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In vitro gastrointestinal digestion impact on stability, bioaccessibility and antioxidant activity of polyphenols from wild and commercial blackberries (*Rubus* spp.)

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Gastrointestinal digestion (GID) is a physiological process that transforms the stability, bioaccessibility and antioxidant activity (AOX) of polyphenols from blackberries (*Rubus* spp.). This study aimed to investigate the effect of the INFOGEST® GID protocol on the phenolic stability, bioaccessibility and AOX of Mexican wild (WB) and commercial (CB) blackberries. After GID, the total phenolic and anthocyanin contents in blackberries decreased by $\geq 68\%$ and $\geq 74\%$, respectively. More than 40 phenolics were identified during GID; most of them degraded completely during digestion. GID had a negative effect on the AOX of both fruits ($>50\%$), but WB showed the highest antioxidant activities, as assessed by the ORAC, DPPH, reducing power and β -carotene bleaching methods. In Caco-2 cells, the cell-based antioxidant activity of digested blackberries ($p < 0.05$) decreased by 48% in WB and by 56% in CB. The capacity to inhibit intracellular ROS decreased by 50% in WB and by up to 86% in CB, after digestion. GID is a complex process that impacts on the bioactive properties of food nutrients, especially phenolics. *In vitro* and cellular AOX of WB polyphenols withstood the gastrointestinal environment better than CB phenolics. The *in vitro* assays results suggest that phenolics from underutilized WB have a higher bioaccessibility and antioxidant capacity than the polyphenols from the most frequently consumed CB. However, whether this corresponds to a better bioaccessibility in humans remains to be determined in future work.

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1. Introduction

Blackberries (*Rubus* spp.) are a large group of fruits with a rich diversity of bioactive polyphenols, such as phenolic acids, flavonoids and tannins.¹ The phenolic profile of blackberries is closely related to biotic and abiotic stimuli, resulting in some important differences between wild species and cultivated

species. In Mexico, wild blackberries and other Rosaceae berries are underutilized seasonal food sources collected only by inhabitants of remote mountainous regions and of minor economic importance, which are consumed as fresh fruits or processed for consumption in traditional dishes and beverages.^{2,3} Wild blackberries, although reported to have a higher phenolic content and biological effects, are scarcely consumed as edible fruits in comparison with commercial blackberries.⁴ Blackberries contain phenolic compounds, which have health promoting effects, such as excellent chemopreventive and chemotherapeutic effects and protection against chronic and non-chronic diseases.^{5,6} However, the health benefits of dietary phenolics are dependent on their bioaccessibility during gastrointestinal digestion (GID).^{7–9}

In blackberries, different hydroxycinnamic and hydroxybenzoic acids are present in free forms, but they are mainly found as blocks of polymeric polyphenols, such as proanthocyanidins and hydrolysable tannins.¹⁰ In contrast, flavonoid derivatives, which have a characteristic C6–C3–C6 (A, B and C rings) carbon skeleton, exhibit several patterns of hydroxylation, acylation, glycosylation and methoxylation of the main structure

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of each phenolic compound.⁹ Also, these substituents may be found in phenolic acid derivatives and polymers, and may explain their biological importance, which is seriously compromised during digestion.^{9,11–13}

In blackberry fruits, polyphenols are the main compounds responsible for fruit sensory properties, and their stability is highly dependent on factors such as light exposure, pH, native chemical structure, sugar conjugation, temperature, metal ions, oxygen, and enzymes.^{14,15} Phenolic compounds are stored in cell structures (e.g. vacuoles and cell walls) of the skin, pulp and seeds of fruits.¹⁵ These compounds are bound to macromolecules, such as proteins and peptides (linking the phenolic group of the phenol ring with the –NH group of peptides) or carbohydrates (linked by α - and β -glycosidic bonds), the dry matter content of which is about 80% and 5%, respectively.^{8,16} It should be noted, however, that these macromolecules undergo marked degradation during GID, becoming bioavailable sugars and small peptides/amino acids.¹⁷

With increasing evidence that supports the bioactive effects of multiple phenolic compounds present in foods, there is a need to develop methods for simulating the physiological conditions, with harmonized requirements and predictable outcomes of *in vivo* GID experiments using non-invasive procedures.¹⁷ Recent data suggest that the transformation of phenolic compounds during GID stages is attributable to several factors, such as pH changes, enzyme activity and concentration, peristaltic movements, and food matrices.^{9,17–19}

Several protocols have been used to study the bioaccessibility of phenolic compounds in red fruits.^{6,11,20,21} To date, however, no clear consensus has been reached on the biotransformation of these phytochemicals. It is crucial to acquire more knowledge in this regard to be able to determine the bioaccessibility and bioactivity of berry phenolics, since we still do not have a clear picture of the mechanisms involved prior to transport through the small intestine epithelium into the bloodstream.^{9,18} Keeping in mind the above considerations, it is necessary to prevent extra degradation of phenolics by external factors (due to atmospheric O₂ and/or the presence of light) during *in vitro* GID protocols.^{11,17} Therefore, the objectives of the present study were (a) to evaluate the bioaccessibility and digestive stability of each phenolic compound after GID, using the INFOGEST® 2.0 protocol; (b) to determine the *in vitro* antioxidant activity of wild (WB) and commercial (CB) blackberry polyphenols during GID; and (c) to assess the antioxidant potential of WB and CB digestates based on their cellular antioxidant activity (CAA) and their capacity to inhibit the synthesis of reactive oxygen species (ROS) in the Caco-2 cell line.

2. Materials and methods

2.1 Biological material and reagents

Ripe wild blackberry (*Rubus liebmannii* Focke) fruits (WB) were collected in El Palmito, Sinaloa, Mexico (23°35'22.2"N, 105°52'11.8"W, 2170 mASL), at the beginning of summer 2018, while commercial *Rubus fruticosus* brambles (CB) were purchased at

a local market. WB and CB fruits were weighed, frozen at –80 °C, and then freeze-dried (FreeZone Dryer System; Labconco, Kansas City, MO, USA) and weighed again. All reagents and enzymes were obtained from Sigma-Aldrich, unless otherwise stated. Commercial standards of cyanidin-3-rutinoside (PHL80577), gallic acid (G7384), chlorogenic acid (C38378), catechin (C1788), vanillic acid (94770), caffeic acid (C0625), epicatechin (E1753), *p*-coumaric acid (C9008), ferulic acid (Y0001013), quercetin (Q4951), and kaempferol (60010) were used for phenolic identification and/or AOX evaluation. Reagents 2,2'-azino-bis(3-ethylbenzothiazoline-6-sulphonic acid) (ABTS, A9941), di(phenyl)(2,4,6-trinitrophenyl)iminoazanium (DPPH, D9132), β -carotene (C9750), 2,2'-azobis(2-amidinopropane)dihydrochloride (AAPH, 440914), potassium persulphate (K₂S₂O₈, 216224), Tween-20 (P7950), linolenic acid (L2376), potassium hexacyanoferrate(III) (K₃Fe(CN)₆, P8131), iron(III) chloride hexahydrate (FeCl₃, 236489), sodium bicarbonate (NaHCO₃, S6297), fluorescein (F46960), and Folin-Ciocalteu's reagent (F9252) were used. Calcium chloride (CaCl₂, 449709) and α -amylase enzymes from porcine pancreas (EC 3.2.1.1, A3176), pepsin from porcine gastric mucosa (EC 3.2.4.3.1, P7012), trypsin from bovine pancreas (EC 3.4.21.1, T0303), α -chymotrypsin from bovine pancreas (EC 3.4.21.1, C4129), lipase from porcine pancreas Type II (EC 3.1.1.3, L3126), and bile salt from porcine bile extract (B8631) were used. Butylated hydroxytoluene (BHT, W218405), Trolox (238813), and cyanidin-3-glucoside chloride (89616 PhytoLab GmbH & Co. KG, Vestenbergsgreuth, Germany) were utilized as external standards in different experiments. For cellular experiments, Gibco Thermo Fisher Scientific (Life Technologies Corp., San Diego, CA, USA) supplied Dulbecco's modified Eagle's medium (DMEM, 11965-092) and Hank's balanced salt solution (HBSS, 14025-092), while fetal bovine serum (FBS, 080450), non-essential amino acids (321-011-EL), L-glutamine (609-065-EL), penicillin-streptomycin (450-201-EL), Dulbecco's phosphate-buffered saline (DPBS, 311-425-CL), dichlorofluorescein diacetate (DCFH-DA, D6883) and Hepes (330-050-EL) were provided by Wisent Bioproducts (Wisent Inc., Saint-Jean-Baptiste, QC, Canada). Hydrogen peroxide (H1009) also was required. Sodium hydroxide (NaOH) and hydrochloric acid (HCl) were used at different concentrations to adjust the pH.

2.2 Simulated *in vitro* gastrointestinal digestion

In vitro gastrointestinal digestion (GID) was performed following the INFOGEST® 2.0 protocol,¹⁷ incorporating specific conditions for the digestion of phenolic compounds, as described by Gil-Izquierdo *et al.*²⁰ and Bermúdez-Soto *et al.*¹¹ Briefly, 20 g of rehydrated WB and CB fruits (with original water proportions) were placed in a 250 mL amber screw cap bottle and homogenized with an Ultra-Turrax® (IKA, Germany) for 30 s. To mimic the oral phase (OP), samples were mixed with simulated salivary fluid (1 : 1, w/v) and adjusted to pH 7.0 with 4 N NaOH, 100 μ L of 0.3 M CaCl₂, and 1500 U mL⁻¹ of α -amylase. The mixture was stirred for 2 min at 350 rpm on a StableTemp stirring-hot plate at 37 °C (Cole-Palmer Instruments Company,



IL, USA). An aliquot of 5 mL was recovered at the end of the OP. After that, the bolus (35 mL) was mixed with simulated gastric fluid (SGF) (1 : 1, v/v), the pH was adjusted to 3.0 with 1 N HCl and 17.5 μL of 0.3 M CaCl_2 , and 2500 U mL^{-1} of pepsin were added. Gastric phase (GP) digestion was carried out for 2 h under constant shaking (100 rpm) at 37 °C using an Innova® 40 incubator (Incubator Shaker Series, New Brunswick Scientific, NJ, USA). The intestinal phase (IP) started with the resulting volume (50 mL), which was mixed with an equal volume of simulated intestine fluid (SIF); the pH was adjusted to 7.0 and then 100 μL of 0.3 M CaCl_2 , trypsin (100 U mL^{-1}), α -chymotrypsin (25 U mL^{-1}), lipase (2000 U mg^{-1} of fat), α -amylase (200 U mL^{-1}), and bile salt (10 mM mL^{-1} SIF) were added. An anaerobic atmosphere was maintained with N_2 flushing during the addition of SGF and SIF, as well as during aliquot sampling every 30 min. Aliquots of 5 mL were recovered every 30 min during the GP and IP phases. Digestates were mixed with pure methanol (1 : 1, v/v) to precipitate enzymes. Then samples were centrifuged at 10 000g for 5 min at 4 °C. Supernatants were filtered with a 0.45 μL nylon filter, freeze-dried and stored in light-protected tubes at -80 °C for further use.

2.3 Total phenolic and anthocyanin content

To assess the total phenolic content (TPC) of various samples, 50 μL of resuspended fruit extracts and digested samples (1 : 50, fruit extract or digestate : distilled water, w/v) were mixed with 100 μL of 70% acetone–1% HCl solution (acetone/water/HCl (70 : 29 : 1, v/v/v) and maintained at room temperature for 2 h.²² After the 2 h period, 100 μL of standard solution (gallic acid 0–500 $\mu\text{g mL}^{-1}$) or 100 μL of the mixture were suspended in 750 μL of Folin–Ciocalteu solution (1 : 10, Folin–Ciocalteu's reagent : water), vortexed and left to stand for 5 min. Then, 750 μL of NaHCO_3 solution (7.5%) was added to the above solution, which was vortexed and centrifuged at 10 000g for 15 min (Benchmark Z206-A Centrifuge, Beckman Coulter, Indianapolis, IN, USA). Finally, 250 μL of each of the aforementioned mixtures were retained in the dark at room temperature, and were read at 750 nm on a microplate reader (BioTek Instruments, VT, USA) equipped with Gen5 software version 3.02 (BioTek Instruments, VT, USA). The results were expressed as μg of gallic acid equivalents (GAE) per mL.

The differential pH method was applied to determine the total anthocyanin content (TAC).²³ Briefly, the pH of 100 μL of WB or CB was adjusted to 1.0 with 1 N HCl, or to 4.5 with 1 N NaOH. Aliquots were measured at 520 nm and 700 nm using a microplate reader (BioTek Instruments, VT, USA) equipped with Gen5 software version 3.02 (BioTek Instruments, VT, USA).

2.4 Characterization of the phenolic compounds

Freeze-dried fruits and digestates of WB and CB (100 mg) were resuspended in water : MeOH (50 : 50, v/v), centrifuged at 15 000 rpm (Z206-A Centrifuge, Beckman Coulter, Indianapolis, IN, USA) and analyzed with the procedure reported by Jiao *et al.*¹² and Iglesias-Carres *et al.*²⁴ using a

1290 UHPLC Infinity II series instrument coupled to a qTOF/MS 6550 system (Agilent Technologies, Palo Alto, CA, USA). Two different methodologies based on UHPLC-ESI-qTOF-MS systems were used to separate, detect, and quantify the non-anthocyanin and anthocyanin phenolic compounds. For the separation of non-anthocyanin compounds, an Acquity HSST3 C18-column (150 \times 2.1 mm, 1.8 μm particle size) (Waters, Milford, MA, USA) was used. The mobile phase consisted of (A) water : acetic acid (99.8 : 0.2, v/v) and (B) acetonitrile. The gradient mode was as follows: 0–0.5 min, 0% B; 0.5–18 min, 0–30% B; 18–21 min, 30–95% B; 21–24 min, 95% B; and 24–25 min, 100–0% B. A post-run time of 6 min was required for column re-equilibration (at 0% B). The flow rate was set at 0.55 mL min^{-1} and column temperature at 45 °C; the injection volume was 2.5 μL for all runs. Electrospray ionization (ESI) operating in negative mode was conducted with a gas temperature at 200 °C and a flow rate of 14 L min^{-1} . A nebulizer gas pressure of 20 psi, sheath gas temperature of 350 °C, sheath gas flow of 11 L min^{-1} and capillary voltage of 3000 V were used. The anthocyanin compounds were separated on an Acquity BEH C18 column (100 mm \times 2.1 mm, 1.7 μm particle size) (Waters) and the mobile phase consisted of water : formic acid (9 : 1, v/v) (A) and acetonitrile (B). The gradient mode was as follows: 0–0.5 min, 0% B; 0.5–5 min, 0–9% B; 5–7 min, 9–15% B; 7–9.5 min, 15–30% B; 9.5–10 min, 30–100% B; 10–12 min, 100% B; and 12–12.1 min, 100–0% B. A post-run time of 5 min was required for column re-equilibration (at 0% B). The flow rate was set at 0.4 mL min^{-1} and column temperature at 25 °C, and the injection volume was 2.5 μL for all runs. ESI operating in positive mode was conducted with a gas temperature of 200 °C and the flow rate was 14 L min^{-1} . A nebulizer gas pressure of 20 psi, sheath gas temperature of 350 °C, sheath gas flow of 11 L min^{-1} and capillary voltage of 3000 V were used. The mass spectra were recorded between m/z 100 and 1000 at 2.5 spectra per s for both methodologies.

Identification of the phenolic compounds was performed by direct comparison with commercial standards or by comparison with the molecular weight for phenolic compounds in the Phenol-Explorer database (<http://www.phenol-explorer.eu>) on the basis of chromatographic behavior, ion molecular mass ($[\text{M} - \text{H}]^-$ or $[\text{M}]^+$) and fragmentation patterns. The calibration curves from commercial standards were used to quantify the corresponding phenolic compounds. For the other compounds, the analysis was semi-quantitative. The behavior of each compound was evaluated, and the calibration curve chosen to calculate its concentration was that of the phenolic compound exhibiting the most similar behavior.²⁴

2.5 Bioaccessibility of WB and CB phenolic compounds

The bioaccessibility index (BI) of each phenolic compound was calculated using the following equation:

$$\text{BI} = \frac{\text{IC}}{\text{FC}} \times 100$$

where IC is the initial concentration of phenolics (concentration in the fruits before GID) and FC is the final concen-



tration of phenolics (concentration in the digestate of the intestinal phase at 120 min).²⁵

2.6 Antioxidant activity

The antioxidant activity (AOX) of phenolics is one of the most relevant properties of these compounds.^{4,12,26,27} The AOX of polyphenols largely depends on the number and position of the hydroxyl groups on the flavylum cation, which gives them their hydrogen-donating capacity. Hydrogen atom transfer (HAT) is one of the mechanisms involved; single electron transfer (SET) ability is another mechanism by which polyphenols stabilize/scavenge free radicals.²⁸ In addition, single-electron transfer followed by proton transfer (SET-PT) has recently been studied.²⁹ Accordingly, it is necessary to evaluate the evolution of the berries' phenolics during digestion to determine the possible antioxidant mechanisms of action and their fate during GID. All undigested and digested aliquots of WB and CB were normalized at 100 mg GAE per mL of sample before *in vitro* AOX assessment.

2.6.1 Oxygen radical antioxidant capacity (ORAC). The ORAC assay was performed following the procedure reported by Ou *et al.*³⁰ The declining fluorescein (0.1 μM) absorbance by AAPH (153 mM) at 37 °C was recorded at 1 min intervals for 60 min at 485/528 nm of excitation/emission using a Synergy HT multi-detection microplate reader (BioTek Instruments, Inc., Winooski, VT, USA). PBS buffer (pH 7.4) was used to dissolve samples at 100 $\mu\text{g mL}^{-1}$ and along with Trolox standard (0–100 $\mu\text{M mL}^{-1}$). The antioxidant capacity (AOX) was estimated using Trolox as the standard curve and the following equation:

$$\text{AOX} = \left[\frac{\text{AUC}_{\text{sample}} - \text{AUC}_{\text{blank}}}{\text{AUC}_{\text{Trolox}} - \text{AUC}_{\text{blank}}} \right] \times k$$

where $\text{AUC}_{\text{sample}}$ is the integrated area under the fluorescence decay curve of each sample solution, $\text{AUC}_{\text{blank}}$ is the integrated area under the fluorescence decay curve of the blank, $\text{AUC}_{\text{Trolox}}$ is the integrated area under the Trolox fluorescence decay standard curve, and k is the dilution factor. Results were expressed as μM of Trolox equivalent (TE) per mL of sample.

2.6.2 DPPH radical-scavenging activity. The DPPH radical-scavenging activity was determined according to the method of Sánchez-Vioque *et al.*³¹ Trolox standard and normalized samples were mixed at a 1 : 1 (v/v) ratio with freshly prepared 0.1 mM DPPH. The microplate was incubated in the dark under agitation (New Brunswick™ Innova® 40/40R Incubator Shaker, Eppendorf, Inc. USA) at room temperature for 30 min. Absorbance was measured at 517 nm on a microplate reader and AOX was calculated using the following equation:

$$\text{RI}(\%) = \left[\frac{\text{ABS}_{\text{sample}} - \text{ABS}_{\text{blank}}}{\text{ABS}_{\text{Trolox}} - \text{ABS}_{\text{blank}}} \right] \times k$$

where RI is the radical inhibition, $\text{ABS}_{\text{sample}}$ is the absorbance of the samples, $\text{ABS}_{\text{blank}}$ is the absorbance of the blank, $\text{ABS}_{\text{Trolox}}$ is the absorbance of the Trolox standard, and k is the dilution factor. Results were expressed as percentage of DPPH radical inhibition (%).

2.6.3 Trolox equivalent antioxidant capacity (TEAC). The Trolox equivalent antioxidant capacity (TEAC) of samples based on the scavenging of $\text{ABTS}^{+\cdot}$ was measured following the method of Re *et al.*³² A solution of 7 mM ABTS reagent was prepared with 40 mg of ABTS salt and 6.95 mg of 2.45 mM $\text{K}_2\text{S}_2\text{O}_8$ solution, which was adjusted to 10 mL with distilled water and maintained in the dark for 16 h at room temperature. The ABTS reagent solution was then diluted with EtOH to obtain an absorbance of 0.80 ± 0.1 at 734 nm. Blank, standards and normalized samples were mixed with ABTS solution (1 : 10, v/v) and vortexed for 10 s, then the absorbance was read at 0 and 6 min at 734 nm. A Trolox curve (0–0.7 mM) was used. AOX was calculated using the following equation:

$$\text{RI}(\%) = \left[\frac{\text{ABS}_{\text{sample}} - \text{ABS}_{\text{blank}}}{\text{ABS}_{\text{Trolox}} - \text{ABS}_{\text{blank}}} \right] \times k$$

where RI is the radical inhibition, $\text{ABS}_{\text{sample}}$ is the absorbance of the samples, $\text{ABS}_{\text{blank}}$ is the absorbance of the blank, $\text{ABS}_{\text{Trolox}}$ is the absorbance of the Trolox standard, and k is the dilution factor. Results were expressed as a percentage of ABTS radical inhibition (%).

2.6.4 β -Carotene bleaching activity. The antioxidant activity assay measuring β -carotene discoloration was used as described by Marco *et al.*³³ A solution of 1 mg mL^{-1} of β -carotene reagent diluted in chloroform and mixed with 200 μL of Tween-20 and 20 μg of linolenic acid was prepared. The resulting mixture was diluted with 20 mL of water previously flushed with O_2 (stock solution). An aliquot of this stock solution was diluted with oxygen-rich water to obtain a β -carotene working solution (absorbance of 1.2–1.3 at 450 nm). The working solution was maintained in an ice bath in the dark before being used. Then 50 μL of normalized samples or blank were mixed with 200 μL of working solution and incubated under agitation at 37 °C, and the absorbance was measured at 0 and 60 min at 450 nm. The following equation was used to calculate the antioxidant activity:

$$\text{DR} = \frac{\text{Ln} \frac{\text{ABS}_{0 \text{ min}}}{\text{ABS}_{60 \text{ min}}}}{60}$$

where DR is the degradation rate for the absorbance at the beginning ($\text{ABS}_{0 \text{ min}}$) and the end of the reaction ($\text{ABS}_{60 \text{ min}}$) for the control and samples. Using the DR, AOX was calculated as a percentage of β -carotene relative inhibition against the blank:

$$\text{AOX}(\%) = \left[\frac{\text{DR}_{\text{blank}} - \text{DR}_{\text{sample}}}{\text{DR}_{\text{blank}}} \right] \times 100$$

where DR_{blank} indicates the degradation of β -carotene in the absence of sample. Results were expressed as percentage of β -carotene bleaching inhibition (%).

2.6.5 Reducing power. The reducing power assay was performed according to the method of Carrasco-Castilla *et al.*³⁴ Briefly, 50 μL of normalized sample were mixed with 50 μL of 0.2 M phosphate buffer (pH 6.6) and 50 μL of fresh (prepared daily) 1% $\text{K}_3\text{Fe}(\text{CN})_6$ solution. The microplate containing the



resulting samples was incubated for 20 min at 50 °C under agitation at 100 rpm (New Brunswick™ Innova® 40/40R Incubator Shaker, Eppendorf, Inc. USA). After that, 50 µL of 10% TCA and 10 µL of 0.1% FeCl₃ were added to each well and the incubation was continued for an additional 10 min at 50 °C under agitation. The absorbance was read at 700 nm and compared with a BHT curve (0.0–0.05 µg µL⁻¹). Results were expressed as the absorbance (Abs) at 700 nm.

2.7 Cellular bioactivity

2.7.1 Caco-2 cell culture. Caco-2 cells were obtained from the American Type Culture Collection (ATCC® HTB-37™, Manassas, VA, USA) and were cultivated in DMEM (supplemented with 10% FBS, 1% non-essential amino acids, 1% L-glutamine, and 1% penicillin–streptomycin). The cells were grown in a 75 cm² flask under a humidified atmosphere (5% CO₂) at 37 °C until 80% confluence was reached.³⁵ Cells were sub-cultured twice a week. Trypan blue dye was used to assess cellular viability.

2.7.2 Caco-2 cytotoxicity assay. Caco-2 cells were seeded in a 96-well flat-bottomed plate (passage number 31), at a concentration of 2.2×10^4 cells per well in 100 µL of DMEM. Following 24 h of incubation (Series 8000DH CO₂ incubator, Thermo Fisher Scientific, Marietta, OH, USA), the samples of a specific concentration (0.1–1000 µg GAE per µL) were added and incubation was continued for an additional 24 h. After that, the 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assay was performed by following the MTT kit's instructions (M2128 Millipore Sigma, Burlington, MA, USA). The culture medium was removed and replaced with fresh medium (100 µL) to the blank (without sample) and sample wells, and MTT solution (10 µL) was added to each well. After an additional incubation step for 4 h in the dark, the MTT formazan produced in wells containing live cells appeared as black, fuzzy crystals on the bottom of the well. Then, 100 µL of isopropanol with 0.04 N HCl was added to the wells and agitated (30 s, 100 rpm at room temperature and in darkness). Finally, within the next hour, the plate was read at 570/630 nm against a blank. The metabolic activity (% MA) of the cells was calculated using the following equation:

$$\%MA = \left[\frac{A_{\text{digestate}}}{A_{\text{control}}} \right] \times 100$$

where $A_{\text{digestate}}$ is the absorbance of the sample exposed to the extract and A_{control} is the absorbance of the control sample (cells without extract exposition, 100% metabolic activity). The results were expressed as the half-maximal effective concentration (EC₅₀) in mg of phenolic compounds per mL of sample (mg mL⁻¹).

2.7.3 Cellular antioxidant activity. The cellular antioxidant activity (CAA) of digested samples was measured according to the method of Kellett *et al.*³⁵ After Caco-2 cells reached confluence (80% confluence), cells were seeded at a density of 6×10^5 cells per mL (passage number 32) in a 96-well black clear-bottomed microplate and incubated until confluence (24–48 h) under the conditions described in the previous section.

Confluence was confirmed using the microscope. The outside wells of the plate were not used. Prior to the experiment with digestates, growth medium was removed and cell monolayers were washed twice with DPBS. Then, 50 µL of treatment medium (FBS free medium) and 25 µM DCFH-DA working solution were added to each well along with 50 µL of the samples (100 µg GAE per mL). Once the DCFH-DA and blackberry samples were added, the cells were placed in the incubator for 1 h at 37 °C. When only intracellular CAA was being measured, wells were washed three times with 100 µL of DPBS. Then, the AAPH radical solution (100 µL, 600 µM AAPH in HBSS) was added to each well. The cells were immediately placed in a microplate reader (BMGLABTECH Inc., Cary, NC, USA), where real-time fluorescence was read at an excitation wavelength of 485 nm and an emission wavelength of 538 nm every minute for 1 h. For the control, 50 µL of DCFH-DA and 50 µL of serum-free culture medium (no samples included) were added to the wells in triplicate. A standard curve with quercetin (0–250 µM) was used. All samples were analyzed in triplicate. For CAA quantification, first, the blank subtraction (wells without samples) was done, followed by integration of the area under the curve of fluorescence *vs.* time at each concentration of sample, as follows:

$$CAA(\%) = 100 - \left[\frac{\int SA}{\int CA} \right] \times 100$$

where $\int SA$ is the integrated area under the sample fluorescence *vs.* time curve and $\int CA$ is the integrated area from the curve. Results were expressed as fluorescence inhibition compared with the quercetin curve standard (%).

2.7.4 Cellular oxidative stress. Oxidative stress in Caco-2 cells was evaluated under the DCFH-DA assay conditions,³⁶ with some modifications. When cells reached 80% confluence, the Caco-2 cell line was grown at a density of 2×10^4 cells per well (passage number 33) in a black flat-bottomed 96-well microplate for 24 h–48 h under optimal conditions until confluence was reached (confirmed using a microscope). The outside wells of the plate were not used. Prior to the experiment, growth medium was removed and cell monolayers were washed twice with 100 µL of DPBS, then cells were incubated for 30 min with 100 µM DCFH-DA and with sample, both diluted in HBSS. The wells were then washed with fresh DPBS solution before 100 µM of H₂O₂ was added. The fluorescence increase ratio (FIR) was evaluated at 0 and 30 min (ex/em: 485/530 nm) on a microplate reader (BMGLABTECH Inc., Cary, NC, USA). The positive control wells contained the cells treated with DCFH-DA and H₂O₂. For the negative control wells, 100 µL of HBSS without H₂O₂ was added to the blank wells. FIR was calculated as:

$$FIR = \left[\frac{F_{30 \text{ min}} - F_{0 \text{ min}}}{F_{0 \text{ min}}} \right] \times 100$$

where $F_{0 \text{ min}}$ is the fluorescence value at 0 min and $F_{30 \text{ min}}$ is the fluorescence at 30 min. FIR was expressed as the percentage (%) of the positive control.



2.8 Statistical analysis

Data were expressed as means \pm SD. Statistical differences were analyzed using the one-way analysis of variance (ANOVA) under Tuckey's test, with Minitab 2013 (Minitab Inc., USA). The differences were considered to be statistically significant at $p < 0.05$.

3 Results and discussion

3.1 *In vitro* GID

Ten aliquots of each blackberry genotype were recovered at different stages during GID: one aliquot for undigested fruit; one aliquot after OP (2 min); 4 aliquots for GP (30, 60, 90 and 120 min); and 4 aliquots for IP (30, 60, 90 and 120 min). In most of the published studies on GID of red fruits,^{13,21,37–41} the digestive procedure was performed in two phases (GP and IP), excluding the first digestive stage (OP). The oral phase is of special importance during the digestion process, since it is during this phase that the plant matrix containing phytochemicals is broken down during the chewing step, the components are exposed to the salivary fluids and to the action of α -amylase, and an anaerobic atmosphere begins. Sigurdson and Giusti⁹ and Brodkorb *et al.*¹⁷ have highlighted the key role that the OP plays in the simulation of gastrointestinal digestion for food matrices with a high carbohydrate content (>80% of dry matter) and in the possible absorption of nutrients that occurs in the oral epithelium.

3.2 Effect of GID on TPC and TAC

The total phenolic content (TPC) of undigested and digested fruits of WB and CB is shown in Fig. 1A. For undigested fruits of WB and CB, the TPC was 127 and 79 mg EAG per mL, respectively, with the TPC being significantly higher for WB ($p < 0.05$) than for CB. Following the OP (2 min), the TPC of WB decreased to 53 mg EAG per mL, which was 58% lower than for the corresponding undigested sample, while the CB sample showed a TPC of 54 mg EAG per mL, which represented a decrease of 31% relative to the undigested sample. There was no significant difference ($p > 0.05$) in the TPC between the two types of blackberries following the OP. For GP_{30 min}, the WB digestates showed values of 44 mg EAG per mL, with no significant difference ($p > 0.05$) when compared to the corresponding GP_{60 min} digestates (42 mg EAG per mL). Digestates obtained after 90 min of GP (GP_{90 min}) had a TPC of 40 mg EAG per mL, which was not significantly ($p > 0.05$) different from the value observed after 30 min or after 60 min. For the same stage, the TPC of CB was 49–52 mg EAG per mL, with no significant difference ($p > 0.05$) between GP_{30 min}, GP_{60 min} and GP_{90 min}. However, these values were statistically higher ($p < 0.05$) than the ones reported for the corresponding WB. During the IP, the TPC of WB showed values of 37–38 mg EAG per mL, with no statistical differences ($p > 0.05$) between IP_{30 min}, IP_{60 min}, IP_{90 min} and IP_{120 min}, while the TPC of the CB digestates varied between 25 and 31 mg EAG per mL, with no significant differences ($p > 0.05$) between them; however,

the TPC values were lower ($p < 0.05$) than those for the WB digestates. These changes in TPC represent a decrease of 70% in WB and 68.35% in CB. This indicates that the phenolic compounds of digested WB are slightly less stable than those found in digested CB.

The quantification of phenolic compounds in the fruits and digestates of blackberries is of great importance, since they are the main group of secondary metabolites in these fruits and they display multiple benefits for human health.^{9,42} Phenolic compounds are released from the plant matrix (mainly from cellular carbohydrates with structural functions) through enzymatic action and pH oscillations during the different digestive stages;^{43–45} however, these compounds are not always stable during and after GID. The TPC of *Rubus* fruit digestates showed a decrease of >41% in commercial raspberry⁴⁵ and >72% in blackberry⁴⁶ following GID. However, in *R. hirsutus*, Chen *et al.*³⁹ found an increase of 26% in TPC at the end of GID. Our results are similar to those reported by Bermúdez-Soto *et al.*¹¹ for commercial blackberries. This may be due to the similar digestive conditions used in these studies (control of external oxidizing agents, *e.g.* light and atmospheric oxygen). As demonstrated by the aforementioned research and the results of the present study, phenolic compounds are highly sensitive to GID conditions. Additionally, the gut microbiota has a key role in phenolic metabolism, breaking down high molecular weight polyphenols to low molecular weight phenolics, which sometimes results in the release of the most bioaccessible and bioactive phenolic compounds with the ability to enter into blood circulation and reach the organs.^{7–9,18,19,21,27,45,47}

The total anthocyanin content (TAC, Fig. 1B) of undigested WB was 0.33 mg EC3G per mL compared with 0.22 mg EC3G per mL for CB; the later value was significantly ($p < 0.05$) lower. Following the OP, TAC values of 0.27 and 0.10 mg EC3G per mL were observed for WB and CB, respectively; these values were significantly different ($p < 0.05$). This represents a decrease of 19% for WB and 54% for CB compared with the undigested samples. During the GP, the TAC of WB varied between 0.16 and 0.18 mg EC3G per mL, and there were no significant differences ($p < 0.05$) between the digestates. The corresponding TAC values varied between 0.18 and 0.19 mg EC3G per mL for CB, with the GP_{90 min} values being statistically higher ($p < 0.05$) than those of GP_{30 min} and GP_{60 min}. For the IP, the TAC of WB varied between 0.08 and 0.09 mg EC3G per mL. In contrast, the TAC of CB decreased sequentially during IP and it was significant ($p < 0.05$) after IP_{90 min}, with values varying between 0.07 and 0.05 mg EC3G per mL. All TAC values of CB during IP were significantly lower ($p < 0.05$) than the corresponding values for WB. At the end of GID, the TAC of the WB digestates decreased by 74% and the TAC of the CB digestates by 75%, when compared with the undigested samples.

Anthocyanins are flavonoid compounds that are found in *Rubus* berries and are mainly responsible for their organoleptic properties. These compounds are highly sensitive to changes in pH and to the presence of enzymes in the matrices.⁹ This



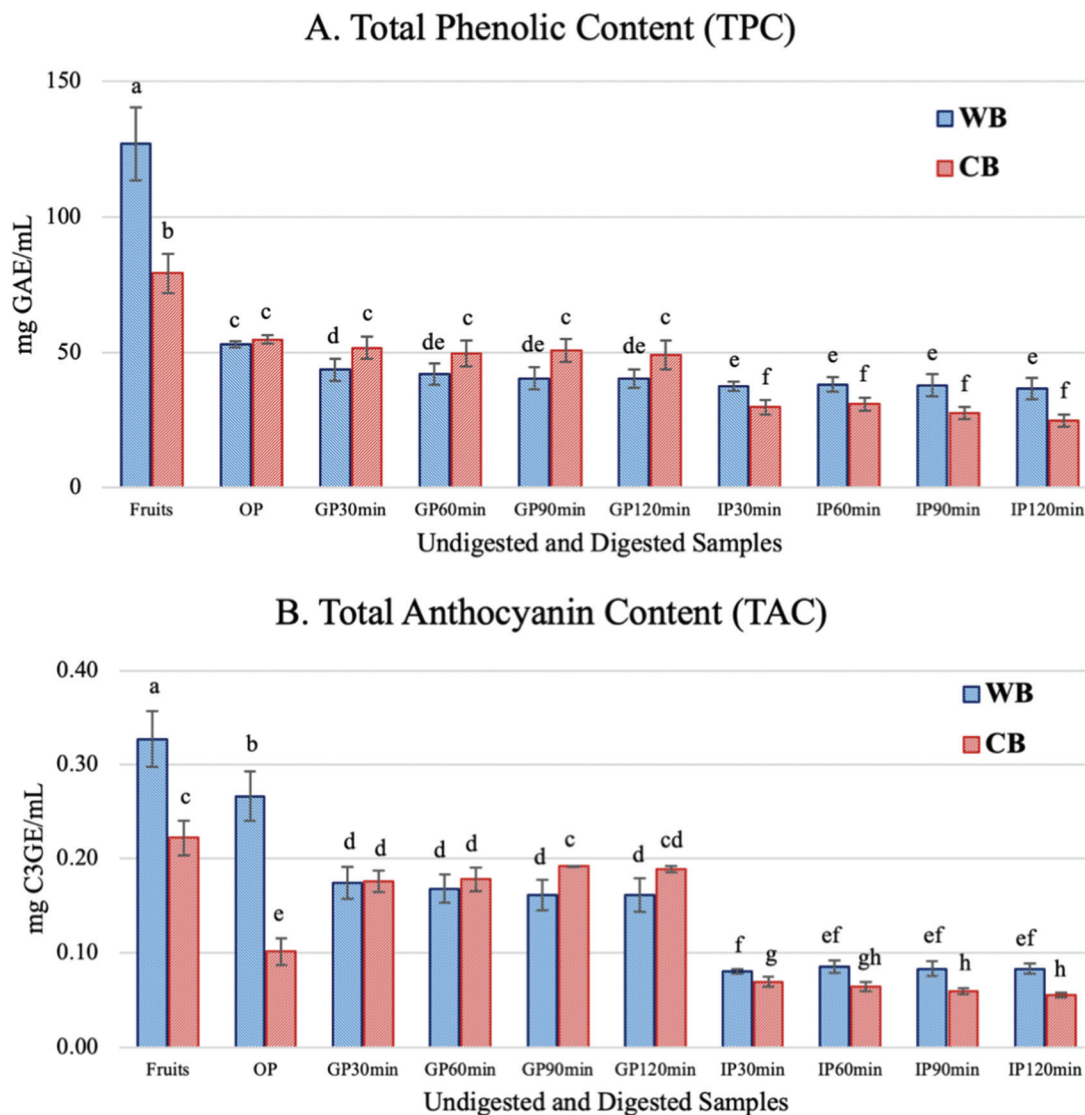


Fig. 1 Total phenolic (A) and anthocyanin (B) contents in undigested and digested WB and CB fruits. WB, wild blackberry; CB, commercial blackberry. mg GAE per mL, mg of gallic acid equivalents per mL; mg C3GE per mL, mg of cyanidin-3-glucoside per mL. Non-equal letters above bars indicate statistical differences ($p < 0.05$).

sensitivity explains the decrease of up to 97% in the TAC observed by Ryu and Koh¹³ for anthocyanin-enriched extracts from *R. occidentalis* after the IP. Similar observations were made by Azofeifa *et al.*⁴⁸ for *R. adenotrichos* juice, with a decrease in TAC estimated at 88%. Correa-Betanzo *et al.*⁴⁹ reported a 90% recovery of polyphenols and/or anthocyanins after gastric digestion of different berries. The release of anthocyanins into the matrix (bolus, chyme or chyle) from the fruit is a gradual process, framed by the different digestive stages, during which the cellular structures of the fruits are exposed to changes in redox potentials, light exposure, transition to an anaerobic atmosphere, and the action of specific enzymes.^{21,50} Considering that, in the current study, the berries were not subjected to an anthocyanin extraction or a semi-purification step, a larger amount of these compounds could be quantified at the end of GID. Another significant

factor in the GID simulation was the use of amber flasks as a digestion container, which reduced exposure of the digestates to light, in addition to maintaining a constant flow of nitrogen gas to maintain anaerobic conditions. This helped reduce interference from external factors during GID with the aim of obtaining TAC and TPC values after GID that were more representative of *in vivo* systems.

3.3 Changes in anthocyanin and non-anthocyanin profile during GID

Anthocyanins are a large subclass of phenolic compounds that are found in many edible fruits and vegetables and that exhibit potential health benefits.^{42,51} The basic chemical structure of anthocyanins has the flavylium ion (2-phenyl-benzopyryl) as its core and consists of two aromatic groups (a benzopyryl and a phenolic ring). Anthocyanins are found in the vacu-



oles of plant cells and provide a wide diversity of colors, such as red, purple or blue, to many types of leaves, flowers and fruits. Acylated and glycosylated aglycones of cyanidin (Cy), delphinidin (Dp), malvidin, pelargonidin (Pg), peonidin (Pn) and petunidin (Pt) are often reported in *Rubus* fruits.^{42,52} Glycosylation (sugar attachment through O-linkages) improves the stability of the anthocyanins and increases their water-solubility.⁵³ Several sugar moieties, from monosaccharides to trisaccharides (e.g. galactose, glucose, rutinose, sambubiose, sophorose, or xylose), have been reported to have the ability to attach to the aglycones. Additionally, anthocyanins can undergo greater molecular substitution in *Rubus* fruits with the addition of acyl or methyl residues to glycosyl moieties.⁵⁴

About 21 acylated and/or glycosylated forms of five aglycones were found in blackberry samples (Table 1). Cy-3-rutinoside (Cy-3-rut) was quantified at 3693 $\mu\text{g mL}^{-1}$ in WB fruits and 1226 $\mu\text{g mL}^{-1}$ in CB fruits, but decreased to 4 $\mu\text{g mL}^{-1}$ in both types of blackberries (BI –99% in both cases). It is important to mention that the Cy-3-rut content increased by about 460% in WB and 314% in CB during the GP. This anthocyanin was the main one found in WB and the second most important one found in CB. In WB samples, cyanidin-3-glucoside (Cy-3-glu) was quantified at 2854 $\mu\text{g mL}^{-1}$ in fruits, but only 8 $\mu\text{g mL}^{-1}$ in digestate IP_{120 min}, a decrease of –99% in BI, while in CB fruits the corresponding value was 4994 $\mu\text{g mL}^{-1}$ and 16 $\mu\text{g mL}^{-1}$ at the end of GID (BI of –99%). This anthocyanin was the most abundant in CB and the second most abundant in WB.

Cy-3-xylosyl-rutinoside (Cy-3-xyl-rut) was quantified at 2215 $\mu\text{g mL}^{-1}$ in WB fruits and at 80 $\mu\text{g mL}^{-1}$ in IP_{120 min} (BI –98%); it was not found in CB samples either before or after GID, only in GP samples (3–27 $\mu\text{g mL}^{-1}$). Cy-3-malonyl-glucoside (Cy-3-mal-glu) was found at a concentration of 466 $\mu\text{g mL}^{-1}$ in WB fruits; it decreased to 3 $\mu\text{g mL}^{-1}$ in IP_{120 min} aliquot (BI < 1%), while in CB fruits it was 160 $\mu\text{g mL}^{-1}$ and 2 $\mu\text{g mL}^{-1}$ (BI < 2%). Cy-3-dioxalyl-glucoside (Cy-3-dioxal-glu) was found at a concentration of 21 $\mu\text{g mL}^{-1}$ in WB fruits and 19 $\mu\text{g mL}^{-1}$ after GID (BI of 90%); in CB samples its content was 613 $\mu\text{g mL}^{-1}$ in fruits and 124 at IP_{120 min} (BI of 20%). Dp-3-(6-*p*-coumaroyl)-glucoside (Dp-3-(6-*p*-cou)-glu) was estimated at 50 $\mu\text{g mL}^{-1}$ in WB fruits and 17 $\mu\text{g mL}^{-1}$ in digested sample (BI of 66%), compared with 83 $\mu\text{g mL}^{-1}$ in CB fruits and 21 $\mu\text{g mL}^{-1}$ in digested CB, which corresponds to a BI of 75%. Pg-3-rutinoside (Pg-3-rut) was found at a concentration of 101 $\mu\text{g mL}^{-1}$ in WB fruits and only 2 $\mu\text{g mL}^{-1}$ after GID, with a BI < 1%; however, it was not detected in CB fruits or digestates. Pt-3-glucoside (Pt-3-glu) was detected at a concentration of 42 $\mu\text{g mL}^{-1}$ in WB fruits; its concentration increased to 264 $\mu\text{g mL}^{-1}$ after GID, for a BI of 629%. In CB fruits, its concentration decreased from 607 $\mu\text{g mL}^{-1}$ to 312 $\mu\text{g mL}^{-1}$ after GID, for a BI of 49%.

Two isoforms of Cy-3-glucosylrutinoside (Cy-3-glu-rut-iso 1 and 2) were found in WB fruits: Cy-3-glu-rut-iso 1 was found at a concentration of 30 $\mu\text{g mL}^{-1}$ and Cy-3-glu-rut-iso 2 at a concentration of 54 $\mu\text{g mL}^{-1}$, respectively. Cy-3-glu-rut isomers were undetected after GID. In CB these anthocyanins were detected only during GP at low concentrations (<12 $\mu\text{g mL}^{-1}$). A similar pattern was observed for other anthocyanins in both

fruits: Cy-3-5-diglucoside isomer 1 (Cy-3-5-diglu-iso 1) was found at a concentration of 6 $\mu\text{g mL}^{-1}$ in WB and 30 $\mu\text{g mL}^{-1}$ in CB; Cy-3-5-diglucoside isomer 2 (Cy-3-5-diglu-iso 2) at a concentration of 3 and 8 $\mu\text{g mL}^{-1}$, respectively, in WB and CB; Cy-3-sophoroside (Cy-3-sop) at a concentration of 16 $\mu\text{g mL}^{-1}$ and 25 $\mu\text{g mL}^{-1}$ in WB and CB fruits, respectively; Cy-3-xylosyl-(feruloyl-glucosyl)-galactoside (Cy-3-xyl-(fer-glu)-gal) at a concentration of 2 $\mu\text{g mL}^{-1}$ in WB and not detected in CB; Cy-3-sambubioside (Cy-3-sam) at a concentration of 103 $\mu\text{g mL}^{-1}$ in WB and 14 $\mu\text{g mL}^{-1}$ in CB, before digestion; Cy-3-xylosyl-galactoside (Cy-3-xyl-gal) at a level of 103 and 14 $\mu\text{g mL}^{-1}$ in WB and CB, respectively; Cy-3-xyloside (Cy-3-xyl) at a level of 14 $\mu\text{g mL}^{-1}$ in WB and 8 $\mu\text{g mL}^{-1}$ in CB; Cy-3-acetyl-glucoside (Cy-3-act-glu) was found at a concentration of 35 $\mu\text{g mL}^{-1}$ in CB, but it was not detected in WB fruits; Dp-3-glucoside (Dp-3-glu) was estimated at 1 $\mu\text{g mL}^{-1}$ in WB and 16 $\mu\text{g mL}^{-1}$ in CB; Dp-3-acetyl-glucoside (Dp-3-act-glu) was quantified at 13 $\mu\text{g mL}^{-1}$ in CB, but it was not detected in WB samples; Pg-3-glucoside (Pg-3-glu) was found at a concentration of 67 $\mu\text{g mL}^{-1}$ in WB and 15 $\mu\text{g mL}^{-1}$ in CB; Pg-3-malonyl-glucoside (Pg-3-mal-glu) was found at a level of 10 $\mu\text{g mL}^{-1}$ in WB, but it was not detected in CB; Pn-3-glucoside (Pn-3-glu) was quantified at 7 $\mu\text{g mL}^{-1}$ in both WB and CB. In all these cases, the BI was 0%.

All concentrations of anthocyanins in WB fruits decreased during the OP; they increased in CB fruits following this digestive step. It is important to consider the role of α -amylase in the cleavage of non-reducing terminal 1–4 linked α -glucose residues; a higher content of carbohydrate concentration in CB could lead to substrate inhibition and interrupt anthocyanin cleavage during OP.⁵⁵ Considering that WB fruits had a higher TAC than CB, anthocyanins may have saturated the salivary amylase, retained them in their carbinol pseudobase/chalcone form, and thus, prevented subsequent enzymatic activity.^{9,56}

Frequently, anthocyanins are the first compounds that come into contact with the hostile environment of the mouth during mastication, which is when the metabolism of plant pigments begins.^{9,57,58} The physicochemical conditions in the oral cavity (pH, electrolytes, temperature, darkness, decreasing O₂ content, enzymes, etc.) have a negative influence on anthocyanin stability, usually resulting in anthocyanin degradation.^{6,59} First, mastication breaks down the macro- and microstructures of fruit cells during bolus formation, which helps to release anthocyanin. Then atmospheric O₂ starts to oxidize the free anthocyanins. The pH level (>7.0) triggers rapid (30–103 s) hydration of the flavylium cation at C2 (ring C), generating a carbinol pseudobase (it can also further equilibrate to the colorless open-ring chalcone form) and a loss of color.^{6,9,59} Deprotonation continues at a pH level of >6.0, forming violet–blue quinoidal and anionic bases, which are fully-conjugated cyclic diones. With the addition of water, two electrons are added to the resonant molecules (chalcone or carbinol pseudobase), increasing their electronegativity at C6 and C8 of ring A.⁹ After chalcone is formed, it undergoes deglycosylation and further chemical degradation.

In addition, enzymatic deglycosylation of the flavylium form generates unstable aglycones, which are then spon-





Table 1 Anthocyanin profile of WB and CB fruits subjected to simulated *in vitro* digestion by the INFOGEST protocol

Anthocyanin	R_f (min)	[M] ⁺ (m/z)	Frag (m/z)	Fruit	Wild blackberry (µg mL ⁻¹)										Commercial Blackberry (µg mL ⁻¹)										BI (%)
					OP	GP ₃₀ min	GP ₆₀ min	GP ₉₀ min	GP ₁₂₀ min	IP ₃₀ min	IP ₆₀ min	IP ₉₀ min	IP ₁₂₀ min	BI (%)	Fruit	OP	GP ₃₀ min	GP ₆₀ min	GP ₉₀ min	GP ₁₂₀ min	IP ₃₀ min	IP ₆₀ min	IP ₉₀ min	IP ₁₂₀ min	
Cy-3-5-diglu-iso 1	3.85	611	449; 287	6	63	61	42	53	53	0	0	0	0	0	30	12	72	68	68	50	0	0	0	0	0
Cy-3-5-diglu-iso 2	4.33	611	449; 287	3	20	25	16	21	22	0	0	0	0	0	8	2	21	22	23	20	0	0	0	0	0
Cy-3-glu-rut-iso 1	4.72	757	287; 611	31	286	352	194	301	298	21	5	6	0	0	nd	0	5	5	6	0	0	0	0	0	0
Dp-3-glu	5.08	465	303	1	30	12	8	9	10	5	0	0	0	0	16	5	102	98	93	92	0	0	0	0	0
Cy-3-sop	5.43	611	287	16	28	75	46	74	67	0	0	0	0	0	25	4	133	147	146	164	0	0	0	0	0
Cy-3-glu-rut-iso 2	5.45	757	287; 611	54	616	675	399	565	562	96	38	70	0	0	nd	0	8	9	8	11	0	0	0	0	0
Cy-3-glu	5.85	449	287	2854	7690	11 693	9413	11 338	11 172	197	53	51	8	<1	4994	184	15 735	16 305	16 111	14 216	222	217	63	16	<1
Cy-3-xyH-(fer-glu)-gal	5.86	919	287	2	4	8	5	10	10	0	0	0	0	0	nd	0	19	25	25	19	0	0	0	0	0
Cy-3-sam	5.95	581	287	103	240	394	342	365	356	8	0	0	0	0	14	1	54	54	53	41	0	0	0	0	0
Cy-3-xyH-gal	5.95	581	287	103	240	390	341	365	361	8	0	0	0	0	14	1	56	54	53	43	0	0	0	0	0
Cy-3-xyH-rut	6.12	727	287	2215	6171	9528	8429	8926	8823	473	319	175	80	<2	nd	0	4	12	3	27	0	0	0	0	0
Cy-3-rut	6.31	595	449; 287	3693	11 994	17 822	14 249	17 161	16 981	635	581	198	4	<1	1226	56	4319	4398	4379	3854	58	49	18	4	<1
Pg-3-glu	6.60	433	271	67	182	348	347	321	312	9	0	0	0	0	15	3	77	81	81	80	5	0	0	0	0
Pg-3-rut	7.00	579	271	101	412	462	710	650	635	40	22	15	2	<1	nd	nd	nd	nd	nd	nd	nd	nd	nd	nd	0
Pn-3-glu	7.17	463	301	7	22	37	23	36	34	0	0	0	0	0	7	1	37	40	39	31	0	0	0	0	0
Cy-3-xyH	7.25	419	287	14	29	55	33	48	49	0	0	0	0	0	8	0	44	50	47	27	0	0	0	0	0
Cy-3-mal-glu	7.58	535	287	466	1313	1916	1366	1721	1693	60	10	9	3	<1	160	30	986	967	953	712	11	11	5	2	<2
Dp-3-(6-p-cou)-glu	7.73	611	303	50	200	267	178	248	248	210	175	108	17	66	83	36	110	120	112	119	68	67	63	21	75
Dp-3-act-glu	7.77	507	303	nd	nd	nd	nd	nd	nd	nd	nd	nd	nd	0	13	0	6	7	6	9	0	0	0	0	0
Pt-3-glu	7.84	491	287	nd	nd	nd	nd	nd	nd	nd	nd	nd	nd	0	35	2	3	11	9	11	0	0	0	0	0
Cy-3-dioxal-glu	8.01	593	287	21	90	150	102	145	143	74	12	9	19	90	613	132	3302	3208	3120	2496	116	118	79	124	20
Pg-3-mal-glu	8.19	519	271	10	31	42	33	37	36	0	0	0	0	0	nd	nd	nd	nd	nd	nd	nd	nd	nd	nd	0

R_f , retention time; min, minutes; Frag, fragment; m/z , relationship mass/charge; OP, oral phase; GP, gastric phase; IP, intestinal phase. BI, bioaccessibility index. Cy-3-5-diglu-iso, cyanidin-3-5-diglucoiside isomer; Cy-3-glu-rut-iso, cyanidin-3-glucosyl-rutinoside isomer; Dp-3-glu, delphinidin-3-glucoside; Cy-3-sop, cyanidin-3-sophoroside; Cy-3-glu, cyanidin-3-glucoside; Cy-3-xyH-(fer-glu)-gal, cyanidin-3-xyloxy-(feruloylglucosyl)-galactoside; Cy-3-sam, cyanidin-3-sambubioside; Cy-3-xyH-gal, cyanidin-3-xyloxy-galactoside; Cy-3-xyH-rut, cyanidin-3-xyloxy-rutinoside; Cy-3-rut, cyanidin-3-rutinoside; Pg-3-glu, pelargonidin-3-glucoside; Pg-3-rut, pelargonidin-3-rutinoside; Pn-3-glu, peonidin-3-glucoside; Cy-3-xyH, cyanidin-3-xyloxy; Cy-3-mal-glu, cyanidin-3-malonyl-glucoiside; Dp-3-(6-p-cou)-glu, delphinidin-3-(6-p-coumaroyl)-glucoside; Dp-3-act-glu, delphinidin-3-acetylglucoside; Pt-3-glu, petunidin-3-glucoside; Cy-3-dioxal-glu, cyanidin-3-dioxalyl-glucoiside; Pg-3-mal-glu, pelargonidin-3-malonyl-glucoiside.

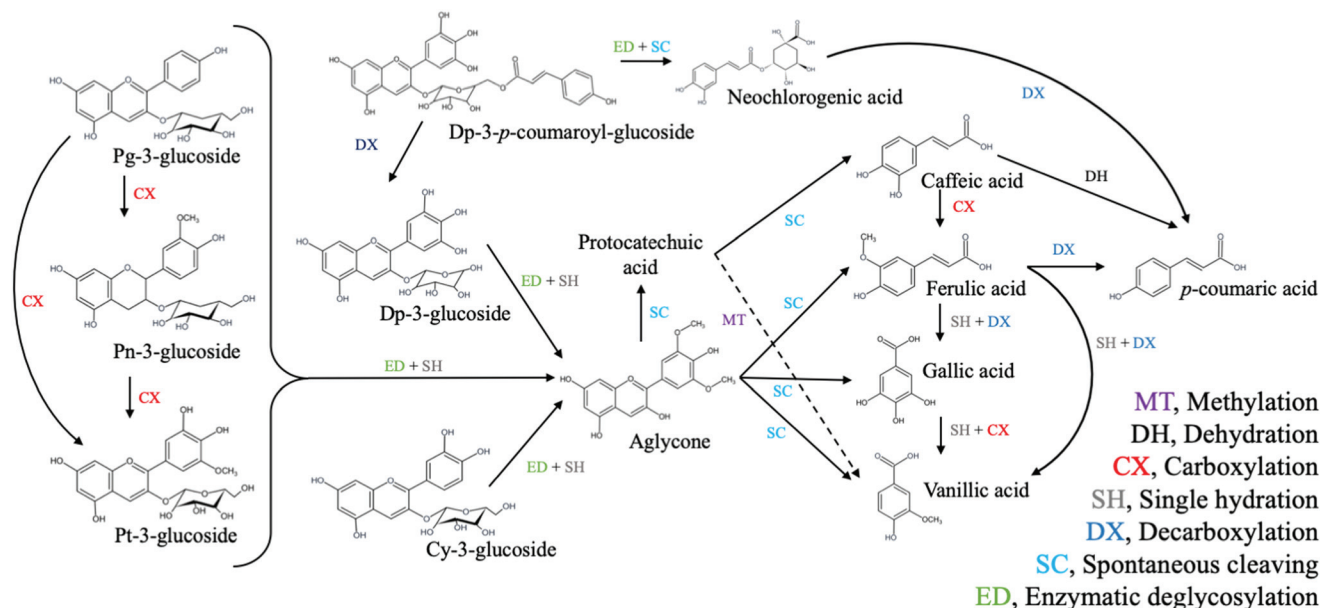


Fig. 2 Polyphenol metabolites implicated in anthocyanin metabolism during GID. Adapted from McGhie *et al.*,⁶⁰ Nurmi *et al.*,⁶¹ de Ferrars *et al.*⁶² and Olivas-Aguirre *et al.*⁵⁸ Cy, cyanidin; Dp, delphinidin; Pg, pelargonidin; Pn, peonidin; Pt, petunidin.

taneously cleaved into phenolic derivatives (Fig. 2). Amylase is the main enzyme synthesized in the oral cavity that contributes to anthocyanin degradation.⁵⁵ The sugar moiety is particularly important to the structural stability of the anthocyanins, and oral glycosidases cleave the glycosidic bond between the aglycone and glycosyl group. Recently, it has been suggested that it is responsible for the glycosylation of anthocyanidins, giving rise to anthocyanins and consecutive derivatives in the mouth.^{9,55} At the same time, amylases release anthocyanins from cell structures as well as large carbohydrates. The most common sub-products of anthocyanin deglycosylation, through spontaneous cleavage of aglycone, are protocatechuic, caffeic, gallic and vanillic acids. The last three phenolics could be produced by other spontaneous cleavage, decarboxylation, carboxylation, single hydration or dehydration processes.^{6,9,57,58,60–62} Since anthocyanins have low bioavailability after GID, the above-mentioned phenolic acids plus *p*-coumaric acid (one of the simplest metabolites of anthocyanins and other phenolic compounds produced during GID) are considered to be bioavailable and bioactive remnants of anthocyanins in the human body.^{58,63}

About 12 flavanols and proanthocyanidins found in the WB and CB samples are listed in Table 2. While the catechin content of WB fruits, initially $48.52 \mu\text{g mL}^{-1}$ and completely degraded after GID, while it remained $5.32 \mu\text{g mL}^{-1}$ in CB after GID (BI of 14%). The isoform of this flavanol, epicatechin, was quantified at $34.51 \mu\text{g mL}^{-1}$ in CB fruits and $5.35 \mu\text{g mL}^{-1}$ in the aliquot of IP_{120 min} (BI of 16%), while in WB samples its content was $412.19 \mu\text{g mL}^{-1}$ before GID and $3.91 \mu\text{g mL}^{-1}$ after GID (BI < 1%). Quercetin (Qc) and kaempferol (Kf) aglycones were detected at levels of 158.54 and $16.87 \mu\text{g mL}^{-1}$ in CB, respectively, and 385.35 and $47.97 \mu\text{g mL}^{-1}$

mL^{-1} in WB, respectively. These two aglycones were degraded after GID in both types of blackberries. In comparison with these compounds, their glycosylated forms seem to be more resistant to GID: Qc-3-glucoside (Qc-3-glu) was found at a concentration of $21159.74 \mu\text{g mL}^{-1}$ in CB fruits and then decreased to $8207.35 \mu\text{g mL}^{-1}$ after GID (BI of 39%). In WB, its content was $1360.17 \mu\text{g mL}^{-1}$ and then $6163.50 \mu\text{g mL}^{-1}$ after IP_{120 min} (an increase in BI of 453%). This flavanol was the major non-anthocyanin phenolic in both types of blackberries. Qc-3-glucuronide (Qc-3-glur) was detected at a level of $4123.47 \mu\text{g mL}^{-1}$ in CB fruits, dropping to $2240.40 \mu\text{g mL}^{-1}$ after GID (BI of 54%); it was quantified at 253.04 and $1522.41 \mu\text{g mL}^{-1}$ in fruits and digested WB (BI of 602%). Qc-3-galactoside (Qc-3-gal) or hyperoside was found at a level of 4847.24 and $1794.07 \mu\text{g mL}^{-1}$ in fruits and digested CB, respectively (BI of 39%). Corresponding concentrations of 198.87 and $1415.59 \mu\text{g mL}^{-1}$ were found in undigested and digested WB (BI of 712%). Kf-3-glucuronide (Kf-3-glur) was quantified at $1675.19 \mu\text{g mL}^{-1}$ in CB fruits and $1570.54 \mu\text{g mL}^{-1}$ after digestion (BI of 94%); its concentration was $63.35 \mu\text{g mL}^{-1}$ in WB fruits and $1195.54 \mu\text{g mL}^{-1}$ after GID (BI of 1887%).

Three isoforms of procyanidin dimers were found in CB fruits: procyanidin dimer iso-2, iso-3 and B2 in concentrations of 54.10, 41.41 and $6.33 \mu\text{g mL}^{-1}$, respectively. In WB fruits, procyanidin dimer iso-1, iso-2, iso-3 and B2 were quantified respectively at 55.24, 236.85, 7.71 and $28.96 \mu\text{g mL}^{-1}$. These procyanidins were totally degraded at the end of GID.

Flavanols and proanthocyanidins have low bioavailability in the human body. Salivary exposure has been found to produce no large changes in these phenolics, which is inconsistent with our findings.⁶⁴ A mild acidic environment is the main factor responsible for oligomeric proanthocyanidin or flavanol

Table 2 Non-anthocyanin profile of WB and CB fruits subjected to simulated *in vitro* digestion by the INFOGEST protocol

Phenolic Compound	R_T (min)	$[M]^-$ (m/z)	Frag (m/z)	Wild blackberry (mg mL ⁻¹)												Commercial blackberry (mg mL ⁻¹)												BI	
				Fruit						OP						Fruit						OP							
				GP ₃₀ min	GP ₆₀ min	GP ₉₀ min	GP ₁₂₀ min	IP ₃₀ min	IP ₆₀ min	IP ₉₀ min	IP ₁₂₀ min	(%)	GP ₃₀ min	GP ₆₀ min	GP ₉₀ min	GP ₁₂₀ min	IP ₃₀ min	IP ₆₀ min	IP ₉₀ min	IP ₁₂₀ min	(%)	GP ₃₀ min	GP ₆₀ min	GP ₉₀ min	GP ₁₂₀ min	IP ₃₀ min	IP ₆₀ min	IP ₉₀ min	IP ₁₂₀ min
Galic acid	2.60	169		1882.38	10 039.35	7674.73	7958.87	7557.92	7554.77	5213.67	4069.60	2171.20	639.02	34	489.34	1041.82	997.62	959.74	1048.13	11684.10	465.60	429.36	335.59	366.22	75				
Neochlorogenic acid	5.53	353		12.62	64.88	102.73	96.00	97.33	99.13	160.00	155.00	100.93	80.00	667	48.21	400.00	102.73	131.57	117.15	189.24	0.00	0.00	0.00	0.00	0				
Procyanidin	6.97	577	289	55.24	1859.48	1065.22	732.63	925.67	1025.68	0.00	0.00	0.00	0.00	0	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0				
Catechin	7.22	289		48.52	1392.75	707.79	654.52	602.51	452.83	289.20	169.21	76.20	0.00	0	38.37	5.71	133.19	101.48	109.09	261.30	43.91	35.52	15.22	5.32	14				
Vanillic acid iso-1	7.40	167		7.23	37.28	44.12	39.56	41.08	39.56	27.39	42.60	48.69	60.86	842	7.71	6.85	10.65	4.56	16.74	9.13	15.22	15.22	9.13	118					
Vanillic acid iso-2	7.60	167		270.35	15.59	17.59	17.05	18.82	19.63	32.26	105.26	57.43	80.97	30	105.03	62.92	86.91	89.46	88.92	75.72	109.73	117.65	141.13	184.27	175				
Caffeic acid	7.77	179		200.00	72.99	66.14	54.43	63.76	64.63	106.98	40.20	45.55	80.00	40	7.59	8.00	14.57	15.80	13.56	14.43	0.00	0.00	0.00	0.00	0				
Procyanidin	7.99	577	289	236.85	4291.99	2620.31	1853.36	2302.07	2505.69	0.00	0.00	0.00	0.00	0	54.10	0.00	164.91	135.59	140.60	326.25	0.00	0.00	0.00	0.00	0				
Procyanidin	8.18	577	289	7.71	63.34	58.47	42.22	51.97	63.34	51.05	0.00	0.00	0.00	0	41.41	5.20	128.30	116.93	141.29	87.70	0.00	0.00	0.00	0.00	0				
Epicatechin	9.06	289		412.19	5945.22	3217.02	2860.19	2696.88	2109.63	568.24	53.69	18.49	3.91	<1	34.51	9.22	187.95	138.51	103.20	95.51	19.20	14.08	10.82	5.35	16				
Procyanidin	9.71	577	289	26.98	437.32	159.03	145.40	124.95	159.03	63.61	0.00	0.00	0.00	0	6.33	5.11	57.93	39.76	43.16	101.10	0.00	0.00	0.00	0.00	0				
Procyanidin B2																													
<i>p</i> -Coumaric acid	9.84	163		14.97	106.96	95.08	73.07	92.44	91.55	206.00	172.55	195.43	192.00	1371	11.17	40.00	27.46	26.95	25.41	26.28	9.86	9.17	11.82	16.00	143				
Ferulic acid	11.21	193		6.95	64.36	51.48	40.16	47.37	55.60	144.16	107.09	111.21	92.67	619	107.60	84.44	96.79	101.94	102.97	75.17	102.97	105.03	102.97	90.61	84				
Qc-3-glu	12.15	463	301	1360.17	4198.95	6431.75	4292.09	6032.56	6225.24	2946.56	4560.35	5184.15	6163.50	453	2159.74	10244.82	34577.32	34674.19	34527.29	30249.04	23325.48	24262.25	21649.95	8207.35	39				
Qc-3-gal	12.18	477	301	253.04	1175.88	1500.80	963.97	1431.00	1474.21	744.58	940.70	1150.12	1522.41	602	4123.47	2931.80	7105.13	7573.82	7505.68	8516.19	4979.41	5072.48	4394.38	2240.40	54				
Qc-3-gal	12.42	289		198.87	725.97	1058.48	717.42	932.32	1028.54	547.42	801.88	868.17	1415.59	712	4847.63	2546.24	9734.83	9705.96	10051.30	8591.88	6119.95	6417.18	5786.37	1794.07	39				
KF-3-glu	13.50	461	285	63.35	298.88	351.79	188.43	336.01	341.58	207.92	358.29	454.82	1195.54	1887	1675.19	1599.31	3018.55	3233.90	3110.45	3406.55	3033.40	3235.75	3012.98	1570.54	94				
Quercetin	17.06	301		358.35	262.45	1003.49	933.42	1091.36	1037.92	28.00	0.00	0.00	0.00	0	158.54	11.28	299.26	305.20	315.89	237.51	0.49	0.00	0.00	0.00	0				
Kaempferol	19.39	285		47.97	22.14	97.00	73.81	103.33	97.00	6.40	2.45	0.00	0.00	0	16.87	0.00	16.87	18.98	23.20	14.76	0.00	0.00	0.00	0.00	0				

K_a, retention time; min, minutes; Frag, fragment; *m/z*, relationship mass/charge; OP, oral phase; GP, gastric phase; IP, intestinal phase. BI, bioaccessibility index. Qc-3-glu, quercetin-3-glucoside; Qc-3-gal, quercetin-3-galactoside; KF-3-glu, kaempferol-3-glucuronide.

R_T , retention time; min, minutes; Frag, fragment; m/z , relationship mass/charge; OP, oral phase; GP, gastric phase; IP, intestinal phase. BI, bioaccessibility index. Qc-3-glu, quercetin-3-glucoside; Qc-3-gal, quercetin-3-galactoside; KF-3-glu, kaempferol-3-glucuronide.

(>2000 Da) depolymerization. This breakdown may be good for the stability of dimeric procyanidins and acylated and aglycones of flavanols,⁶⁴ which could explain the increase in the concentration of these proanthocyanidins and flavanols during the GP. However, following the addition of pancreatic enzymes during the IP, their degradation is promoted, and glycosylated flavanols are progressively degraded, resulting in stable final products like phenolic acids, including gallic, ferulic, vanillic and hydroxybenzoic acids.⁶⁵

The six phenolic acids profiled in both types of blackberries are displayed in Table 2. Phenolic acids are the major metabolites of tannins, anthocyanins, flavanols and other polyphenol metabolism before absorption by the gut epithelium or their total degradation.⁶³

The gallic acid content was quantified at 489.34 $\mu\text{g mL}^{-1}$ and 366.22 $\mu\text{g mL}^{-1}$ in CB fruits and the digestate IP_{120 min}, respectively (BI of 75%). In WB, this phenolic acid was found at levels of 1882.38 and 639.02 $\mu\text{g mL}^{-1}$ in undigested and digested fruits, respectively (BI of 34%). Gallic acid is a hydroxybenzoic acid present in *R. liebmannii* in the form of hydrolysable (gallotannins and ellagitannins) and condensed (proanthocyanidins) tannins.⁶⁶ Despite the fact that gallic acid is unstable under alkaline conditions, its content increased in both types of fruit following the OP, possibly due to its liberation from the matrix by α -amylase. Gallic acid is released into the digestive matrix from metabolized proanthocyanidins, gallotannins and ellagitannins during degeneration of the cell structures of fruits, such as vacuoles and cell walls, by mechanical action, but particularly by enzymatic activity *via* non-covalent interactions between gallic acid- α -amylase.⁶⁶ Gallic acid could be a product of tannin hydrolysis, spontaneous cleavage of the cyanidin aglycone, or a single hydration + carboxylation of ferulic acid.^{6,62,67} The short time of exposure to mouth conditions could result in irreversible changes to gallic acid molecules, including enzymatic hydrolysis, depolymerization and deglycosylation,^{9,68} which are more evident than in the GP, since proteases (pepsin) and low pH (≤ 3) may favor the release of this phenolic acid and maintain its chemical stability in the food matrix.⁶⁹ Agudelo *et al.*⁷⁰ also reported an increase in gallic acid content during mouth and stomach simulation of the digestion of the Andean berry (*Vaccinium meridionale*). This is one of the most bioaccessible, bioactive and bioavailable phenolics in this fruit.

Neochlorogenic acid was not detected in CB samples after GID but it was found at a level of 48.21 $\mu\text{g mL}^{-1}$ in fruits. In WB, it was quantified at 12.62 $\mu\text{g mL}^{-1}$ and 80.00 $\mu\text{g mL}^{-1}$ in fruits and digestate, respectively (BI of 667%). Neochlorogenic acid is an isomer of chlorogenic acid, a hydroxycinnamic acid derived from caffeic acid; it is commonly found in many berry fruits.^{27,47,71} It is also a metabolite of acylated anthocyanins, specifically Dp-3-*p*-cou-glu, resulting from enzymatic deglycosylation + spontaneous cleavage.^{9,68} In contrast to the results found in the present study for CB digestates, the neochlorogenic acid content was found to decrease by 28% in chokeberries (*Aronia melanocarpa*)¹¹ and by 55% in blueberries (*Vaccinium sp.*)¹² after GID digestion; however, it was not

totally degraded, as was the case in digested WB fruits. In black mulberry (*Morus sp.*) wine, the chlorogenic acid content decreased by 43% during GID digestion.⁷² The non-detection of neochlorogenic acid in commercial blackberry IP digestates could be due to its isomerization under pancreatic incubation conditions, not just its degradation.¹¹

Two isomers of vanillic acid (iso-1 and iso-2) were found in both types of blackberries: vanillic acid iso-1 was quantified at 7.71 $\mu\text{g mL}^{-1}$ and 9.13 $\mu\text{g mL}^{-1}$ in undigested and digested CB fruits, respectively (BI of 118%). Its content was 7.23 $\mu\text{g mL}^{-1}$ in WB fruits and 60.86 $\mu\text{g mL}^{-1}$ after GID (BI of 842%). The concentration of vanillic acid iso-2 was 105.03 $\mu\text{g mL}^{-1}$ and 184.27 $\mu\text{g mL}^{-1}$ in CB before and after GID, respectively (BI of 175%). In WB samples, it was found at a level of 270.35 $\mu\text{g mL}^{-1}$ in fruits and 80.97 $\mu\text{g mL}^{-1}$ in the digestate IP_{120 min} (BI of 30%). Vanillic acid isoforms are derivatives of the hydroxybenzoic acid structure that is present in berry fruits or digested products in its native form, or as a metabolite from anthocyanins, flavanols or hydroxycinnamic acids.^{47,61} Vanillic acid isomers may be produced by spontaneous cleavage of anthocyanidins (like cyanidin), methylation of protocatechuic acid and/or single hydration + carboxylation of gallic acid.^{60,61} During *in vitro* GID of blueberries (*Vaccinium myrtillus*), the content of vanillic acid decreased from the 0.08 mg g⁻¹ fresh weight level until it was undetectable.⁶⁸ Nurmi *et al.*⁶¹ found an increase in homovanillic (+23%) and vanillic acids (+57%) in a mixed extract of blueberries (*Vaccinium myrtillus*) and lingonberries (*Vaccinium vitis-idaea*) after *in vivo* metabolism. Vanillic acid isoforms are recognized as products of protocatechuic acid metabolism, which in turn is produced by the hydrolysis of anthocyanins; this may explain the increase in the concentration of this phenolic acid at the end of GID.⁶²

Caffeic acid was quantified at 7.59 $\mu\text{g mL}^{-1}$ in CB fruits, but it was not detected after GID. In contrast, in WB it was found at a concentration of 200.00 and 80.00 $\mu\text{g mL}^{-1}$ before and after digestion of the fruits, respectively (BI of 40%). Caffeic acid is a hydroxycinnamic acid, the main metabolites of which are vanillic acid isomers, which could be produced by spontaneous cleavage of protocatechuic acid.^{47,58} The metabolism of this phenolic compound could contribute to the increase in vanillic acid isoforms. In the Andean berry, the caffeic acid content decreased by 74% after the IP⁷⁰ (a bigger decrease than in CB digestates, but smaller than in WB samples). An experiment with blueberry-lingonberry extract showed that <10% of ingested caffeic acid was recovered. Additionally, its intact chemical structure showed low bioavailability and bioaccessibility in comparison with its metabolites: 3-caffeoylquinic, 4-caffeoylquinic, ferulic and/or *p*-coumaric acids.⁷³

Ferulic acid was found at a concentration of 107.60 $\mu\text{g mL}^{-1}$ in CB fruits and at 90.61 $\mu\text{g mL}^{-1}$ after digestion (BI of 84%), compared with 6.95 $\mu\text{g mL}^{-1}$ in WB fruits and 92.67 after GID (BI of 619%). It is known that ferulic acid moieties can stabilize anthocyanins, which can generate co-pigmentation in the matrix even at pH 3.0–4.0 (such as in fruits or during the GP);^{59,62} however, this compound is also an intermediate



product of the catabolism of these flavonoids, the spontaneous cleavage of the B-ring of cyanidin aglycone and protocatechuic acid, or the spontaneous carboxylation of caffeic acid in the gastrointestinal tract.^{6,9,57,62}

In CB, *p*-coumaric acid was found at levels of 11.17 and 16.00 $\mu\text{g mL}^{-1}$ in fruits and digested samples (BI of 143%), whereas in WB its concentration was 14.97 and 192.00 $\mu\text{g mL}^{-1}$ in fruits and digested samples (BI of 1371%). *p*-Coumaric acid, a hydroxycinnamic acid, is found in blackberry fruits in moderate concentrations, but its content increases with the metabolism of cyanidin anthocyanins during GID, for example, *via* dehydration of caffeic acid and/or decarboxylation of ferulic and neochlorogenic acids.^{6,58,60,74} In blueberry fruits, Gapski *et al.*⁷³ observed that this phenolic compound was not reported after *in vivo* GID. It has been found in the bloodstream after berry consumption, which may be due to the metabolism of free phenolic acids, such as caffeic or ferulic acids, or degradation of flavonoids, including cyanidin and quercetin derivatives.^{75,76}

Polyphenols from red fruits are released into the digestive matrix and metabolized by the above-mentioned processes; however, it should be noted that other factors, such as oral, stomach and gut microbiota, and absorption during the three GID steps, have a strong influence on polyphenol bioaccessibility and bioavailability.⁷⁷ There is no consensus on the bioavailability of phenolic compounds with high molecular weight ($>450 \text{ g mol}^{-1}$), such as anthocyanins, flavanols and tannins. Native high molecular weight polyphenols are undetected or detected in very low amounts after digestion in plasma, tissues or urine.⁷⁸ This is because these phytochemicals are recognized as compounds with variable bioaccessibility and low bioavailability.⁷⁷ However, their metabolites are frequently found in body fluids in concentrations that do not correspond with their contents in fruits.⁷³

The digestion conditions are one of the main factors responsible for the biotransformation of these compounds; however, gut microbiota, interactions with other macro- and micro-compounds in the food matrix, age, health conditions, sex, and other factors are implicated in the profiling of polyphenol intake.⁷⁸ Thus, digestion increases the amount of phenolic metabolites (low molecular weight phenolics), enhancing their bioaccessibility, which could have a direct effect on their bioavailability and bioactivity.⁷⁹ More studies are required to better understand the *in vivo* effects of digestion and metabolism of phenolics and other phytochemicals.

3.4 *In vitro* antioxidant activity

The antioxidant capacity (AOX) of fruits is directly related to their chemical profile. Phenolics are the most important antioxidant phytochemicals in *Rubus* fruits.⁴² Phenolic anti-radical activity follows the HAT or SET mechanisms, or both, depending on many factors (*e.g.*, pH of the matrix and chemistry of the antioxidant agents).²⁶ The results of five *in vitro* bioactivity assays are summarized in Table 3.

3.4.1 ORAC. The results obtained with the ORAC assay showed higher AOX in WB fruits compared with CB fruits

Table 3 *In vitro* antioxidant activity of WB and CB fruits subjected to simulated *in vitro* digestion by the INFOGEST protocol

Bioactivity assay	Sample/digestates											
	BS	Fruits	OP	GP _{30 min}	GP _{60 min}	GP _{90 min}	GP _{120 min}	IP _{30 min}	IP _{60 min}	IP _{90 min}	IP _{120 min}	
ORAC	WB	4219.20 ± 111.70 ^a	2560.15 ± 47.85 ^b	1909.70 ± 37.30 ^d	1762.4 ± 12.35 ^e	1891.55 ± 60.30 ^d	2035.55 ± 37.05 ^c	2041.25 ± 32.95 ^c	2060.60 ± 159.00 ^c	2079.35 ± 46.95 ^c	2006.75 ± 55.85 ^c	
($\mu\text{M TE per mL}$)	CB	1902.55 ± 35.28 ^d	987.51 ± 31.28 ^{hi}	1234.76 ± 2.15 ^f	1231.12 ± 0.96 ^f	1233.51 ± 5.75 ^f	1233.20 ± 8.04 ^f	1089.49 ± 13.34 ^g	1036.51 ± 10.05 ^h	972.87 ± 20.44 ⁱ	829.57 ± 36.31 ^j	
DPPH RI (%)	WB	89.37 ± 0.00 ^a	61.20 ± 1.09 ^c	63.25 ± 6.89 ^c	60.82 ± 3.99 ^c	70.04 ± 3.52 ^b	61.01 ± 0.54 ^c	43.79 ± 6.97 ^e	47.12 ± 5.40 ^e	40.33 ± 8.90 ^{ef}	43.15 ± 3.26 ^{ef}	
	CB	39.50 ± 3.80 ^{ef}	42.77 ± 1.09 ^e	56.34 ± 7.70 ^{cd}	45.33 ± 9.23 ^e	52.56 ± 1.63 ^d	42.96 ± 4.89 ^e	26.63 ± 7.68 ^f	34.31 ± 9.06 ^f	41.87 ± 12.2 ^{ef}	41.61 ± 6.52 ^{ef}	
ABTS RI (%)	WB	45.29 ± 4.81 ^a	25.58 ± 3.32 ^f	42.45 ± 3.54 ^{ab}	41.29 ± 3.21 ^{ab}	39.87 ± 4.06 ^b	38.36 ± 3.01 ^{bc}	31.32 ± 3.24 ^{de}	31.38 ± 2.06 ^d	28.96 ± 2.19 ^c	26.96 ± 1.99 ^{ef}	
	CB	40.27 ± 3.59 ^b	23.05 ± 2.10 ^{fg}	35.36 ± 1.36 ^c	36.89 ± 2.87 ^{bc}	35.87 ± 2.00 ^c	32.56 ± 3.51 ^d	24.22 ± 2.17 ^{fg}	25.23 ± 2.38 ^f	22.73 ± 1.71 ^g	21.45 ± 2.11 ^g	
β -Carotene	WB	93.86 ± 11.01 ^a	64.93 ± 1.12 ^d	38.61 ± 9.26 ^h	38.11 ± 10.70 ^h	39.18 ± 6.52 ^h	35.90 ± 2.40 ^j	64.58 ± 7.82 ^{cde}	73.99 ± 5.88 ^b	69.36 ± 1.73 ^c	54.53 ± 3.66 ^{gh}	
Inhibition (%)	CB	61.35 ± 1.01 ^e	45.22 ± 11.95 ^h	45.22 ± 11.95 ^h	46.18 ± 5.08 ^h	37.26 ± 5.45 ^h	24.10 ± 3.25 ^j	73.18 ± 3.92 ^b	67.13 ± 3.58 ^{cd}	64.29 ± 4.37 ^d	58.11 ± 0.16 ^f	
Reducing power	WB	0.82 ± 0.05 ^a	0.55 ± 0.01 ^b	0.45 ± 0.03 ^a	0.39 ± 0.04 ^c	0.43 ± 0.01 ^d	0.43 ± 0.00 ^d	0.46 ± 0.01 ^d	0.44 ± 0.02 ^d	0.50 ± 0.02 ^c	0.49 ± 0.00 ^e	
(ABS 700 nm)	CB	0.40 ± 0.00 ^e	0.45 ± 0.01 ^d	0.35 ± 0.00 ^g	0.33 ± 0.01 ^h	0.36 ± 0.00 ^f	0.33 ± 0.00 ^h	0.40 ± 0.05 ^e	0.39 ± 0.00 ^e	0.31 ± 0.02 ⁱ	0.34 ± 0.01 ^h	

BS, blackberry species; WB, wild blackberry; CB, commercial blackberry. OP, oral phase; GP, gastric phase; IP, intestinal phase. $\mu\text{M TE per mL}$, μM of Trolox equivalents per mL; RI, radical inhibition. Mean \pm s.d. ($n = 3$), equal letters on the same line indicate a non-statistical difference ($p < 0.05$).



(4219.20 and 1902.55 μM TE per mL, respectively). Following the OP, the AOX of both types of fruit was drastically decreased. Afterward, it increased again in CB, reaching a constant value until GP_{120 min}. The value for WB at GP_{120 min} was lower ($p < 0.05$) than that for the undigested fruits; it was also lower following the OP, at 2035.55 μM TE per mL. After that, the AOX for WB did not change, whereas it declined to 829.57 μM TE per mL in CB at the end of GID. The results showed statistically ($p < 0.05$) higher values in WB compared with those in CB at every single GID stage. The AOX of IP_{120 min} digestates for WB and CB decreased by 52% and 56%, respectively, compared with the values for undigested fruits. The ORAC assay measures the capacity of an antioxidant to scavenge $\text{H}_2\text{O}_2^{\cdot}$ produced by the AAPH radical generator. Free phenolic acids and other metabolites of anthocyanins, flavanols and tannins from berries have a high antioxidant effect on $\text{H}_2\text{O}_2^{\cdot}$, which is one of the major radicals produced in the human body and may cause cellular damage to membrane lipids, DNA and other valuable macromolecules.^{42,48} Phenolic acids and flavonoids with free OH groups are free-radical scavengers that are able to interrupt oxidation chains *via* the HAT mechanism. These phenolics inhibit the degradation of fluorescence after being mixed with an azo-initiator (AAPH), which generates ROO^{\cdot} at body temperature.²⁹

Undigested samples of WB had AOX similar ($>3800 \mu\text{M}$ TE per mL) to that reported by Sánchez-Velázquez *et al.*^{16,66} in wild blackberries (*R. liebmannii* and *R. palmeri*) collected in Sinaloa during the 2014 season, but higher than that of CB. Fresh fruits of wild *R. croceacanthus* and *R. sieboldii* showed ORAC values of $>4000 \mu\text{mol}$ TE per mL, closer to that of WB, but in commercial fruits of *R. idaeus* ORAC showed a value of $\sim 2000 \mu\text{mol}$ TE per mL,⁸⁰ similar to that of CB. Cerezo *et al.*⁸¹ observed that the AOX of the Camarosa strawberry (*Fragaria* \times *ananassa*) as determined by the ORAC assay decreased by 43% after GID. Tavares *et al.*⁴⁶ reported a decrease of 83.1% in the AOX of cultivated blackberries after GID. However, in *R. adenotrichos*, no statistical difference was found between undigested and digested juice.⁴⁸ Alternatively, Chiang *et al.*⁸² found that, during digestion of European currant fruits, antioxidant capacity increased by up to 11%. With respect to GID of black elder (*Sambucus nigra*), Zhou⁸³ observed a 19% decrease in the antioxidant capacity as evaluated by ORAC; however, the OP was omitted in this model. As for the black elderberry, our results showed a decrease of 52.43% in the antioxidant capacity of WB and a decrease of 56.39% in CB as assessed by the ORAC method. Despite the fact that numerous studies, in which commercial and wild fruit matrices were subjected to *in vitro* digestion, have shown that digestion improves their antioxidant activity,^{48,82,84,85} more research is required to harmonize the digestion methods, to evaluate the impact during different digestive stages, and to determine which compounds are responsible for the AOX activity found.

3.4.2 DPPH scavenging activity. The inhibition rate (IR) of DPPH radicals in WB fruits was 89.37%, compared to 39.50% for CB fruits. The inhibition rate in WB decreased to 61.20% following the OP and it was not statistically different at

GP_{120 min}. On the other hand, the value for CB fruits did not change during OP as it did in the undigested fruits; however, it increased to 56.34% at GP_{30 min} and then decreased to 42.96% at GP_{120 min}. The AOX of WB at IP_{30 min} was 43.79% and there was no statistically significant change ($p < 0.05$) during the IP. However, the AOX values for CB increased ($p < 0.05$) from 26.63% to 41.61% between IP_{30 min} and IP_{120 min}.

After GID, the AOX of WB fruits was less than 50% of the value for undigested fruits. Despite the variations in the IR during GID of CB, no statistical difference ($p < 0.05$) was observed between the undigested fruits and the final GID digestate. The DPPH radical antioxidant method is a popular method for determining antioxidant activity; this involves monitoring the decrease in the initial absorbance of the radical at 517 nm.²⁸ DPPH $^{\cdot}$ is easily reduced by the action of flavonoids (like quercetin) *via* HAT or sequential proton loss electron transfer (SPLET) mechanisms, depending on the polarity of the medium. Independently of this, at the end of the reaction, the structure of the flavonoid is stabilized after the loss of one or two H^+ from the OH group at positions A7, B4' or B5' and subsequent opening of the C ring.²⁹ The A7-OH group has a key role as the main molecular site of ionization and electron (e^-) transfer, following a SPLET mechanism in flavonoids.^{86–88} DPPH $^{\cdot}$ is quenched after a H^+ or electron (e^-) is accepted from hydroxycinnamic acids to become DPPH₂. Low molecular weight phenolic acids can easily remove one DPPH radical from the medium, but their polymers or flavonoids may scavenge more than two of these radicals through the strong density of the rings A and B. However, an additional mechanism may be present in the DPPH $^{\cdot}$ quenching chain, specifically single-electron transfer followed by proton transfer (SET-PT).⁸⁹ However, very few data are available on this mechanism for polyphenols in a digestive matrix.

At a concentration of $100 \mu\text{g mL}^{-1}$, ethanolic extracts from wild fruits of *R. alpestris*, *R. fraxinifolius* and *R. moluccanus* showed DPPH $^{\cdot}$ inhibition of 95%, 59% and 86%, respectively,⁹⁰ where *R. alpestris* and *R. moluccanus* had an AOX similar to that of undigested WB, but all of them were considerably higher than that of CB. By digesting an extract rich in polyphenols from the Chilean strawberry, Thomas-Valdés *et al.*⁸⁵ reported that the antioxidant capacity evaluated by DPPH increased by 27.15% when subjected to GID, using the INFOGEST protocol. Azofeifa *et al.*⁴⁸ also reported an increase of 3.77% in the IR of digestates from *R. adenotrichos*. However, in a study using pulp from *Vitis vinifera*, Corrêa *et al.*⁹¹ found no significant changes between the undigested samples and after GID. Zhou⁸³ observed a decrease in phenolics and anthocyanins during GID of black elderberry (*Sambucus nigra*) and a corresponding decrease of more than 75% in antioxidant activity as assessed by DPPH.

3.4.3 ABTS scavenging activity. The inhibition rate (IR) of ABTS radicals in WB and CB fruits showed statistically different ($p < 0.05$) values of 45.29% and 40.27%, respectively. During the OP, inhibition was significantly decreased to 25.58% and 23.05% for WB and CB, respectively, but these values were not statistically different ($p < 0.05$). The result rep-



resents a decrease of >40% of inhibition of ABTS. Nevertheless, at GP_{30 min} the IR increased again to a value similar to that obtained for undigested WB and it remained stable until GP_{90 min}, and subsequently decreased to 38.36% at GP_{120 min}. The IR of CB samples was not statistically affected during the GP. However, during the IP, the AOX for both fruits decreased, reaching 26.96% and 21.45% in WB and CB, respectively, at the end of GID. Both the HAT and SET mechanisms are reported in the ABTS method because ABTS solubility is not influenced by the pH of the medium. Therefore, the use of this assay alone to determine the possible antioxidant mechanisms of the assayed compounds is not recommended.⁹² ABTS is oxidized by oxidants to its radical cation (ABTS^{•+}), thus, AOX is measured through the ability of test compounds to decrease the color by reacting directly with the ABTS.²⁹

Abu Bakar *et al.*⁹⁰ evaluated the TEAC in ethanolic extracts of wild fruits of *R. alpestris*, *R. fraxinifolius* and *R. moluccanus* and found an AOX of 0.73, 0.75 and 0.79 mg of ascorbic acid equivalent per g of dry sample, respectively. The authors attributed this to the high content of phenolic compounds, similar to that found in WB and CB. In their work, Mihailović *et al.*⁹³ found an antioxidant capacity of 39.5 µM TE per g for a fresh wild raspberry extract rich in polyphenols, flavonoids and anthocyanins that was subsequently subjected to GID. However, the contribution of these phenolics to AOX was not reported. During the GID of commercial raspberry, Gião *et al.*³⁸ did not find a significant difference in AOX, as evaluated by the ABTS method, between the undigested sample and the value measured after OP. However, a significant decrease of >70% was observed for AOX during the GP (relative to the OP). Following the GP stage, the AOX values increased to a value higher than that before GID, possibly because phenolics are more stable and still exhibit their AOX at an acidic pH.^{26,48} Using INFOGEST, a study on a polyphenol-rich extract from the Chilean strawberry (*Fragaria chiloensis*) found that the antioxidant capacity was reduced by 47.66% when it was subjected to GID.⁸⁵ In this study, we obtained maximum antioxidant capacity values for undigested samples and during the gastric stage, a finding that is attributable to the stability of anthocyanins and other polyphenols that are stable under the digestive conditions of the stomach, but are easily degraded in the small intestine. This could explain the decrease in antioxidant capacity of up to 40.47% in WB and 46.73% in CB, as assessed by ABTS at the end of the IP. The use of both the ABTS and ORAC assays is recommended for measuring the AOX of phenolics in intestinal digestates at a pH of 7. However, it is necessary to use other AOX methods as well to confirm the radical scavenging activity.^{26,29}

3.4.4 β-Carotene bleaching activity. The bleaching rate of β-carotene was 93.86% and 61.35% for WB and CB fruits, respectively. These values decreased during the OP (to 64.93% in WB and 56.13% CB) and during the GP, reaching 35.90% in WB and 24.10% in CB at GP_{120 min}. Following the GP, a major increase was observed for CB at IP_{30 min} ($p < 0.05$), and it was slightly higher than the corresponding value for WB at IP_{30 min} and higher than that at IP_{120 min}. When β-carotene is exposed

to O₂, it starts a chain of reactions (a series of rotations and vibration interactions with the solvent) that result in a yellowish-orange medium.²⁹ Phenolics incorporated into β-carotene–O₂ water may prevent the oxidation of β-carotene, and thus, its AOX may be estimated according to a HAT mechanism.²⁸

The β-carotene bleaching activity found in 10 µg TPC per mL of extracts from nine Brazilian blackberry cultivars ranged from 30 to 70%.⁹⁴ Undigested WB showed a higher β-carotene bleaching activity than these nine *Rubus* varieties, but the results of CB agree with these genotypes. The O₂[•] inhibition of different fruits of domesticated blackberry genotypes was 40–74%,⁹⁵ which was lower than the value reported for WB, but similar to the value reported for CB. Corrêa *et al.*⁹¹ observed a 36% increase in the ability to inhibit O₂[•] during the digestion of *Vitis vinifera* pulp. Our results showed a final decrease of 41.77% in the ability to inhibit O₂[•] in WB and only 5.28% in CB. The final AOX value as obtained by using this method for various food materials subjected to GID has been reported and compared against other antioxidant capacity protocols;⁹⁶ however, they were not used to compare the AOX values of the sample before and after GID. Since the β-carotene bleaching method employs the O₂[•] radical, it can be considered an antioxidant evaluation protocol similar to that occurring under physiological redox conditions.²⁸

3.4.5 Reducing power. The reducing power (RP) of both types of fruit was affected during GID. An Abs of 0.82 was observed in WB fruits, compared with 0.40 in CB. Following the OP, the Abs decreased to 0.55 in WB and 0.45 in CB. WB showed values of 0.45 between GP_{30 min} and GP_{120 min}, with no significant difference ($p < 0.05$) during this stage. The Abs of CB showed a statistically significant decrease to 0.33. At the end of the IP, the value for WB was 0.49, which was higher than the values observed during the GP. CB did not show a statistically significant difference ($p < 0.05$) between GP_{120 min} and IP_{120 min}. The reducing power decreased by 56% in WB digestates compared with undigested samples; the RP of WB decreased by 40% during GID. CB showed a decrease of 15% at the end of GID. The AOX of WB was statistically ($p < 0.05$) higher than that of CB.

The RP method measures the ability of an antioxidant (or any other molecule) to donate an electron to a radical. It is based on the reduction of a ferroin analog, the Fe³⁺ complex of tripyridyltriazine Fe³⁺.²⁸ The Fe³⁺ reducing power is a useful method that may emulate the ability of ingested phenolics to modulate the redox tone in plasma and tissues through the HAT mechanism.²⁹ An ethanolic extract from fruits of *R. ellipticus* (100–200 mg L^{−1}) showed an Abs of 0.43–1.11.⁹⁷ It is important to mention that WB showed values similar to the highest concentrations of *R. ellipticus* (150–200 mg L^{−1}), and CB had barely less RP than the lowest tested amount of *R. ellipticus* fruit extract. In contrast with our findings, a study on a pomace of grapes (*Vitis vinifera*) showed that the reducing power increased by 36% after GID.⁹¹ The difference in results may be attributable to the fact that the digestive conditions differed between the two studies and to the use of an extract from grapes in the other study.



3.5 Cellular bioactivity

Many studies suggest that *in vitro* assays on food antioxidants need to be supported by cellular assessment to elucidate the physiological anti-radical activity in the human body.^{35,36} Cellular bioactivity results for berry digestates indicate that the phenolics in these fruits contribute to enhanced CAA, inhibition of oxidative stress, apoptosis, aging, or neurological diseases.^{29,46,57,71,98} The cytotoxicity, CAA, inhibition of ROS-synthesis effects on Caco-2 cells of digested and undigested WB and CB samples are summarized in Table 4.

3.5.1 Cytotoxicity in Caco-2 cells. The cytotoxicity of the undigested samples of WB and CB showed mean-maximum effective concentration (EC₅₀) values of 0.50 mg GAE per mL and 1.20 mg GEA per mL, respectively, with significant differences ($p < 0.05$) between them (Table 4). However, with GID, the EC₅₀ was 1.14 mg GAE per mL for the WB and 1.55 mg GAE per mL for the CB, an EC₅₀ increase of 56% and 23%, respectively, indicating a lower level of cytotoxicity than in undigested fruits. HepG2 cells, when exposed to undigested and digested blueberry (*Vaccinium angustifolium*) samples, showed a cytotoxicity decrease of 85%.¹² In digested Chinese bayberry (*Myrica rubra*) extracts, the cytotoxicity was 44–60% lower than that in undigested extracts.⁹⁸ These studies are in agreement with our results, given that an increase in some free phenolic acids was observed following GID. This means that the degradation of large phenolics (e.g. tannins and flavonoids) and the increase in small phenolics (e.g. phenolic acids) in “chyle” during GID, may favor an increase in cytotoxicity in both Caco-2 and HepG2 cell lines.

3.5.2 Cellular antioxidant activity. The undigested samples of WB and CB showed a fluorescence inhibition rate of 85.88% and 78.70%, respectively, compared with the quercetin control; these values are statistically different ($p < 0.05$) from each other (Table 4). The IP_{120 min} samples indicated fluorescence inhibition of 58.39% in WB and 43.49% in CB, with significant differences ($p < 0.05$) between both digestates and the undigested samples. In summary, the GID of WB and CB samples produced 32% and 45% less CAA, respectively, than for undigested fruits.

As reflected in the cytotoxicity of the blueberry to HepG2 cells, Jiao *et al.*¹² reported a decrease of 85% of CAA at the end of GID; they also reported a strong correlation with the TPC

and cytotoxicity. These results are consistent with those reported by Huang *et al.*⁹⁸ and Boaventura *et al.*⁹⁶ for *Myrica rubra* and *Ilex paraguariensis*, respectively. According to Huang *et al.*,⁹⁸ the CAA of Chinese bayberry pomace improved after GID. However, in contrast with these authors' findings, we observed inhibition in CAA with GID, maybe because we used whole fruits rather than a fruit extract. During digestion, phenolics are metabolized and they can exhibit bioactivity while interacting with the gut epithelial surface.⁵⁸

The CAA assay evaluates the ability of an antioxidant to interrupt the reaction between ROO· (generated by AAPH) and a target probe.³⁵ It has gained importance in recent years because it provides more relevant information on antioxidant mechanisms in experimental cells lines than in *in vitro* (chemical) AOX assays. This method considers some aspects related to the take up, metabolism and location of antioxidants within cells, making it a potent tool for comparison in conjunction with *in vivo* AOX testing.³⁵ Nevertheless, it is recommended to highlight the physiological relevance of CAA over other cellular and *in vitro* AOX assays.

3.5.3 Cellular oxidative stress. Intracellular oxidative damage related to the formation of reactive oxygen species (ROS) in the undigested samples of WB and CB showed an inhibition of 72.53% and 38.87%, respectively, with significant differences ($p < 0.05$) between these values (Table 4). The values reported at IP_{120 min} indicated an inhibition of 36.95% in WB and only 5.37% in CB, with a significant difference between them ($p < 0.05$). This means that, after GID, the capacity to decrease ROS formation in Caco-2 cells of blackberry digestates was drastically inhibited, by about 50.05% in WB and by up to 86.14% in CB. Also, although the capacity to inhibit the formation of intracellular ROS decreased after GID, the pattern was not the same in WB and CB. Arango-Varela *et al.*⁹⁹ observed a reduction in ROS of 63–78% in samples of the Andes berry (*Vaccinium meridionale*), which was attributed to the presence of phenolics with different molecular weights. According to this, anthocyanidins (like cyanidin chloride) have greater ROS-synthesis inhibitory activity than small phenolics (like gallic acid). Nevertheless, the inhibition of ROS-synthesis by digested samples rich in phenolics is still limited.

The cellular oxidative stress assay evaluates the ability of an antioxidant to prevent the formation of oxidative sub-products in cells when they are exposed to a stress inductor (*i.e.*

Table 4 Cellular bioactivity of WB and CB fruits subjected to simulated *in vitro* digestion by the INFOGEST protocol

Cellular assay	Before GID		After GID	
	WB	CB	WB	CB
Caco-2 cells				
Cytotoxicity (EC ₅₀ , mg GAE per mL)	0.50 ± 0.04 ^c	1.20 ± 0.07 ^b	1.14 ± 0.08 ^b	1.55 ± 0.07 ^a
CAA ^a (%)	85.88 ± 7.30 ^a	78.70 ± 2.61 ^b	58.39 ± 2.80 ^c	43.49 ± 0.00 ^d
FIR ^a (%)	72.53 ± 6.71 ^a	38.87 ± 4.23 ^b	36.95 ± 2.62 ^b	5.37 ± 1.06 ^c

WB, wild blackberry; CB commercial blackberry. mg GAE per mL, mg of gallic acid equivalents per mL; CAA, cellular antioxidant activity; FIR, fluorescence increase ratio. ^a % inhibition of fluorescence (100 µg GAE per mL). Mean ± s.d. ($n = 3$), equal letters on the same line indicate a non-statistical difference ($p < 0.05$).



H₂O₂).³⁶ Cellular oxidative stress has been linked to various degenerative diseases.³⁶ Phenolics from berries and other plant sources help to prevent the formation of intracellular ROS in the Caco-2 cell model, as well in RAW 264.7 macrophages.^{56,99,100} Small phenolic acids, like those found in blackberry digestates, reduce oxidative stress induced by ROS in Caco-2 cells.^{101,102} Therefore, the metabolites of these anthocyanins and flavanols may play a key role in radical scavenging activity at the intestinal epithelium by direct interaction between ROS and phenolic acids. However, phenolic metabolites might upregulate the activation of the Nrf2/Keap1-ARE signaling pathway, an endogenous cellular antioxidant system related to the production of self-antiradical cell enzymes like glutathione reductase and oxidase in gut epithelial cells.¹⁰²

4. Conclusions

Wild blackberries are underutilized fruits with a high content and large diversity of polyphenols. Despite the fact that WB samples had 70% less TPC than CB at IP_{120 min} (68% less), WB still showed higher TPC than CB in the final digestate. Similar behavior was observed for TAC, where the TAC of WB was 75% lower at IP_{120 min}, but statistically ($p < 0.05$) higher than that of CB with 72% lower TAC at the end of GID. Several anthocyanins resisted the hostile environment of the gastrointestinal tract during digestion, including Cy-3-glu, Cy-3-rut, Cy-3-xyl-rut, Dp-3-(6-*p*-cou)-glu and Pt-3-glu, as well some flavanols such as epicatechin, Qc-3-glu, Qc-3-glur and Kf-3-glur. In contrast, the concentrations of many phenolic acids, such as gallic, vanillic, *p*-coumaric and ferulic acids, increases following digestion, which suggests that the catabolism of anthocyanins, flavanols and tannins may result in an increase in free phenolic acids in the digestive matrix.

The ORAC and ABTS assays showed that WB IP_{120 min} digestate had an AOX of >50% after GID, compared with an AOX of <50% for CB. The DPPH results also indicated that AOX was >50%; however, after GID, AOX values were statistically ($p > 0.05$) similar for undigested CB and digested WB. The β -carotene bleaching activity and reducing power methods showed a less significant decrease in CB than in WB IP_{120 min} digestates. However, the cytotoxicity of WB to Caco-2 cells was more inhibited than that of CB after GID. At the same time, the CAA of digested blackberries showed a statistically significant decrease ($p < 0.05$) of 48% in WB and 56% in CB following GID. Finally, the inhibition of the ROS-synthesis capacity in Caco-2 cells decreased by 50% in WB IP_{120 min}, and up to 86% in CB IP_{120 min}. WB polyphenols exhibited similar bio-transformations to those of CB phenolics during GID. However, certain particularities, such as the higher increase in many antioxidant free phenolic acids, suggest that WB provide better antioxidant effects than CB. The results also indicate that bioaccessible phenolics from WB may have a higher bio-availability and bioactivity in the human body than CB polyphenols. Therefore, WB could be considered an emerging food with potential health benefits.

Author contributions

Conceptualization: O.A.S.-V. and A.J.H.-Á.; methodology: O.A.S.-V. and M. Mulero; investigation: O.A.S.-V. and M. Mulero; data curation: O.A.S.-V. and Y.A.; writing – original draft: O.A.S.-V. and A.J.H.-Á.; supervision: E.O.C.-R. and A.J.H.-Á.; writing – editing: E.O.C.-R., M. Mondor and A.J.H.-Á.; validation: M. Mondor. and Y.A.; resources: Y.A.

Conflicts of interest

The authors declare that they have no known competing financial interests or personal relationships that could have impacted on the work reported in this paper.

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