Effect of β-carotene-rich tomato lycopene β-cyclase (tlcy-b) on cell growth inhibition in HT-29 colon adenocarcinoma cells

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Lycopene β-cyclase (tlcy-b) tomatoes, obtained by modulating carotenogenesis via genetic engineering, contain a large amount of β-carotene, as clearly visible by their intense orange colour. In the present study we have subjected tlcy-b tomatoes to an in vitro simulated digestion and analysed the effects of digestate on cell proliferation. To this aim we used HT-29 human colon adenocarcinoma cells, grown in monolayers, as a model. Digested tomatoes were diluted (20 ml, 50 ml and 100 ml/l) in culture medium and added to the cells for different incubation times (24 h, 48 h and 72 h). Inhibition of cell growth by tomato digestate was dose-dependent and resulted from an arrest of cell cycle progression at the G0/G1 and G2/M phase and by apoptosis induction. A down-regulation of cyclin D1, Bcl-2 and Bcl-xl expression was observed. We also found that heat treatment of samples before digestion enhanced β-carotene release and therefore cell growth inhibition. To induce with purified β-carotene solubilised in tetrahydrofuran the same cell growth inhibition obtained with the tomato digestate, a higher amount of the carotenoid was necessary, suggesting that β-carotene micellised during digestion is utilised more efficiently by the cells, but also that other tomato molecules, reasonably made available during digestion, may be present and cooperate with β-carotene in promoting cell growth arrest.

β-Carotene-rich tomatoes: In vitro digestion: HT-29 cells: Cell cycle: Apoptosis

Retinol (vitamin A₂) and its metabolites are essential in a wide range of physiological regulatory processes(13). At the molecular and cellular level, this regulation involves the control of cell proliferation and differentiation through retinoid-dependent effects on gene expression(13). Vitamin A deficiency is still a public health problem worldwide(3,4). Since animals are not able to synthesise retinoids de novo, they must derive them from dietary vitamin A and provitamin A carotenoids. β-Carotene is the most potent precursor of vitamin A, and therefore efforts to manipulate the genetic background of plants widely used as food to produce β-carotene have received considerable attention(5–7). Tomato fruit is characterised by the high content of lycopene that, although not a vitamin A precursor, can be readily cyclised to β-carotene by the plant enzyme lycopene β-cyclase. Some authors(8) have succeeded in modulating the carotenogenesis in tomato fruit to induce accumulation of β-carotene. Tomato plants were transformed with tomato lycopene β-cyclase (tlcy-b) cDNA under the control of cauliflower mosaic virus 35S promoter, and acquired the ability of converting almost all lycopene contained in the fruits into β-carotene. β-Carotene is not only a vitamin A precursor, but performs in animals other functions related to its chemical structure, mainly acting as an antioxidant(9,10). The consumption of a diet rich in fruit and vegetables, and therefore in carotenoids, has been associated with a decreased risk of certain types of cancer and cardiovascular disease(11,12). Moreover, tomatoes contain not only lycopene, but also other carotenoids, vitamins and other molecules that, possibly in combination, can contribute to their positive effects on health.

Human intestinal cells cultured in vitro represent very useful models to study the effects of carotenoids on the barrier between the external world and the inside of the human organism(13). In a previous study(14) we utilised an in vitro model consisting of the adenocarcinoma colon cells HT-29 and HCT-116, that still exhibit several morphological and biochemical characteristics of intestinal cells and that accumulate carotenoids, to investigate the potential mechanisms underlying the anti-tumoural effects of tomatoes reported in the literature(15,16). In order to deliver tomatoes to cells in culture, an in vitro digestion procedure mimicking the physiological in vivo function has been adapted and utilised(17). The resulting digestate was then added to the cell cultures at different concentrations and for various lengths of times.

The aim of the study reported in the present paper was to utilise HT-29 human colon adenocarcinoma cells cultured in monolayers to investigate the effects of the β-carotene-rich...
tomato *tlcy-b* digestates on cell growth and on the expression of genes that regulate the cell cycle and apoptosis, in order to understand the molecular mechanisms that govern cell growth inhibition by β-carotene liberated during digestion together with other molecules from the food matrix represented by the β-carotene-rich *tlcy-b* tomato.

The results reported demonstrate that the molecules liberated from digested *tlcy-b* tomatoes by *in vitro* simulated digestion inhibit the growth of HT-29 colon cancer cells by interfering with cell cycle progression and apoptosis, in particular by modulating the expression of some key regulator proteins. To induce the same rate of cell growth inhibition with purified β-carotene solubilised in tetrahydrofuran, HT-29 cells must be treated with a higher amount of β-carotene; it is therefore evident that β-carotene provided as *tlcy-b* digestate is more active in the anti-proliferative action on HT-29 cells than purified β-carotene, either for the efficient micellarisation that takes place during digestion, and for the presence and cooperation of other molecules(18).

**Materials and methods**

**Preparation of tomato samples and in vitro simulated digestion**

The GM *tlcy-b* tomato was obtained by transformation of the Red Setter tomato with *Agrobacterium tumefaciens*, harbouring the tomato lycopene β-cyclase (*tlcy-b*) cDNA, under the control of cauliflower mosaic virus 35S promoter. The transgenic tomato fruits acquired the ability to convert almost all lycopene into β-carotene, and to increase the total carotenoid content(8). To prepare samples, *tlcy-b* frozen tomatoes were chopped and homogenised with a B-400 Buchi mixer (Buchi Labortechnik AG, Flawil, Switzerland).

All manipulations with tomato samples were performed under subdued lighting and in amber glass bottles to minimise the destruction of carotenoids. Samples of 1 g homogenised tomato were mixed with 1:8 ml saline (140 mM-NaCl, 5 mM-KCl, 150 μM-butylated hydroxytoluene in tetrahydrofuran), and then hand-homogenised in a Teflon/glass Potter homogeniser. *In vitro* simulated digestion was performed according to Garrett et al. (17) with modifications. Briefly, after homogenisation, samples were acidified to pH 2.0 with 1 M-HCl before the addition of 50 μl of pepsin, from porcine stomach mucosa (0.2 g pepsin in 5 ml of 0.1 M-HCl), and samples were incubated in a shaking water-bath for 60 min at 37°C.

After gastric digestion, the pH was raised to 6.9 with 1 M-NaHCO₃. Intestinal digestion was simulated by the addition of 200 μl of pancreatin–bile solution from porcine pancreas (0.45 g porcine bile extract and 0.075 g pancreatin in 37.5 ml of 0.1 M-NaHCO₃) and incubated in a shaking water-bath at 37°C for 120 min. The pH of the samples was then adjusted to 7.5. A sample of homogenate was subjected to heat treatment before gastric digestion by boiling for 15 min in a water-bath.

At the end of the digestion procedure, samples were centrifuged at 12 000 rpm for 30 min at 4°C in a Sorvall SS-34 angle rotor, and the supernatant fractions were collected and stored at −80°C.

**Cell culture**

HT-29 human colon adenocarcinoma cells (American Type Culture Collection (ATCC), Rockville, MD, USA) were grown in minimum essential medium. Cells were maintained in log phase by seeding them twice per week at the density of 3 × 10⁴ cells/ml at 37°C under a 5% CO₂–air atmosphere. The medium was supplemented with 10% (v/v) fetal calf serum (Flow, Irvine, North Ayrshire, UK) and 2 mM-glutamine and was not further replaced throughout the experiment. Experiments were routinely carried out on triplicate cultures. After the incubation, cells were harvested and quadruplicate haemocytometer counts were performed. The trypan blue dye exclusion method was used to evaluate the percentage of viable cells.

The Caco-2 cell line was obtained from the ATCC. The cells were grown and maintained in Dulbecco’s modified Eagle’s medium (DMEM) without pyruvate, containing 25 mm-glucose and NaHCO₃ (3.7 g/l) (DMEM; Biochrom, KG, Berlin, Germany), supplemented with 4 mM l-glutamine (Euroclone, Wetherby, West Yorks, UK) and 1% non-essential amino acids (Euroclone). Cells were left to differentiate for 15 d after confluence. Differentiated Caco-2 cells were incubated for 24 h in the presence or in the absence of *tlcy-b* tomato digestate (100 ml/l). Cells were then removed from the dishes, stained with trypan blue and counted under a microscope for viable and dead cells.

**Cell growth-inhibition assay**

Tomato digestates were serially diluted in cell culture medium as reported in the figure legends. Cells were seeded in twenty-four-well plates with 3 × 10⁴ cells/well and maintained for 24 h before any treatment to facilitate their adhesion on the well. The control group consisted of cells treated with the same amount of digestion mixture, for example, proteolytic and lipolytic enzymes without tomato, diluted with culture medium and of cells without any treatment. Since no differences in terms of viability, cell cycle, caspase activity and cell cycle-related proteins were found between the two control groups, control cells are referred to cells without any treatment. At the time indicated, cells were removed from the wells, stained with trypan blue and counted under a microscope for viable and dead cells.

**Cell cycle analysis**

Cell cycle distribution was analysed by flow cytometry, as previously described(19). Samples of 10⁶ cells were harvested by centrifugation, washed in PBS and fixed with ice-cold 70 % ethanol. The cells were incubated at 4°C for 30 min and then centrifuged at 2500 g for 10 min. The pellet was resuspended in 0.5 ml PBS and 0.5 ml DNA-Prep stain (Coulter Reagents, Miami, FL, USA), containing RNAse (1 g/l) and propidium iodide (50 g/l). All samples were incubated for 30 min in the dark at 4°C. The DNA content of cells stained with propidium iodide was measured with a FACS instrument (EPICS XL-MCL Flow Cytometer; Coulter Electronics, Miami, FL, USA), by using Multicycle AV software (Phoenix Flow Systems, San Diego, CA, USA).
Apoptosis detection by the terminal deoxynucleotidyl transferase-mediated dUTP nick end labelling method

The percentage of apoptotic cells was determined by terminal deoxynucleotidyl transferase-mediated dUTP nick end labelling (TUNEL). Briefly, cells were centrifuged, fixed with acetone, and incubated for 5 min with the hybridisation buffer (Boehringer-Mannheim, Germany). Then 2-5 units of terminal deoxynucleotidyltransferase (Tdt) and 100 pmol biotin–dUTP in hybridisation buffer were added and incubated for 1 h at 37°C. Thereafter, cells were incubated with streptavidin–biotin–peroxidase complex for 30 min at room temperature. The sites of peroxidase binding were detected with diaminobenzidine. The percentage of TUNEL-positive apoptotic cells (labelling index, LI %) was counted at x 40 magnification. In the absence of Tdt, no unspecific staining was observed. For each slide, three randomly selected microscopic fields were observed and at least 100 cells per field were evaluated.

Caspase-3 activity assay

The extent of apoptosis was also measured by the caspase-3 activity assay as previously described (20). Briefly, after a 24 h treatment, cells (2 × 10⁶) were lysed in 50 mM-2-amino-2-hydroxymethylpropene-1,3-diol–HCl buffer (pH 7.5) containing 0·5 mM-EDTA, 0·5% IGEPAL® Ca-630 and 150 mM-NaCl. Then the cell lysate was incubated with 50 μM-fluorogenic substrate, N-acetyl-Asp-Glu-Val-Asp-(7-amino-4-methylcoumarine) (Ac-DEVD-AMC; Alexis Biochemicals, San Diego, CA, USA), in a reaction buffer (10 mM-HEPES (pH 7.5) containing 50 mM-NaCl and 2·5 mM-dithiothreitol) for 120 min at 37°C. The release of AMC was measured with excitation at 380 nm and emission at 460 nm using a fluorescence spectrophotometer.

Western blot analysis

Cells (10 × 10⁶) were harvested, washed once with ice-cold PBS and gently lysed for 30 min in ice-cold lysis buffer (1 mM-MgCl₂, 350 mM-NaCl, 20 mM-HEPES, 0·5 mM-EDTA, 0·1 mM-ethylene glycol tetra-acetic acid, 1 mM-dithiothreitol, 1 mM-Na₂P₂O₇, 1 mM-phenylmethanesulfonyl fluoride, 1 mM-aprotinin, 1·5 mM-leupeptin, 1 mM-Na₃VO₄, 20% glycerol, 1 mM-MgCl₂, 350 mM-NaCl, 20 mM-HEPES, 0·5 mM-EDTA, 0·1 mM-ethylene glycol tetra-acetic acid, 1 mM-dithiothreitol, 1 mM-Na₂P₂O₇, 1 mM-phenylmethanesulfonyl fluoride, 1 mM-aprotinin, 1·5 mM-leupeptin, 1 mM-Na₃VO₄, 20% glycerol, 1% NP40). Cell lysates were centrifuged for 10 min at 4°C (10,000 g) and the supernatant fractions were used for Western Blot analysis. The anti-cyclin D1 (clone 72-13G; catalogue no. SC-126), anti-Bcl-2 (clone: Bcl-2/100/D5; catalogue no. SC-1041) and anti-p53 (clone DO-1; catalogue no. SC-126) monoclonal antibodies were purchased from YLEM (Rome, Italy). Blots were purchased from Santa Cruz Biotechnology (Santa Cruz, CA, USA). The anti-Bcl-2 (clone: Bcl-2/100/D5) monoclonal antibody was purchased from YLEM (Rome, Italy). Blots were washed with PBS and exposed to horseradish peroxidase–labelled secondary antibodies (Amersham Pharmacia Biotech, Arlington Heights, IL, USA) for 45 min at room temperature. Immunocomplexes were visualised by the enhanced chemiluminescence detection system (ECL). Western Blotting

Analysis System; ECL, Amersham, Bucks, UK) and quantified by densitometric scanning.

β-Carotene extraction and assay

β-Carotene was extracted with 1 volume of methanol and 3 volumes of hexane–diethyl ether (1:1) from 10 × 10⁶ cells after the different times of treatment and analysed by HPLC, as described previously (21).

Samples were dissolved in methanol and a 20 μl sample was analysed by reverse-phase HPLC with spectrophotometric detection on a Perkin-Elmer LC-295 detector at 450 nm. The column was packed with Alttech C18 Adsorbosphere HS material, 3 μm particle size, in a 15 × 0·46 cm cartridge format (Alttech Associates, Deerfield, IL, USA). A 1 cm cartridge precolumn, containing 5 μm C18 Adsorbosphere packing was used. Analyses were done by gradient elution; the initial mobile phase was 85% acetonitrile–15% methanol, with the addition of 30% 2-propanol at 8 min. Ammonium acetate (HPLC grade, 0·01%) was added to the initial mobile phase.

Statistical analysis

Three separate cultures per treatment were utilised for analysis in each experiment. Values are presented as mean values with their standard errors. One-way ANOVA was adopted to assess differences among the concentrations. When significant values were found (P<0·05), post hoc comparisons of means were made using Fisher’s test. Differences were analysed using Minitab Software (Minitab, Inc., State College, PA, USA). Multifactorial two-way ANOVA was adopted to assess any differences among the treatments and the times (Fig. 1(B), Fig. 6). When significant values were found (P<0·05), post hoc comparisons of means were made using the honestly significant differences test.

Results

The amount of β-carotene transferred to the aqueous fraction during the digestion procedure (8·7 (SEM 0·8) μg/g fresh weight) was less than one tenth of the amount of β-carotene originally contained in the homogenate (140·0 (SEM 14·5) μg/g fresh weight). Moreover, heat treatment of tomato homogenate before in vitro digestion induced, as demonstrated also in other studies (21), a three-fold increase in the carotenoid content, reaching the value of 22·0 (SEM 0·3) μg/g fresh weight. This is in agreement with other findings according to which processing of vegetable foods before consumption enhances to a lesser or greater extent – depending on treatment and on carotenoid type – carotenoid release (22).

Fig. 1 shows the effects of tclt-l tomato digestate on the growth of HT-29 cells, as measured by direct cell counting. Fig. 1(A) shows the effects of varying digestate concentrations in cells treated for 24 h and Fig. 1(B) shows the effects of 100 ml/l digestate in cells treated for different periods of time. tclt-l Tomato digestate inhibited the growth of HT-29 cells in a dose-dependent manner. After 24 h of incubation, a cell growth inhibition of about 10, 25 and 55% was observed following the addition of tclt-l tomato digestate to the culture medium at the concentrations of 20, 50 and
100 ml/l, respectively (Fig. 1(A)). A clear inhibition of cell growth was evidenced after 24 h of incubation following the addition of digestate at the concentration of 100 ml/l, which was not significantly modified prolonging the incubation time until 72 h (Fig. 1(B)). The effect seems to be specific for tumour cells, since differentiated Caco-2 cells were not affected in cell viability by addition of digestate at the concentration of 100 ml/l, which growth was evidenced after 24 h of incubation following the interaction was significant ($P<0.05$).

Since it has been previously reported that carotenoids act as strong apoptotic inducers in tumour cells, we examined whether the reduction in cell number was associated with apoptosis induction.

Apoptosis induction by tcly-b tomato digestate was studied by analysing the percentage of TUNEL-positive cells (Fig. 2(A)) and the activation of caspase-3, one of the most important cell death executioners for apoptosis (Fig. 2(B)). We found that a 24 h treatment with the digestate resulted in both an increase in the percentage of TUNEL-positive cells and in a strong dose-dependent increase in 7-amido-4-methylcoumarin fluorescence, indicative of the activation of caspase-3 in HT-29 cells.

Consistent with these results, analysis of the cell cycle progression showed by the accumulation of cells in both the G0/G1 and G2/M phase and by a concomitant decrease in the percentage of cells in the S phase.

To explore the effects of tcly-b tomato digestate on apoptosis-regulating proteins, we examined the expression of Bcl-2 and Bcl-xl, which are known to suppress programmed cell death, and that of Bax, which promotes it, in HT-29 cells (Fig. 3). Treatment with tomato digestate significantly reduced the expression of both Bcl-2 and Bcl-xl in a dose-dependent manner. In contrast, no significant changes in the expression of Bax were found under the same conditions.

Fig. 4 shows the effect of increasing β-carotene concentrations on cell growth (Fig. 4(A)), caspase-3 activation (Fig. 4(B)), and Bcl-2 (Fig. 4(C)) and Bcl-xl (Fig. 4(D)) content following a 24 h treatment. Carotenoid treatment inhibited cell growth and induced caspase-3 activation in a dose-dependent manner. Such effects were also accompanied by a decreased Bcl-2:actin and Bcl-xl:actin ratio.

When the activity of tcly-b tomato digestate, containing in our experimental conditions β-carotene at the concentrations of 0.08, 0.16 and 0.8 μM corresponding to 20, 50 and 100 ml/l, respectively, was compared with that of purified β-carotene at concentrations ranging from 0.5 to 10 μM (Fig. 5), the growth-inhibitory effect of the digestate was remarkably higher than that of the purified β-carotene, as highlighted by the different percentage of cell growth inhibition.

As shown in Fig. 6, HT-29 cells incorporated and/or associated β-carotene linearly. A time-dependent linear increase in β-carotene content was observed in cells treated for 24 h with both 100 ml/l digestate and purified β-carotene at the concentration of 0.8 μM, which corresponds to the digestate concentration of 100 ml/l.
carotenoid amount. Purified β-carotene was incorporated and/or associated to a greater extent than β-carotene in the digestate. The cellular carotenoid amount increased in a dose-dependent manner after the addition of 20, 50 and 100 ml/l digestate, respectively, for 24 h (data not shown).

**Discussion**

In the present study we investigated the growth-inhibitory effect of the β-carotene-rich tlc-y-b tomato, obtained by the modulation of carotenogenesis in tomato fruits via genetic engineering(9), on human intestinal cancer cells cultured in vitro. Colon cancer is one of the most common causes of death in Western populations(24), epidemiological and experimental data have shown that a high dietary intake of fruit and vegetables rich in carotenoids is associated with a lower risk of colon neoplasia(25-27). Although some human intervention trials failed to demonstrate the prevention of colon cancer by β-carotene supplements(28-30), other interventions showed a significant protective effect of β-carotene(31), a reduced rate of colon cell proliferation in patients with adenomatous polyps(32) and protective effects against colon carcinogenesis in animal models(33,34). More recently, growth-inhibitory effects by carotenoids in human colon cancer cell lines were also reported by our group(35,36).

The model system utilised in the present study consisted of HT-29 human adenocarcinoma colon cells treated with different amount of tlc-y-b tomato subjected to an in vitro simulated digestion(12). During the in vitro digestion, thanks to the action of pancreatic enzymes and bile salts, at specific pH conditions, bioactive molecules are transferred from the food matrix to the aqueous micellar fraction, becoming available for cell uptake. In vitro simulated digestion has been proven to represent a very useful and efficient approach to deliver molecules to cells in culture, quite similar to the physiological process that takes place in vivo in the gastrointestinal tract. This method has been utilised for assessing bioaccessibility and bioavailability of Fe(37), amino acids(38), carotenoids from baby food meals(17), and of other molecules. It has also been recently applied by us(14) to the study of the effects of tomato lycopene on the growth of colon cancer cells. We have shown that ripe tomato, containing essentially lycopene, is an anti-proliferative agent not only in prostate cancer cells(39), but also in colon cancer cells such as HT-29. In fact, tomato digestate inhibited cell cycle progression through the decrease of cyclin D1 expression and induced apoptosis through the modulation of Bcl-2 and Bcl-xl proteins. This effect seems to be specific for colon cancer cells since, as it has been reported in the Results section, differentiated Caco-2 cells, representing a suitable model of absorbing intestinal cells(23), are not affected in their viability by digestate in the range of concentrations used in the present study. This observation is in agreement with previous findings, comparing the effects of β-carotene in normal and tumour cells(40). In particular, in a previous study, we reported that β-carotene was much more potent in inhibiting the growth and altering the viability in undifferentiated than in dimethyl sulfoxide-differentiated HL-60 cells(41).

Lycopene and β-carotene exert a significant antioxidant activity, but their anti-proliferative action, tested in different cell types, has been attributed also to other specific molecular processes to which these molecules take part(42). For instance, it has been demonstrated in prostate cancer cells that lycopene regulates the expression of specific proteins, such as connexin 43, involved in proper cell—cell communication and therefore in the control of cell proliferation(43). β-Carotene is the main precursor of vitamin A, a well-characterised morphogen that, once metabolised to retinoic acid, modulates the expression of hundreds of genes involved in embryogenesis, cell growth and differentiation(44).

**Table 1. Effect of tomato lycopene β-cyclase (tlcy-b) tomato digestate on cell cycle distribution in HT-29 cells (%)**

(Mean values of three different experiments with their standard errors)

<table>
<thead>
<tr>
<th>Phase...</th>
<th>G0/G1</th>
<th>S</th>
<th>G2/M</th>
<th>Sub-G1</th>
</tr>
</thead>
<tbody>
<tr>
<td>Treatment</td>
<td>Mean</td>
<td>SEM</td>
<td>Mean</td>
<td>SEM</td>
</tr>
<tr>
<td>Control</td>
<td>47.0±5</td>
<td>3.0</td>
<td>41.0±5</td>
<td>2.9</td>
</tr>
<tr>
<td>T (20 ml/l)</td>
<td>47.8±5</td>
<td>2.9</td>
<td>39.5±5</td>
<td>3.0</td>
</tr>
<tr>
<td>T (50 ml/l)</td>
<td>53.3±5</td>
<td>3.5</td>
<td>27.8±5</td>
<td>3.0</td>
</tr>
<tr>
<td>T (100 ml/l)</td>
<td>58.5±5</td>
<td>4.2</td>
<td>15.3±5</td>
<td>2.5</td>
</tr>
</tbody>
</table>

Control, vehicle control-treated cells; T, tlc-y-b tomato digestate-treated cells.

a,b,c Mean values within a column with unlike superscript letters were significantly different (P<0.05; Fisher’s test).
We have investigated in the present study the mechanisms involved in the growth-inhibitory effects of molecules, mainly β-carotene, released from the *tly*-b tomato during the *in vitro* digestion treatment and probably presented to the cells in micelles(45). Only a small amount of β-carotene, less than 10%, is released during digestion; in agreement with the observation of other authors(46), this amount increases more than three times when the tomato homogenate is subjected to heat treatment before digestion.

The mechanisms by which β-carotene inhibits the growth of colon cancer cells is still not completely understood. It was shown that apoptosis is involved, since this carotenoid interferes with cell cycle progression(20). In the present paper we have investigated and elucidated in HT-29 human colon adenocarcinoma cells the effects of *tly*-b tomato digestate on individual steps of the cell cycle and some key proteins involved in this activity.

Since it has been previously reported(41) that carotenoids are strong apoptotic inducers in cancer cells, we investigated whether the inhibition of proliferation caused by digested *tly*-b tomato was associated with induction of apoptosis. By both methods utilised, TUNEL-positive cells and activation of caspase-3, it was demonstrated that *tly*-b tomato digestate indeed induced apoptosis. In parallel, the expression of Bcl-2 and Bcl-xl, inhibitors of programmed cell death, was decreased. On the other hand, the pro-apoptotic protein Bax was not affected by carotenoid treatment. It is well known that the ratio of the anti-apoptotic:pro-apoptotic Bcl-2 family proteins, more than the level of the individual proteins, is extremely important in modulating the apoptotic process(47). It has been demonstrated that β-carotene is able to decrease the expression of Bcl-2 and Bcl-xl in colon cancer cells(20,48) and to diminish that of Bcl-2 in HL-60 cells(41). Such effects were strictly related to apoptosis induction and to reactive oxygen species production by the carotenoid. This finding is particularly interesting in the light of the data supporting a role for Bcl-2 in an antioxidant pathway.

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**Fig. 4.** Effect of β-carotene on cell growth (A), caspase-3 activation (B), Bcl-2 (C) and Bcl-xl (D) content in HT-29 cells following a 24 h treatment. Values are the means of three different experiments, with standard errors represented by vertical bars. *ABC* Mean values with unlike letters were significantly different (*P*<0.05; Fischer's test).

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**Fig. 5.** Comparison of β-carotene (●) and tomato lycopene β-cyclase (*tly*-b) tomato digestate (■) ability in inhibiting HT-29 cells growth after 24 h treatment. Concentrations of *tly*-b tomato digestate of 20, 50 and 100 ml/l correspond to β-carotene concentrations of 0.08, 0.16 and 0.8 μM, respectively.

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**Fig. 6.** β-Carotene incorporation and/or association in HT-29 cells treated with tomato lycopene β-cyclase (*tly*-b) tomato digestate (■) and purified β-carotene (●) for 24 h. The concentration of digestate in cultured medium was 100 ml/l and that of the purified carotenoid was 0.8 μM. Values are the means of three different experiments, with standard errors represented by vertical bars. *Mean value was significantly different from that of the digestate-treated cells (*P*<0.005; Tukey’s test).
whereby this protein prevents programmed cell death by decreasing the formation of reactive oxygen species and lipid peroxidation products (69).

In addition, data from other studies suggest that carotenoids modulate pro-apoptotic Bcl-2 family proteins, including Bad (50), Bid (19) and Bax (51) in different experimental models. In particular, it has been recently shown that the increase in Bax expression and in apoptosis induction caused by cigarette tar was prevented by the addition of β-carotene in Rat-1 fibroblasts as well as in different tumour cell lines (51). On the other hand, an increased expression of Bax protein by β-carotene has been recently reported in U-937 cells and human umbilical vein endothelial cells by micro-array analysis (52, 53).

Since apoptosis is the end point of some epithelial colonic cell differentiation pathways, a process that induces apoptosis should also reduce proliferative signals. In fact, the inhibition of growth and the increased expression of Bcl-2 and Bcl-xL was associated with the slowdown of cell cycle progression at the G0/G1 and G2/M phase.

HT-29 cells accumulate β-carotene in a linear dose-dependent way (Fig. 6). It is crucial to remark that the growth-inhibitory action of 1,3c-cb tomato digestate is active at a concentration much lower than that of purified β-carotene, although purified β-carotene was incorporated to a greater extent into the cells. Tomatoes contain several bioactive compounds, mainly lycopene but also other carotenoids such as β-carotene, phytotoxins, phytotoflueins, their metabolites and oxidative products, as well as vitamin E, vitamin C and polyphenols (10). 1,3c-cb Tomatoes are engineered to be very rich in β-carotene; however, as underlined above, other molecules, including a very small amount of lycopene, are present that are probably released by digestion and that can contribute in cooperation with β-carotene to the effects described.

It is also important to stress that micellisation is a more physiological method to solubilise and deliver molecules, in particular lipophylic molecules, compared with the use of a solvent. In conclusion, the results obtained with this method point out that the whole digested food, containing different bioactive molecules at different relative concentrations, their metabolites, some minor components and the matrix itself, provides a unique mixture with a specific function in maintaining or protecting health.

In conclusion, simulated digestion is confirmed as a valid method to perform in vitro nutritional studies. The next crucial effort will be the analysis of the complete pattern of molecules involved in the effect observed and their interaction.

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D. B. and P. P. performed most of the experiments and contributed equally to the discussion and to writing the manuscript; R. S. performed HPLC determinations; A. B. performed the cytotoxicity assays; F. C. produced the 1,3c-b tomato and provided the homogenates; G. M., as coordinator of the project ‘GMO in Agriculture’, participated in the research planning and discussion of results; S. G. coordinated the research and contributed to the planning, realisation and discussion of the work.

P. P. and D. B. contributed equally to the present study. There is no conflict of interest that should be disclosed.

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