Carlos, Ayala-Zavala, & González-Aguilar, 2011). At the same time, the chromoplast structure provides a stable environment for carotenoids, preventing their complete release and subsequent solubilisation, affecting both the stability and bioaccessibility (Schweiggert, Steingass, Heller, Esquivel, & Carle, 2011).

Our results show direct evidence of barriers and therefore support the existing hypothesis, since lower lycopene bioaccessibility values were obtained in fractions where cell walls and chromoplasts were present as structural barriers. However, interesting differences were observed between fractions obtained from the two matrices (red carrots and red tomatoes), thus indicating that the matrix in which lycopene is embedded plays an important role.

For red carrots, the largest increase of lycopene bioaccessibility was obtained after the removal of the cell wall (i.e., the chromoplast fraction vs. the cell cluster fractions). The difference ($P < 0.05$) in lycopene *in vitro* bioaccessibility between cell clusters and the chromoplast fraction was significant. Lycopene *in vitro* bioaccessibility in red carrots was further increased after removal of the chromoplast barrier (lycopene solubilised in the oil phase), where a significant difference ($P < 0.05$) of the lycopene *in vitro* bioaccessibility was found between the carotenoid-enriched oil fraction and the three other fractions.

In the case of red tomatoes, the enhanced bioaccessibility was more pronounced when lycopene was solubilised in the oil fraction, rather than when cell walls were removed, as in red carrots. Statistical analysis showed that there was no significant difference ($P < 0.05$) between the lycopene *in vitro* bioaccessibilities in cell clusters and in chromoplast fractions of red tomatoes. Nevertheless, the lycopene bioaccessibility in the oil fraction was significantly higher ($P < 0.05$) than those in the other three fractions.

Further analysis showed that, for red carrots, cell walls and chromoplast substructure constitute important barriers against lycopene solubilisation while, in red tomatoes, the chromoplast substructure becomes the strongest obstacle hindering lycopene release during digestion. Therefore, it can be concluded that the cell wall is the most important barrier against lycopene bioaccessibility in case of red carrots, since a threefold increase of the lycopene bioaccessibility was obtained after isolation of the chromoplast fraction. The chromoplast substructure is also an important barrier against lycopene bioaccessibility, as a significant improvement (but to a lower extent) was found after isolating the carotenoid in the enriched oil fraction. For red tomatoes, however, the most important barrier is associated with the chromoplast substructure, as the solubilisation of lycopene in the enriched oil fraction was the most important step for increasing its bioaccessibility.

These observations clearly indicate that differences in cell wall composition and chromoplast substructure among the matrices

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Table 1

Carotenoid concentration ($\mu\text{g/g}$ fraction) (mean \pm standard deviation) in each fraction of red carrots, red tomatoes, orange carrots and orange tomatoes, respectively.

	Lycopene in red carrots ($\mu\text{g/g}$)	Lycopene in red tomatoes ($\mu\text{g/g}$)	β -Carotene in orange carrots ($\mu\text{g/g}$)	ζ -Carotene in orange tomatoes ($\mu\text{g/g}$)
Carotenoid-enriched oil fraction	3.04 ± 0.05	2.43 ± 0.05	4.39 ± 0.28	0.24 ± 0.01
Chromoplast fraction	45.8 ± 1.81	36.2 ± 0.98	25.8 ± 11.46	3.35 ± 0.13
Small cell cluster fraction	46.3 ± 0.51	10.9 ± 0.45	21.1 ± 4.95	4.89 ± 0.12
Large cell cluster fraction	27.8 ± 6.10	45.1 ± 1.25	6.96 ± 0.43	3.96 ± 0.57

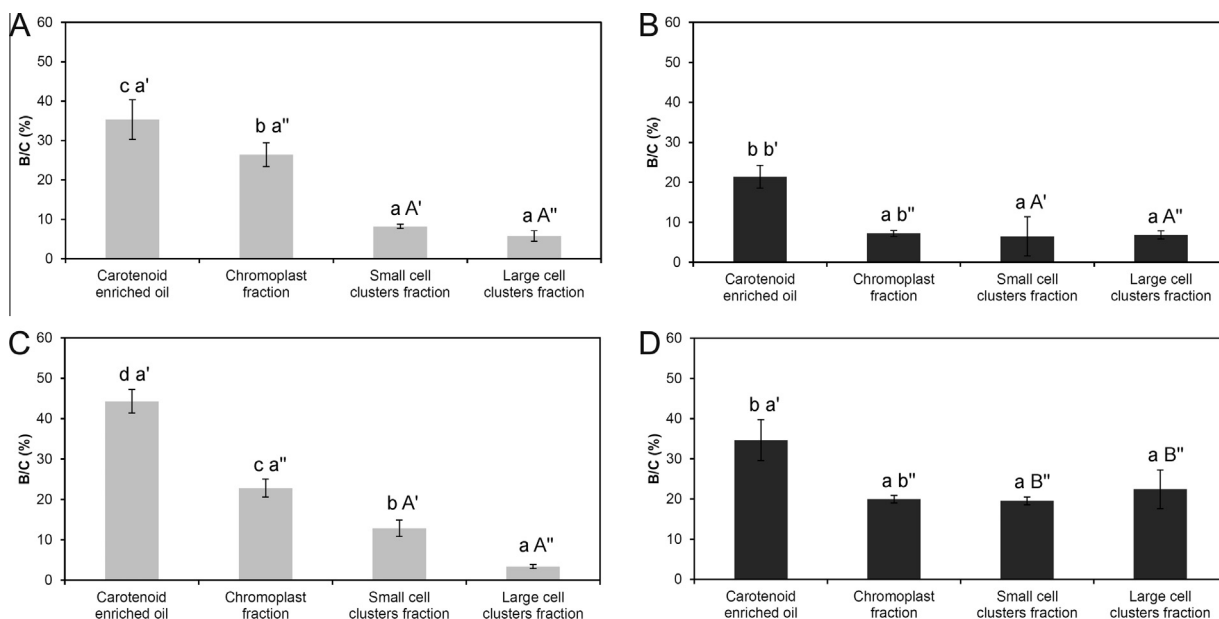


Fig. 2. Percentage *in vitro* bioaccessibility (B/C) of lycopene (mean \pm standard deviation) in fractions of red carrots (A) and of red tomatoes (B), β -carotene (mean \pm standard deviation) in fractions of orange carrots (C) and ζ -carotene (mean \pm standard deviation) in fractions of orange tomatoes (D). Lower case different letters represent significant differences ($P < 0.05$) among bioaccessibility values in fractions of the same matrix. Significant differences ($P < 0.05$) among bioaccessibility values in carotenoid-enriched oil, chromoplast, small cell cluster and large cell cluster fractions of the four matrices are represented by lower case different prime letters, lower case double prime letters, upper case prime letters and upper case double prime letters, respectively.

are important factors determining carotenoid bioaccessibility. This is in line with previous suggestions that pectin composition and the presence of other polysaccharides (which vary according to the matrix) in the cell wall influence the bioaccessibility of carotenoids by interacting differently with the target compounds (Castenmiller & West, 1998; Epriliati, D'Arcy, & Gidley, 2009; Parada & Aguilera, 2007).

In the literature, some important differences between carrot and tomato cell wall material, which may be related to differences in carotenoid bioaccessibility, can be found. For example, Jeffery, Holzenburg, et al. (2012) recently reported that the cell wall in carrots is very fibrous and compact and pectin may reduce cell wall porosity. In tomato, cell walls are thinner and less fibrous than in carrots, and present higher porosity due to their tendency to lose cellular adhesion. These suggestions are in line with our results. The cell walls in red carrots are fibrous and compact, therefore acting as an important barrier against lycopene bioaccessibility. By contrast, red tomato cell walls, due to their higher porosity, do not determine lycopene bioaccessibility, and the chromoplast substructure becomes the main barrier.

When the absolute values for the lycopene bioaccessibility between red carrot and red tomato fractions are compared, it can be observed that, for red carrot chromoplast fractions, the lycopene bioaccessibility was significantly higher ($P < 0.05$) than that for the fractions of red tomato (Fig. 2A and B). Fig. 3 represents the microscopic observations of chromoplast fractions of different matrices in this study. Red carrot (A) and red tomato (B) chromoplasts

contained crystalline carotene substructures, which have been observed previously too by Schweiggert et al. (2012), Jeffery, Holzenburg, et al. (2012), Kim, Rensing, Douglas, and Cheng (2010) and Shi and Le Maguer (2000). Among the different chromoplast morphologies, containing globular, tubular, reticulotubular, membranous and crystalline substructures, chromoplasts containing carotenoids in the solid crystalline state are known to result in the poorest bioaccessibility (Fleshman et al., 2011). However, although both matrices present the same chromoplast morphology, differences in lycopene *in vitro* bioaccessibility of the two chromoplast fractions indicate that, besides the crystalline state of the carotenoids, the chromoplast sub-organisation, as well as other matrix effects, probably also have an important effect on the subsequent solubility and incorporation into the micelles. Schweiggert et al. (2011) already indicated that the release of lycopene is different between several fruit matrices because of its different association with other chromoplast substructures, such as the thylakoid membranes and its localisation within the organelle. In the case of the two studied matrices studied here, the carrot root chromoplasts accumulate carotenoids inside the lumina of the thylakoid membranes while, in tomato fruits, they accumulate in membrane-shaped structures as carotenoid-protein complexes or are membrane bound (Egea et al., 2010; Shi & Le Maguer, 2000). Such characteristics may explain the differences in lycopene bioaccessibility among chromoplast fractions of red carrots and red tomatoes.

In the carotenoid-enriched oil fraction, the lycopene *in vitro* bioaccessibility value was highly improved for both matrices.

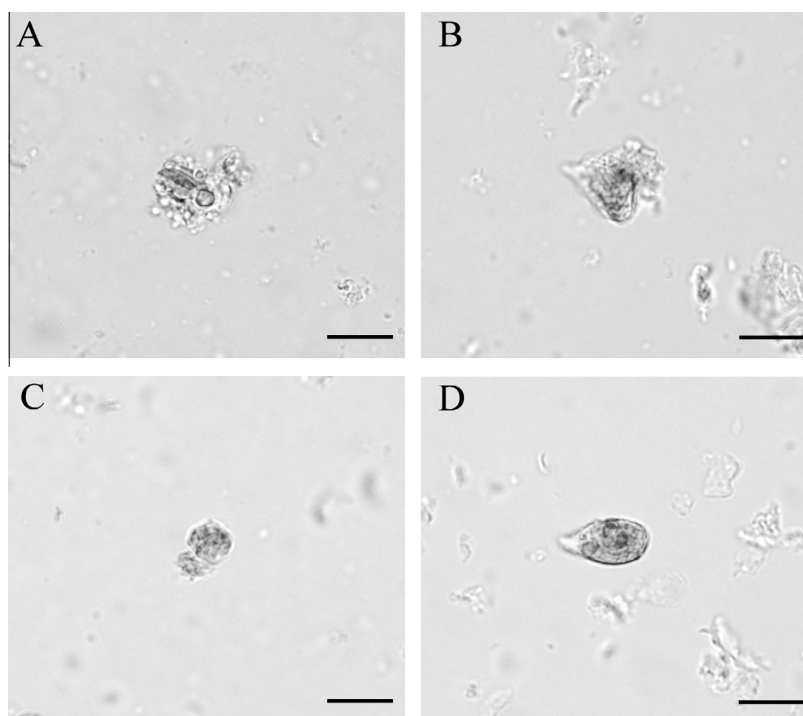


Fig. 3. Isolated chromoplasts from red carrot (A), red tomato (B), orange carrot (C) and orange tomato (D). Scale bars = 10 μm .

Crystalline structures of carotenoids found in tomatoes and carrot roots can be dissolved in oil, resulting in a higher bioaccessibility (Colle et al., 2012; Fernández-García, Mínguez-Mosquera, & Pérez-Gálvez, 2007; Vásquez-Cañedo, Heller, Neidhart, & Carle, 2006). Similar to the chromoplast fraction, the lycopene bioaccessibility was higher in the carotenoid-enriched oil isolated from red carrots, compared to red tomatoes. Solubilisation of carotenoids into the oil phase is greatly dependent on the polarity of the molecule (Nguyen, Francis, & Schwartz, 2001; Tyssandier, Lyan, & Borrel, 2001). Lycopene is a highly lipophilic molecule which, in order to be isolated from the matrix, has to be transferred first from the innermost physical barrier (chromoplast), across the cell wall and move through the aqueous environment surrounding the cells, which might be considered an additional physical barrier for carotenoid incorporation into the oil phase. Considering, the same type of carotenoid within different matrices, differences in matrix characteristics, previously described, may influence lycopene transfer through the chromoplast substructure and cell wall into the oil phase.

3.3. β -Carotene *in vitro* bioaccessibility in fractions of orange carrots

Orange carrot fractions (carotenoid-enriched oil, chromoplasts, small and large cell clusters) were prepared and the corresponding β -carotene *in vitro* bioaccessibility was measured.

Fig. 2C shows the values of the β -carotene *in vitro* bioaccessibility. An increased β -carotene bioaccessibility with a decreasing level of barriers can be observed. There was a significant difference ($P < 0.05$) in β -carotene *in vitro* bioaccessibilities among the four fractions.

Cell walls and chromoplast substructure were confirmed to be important physical barriers that hamper the release of β -carotene from the matrix, a behaviour which was also observed in red carrots. Once β -carotene is dissolved in the carotenoid-enriched oil fraction, the percentage of *in vitro* bioaccessibility increased six times compared to the value obtained for the large cell cluster

fraction. In the latter fraction, cell walls and chromoplast substructures were impeding the β -carotene solubilisation and incorporation into micelles.

Similar to the lycopene-rich chromoplasts from red carrots, chromoplasts from orange carrots, which contain β -carotene as the main carotenoid, showed a crystalline carotene substructure (Fig. 3C). Kim et al. (2010), Schweiggert et al. (2012), Jeffery, Holzenburg, et al. (2012) and Hornero-Méndez and Mínguez-Mosquera (2007) came to similar conclusions for orange carrots. The crystalline substructure, in the chromoplasts from orange carrots, is an important characteristic that determines the low β -carotene bioaccessibility. As mentioned in the previous section, from the different types of chromoplast morphologies, the crystalline carotene substructure presents the lowest carotenoid bioaccessibility.

When the β -carotene bioaccessibility in orange carrots is compared to the lycopene bioaccessibility in red carrots (same matrix, different type of carotenoid) in each matrix, no significant differences ($P < 0.05$) were found. This might indicate that the structural matrix of the two types of carrots is affecting the lycopene and β -carotene *in vitro* bioaccessibility in a similar way. Interestingly in this case, the differences of the molecular structure, and hence polarity, between lycopene and β -carotene in carrots do not seem to be a major factor affecting solubilisation into the micelles.

3.4. ζ -Carotene *in vitro* bioaccessibility in fractions of orange tomatoes

Orange tomato fractions (carotenoid-enriched oil, chromoplasts, small and large cell clusters) were prepared and ζ -carotene *in vitro* bioaccessibility was measured. Fig. 2-D shows the results of the ζ -carotene *in vitro* bioaccessibility for the fractions of orange tomatoes. No significant differences ($P < 0.05$) for the ζ -carotene *in vitro* bioaccessibility between chromoplast, small and large cell clusters fractions could be observed. Once ζ -carotene was solubilised in the oil fraction (absence of cellular material and chromoplast substructures), there was a significant enhancement ($P < 0.05$) of its bioaccessibility.

Similar to what was observed for the lycopene bioaccessibility in red tomatoes, the results suggest that the chromoplast morphology is more important in determining the ζ -carotene bioaccessibility than is the cell wall itself. In contrast to the chromoplast barrier, the removal of the cell walls did not result in a significant improvement of ζ -carotene bioaccessibility in orange tomato fractions.

It was observed by Rosso (1967) that chromoplast morphologies from different members of the family *Solanaceae* (to which tomatoes belong) are not the same. Using electron microscopy, chromoplast substructures from the Golden Jubilee tomatoes (with ζ -carotene and pro-lycopene as the major carotenoids) appeared as spheroidal globules and crystals. From our light microscopy observations, globules of various sizes in orange tomato chromoplasts were observed (Fig. 3D). The globules clearly differed from the high amount of crystals observed in red tomatoes (Fig. 3B). This structural difference may explain why the ζ -carotene *in vitro* bioaccessibility, in all fractions of orange tomatoes, was significantly higher ($P < 0.05$) than that for lycopene in fractions from red tomatoes. Probably, the crystalline substructures of the lycopene-rich chromoplasts from red tomatoes confer a high resistance to lycopene solubilisation and subsequent incorporation into micelles. It has been previously reported by Fleshman et al. (2011) that melon chromoplasts with globular substructures show a higher β -carotene bioaccessibility than do carrot root chromoplasts with crystalline substructures. Jeffery, Turner, et al. (2012) similarly concluded that globular carotenoids are the most prone to solubilisation during digestion. Schweiggert et al. (2012) also detected a higher bioaccessibility of β -carotene in mango and papaya, in which carotenoids are deposited in lipid elements in the chromoplasts, compared to tomatoes and carrots, where carotenoids are encountered as solid crystalline substructures.

The observations in orange tomatoes support the hypothesis that the chromoplast morphology and its substructures are important determinants of the carotenoid bioaccessibility. Additionally, differences in cell wall composition between orange and red tomatoes might play a role.

4. Conclusion

Several previous studies have investigated carotenoid bioaccessibility in different fruit- and vegetable-based systems. However, there is a remaining problem of the detailed understanding of carotenoid bioaccessibility in relation to (natural and/or process-induced) structural physical characteristics of the matrix. Our experiment was a targeted study, using specific separate fractions, in which the most important barriers governing carotenoid bioaccessibility (cell walls/membranes and chromoplast structure/organisation) were systematically considered. This allowed improved insight of how the different levels of carotenoid physical bioencapsulation affected their release, solubilisation and incorporation in micelles during digestion. The results of this experiment show an inverse correlation between the levels of carotenoid bioencapsulation and the carotenoid *in vitro* bioaccessibility. Observations of the bioaccessibility of different carotenoids in fractions of different matrices emphasise the importance of the type of carotenoid (although to a limited extent in this study) and the matrix-related effects (cell wall material composition and chromoplast morphology and substructure organisation) on carotenoid solubilisation and subsequent incorporation into the micelles.

It becomes evident that, depending on the food matrix considered, specific barriers are important. In the case of carrots (red and orange), the chromoplast substructure is an important barrier for carotenoid bioaccessibility, but the cell wall appears to be the most important barrier for obtaining a more significant improvement. For tomatoes (red and orange) the chromoplast substructure

proves to be the principal barrier affecting carotenoid bioaccessibility. Higher ζ -carotene *in vitro* bioaccessibility in fractions of orange tomatoes than in fractions of the other matrices studied, supports the hypothesis that crystalline chromoplast substructures (observed in red carrots, red tomatoes, and orange carrots) result in a lower carotenoid bioaccessibility.

Nevertheless, in order to obtain the highest increase in carotenoid bioaccessibility in the four matrices, it is necessary to solubilise carotenoids in the oil phase. Therefore, detailed knowledge of how to improve the transfer of carotenoids to the oil phase of food systems during processing and/or digestion is of key importance for developing foods with enhanced levels of bioaccessible carotenoids.

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