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Betanin-enriched red beetroot (*Beta vulgaris L.*) extract induces apoptosis and autophagic cell death in MCF-7 cells

Short title: *In vitro* antitumor activity of betacyanins

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ABSTRACT

Recent studies have pointed out the preventive role of beetroot extracts against cancers and their cytotoxic activity on cancer cells. Among many different natural compounds, these extracts contained betanin and its stereoisomer isobetanin which belong to the betalain group of highly bioavailable antioxidants. However, a precise identification of the molecules responsible for this tumor-inhibitory effect was still required. We isolated a betanin/isobetanin concentrate from fresh beetroots, corresponding to the highest purified betanin extract used for studying anticancer activities of these molecules. The cytotoxicity of this betanin-enriched extract was then characterized on cancer and normal cells and we highlighted the death signalling pathways involved. Betanin/isobetanin concentrate significantly decreased cancer cell proliferation and viability. Particularly in MCF-7-treated cells, the expressions of apoptosis-related proteins (Bad, TRAILR4, FAS, p53) were strongly increased and the mitochondrial membrane potential was altered, demonstrating the involvement of both intrinsic and extrinsic apoptotic pathways. Autophagosome vesicles in MCF-7-treated cells were observed, also suggesting autophagic cell death upon betanin/isobetanin treatment. Importantly, the betanin-enriched extract had no obvious effect towards normal cell lines.

Our data bring new insight to consider the betanin/isobetanin mix as therapeutic anticancer compound, alone or in combination with classical chemotherapeutic drugs, especially in functional p53 tumors.

Keywords: betanin concentrate, isobetanin, betalains, anticancer activity, 2D and 3D cultures

INTRODUCTION

With surgery and radiotherapy, chemotherapy is a major treatment for cancer. Chemical agents act on cancer cells at different levels and trigger cell destruction *via* the induction of an apoptotic program. However classic chemotherapeutic agents often target rapidly dividing cancer and normal cells, leading to side effects for the patients (Rao *et al.*, 2013). As the cancer incidence in worldwide tends to increase, the development of new, safer and efficient anticancer agents is a major challenge.

Certain plant-derived compounds exhibit anticancer activities and have attracted interest in preventing and treating cancers with over 60% of currently used anticancer agents derived from natural sources (Cragg and Newman, 2005; Gupta and Prakash, 2014). Beetroots contain both red (betacyanins) and yellow pigments (betaxanthins) known collectively as betalains, which constitute a class of highly bioavailable natural antioxidant pigments (Kanner *et al.*, 2001; Tesoriere *et al.*, 2004 and 2013). In the red beetroot species, the predominant betacyanins are betanin (Figure 1A), which represents between 75 and 90 % of the total pigments of red beetroot (Henry, 1996), and isobetanin, its C15 stereoisomer. In addition to the powerful antioxidant properties of betalains, studies have pointed out their preventive role against cancers (Kapadia *et al.*, 1996 and 2003; Lechner *et al.*, 2010). Nevertheless these studies were based on animal diet with red food coloring (E162) and other standard beetroot extracts, whose betanin content does not exceed 1.2%. Antitumoral activity of betalains was also evaluated *in vitro* on several cancer cell lines, but data obtained by the authors were disparate according to the cell lines and the cytotoxicity assays used (Reddy *et al.*, 2005; Sreekanth *et al.*, 2007; Paluszczak *et al.*, 2010). Moreover, the betanin content or the

purity degree of beetroot extracts used have never been precisely defined. In order to dispel any ambiguity, the characterization of the betanin effect on tumor cells requires a highly purified betanin concentrate with a precisely known composition.

During preclinical testing, *in vitro* analyses are performed to select compounds with potential antitumor activity before *in vivo* studies in relevant animal models. Cell culture approach is therefore of the highest importance for the initial screening of molecules. However cell lines used are conventionally cultured as monolayers (2D culture) and do not mimic the phenotype of cancer tissues due to the lack of cell-cell and cell-extracellular matrix interactions. 3D models (cell spheroids or aggregates) are obtained by preventing cells from attaching to the culture substrata. Based on their ability to mimic tissue-like structures more effectively than 2D cell cultures, the use of 3D *in vitro* models is believed to be a requisite mean into anticancer drug development (Breslin and O'Driscoll, 2013).

This study was carried out to clarify the antitumor effect of a betanin-enriched beetroot dried extract. We used both 2D and 3D cultures to assess the cytotoxic activity of betanin on cancer and normal cells. Our results showed that the betanin concentrate induces cancer cell growth inhibition, associated with an apoptotic cell death and an autophagic activity increase in MCF-7 cells, but has no obvious effect towards normal cells.

MATERIALS AND METHODS

1. Betanin extraction and purification

Grinded red beetroots (local market) were extracted into ethanol/water (80/20 v/v) solvent at a solid/liquid ratio of 1/3 (g/mL) for 1 h under continuous mechanical stirring. The solid material was separated from the macerate by centrifugation at 12 000 g for 15 min at 4°C followed by a filtration on a polypropylene membrane filter (0.2 µm, Pall Corporation). The filtrate was concentrated by ethanol evaporation under reduced pressure at 30°C and then enriched in betanin and isobetanin by adsorption chromatography (Stintzing *et al.*, 2002). A glass column was packed with 1 L of activated polymeric adsorbent (Amberlite XAD-16, Sigma-Aldrich, France). The pH of beetroot extract was adjusted to 3 with hydrochloric acid (HCl). 100 mL of extract was passed through the resin column at 1 Bed Volume (BV) per hour. After adsorption of the betacyanins, the other extract constituents were eluted using 3 BV of deionized water (resistivity > 18.2 mΩ.cm⁻¹, MilliQ plus unit, Merk-Millipore) acidified to pH 3 at 2 BV/h. Betanin and isobetanin were eluted with ethanol at 0.5 BV/h. The collected fraction was concentrated by ethanol evaporation under vacuum at 30°C and freeze-dried. The obtained betanin/isobetanin-enriched powder was stored at -80°C.

2. Extract characterization

Betanin quantification. All betacyanin/betaxanthin quantifications were performed using the multi-component photometric method described by J.H. von Elbe (von Elbe *et al.*, 2001). The pigment concentration calculations were based upon the absorptivity

values: $A^{1\%}$ 1120 for betanin and isobetanin (at 538 nm) and 750 for betaxanthins (at 477 nm).

Protein and total carbohydrate quantifications. Total carbohydrate amount present in extracts was determined using the DuBois method (DuBois *et al.*, 1956). Glucose was used as a standard. The protein quantification was performed using the Lowry method (Lowry *et al.*, 1951). The absence of pigment interferences with these two colorimetric methods was controlled.

HPLC/ESI-MS analysis. The HPLC system (Infinity 1290, Agilent Technologies, France) was equipped with diode array detector coupled with a Q-TOF micro hybrid quadrupole time of flight mass spectrometer (Agilent 6538, Agilent technologies, France). HPLC analyses were performed using an analytical scale (150 mm x 2.1 mm i.d.) hypersil gold C18 reversed phase column with a particle size of 3 μm (Thermo Scientific, France). Eluents A and B (LC-MS grade) consisted of 100% water with 0.2 % (v/v) formic acid and 100 % acetonitrile respectively. Betalains separation was achieved within 40 min at 20°C at a flow rate of 0.3 mL/min. The elution profile was: 0-3 min 100% A, 3-21 min 0-13% B in A (linear gradient), 21-24 min 13 % B, 24-30 min 13-50% B (linear gradient), 30-33 min 50-100 % B, 33-40 min 100 % B. Detection wavelengths were 477 nm (betaxanthins and betacyanins) and 538 nm (betacyanins only). Positive ion electrospray mass spectra were acquired by scan mode, consisted of scanning from m/z 50 to 1700 at electrospray voltage 3800 V, fragmentor voltage 140 V. Nitrogen was nebulized at 12.0 L/min under 45.0 psi and heated at 350°C. Betanin and isobetanin structural identities were confirmed by tandem mass (relative collision energy of 15 eV) spectroscopy and NMR studies. The ^1H NMR spectrum of betanin was

recorded in D₂O (data not shown) and corresponds to the previously published data (Stinzting *et al.*, 2004).

3. Cell culture and treatment

The mouse melanoma cell line B16F10 was a generous gift from Dr L. Larue (Institut Curie, France). Human breast cancer lines (MCF-7 ATCC[®] HTB-22, MDA-MB-231 ATCC[®] HTB-26), human colorectal cells (HT-29 ATCC[®] HTB-38) and normal human fibroblasts (MRC-5 ATCC[®] CCL-171) were obtained from the American Type Culture Collection. Human umbilical vein endothelial cells (HUVEC) were purchased from PromoCell.

Cells were cultured as monolayers in RPMI 1640 (B16F10), MEM (MCF-7, MDA-MB-231, MRC-5), DMEM (HT-29) or M199 (HUVEC). All cell culture media (Gibco, Invitrogen, France) were supplemented with 10% fetal bovine serum (FBS, Gibco, Invitrogen), 2 mM L-glutamine (Gibco, Invitrogen), penicillin (100 µg/mL, Gibco, Invitrogen) and streptomycin (100 µg/mL, Gibco, Invitrogen). All cell lines were maintained at 37°C in an air atmosphere of 10% CO₂ (B16F10) or 5% CO₂ (MCF-7, MDA-MB-231, HT-29, HUVEC, MRC-5).

Nunclon[®] polystyrene plates (tPS) were used for 2D cultures. Polyhydroxyethylmethacrylate (polyHEMA, Sigma Aldrich) coated polystyrene plates were prepared as previously described (Velzenberger *et al.*, 2008) and used for 3D cultures. Cells were seeded at 10000 cells/cm² without (controls) or with increasing concentrations of betanin/isobetanin (from 10 to 40 µM), and cultured for 24 or 48 hours.

4. Proliferation assay and cell cycle analysis

In 2D cultures, cells were removed by incubation for 5 min at 37°C in 0.25% trypsin + 1 mM EDTA solution (Gibco, Invitrogen). In 3D cultures, aggregates were collected by centrifugation (5 min, 200g) and dissociated using trypsin-EDTA solution. Trypsin reaction was stopped by adding half a volume of FBS, and cells were counted in a Malassez hemocytometer using trypan blue exclusion assay. The proliferative index was estimated as the ratio between viable cell counts 48 hour post-seeding and number of seeded cells. Each proliferative index was normalized to this one obtained for control cells and data were expressed as percentages.

For cell cycle analysis, harvested cells were washed twice in PBS containing 5 mM EDTA, and fixed for 45 min at 4°C in 1 mL 75% ethanol in PBS with 5 mM EDTA. Cell lines were washed twice and suspended in PBS, 5 mM EDTA containing 0.1% TritonX100 (Sigma Aldrich) mixed with 40 µg RNase A (Sigma Aldrich) and 25 µg propidium iodide (PI, Sigma Aldrich) and incubated for 15 min protected from light. The stained samples were analysed in Gallios flow cytometer (Beckman Coulter, France). Histograms were analysed using Wincycle software (PHOENIX flow systems, USA).

5. Apoptosis detection and quantification

To assess early stage of apoptosis, cells cultured in 2D and 3D conditions were treated with 30 µM betanin/isobetanin mix for 24 hours and labeled with Annexin-V-FITC (Beckman Coulter). For each cell line, a positive control for apoptosis was tested (cells treated with 1 µM staurosporine). The cytometer was adjusted according to the Annexin-V positive cells of these controls. According to the manufacturer's

instructions, cells were washed with PBS and suspended in the binding buffer at 5.10^5 to 5.10^6 cells/mL. 1 μ L of Annexin-V FITC solution and 5 μ L of PI solution were added to 100 μ L of cell suspension. Cells were incubated for 15 min on ice in the dark, and then diluted by adding 400 μ L of binding solution. The Annexin-V positive/IP negative cell populations were detected with a Gallios flow cytometer and were representative of apoptotic cells.

To detect DNA fragmentation, an Apostain binding assay (AbCys) was used. MCF-7 and HUVEC cells were treated with betanin/isobetainin (30 μ M) or staurosporine (1 μ M) for 48 hours, fixed with ice-cold methanol 80% (v/v) for 30 min, washed and suspended in 250 μ L formamide for 10 min at 75°C and for 5 min at room temperature (RT). Cells were then incubated for 45 min at RT with a mouse monoclonal antibody to single-stranded DNA (ssDNA, diluted 1:10 in PBS containing 5% FBS). Bound specific antibodies were revealed by incubation with Cy3-conjugated anti-mouse antibody (diluted 1:200 in PBS containing 1% milk, Jackson ImmunoResearch) for 30 min at RT. Each sample was counterstained with DAPI (4',6'-diamidino-2-phénylindole, 1 μ g/mL, Sigma-Aldrich). The cells were then cytocentrifuged (Shandon Cytospin®) and each slide was mounted in mowiol and examined by epifluorescence microscopy (Leica DMI6000). Ten fields were randomly chosen from each slide and the percentage of apoptotic cells in each picture was calculated as the ratio of the labeling cells to the total number of cells.

6. Apoptotic signaling pathway analysis

Human Apoptosis arrays (Proteome Profiler™ Array, R&D Systems) were used to simultaneously detect the relative levels of expression of 35 apoptosis-related proteins.

MCF-7, MDA-MB-231 and HUVEC cells were treated with betanin/isobetanin-enriched extract for 48 hours, rinsed with PBS and lysed for 30 min at 4°C using lysis buffer provided by the manufacturer. Lysates were then centrifuged at 14 000 g for 5 min and supernatants were harvested. Protein concentrations for each sample were determined using the Bradford method (Coomassie Protein Assay Reagent, Pierce, Interchim). 400 µg of cell lysates were incubated on each array overnight at 4°C. Arrays were incubated for 1h at RT with a reconstituted detection antibody cocktail, and then with a streptavidin-HRP solution for 30 min. Arrays were revealed with a Chemireagent Mix using ChemiDoc (Biorad), and each spot was quantified with Quantity One software (Biorad).

To detect change in the mitochondrial membrane potential, a MitoCapture™ dye (BioVision) was used. MCF-7 cells were treated with betanin/isobetanin mix (15 and 30 µM) for 24 hours, harvested and then resuspended in 1 mL diluted MitoCapture™ reagent provided by the manufacturer. Cells were incubated for 20 min at 37°C in a 5% CO₂ incubator. After centrifugation (500 g), the supernatant was discarded, the cells were suspended in 1 mL of pre-warmed incubation buffer and analyzed by epifluorescence microscopy.

7. Autophagy assay

Autophagy was detected using the Autophagy Detection Kit (Abcam). Briefly, MCF-7 cells were grown 24 hours on glass slides, and then were treated with betanin/isobetanin (15 and 30 µM) or with an autophagy inducer, rapamycin (500 nM provided by the manufacturer) for 24 hours. Medium was removed and cells were washed with a buffer supplied by the manufacturer. Cells were incubated with the dye (diluted at 1:500 in

assay buffer) and DAPI for 20 min at 37°C. Cells were carefully washed and slides were examined by epifluorescence microscopy.

8. Statistical analysis

All statistical evaluations were performed using GraphPad InStat software. Continuous variables are expressed as means \pm standard deviation. As most of the continuous values measured had a non-Gaussian distribution, non-parametric Dunn and Kruskal-Wallis tests were used for comparisons. A value of $p < 0.05$ was taken as significant.

RESULTS

1. Betanin/isobetanin concentrate characterization

Figure 1B shows the chromatogram corresponding to the pigment-enriched red beetroot dried extract obtained after purification by adsorption chromatography. The HPLC profile monitored at 477 nm combined with electrospray mass spectrometry in positive mode allows the identification of the betaxanthin and betacyanin molecules contained in this final extract (Stintzing *et al.*, 2004). Only two betalains were detected and identified as betanin ($t_R = 13.05$ min, $[M+H]^+ = 551$) and isobetanin, the 15R stereoisomer of betanin ($t_R = 14,51$ min, $[M+H]^+ = 551$) (Kujala *et al.*, 2002; Stintzing *et al.*, 2004). The betanin and isobetanin structures were confirmed by mass tandem spectroscopy. The positive ion spray mass spectrum (Figure 1C) shows the daughter ion produced by fragmentation of the parent ion of m/z of 551 $[M+H]^+ = 551$ assigned to betanin or isobetanin. The fragment ion at the mass charge (m/z) of 389 indicated that this ion is

obtained by glucose loss and corresponds to the protonated aglycones [betanidin + H]⁺ or [isobetanidin + H]⁺ (Castellanos-Santiago and Yahia, 2008).

All the betaxanthins were eluted during beetroot extract purification on polymeric resin while the betacyanins remained absorbed. Total carbohydrate quantification reveals that they are totally eliminated at this purification step, no residual complex or simple carbohydrates could be detected in the final betanin concentrate. As betanin and isobetanin differ only by the absolute configuration of their C15 chiral center, we did not distinguish them for determination of the total pigment amount in the purified extract. The total betanin/isobetanin content in the dried concentration was evaluated at 80% by photometric quantification. The remaining 20% was attributed to residual proteins by total protein content determination. Based on HPLC peak area values, the betacyanin composition was estimated at 64% of betanin and 36% of isobetanin. In the rest of the article, all the concentrations given correspond to the total betanin/isobetanin (Bet./IsoBet.) concentrations applied to the cells.

2. Effect of betanin/isobetanin on cell proliferation and morphology

We used both cancer (B16F10, MCF-7, MDA-MB-231, HT-29) and normal (HUVEC, MRC-5) cells to assess the biological effect of betanin extract. All cells were cultured as monolayer (2D) on tPS and as aggregates (3D) on anti-adhesive polyHEMA substratum, and then treated with betanin-enriched extract for 48 hours.

As shown in Figure 2A, Bet./IsoBet. treatment significantly decreased the proliferation of B16F10 and MCF-7 cells on both surfaces. The half maximal inhibitory concentration (IC₅₀) was determined for each cell lines, and was about 25 μM for B16F10 and MCF-7 cells. No viable cells were observed with 40 μM Bet./IsoBet. (data

not shown). MDA-MB-231 and HUVEC cells were also sensitive to betanin-enriched extract, but to a lower extent (IC₅₀ value was 35 μ M) (Figure 2B). The cell viabilities measured 48 hour post-treatment with 40 μ M Bet./IsoBet. decreased weakly: 35.6% \pm 17.2 (2D) and 52.0% \pm 19.2 (3D) for MDA-MB-231 and 75.2% \pm 26.9 (2D) and 71.0% \pm 27.4 (3D) for HUVEC cells (data not shown). For the same concentration range, HT-29 and MRC-5 cells were not sensitive to betanin concentrate: at 40 μ M, cell proliferation was not significantly decreased compared to controls (Figure 2C). Viabilities of HT-29 cells treated with 40 μ M Bet./IsoBet. for 48 hours were 95.2% \pm 5.3 (2D) and 96.6% \pm 5.2 (3D) (data not shown). For MRC-5 cells, viabilities remained high with betanin/isobetanin treatment: 87.7% \pm 9.3 (2D) and 76.8% \pm 12.8 (3D).

Cell morphologies were examined (Figure 2D): MCF-7 and HUVEC control cells cultured on tPS spread well and colonized the culture dishes, while cells on polyHEMA had a rounded aggregated morphology. Betanin-enriched extract reduced MCF-7 cell spreading on tPS and MCF-7 cell aggregation on polyHEMA, whereas a moderate effect of betanin/isobetanin concentrate was observed on HUVEC cells cultured in 2D and 3D conditions. No morphological change was observed for the HT-29 and MRC-5 cells treated with Bet.IsoBet. (data not shown).

Thus we identified three groups of cell lines with different sensitivities to betanin/isobetanin: B16F10 and MCF-7 as highly sensitive (no viable cells with 40 μ M Bet./IsoBet.); MDA-MB-231 and HUVEC as weakly sensitive (cell viabilities from 35% to 70% with 40 μ M Bet./IsoBet.); HT-29 and MRC-5 as resistant (cell viabilities about 90% with 40 μ M Bet./IsoBet.).

3. Effect of betanin/isobetanin concentrate on cell cycle distribution

To better understand betanin/isobetanin concentrate effect, we focused on **three human cell lines that represent three different and specific phenotypes: p53 wild-type cancer cell line MCF-7, p53-mutated cancer cell line MDA-MB-231 and normal HUVEC cells.**

The cell cycle progression was studied 48 hour post-seeding without (control) or with 30 μ M Bet./IsoBet. (Table 1). For each cell line, the cell cycles of control cultures on tPS and polyHEMA were significantly different: there were more cells in G1 phase and fewer in the S and G2/M phases in 3D cultures.

Betanin/isobetanin-enriched extract significantly decreased the percentage of MCF-7 cells in G1 phase (2D and 3D conditions) and increased percentage of cells in **S phase (2D culture) or G2/M phase (3D culture)**. For MDA-MB-231 and HUVEC cell lines, betanin/isobetanin concentrate decreased cell number in S phase (2D). No obvious effect of Bet./IsoBet. was observed on the 3D MDA-MB-231 cell cycle repartition. Betanin/isobetanin treatment decreased the 3D HUVEC cell number in G1 phase and increased the percentage of cells in S and G2/M phases.

4. Characterization of MCF-7 cell death induced by betanin/isobetanin concentrate

As MCF-7 cell viabilities strongly decreased during Bet./IsoBet. treatment, we analyzed the nature of cell death induced by the pigments. First we checked apoptosis induction and quantification using Annexin-V-FITC labeling and cell analysis by flow cytometry (Table 2, Figure 3A). Our results showed that betanin/isobetanin treatment significantly increased the percentage of Annexin-V positive/PI negative MCF-7 cells, as the positive control staurosporine. MCF-7 3D cell culture promoted phosphatidylserine externalization, and the percentage of Annexin-V positive cells was found to be

increased with betanin/isobetanin or staurosporine. MDA-MB-231 cells cultured in 2D or 3D with Bet./IsoBet. were not strongly labeled with Annexin-V-FITC probe (Table 2). Interestingly, HUVEC cells were found to be non-apoptotic when cultured in 2D and treated with Bet./IsoBet., whereas staurosporine treatment induced significantly apoptosis (Table 2, Figure 3A). Anchorage-independent culture of HUVEC cells triggered Annexin-V labeling and the percentage of apoptotic cells was slightly increased with betanin/isobetanin treatment. To confirm apoptosis induction in MCF-7 cells treated with betanin/isobetanin concentrate, we analyzed DNA fragmentation using an antibody recognizing single-stranded DNA. After culture for 48 hours with betanin/isobetanin (30 μ M) or staurosporine (1 μ M), about 80% of MCF-7 cells were labelled (Figure 3B). Only 1.84% of HUVEC cells treated with betanin-enriched extract were found to be positive (Figure 3B), whereas 29.03% were labeled with the positive control staurosporine.

Proteome profiler arrays allowed us to detect the expression levels of apoptosis-related proteins in MCF-7, MDA-MB-231 and HUVEC cells treated with 30 μ M Bet./IsoBet. Typical results of obtained arrays are shown in Figure 4A. The most important differences in protein expressions were found in sensitive MCF-7 cells: we observed a strong induction of Bad, TRAILR1/DR4, Fas/TNFRSF6/CD95 and phospho-p53 (S392) proteins in betanin-treated cells compared to control cells (Figure 4B, Table I supplementary data). No significant variation in protein expression was observed in MDA-MB-231 cells. In HUVEC cells, Bet./IsoBet. treatment induced a moderate overexpression of p21 and p27 proteins (Table I supplementary data).

The effect of betanin/isobetanin on the mitochondrial membrane potential was checked in MCF-7 cells using MitoCapture dye (Figure 5). Our data showed an increase in green

fluorescence in betanin/isobetanin-treated cells, in a same extent than positive control (curcumin 50 μ M).

To assess another cell death, autophagy, MCF-7 cells treated with Bet./IsoBet. or rapamycin (an autophagy inducer) were analyzed for lysosomal vacuole formation (Figure 6). We observed positive labeling in MCF-7-treated cells, indicating an autophagic activity increase.

DISCUSSION

We described that a highly betanin/isobetanin-enriched concentrate produced from red beetroots inhibits cancer cell proliferation and induces MCF-7 cell death, but has no obvious effect towards normal cells.

Betanin and isobetanin are the most predominant betalains in red beetroot. We were able to obtain a pigment-enriched dried extract from beetroots which contains only betanin/isobetanin as coloring agents. This concentrate consists of 80 % of betanin/isobetanin mixture (of which betanin accounts for 64% and isobetanin for 36%) and corresponds, to our knowledge, to the highest purified betanin extract used for studying its anticancer activities. To better evaluate the cytotoxicity of the betanin-enriched extract, we used two different culture conditions: classical monolayer culture (2D) and cells cultured as aggregates on anti-adhesive substratum (3D) (Velzenberger *et al.*, 2008). These 3D culture models are now commonly used to study new anticancer agents because they are believed to bridge the gap between *in vitro* 2D assessment and animal models (Breslin and O'Driscoll, 2013). Cell cytotoxicity analysis revealed that p53 wild-type cancer cell lines (B16F10, MCF-7) were highly sensitive to 40 μ M

betanin/isobetanin mix (assessed by proliferation inhibition and low cell viabilities), whereas cancer cell lines expressing mutated p53 were less (MDA-MB-231) or not sensitive (HT-29) for the same concentration range. The betanin-enriched extract effect was similar whatever the culture conditions, 2D or 3D. This shows that betanin/isobetanin concentrate inhibits aggregated cancer cell proliferation, a cell structure known as more resistant to apoptosis induction (Hirschhaeuser *et al.*, 2010). In this way, B16F10 and MDA-MB-231 are metastatic cell lines, resistant to anoikis when cultured under anchorage-independent conditions because of the activated ERK signaling pathway (Goundiam *et al.*, 2010; Fukazawa *et al.*, 2002). For these reasons, identifying an efficient molecule on anoikis resistant cancer cells is a very hopefully result. Cell proliferation inhibition by betanin/isobetanin was confirmed with affected cell cycle. In MCF-7 cells, betanin/isobetanin extract decreased the G1 and G2/M cell number and promoted S phase increase, a behavior previously described when MCF-7 cells were treated with cell growth inhibitors as resveratrol and riproximin (Joe *et al.*, 2002; Pervaiz *et al.*, 2015). Betanin/isobetanin extract may downregulate the expression of cell cycle regulators (cyclin B1, cyclin A2) resulting in G2 phase entry inhibition and S phase arrest, as act resveratrol and riproximin. 2D cultured MDA-MB-231 cells were blocked in G1 phase during treatment, but the molecules did not significantly affect the cell cycle repartition when cells were cultured as aggregates, probably because of the G1 arrest induced by the culture configuration.

Kapadia *et al.* (2011 and 2013) have previously evaluated the cytotoxic effect of a red beetroot extract in MCF-7 cell line and the IC50 value they obtained was 600 μM (after 72 hours of exposure). Reddy *et al.* (2005) also observed a growth inhibition of MCF-7 cells treated with a betanin concentrate for 48 hours (IC50 value was 294 μM). Our data

clarify these previous studies. The betanin purification process we applied to the crude beetroot extract, allowed us to obtain a significant MCF-7 growth inhibition associated with cell death for very low concentrations (below 40 μM).

As MCF-7 cell viability strongly decreased when treated with betanin/isobetanin mix, the nature of cell death was analyzed. Using different methods, we showed that betanin treatment induced apoptosis in 2D MCF-7 cells. The expressions of apoptosis-related proteins (Bad, TRAILR4, FAS, phosphorylated p53) were strongly increased and the mitochondrial membrane potential was clearly altered. Taken together, all these data suggest both mitochondrial and death-receptor pathway involvement, and a p53-dependent response in MCF-7 cells treated with the betanin/isobetanin concentrate. Betanin extract (30 μM) only induced a moderate apoptosis in aggregated MCF-7 cells, a 3D configuration often promoting cell death resistance. Recently, Gong *et al.* (2015) have compared the cytotoxicity effect of the doxorubicin on MCF-7 cells cultured as monolayer or spheroids. Their data show that 3D cultured MCF-7 cells were less sensitive than their 2D counterpart. According to our data on cell proliferation and viability, we would probably observe an apoptosis induction on 3D MCF-7 cells with higher concentrations of betanin/isobetanin mix.

Intrinsic pathway activation upon betanin exposure was previously observed by Sreekanth *et al.* (2007) in K562 cells with mitochondrial membrane potential decrease and cytochrome c release. A significant activation of caspase 9 and effectors (caspases 3 and 7) was also reported in lung cancer cells treated with 400 μM betanin for 48 hours (Zhang *et al.*, 2013). Our results confirm these previous studies about the mitochondrial pathway involvement. Moreover, we show that betanin/isobetanin mix induces upregulation of death receptors and p53 activation, as curcumin does on MCF-7 cells

(Choudhuri *et al.*, 2002; Mohankumar *et al.*, 2014). P53 is a well-known important actor in response to cellular stresses (Goh *et al.*, 2011). The mutated p53 status in MDA-MB-231 and HT-29 cells (Bartek *et al.*, 1990; Rodrigues *et al.*, 1990) could explain in part the weak betanin cytotoxicity for these cell lines. Particularly, HT-29 cells express a mutant p53 protein which gains function, promoting cell proliferation and chemoresistance (van Oijen and Slootweg, 2000). In this way, Arafa *et al.* (2013) have recently compared HT-29 and MDA-MB-231 sensitivity to novel quinoline-based compounds. They found that the HT-29 cell line was more refractory to the cytotoxic activity of most compounds than MDA-MB-231 cells. In our work, HT-29 cells were also resistant to betanin/isobetatin effect, whereas molecules slowed down MDA-MB-231 cell proliferation without inducing apoptosis. Therefore low concentrations of betanin-enriched extract appear to be efficient on a wild-type p53 cancer cells to induce cell death (MCF-7) and inhibit a p53-mutated cancer cell proliferation (MDA-MB-231), but have no effect on another p53-mutated cell line (HT-29).

As some natural-derived compounds have already been shown to induce cell death *via* an autophagic process (Zhang *et al.*, 2012), we investigated and revealed lysosomal vacuole formation in MCF-7 cells upon betanin treatment. We are the first to describe that betanin/isobetatin mix also induces an autophagic response in cells. The identification of molecules that target autophagy is believed to be a realistic prospect for cancer therapy (Bincoletto *et al.*, 2013).

To fully characterize the biological effect of betanin, we also evaluated its cytotoxicity on normal cells. Importantly, normal cell lines (HUVEC, MRC-5) were found to be weakly or not sensitive to betanin C15 stereoisomer mix for the same concentration range used on cancer cells. Particularly, HUVEC endothelial cells slowed down their

proliferation rate upon betanin exposure, remaining viable for higher concentrations than MCF-7 cells. When cultured as monolayer and treated with the betanin/isobetanin extract, HUVEC cells accumulated in the G1 phase and the percentage of cells in the DNA replicative phase decreased, confirming the cell proliferation data. 3D betanin treated-HUVEC cells progressively accumulated in the S and G2/M phases. Moreover the betanin/isobetanin treatment triggered an overexpression of p21 and p27 proteins, cyclin-dependent kinase inhibitors involved in cell cycle arrest. However, the HUVEC cell proliferation inhibition by the betanin concentrate was not associated with apoptosis or autophagy induction: this suggests that undesired side effects could be limited in therapies integrating these molecules. Indeed, the preliminary data we performed on the non-tumorigenic MCF-12F cell line confirmed the harmless interaction of the molecules on normal epithelial breast cells (data not shown).

In conclusion, we reported that a purified betanin/isobetanin concentrate produced from red beetroots is cytotoxic at low concentrations for cancer cells expressing functional p53, but have no or moderate effect on normal cells, suggesting limited *in vivo* side effects. In p53 wild-type cancer cells, betanin/isobetanin extract induces apoptosis (*via* extrinsic and intrinsic activation pathways) and autophagy. Moreover, betanin-enriched extract inhibits some p53 mutated cancer cell growth, without inducing apoptosis. Our data bring new insight to consider betanin/isobetanin as therapeutic anticancer molecules alone or in combination with classical chemotherapeutic drugs, especially in functional p53 tumors.

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Tables

Table 1: Cell cycle distribution of cells grown on tPS or polyHEMA in the presence or absence (Control) of betanin/isobetatin extract.

			Control	Bet./IsoBet. (30 μ M)
MCF7	2D	G1	54.7% \pm 1.6	49.1% \pm 4.2*
		S	26.4% \pm 2.8	34.2% \pm 9.8
		G2/M	18.8% \pm 4.1	16.7% \pm 5.5
	3D	G1	77.0% \pm 2.6	68.2% \pm 1.8**
		S	12.4% \pm 3.2	17.8% \pm 2.2
		G2/M	10.6% \pm 0.7	14.0% \pm 0.7***
MDA-MB231	2D	G1	62.3% \pm 1.9	67.4% \pm 1.2
		S	24.8% \pm 1.6	19.8% \pm 0.9*
		G2/M	12.9% \pm 0.8	13.5% \pm 2.2
	3D	G1	74.6% \pm 10.0	72.4% \pm 10.5
		S	22.9% \pm 11.6	21.7% \pm 13.2
		G2/M	2.5% \pm 2.9	5.8% \pm 2.7
HUVEC	2D	G1	51.2% \pm 2.4	58.4% \pm 1.9
		S	40.0% \pm 1.5	30.2% \pm 2.5*
		G2/M	8.8% \pm 1.4	11.7% \pm 1.6*
	3D	G1	90.1% \pm 1.4	80.9% \pm 4.5*
		S	5.5% \pm 1.2	13.1% \pm 4.6
		G2/M	4.3% \pm 1.3	5.9% \pm 1.2

Results are means \pm SD of three experiments each performed in triplicate. Asterisks indicate significant differences in cell cycle distribution of treated and control cells (Dunn test): *, $p < 0.05$; **, $p < 0.01$; ***, $p < 0.001$.

Table 2: Annexin V labeling of MCF-7, MDA-MB231 and HUVEC cells treated with betanin/isobetatin or staurosporine.

		Control	Bet./IsoBet. (30 μ M)	Staurosporine (1 μ M)
MCF7	2D	0.59% \pm 1.41	23.24% \pm 2.72 *	18.75% \pm 11.19 *
	3D	11.0% \pm 6.91	17.29% \pm 6.25	29.41 % \pm 14.51 *
MDA-MB231	2D	0.29% \pm 0.04	0.71% \pm 0.15	13.45% \pm 2.52 ***
	3D	1.94% \pm 0.5	4.25 % \pm 1.40*	1.79 % \pm 0.68
HUVEC	2D	1.44% \pm 0.72	4.05% \pm 0.86	56.99% \pm 10.37 ***
	3D	25.06% \pm 5.62	38.72% \pm 7.8	18.13% \pm 7.81

Results are means \pm SD of two experiments each performed in triplicate. Asterisks indicate significant differences in Annexin-V labelling of treated and control cells (Dunn test): *, $p < 0.05$; *** $p < 0.001$.