Opuntia ficus-indica L. Mill Residues—Properties and Application Possibilities in Food Supplements

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Featured Application: In this paper we developed tablets made from Opuntia ficus-indica residues, which can be used as a product of interest for the food supplement industry.

Abstract: Recently, industry has been focusing on the development of new products made from food by-products in order to reduce and take advantage of food wastes. The objective of this study was to evaluate tablet formulations developed by mixing two commercial excipients, microcrystalline cellulose (M) and α-lactose-monohydrate (L), added with powder from residues (mesocarp and pericarp) of green and red (G and R) cactus pear fruit (Opuntia ficus-indica L. Mill), having the following formulations: green with microcrystalline cellulose (GM), green with lactose (GL), red with microcrystalline cellulose (RM), and red with lactose (RL). The results showed lower disintegration times for the tablets with microcrystalline cellulose. The fiber functional properties presented good values for lipid and water holding capacity. There was a higher total phenolic content (TPC) in formulations with green cactus pear residue powder with microcrystalline cellulose and lactose (GM and GL, respectively), but the DPPH and ferric reducing/antioxidant power (FRAP) values were higher in the formulations with red cactus pear residues (RM and RL), while ABTS values were similar among all formulations. In conclusion, tablets made from Opuntia residues are proposed as a product of interest for the food supplement industry because of the good quality parameters and the functional and antioxidant properties that they provide.

Keywords: Opuntia ficus-indica; residues; tablet; functional properties; antioxidants
1. Introduction

Dietary supplements are important sources of nutrients widely used by the population, and they play an essential role in disease prevention [1]. There are many kinds of dietary supplements, such as vitamins, minerals, herbs, and oils, among others. A dietary supplement is defined “as a product that contains one or more concentrated nutrients [2], with the objective of increasing the intake of some nutrients in the daily diet”. These supplements are regulated by the Food and Drug Administration (FDA) as food and not as drugs [3]. Nowadays, the supplement industry is growing fast since people expect to find ready-to-eat products from natural sources such as fruits, vegetables, or herbs with bioactive compounds which promote health [4]. However, several studies have reported that different agro-food by-products such as fruit and vegetable residues, including Opuntia ficus-indica L. Mill residues, possess bioactive compounds with a wide spectrum of pharmacological properties, such as neuroprotective, antidiabetic, antioxidant, anti-inflammatory, hepatoprotective, and anticancer activities, and can be used as raw materials for the development of nutraceuticals, medicines, food, and cosmetics useful to the prevention or control of diseases [5,6].

In this context, the cactus pear fruit (Opuntia ficus-indica L. Mill.) is a product that has been recognized for the health benefits that it can provide, and it has also drawn scientific interest for several years. These species are grown in Mexico and are arborescent plants with elliptic cladodes. Approximately >90% of the production of the fruit is of the yellow or green cultivars; the red type is less abundant (7–8%) [7]. Cactus pear fruit is generally consumed fresh, but it has also the potential to be processed into products like juices or jam [8]. The fruit has juicy pulp with numerous seeds immersed in it, surrounded by a thick peel with small spines [9]. Consumption and processing generate residues such as peels and seeds, which are nonedible portions of the fruit and represent 37% to 67% of the total weight [10]. These residues constitute a problem for environmental pollution due to the high amount of wastes produced and the associated fermentation phenomena [11].

The nonedible portion constituting the peel comprises two fractions, the mesocarp and pericarp, which are rich in antioxidants, soluble sugars, and dietary fiber, as well as trace elements such as calcium, magnesium, potassium, and manganese [12,13]. It has been reported that cactus pear peel has twice the dietetic fiber content of the fruit pulp, with a high proportion of insoluble fiber which provides beneficial effects for human health, such as lipid-lowering and hypoglycemic actions [14,15]. In addition, dietary fiber can be linked to antioxidant compounds like polyphenols, flavonoids, or phenolic acids, becoming bioactive in the human intestine once it is released from the food matrix [16]. Different products have been developed with cactus pear residues, such as yogurts, snacks, and margarine [17–19], and some other fruit wastes (pineapple, banana, or orange peels) have been used for the creation of dietary supplements or products for the pharmaceutical industry [20–22].

The drugs or food supplements are introduced into the body through the mouth and are absorbed in the digestive tract; among other dosage forms such as liquids, emulsions, or powders, tablets provide a convenient and easy method of administration [23]. The tablet formulations must observe specific quality parameters that allow the characteristics of the product to be preserved without physical changes during transportation and packaging, as well as evaluation of the dissolution behavior in the human body [24]. It has been widely reported that polymers isolated from fruit or vegetables residues can be used as excipients; however, there has been little research in which the residues are combined with an excipient in the same formulation. In this context, the aim of the present study was to evaluate the quality parameters (uniformity, breaking force, friability, disintegration) and the functional (lipid and water holding capacity) and antioxidant properties (total phenolic content, antioxidant activity by ABTS, DPPH, and ferric reducing/antioxidant power (FRAP)) of tablet formulations developed from cactus pear residues.
2. Materials and Methods

2.1. Sample Preparation

Two varieties of cactus pear fruits were selected, green and red *Opuntia ficus-indica* L. Mill, which were provided by the Mexican Association CoMeNTuna (Consejo Mexicano del Nopal y la Tuna) in Hidalgo, Mexico. The cactus pear fruits were manually harvested (latitude of 20° 16′ 12″ N, longitude 98° 56′ 42″ W, and altitude of 2600 m above sea level) during September 2015 and were selected in agreement with the Mexican legislation for nonindustrialized food products relating to the human use of cactus pear (*Opuntia* spp.) fruit, being of firm consistency, clean, free of foreign matter, and free of damage caused by pests or diseases in addition to presenting a state of commercial maturity, determined by the observation of the sinking of the fruit receptacle [25]. The fruits were divided into three batches, washed, and manually peeled; the residues (mesocarp and pericarp) were separated, then frozen to −32 °C and lyophilized (VWR26671-581 Labconco, Kansas City, MO, USA) for 96 h at −55 ± 1 °C under a vacuum of 0.040 mbar. Samples were milled (Blender, 38BL52, LBC10, Waring Commercial, Torrington, CT, USA) and sieved to a particle size of <500 µm.

**Tablet Formulation**

Formulations were produced as follows: Two commercial excipients, microcrystalline cellulose (Avicel® PH200 FMC Corp., Spain) and α-lactose-monohydrate (Tabletose® 80 Meggle Milchindustrie GMBH and Co. KG, Wasserburg, Germany), were used at 58%, mixed with 40% of cactus pear powder, and four formulations were thus developed: green cactus pear residue with microcrystalline cellulose (GM), green cactus pear residue with lactose (GL), red cactus pear residue with microcrystalline cellulose (RM), and red cactus pear residue with lactose (RL). Additionally, 1% of talcum powder and 1% of magnesium stearate (Panreac-Montplet and Esteban, Barcelona, Spain) were added as lubricant agents to improve the tablet quality. All components were mixed in a homogenizer (V-mill type MV-6; Turu Grau S.A, Tarrasa, Spain) for 15 min and compacted using an eccentric tableting machine (Bonnals B-40, Barcelona, Spain). These formulations were established according to preliminary tests where the excipients were added until complete compaction was reached.

2.2. Quality Parameters

2.2.1. Mass Uniformity, Thickness, and Diameter

The uniformity analysis was carried out by randomly selecting 10 units of each formulation and measuring their weight (g), thickness (mm), and diameter (mm) using a tablet tester (WHT-1 Pharma Test, Hainburg, Germany) [26].

2.2.2. Tablet Breaking Force

The tablet breaking force is defined as the amount of force required to break a tablet under a diametrical compression test. This was determined using a durometer (PT-B 311 Pharma Test, Hainburg, Germany), applying pressure until the tablets broke. The results are expressed in Newtons (N) [27].

2.2.3. Friability

Friability is the tendency of a tablet to lose particles due to abrasion, friction, or mechanical shock. This was determined according to the American Pharmacopoeia Commission (USP 1217) standard using a friability tester (ERWEKA, TA-UZ 127, Heusenstamm, Germany). Ten undusted and weighed tablets were subjected to 100 drum rotations; then, the tablets were removed from the drum, the dust...
was brushed off, and the tablets were weighed again [28]. Friability is expressed as the percentage of mass lost, calculated using Equation (1):

\[
\% \text{ Friability} = \left( \frac{\text{starting weight} - \text{weight after test}}{\text{starting weight}} \right) \times 100.
\]

(1)

2.2.4. Disintegration

This test is used to analyze the disintegration of tablets or capsules within the prescribed time when placed in a liquid medium. Six tablets were tested using a disintegrator (ERWEKA, DT-60, Heusenstamm, Germany) with a medium of 0.1 N hydrochloric acid at 37 ± 1 °C at 50 rpm. The test was finished when no residue was observed in the bottom of the disintegrator glass, and the disintegration time (min) was measured [29].

2.3. Scanning Electron Microscopy (SEM)

Micrographs were taken to establish structural comparisons between formulations. The morphology of the cactus pear formulations was evaluated by SEM. Samples were deposited on a silicon wafer and set into a sputter coater (Denton Vacuum Desk V, USA) for a thin-layer gold coating, applying 20 mTorr and 20 mA for a duration of 4 min. Samples were observed under a scanning electron microscope (JEOL JSM-6300, USA) at 250× magnification.

2.4. Functional Properties: Lipid and Water Holding Capacity

The lipid holding capacity (LHC) and water holding capacity (WHC) were measured by mixing 250 mg of the tablet powder with 10 mL of soybean oil or water in a centrifuge tube, shaking the mixture for a duration of 10 min, and leaving it at room temperature for 24 h prior to centrifugation (Hamilton Bell, VanGuard V6500 USA) at 3200 rpm for 10 min. The supernatants were decanted and the pellets were weighted. The amount of lipids or water absorbed by the powder is reported as grams of lipids or water retained per gram of sample on a dry basis (g/g db) [30].

2.5. Color Measurement

Color was measured by placing milled tablets on a circular white plate and using a colorimeter (Minolta CM-80 500 SM-508D, Japan), using the D65 illuminant with an angle of observation of 10°. Color was recorded using the CIE-L*a*b* values, where L* indicates lightness (L* values of 0 and 100 indicate black and white, respectively), a* is the axis of chromaticity between green (−) and red (+), and b* is the axis between blue (−) and yellow (+). Numerical values of L*, a*, and b* were used to obtain the Hue angle (h°) \( h^\circ = \tan^{-1} (b/a) \) and chroma (\( C = \sqrt{a^2 + b^2} \)) parameters [31].

2.6. Antioxidant Properties

2.6.1. Extraction of Antioxidants

To determine the antioxidant properties (total phenolic content (TPC), ABTS, DPPH, and FRAP) aqueous extraction was carried out following the methodology described by Serrano and Saura-Calixto [16]. Briefly, 250 mg of tablets was triturated and placed in tubes, 10 mL of methanol/water (50:50, v/v) was added, and then the tubes were shaken (LabTech LSI-3016A, Korea) at 60 rpm for 1 h at room temperature. After this, the tubes were centrifuged at 3200 rpm for 20 min (Hamilton Bell VanGuard 6500, Montvale, NJ, USA) and the supernatant was recovered to perform a second wash in the pellet, adding 10 mL of acetone/water (70:30, v/v). The shaking and centrifugation processes were repeated, and both the methanolic and acetonic supernatants were combined and graduated to 25 mL using a 1:1 methanol/water and acetone/water solution.
2.6.2. Total Phenolic Content (TPC)

The total phenolic content was determined following the Folin–Ciocalteu method according to Stintzing et al. [32]. An aliquot (100 µL) was mixed with 500 µL of diluted Folin–Ciocalteu reagent (1:10). Then, 400 µL (7.5%) of sodium carbonate was added, and the mixture was incubated for 30 min at room temperature. The absorbance of the mixture was measured at 765 nm (Power Wave XS UV-Biotek, USA). For the standard curve, gallic acid was used as the reference standard, with concentrations of 0, 100, 200, and 300 mg of gallic acid/L. The results are expressed as milligrams of gallic acid equivalent per 100 grams on a dry basis (mg GAE/100 g db).

2.6.3. Antioxidant Capacity by ABTS•+

The ABTS analysis was performed according to Kuskoski et al. [33], with some modifications. The radical cation 2,2 azino-bis (3-ethylbenzothiazoline-6-sulfonic acid) diammonium salt (ABTS•+) was produced by reacting 7 mM ABTS stock solution with 2.45 mM potassium persulfate under dark conditions at room temperature for a duration of 16 h before use. The ABTS•+ solution was diluted with deionized water until it reached an absorbance of 0.7 ± 0.1 at 754 nm. An aliquot of 100 µL of sample was added to 900 µL of ABTS solution, and the absorbance at 754 nm (Power Wave XS UV-Biotek, USA) was measured after 7 min of incubation at room temperature. The standard curve was linear with concentrations of 0, 60, 120, 180, 240, and 300 µmol of Trolox/L, and the antioxidant capacity is expressed as micromoles of Trolox equivalents per 100 grams on a dry weight basis (µmol TE/100 g db).

2.6.4. Antioxidant Activity by DPPH•

Antiradical activity was analyzed using the 1,1-diphenyl-2-picrylhydrazyl (DPPH•) radical as described by Aguirre et al. [34]. An ethanolic solution (7.4 mg/100 mL) of the stable DPPH radical was prepared. Then, 100 µL of extract was taken into vials and 500 µL of DPPH solution was added; the mixture was left to stand for 1 h at room temperature. Finally, the absorbance was measured at 520 nm using a microplate reader (Power Wave XS UV-Biotek, USA). A standard curve was constructed with concentrations of 0, 50, 100, 200, and 300 µmol of Trolox/L. The inhibition percentage of the DPPH• cation was calculated using Equation (2):

\[
\% \text{ Inhibition DPPH} = \left(\frac{Ac - As}{Ac}\right) \times 100
\]

where Ac is the absorbance of the control and As is the absorbance of the sample.

2.6.5. Ferric Reducing/Antioxidant Power (FRAP)

The FRAP assay was performed according to the method described by Zafra et al. [35]. The stock solutions included 300 mM acetate buffer (sodium acetate trihydrate C\(_2\)H\(_3\)NaO\(_2\)\(_3\)H\(_2\)O and glacial acetic acid C\(_2\)H\(_4\)O\(_2\)), pH 3.6, 10 mM; TPTZ (2,4,6-tripyridyl-s-triazine) solution in 40 mM HCl; and 20 mM FeCl\(_3\).6H\(_2\)O solution. The fresh solution prepared by mixing 25 mL acetate buffer, 2.5 mL TPTZ solution, and 2.5 mL FeCl\(_3\).6H\(_2\)O solution was warmed at 37 °C before use. The sample (30 µL) was mixed with 90 µL of distilled water and 900 µL of the FRAP solution for 10 min in the dark. A quantity of 5 M ferrous sulfate (FeSO\(_4\)) was used as the standard for the calibration curve (concentrations of 0, 200, 400, 600, 800, and 1000 µmol of FeSO\(_4\)). Absorbance was measured at 593 nm (Power Wave XS UV-Biotek, USA), and the results are expressed as micromoles of Fe(II) per 100 grams on a dry basis (µmol Fe(II)/100 g db).

2.7. Statistical Analysis

All determinations were performed in triplicate, and the results presented correspond to the average of the triplicate ± standard deviation. Statistical analyses were conducted using SPSS 23.
(SPSS® for Windows™ Chicago Illinois), and comparisons between samples were carried out by applying a multivariate analysis of variance (ANOVA). Differences between means were ranked by Tukey’s comparison test at a significance level of \( p < 0.05 \).

3. Results and Discussion

3.1. Quality Parameters

The tablet form represents two-thirds of the worldwide pharmaceutical products commercialized and is one of the most-consumed administration types of natural extracts and drugs [36]. In the pharmaceutical manufacturing process, quality evaluations (of uniformity, hardness, friability, and disintegration, among others) are performed to avoid variation from the pharmacopoeia limits [37]. The quality parameters of the cactus pear residue tablets are shown in Table 1.

### Table 1. Quality and functional parameters of cactus pear residue tablet formulations.

|                  | GM       | GL       | RM       | RL       |
|------------------|----------|----------|----------|----------|
| Weight (g)       | 0.7 ± 0  | 0.9 ± 0  | 0.8 ± 0  | 1 ± 0    |
| Thickness (mm)   | 4.6 ± 0  | 4.8 ± 0  | 4.6 ± 0  | 5.6 ± 0  |
| Diameter (mm)    | 14 ± 0   | 14 ± 0   | 14 ± 0   | 14 ± 0   |
| Breaking force (N)| 121 ± 8  | 162 ± 29 | 164 ± 29 | 96 ± 13  |
| Friability (%)   | 0.3 ± 0  | 0.1 ± 0  | 0.3 ± 0  | 0.1 ± 0  |
| Disintegration (min)| 2 ± 0   | 66 ± 2   | 19 ± 1   | 67.3 ± 4 |
| LHC (g/g)        | 5.3 ± 0  | 4.5 ± 0  | 4.5 ± 0  | 3.8 ± 0  |
| WHC (g/g)        | 5.5 ± 0  | 4.2 ± 0  | 4.6 ± 0  | 3.6 ± 0  |

Mean values ± standard deviation (\( n = 3 \)). *–d* Superscript letters indicate significant differences \( (p < 0.05) \) between tablet formulations. GM: green cactus pear residue with microcrystalline cellulose; GL: green cactus pear residue with lactose; RM: red cactus pear residue with microcrystalline cellulose; RL: red cactus pear residue with lactose. LHC: lipid holding capacity; WHC: water holding capacity.

The variations in these quality parameters depend directly on the pressing force and the punches selected in the tablet compression machine [38]. The average weight of the tablets ranged from 0.7 to 1 g, RL tablets were significantly heavier, and weight variations from the average were of 0.01% to 0.05%; all tablets were within the tolerance limits for mass variation (<5%) established by the United States Pharmacopoeia Commission [26]. The weight of the cactus pear residue tablets was higher than those reported for tablets made in other studies, such as those containing orange peel pectin and banana peel (0.4 and 0.2 g, respectively) [21,39]. The thickness values were between 4.6 and 5.6 mm, and the RL tablets were significantly thicker \( (p < 0.05) \) than the other formulations.

Tablet diameter did not differ among the formulations (14 mm) and could be considered a good size. According to the FDA, a size of >22 mm could propitiate a prolonged esophageal transit time, difficulty swallowing, or even injury to the esophagus [40]. In addition, it has been reported that the ideal size for tablets is between 7 and 8 mm [41].

Evaluation of the hardness parameter is relevant because it is directly related to the resistance against mechanical impacts due to handling, manufacturing, packaging, and transport, and it is also related with tablet disintegration [42]. The breaking force of the cactus pear residue tablets ranged from 96 to 164 N, with the values for GL and RM being higher. The cactus pear residue tablets presented values higher than those reported in tablets made with orange peel pectin (17 to 29 N) [22] and banana peel (19 to 22 N) [21].

The percentage of friability indicates the amount of loose particles on the tablet due to mechanical shock. In all formulations, the loss of mass was below 1%, and according to Porter et al., tablets with friability values of <1% are good, while high friability values lead to unacceptable loss of drug content during processing, storage, and handling [43]. The tablets with microcrystalline cellulose (GM and RM) had higher friability (both 0.3%) in comparison with the lactose tablets (0.1% for GL and RL). These differences could be attributed to the greater resistance to impacts and cohesion force...
of α-lactose agglomerates, while cellulose tablets are characterized by a lower compressibility and crushing strength since cellulose crystals are softer, weaker, and less elastic [24,44].

Disintegration is the first step in the bioavailability process, used to evaluate the behavior of the product through the gastrointestinal tract [45]. The formulations with microcrystalline cellulose had lower disintegration times (1.9 and 19.3 min for GM and RM, respectively), while lactose tablets presented greater times (66 and 67 min for GL and RL, respectively). These differences may be related to the low compressibility of microcrystalline cellulose and its rapid swelling in a humid environment. However, if there is erosion on the tablets, it may explain why tablets of similar mass show little difference in the dissolution profiles [46]. In comparison with other studies, slower disintegration for tablets with lactose derivatives is reported [44]. Other studies on tablets made with pectin from citrus peel exhibited disintegration times of between 12 and 15 min [39], and the times for banana peel tablets were 8–10 min [21]. These differences can be explained by the geometry playing an important role—tablets with larger surface area have faster drug release and disintegration [46]—and the variation in particle size [45]. In this context, the GL tablet presented greater resistance to breakage, less friability, and a longer disintegration time compared to the other excipients. These differences can be explained by the influence of the different excipients analyzed and their physical and chemical properties, given that several researchers have reported that the excipients are a critical factor in the development of new pharmaceutical formulations with greater pharmacological functionality (disintegration, dissolution, release, and bioavailability of the drug) [47,48].

3.2. Scanning Electron Microscopy (SEM)

SEM images of the cactus pear powder and formulations are shown in Figure 1. The structure of the green and red cactus pear powders (Figure 1A,D) without excipients was continuous but with irregular borders. Images of the GM and RM tablets (Figure 1B,E) also showed the structure of the added excipients. The typical structure of microcrystalline cellulose (labeled “a” in the figure) was observed as elongated filamentary structures with angular borders [49], while the lactose structures (labeled “b”) were rather small, spherical and without angular edges. The structure of the matrix by itself, in combination with the structure of an excipient, influences the behavior of the sample; for example, the filamentous structures of the microcrystalline cellulose give greater force against abrasion, while the spherical lactose structures provide greater cohesion between the particles, allowing longer disintegration times [50–52].

3.3. Functional Properties: Lipid and Water Holding Capacity

In previous studies performed by Bensadón et al., on the same raw material of cactus pear wastes used in this work, the amount of dietary fiber was reported as follows: for the green residues, 34.9 g/100 g insoluble fiber and 7.9 g/100 g soluble fiber; for the red residues, 19.3 g/100 g insoluble fiber and 8.1 g/100 g soluble fiber [53]. Thus, the fiber content of each tablet is 0.24 g with green residues and 0.15 g with red residues, approximately. According to the recommended daily intake of fiber for adults (25–30 g/day) [54], in the present study, the daily portion of consumption would be six tablets per day; therefore, the tablets would cover 5.7% (green tablets) and 2.6% (red tablets). Although the content of fiber is low, the impact in terms of the physiological benefits and in the food industry could be important. According to Cardenas et al., the fiber in the cactus pear fruit is characterized by a high insoluble dietary fiber content and the presence of mucilages with high molecular weight which interact in an entangled network, able to hydrate and increase the viscosity of the liquid phase in the food system [55]. Therefore, the physicochemical properties of these samples could be affected in the presence of water.
consumer choices and preferences [31]. The significance in certain disorders of colonic function, obesity, and lipid blood concentrations [59].

Leads to the hydration of the intestinal content for soft and easy-to-pass stools, and LHC has clinical significance in certain disorders of colonic function, obesity, and lipid blood concentrations [59]. Other studies on fibers from fruit by-products have reported good water and lipid retention. For example, a fiber supplement from lemon peel presented 5 g/g and 7 g/g WRC and LRC, respectively, while pineapple peel presented 11 g/g and 6 g/g WRC and LRC, respectively [20,58]. However, these properties have not been reported in other dietary supplements in solid dosage forms, though they are physiologically important, since WHC leads to the hydration of the intestinal content for soft and easy-to-pass stools, and LHC has clinical significance in certain disorders of colonic function, obesity, and lipid blood concentrations [59].

3.4. Color

The four formulations presented a heterogeneous aspect and characteristic color corresponding to the cactus pear residue variety used (Figure 2). Color is an important quality attribute that influences consumer choices and preferences [31].

For the LHC and WHC parameters presented, the GM formulation had the highest values, in contrast with the RL formulation, which had the lowest values for both parameters (Table 1). The lipid and water holding capacity may be attributed to the high amount of insoluble fiber in the cactus pear residue, which is capable of trapping oil and water particles in its structure [56,57]. The structure of the microcrystalline cellulose fiber favors retention [44], and this could explain the faster disintegration of these tablets compared to the lactose formulations (Table 1). Other studies on fibers from fruit by-products have reported good water and lipid retention. For example, a fiber supplement from lemon peel presented 5 g/g and 7 g/g WRC and LRC, respectively, while pineapple peel presented 11 g/g and 6 g/g WRC and LRC, respectively [20,58]. However, these properties have not been reported in other dietary supplements in solid dosage forms, though they are physiologically important, since WHC leads to the hydration of the intestinal content for soft and easy-to-pass stools, and LHC has clinical significance in certain disorders of colonic function, obesity, and lipid blood concentrations [59].

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For the LHC and WHC parameters presented, the GM formulation had the highest values, in contrast with the RL formulation, which had the lowest values for both parameters (Table 1). The lipid and water holding capacity may be attributed to the high amount of insoluble fiber in the cactus pear residue, which is capable of trapping oil and water particles in its structure [56,57]. The structure of the microcrystalline cellulose fiber favors retention [44], and this could explain the faster disintegration of these tablets compared to the lactose formulations (Table 1). Other studies on fibers from fruit by-products have reported good water and lipid retention. For example, a fiber supplement from lemon peel presented 5 g/g and 7 g/g WRC and LRC, respectively, while pineapple peel presented 11 g/g and 6 g/g WRC and LRC, respectively [20,58]. However, these properties have not been reported in other dietary supplements in solid dosage forms, though they are physiologically important, since WHC leads to the hydration of the intestinal content for soft and easy-to-pass stools, and LHC has clinical significance in certain disorders of colonic function, obesity, and lipid blood concentrations [59].

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The color parameters are shown in Table 2. Green tablets had the highest values ($p < 0.05$) for luminosity ($L^*$), having a brighter color than red tablets. For the green tablets, the $a^*$ values were negative and $b^*$ values were positive, which places them in the green-yellow quadrant, while the red tablets presented positive values for $a^*$ and $b^*$, corresponding to the red-yellow quadrant.

Table 2. Color parameters of the cactus pear residue tablet formulations.

|       | GM    | GL    | RM    | RL    |
|-------|-------|-------|-------|-------|
| $L^*$ | 49 ± 0 | 48 ± 1 | 40 ± 1 | 41 ± 0 |
| $a^*$ | −2 ± 0 | −2 ± 0 | 7 ± 0  | 9 ± 0  |
| $b^*$ | 16 ± 0 | 15 ± 0 | 4 ± 0  | 4 ± 0  |
| $h^*$ | 97 ± 0 | 98 ± 0 | 31 ± 1 | 25 ± 0 |
| $C^*$ | 16 ± 0 | 15 ± 0 | 9 ± 0  | 11 ± 0 |

Mean values ± standard deviations ($n = 3$). a–d Superscript letters indicate significant differences ($p < 0.05$) between tablet formulations. GM: green cactus pear residue with microcrystalline cellulose; GL: green cactus pear residue with lactose; RM: red cactus pear residue with microcrystalline cellulose; RL: red cactus pear residue with lactose. $L^*$ values of 0 and 100 indicate black and white, respectively, $a^*$ is the axis of chromaticity between green (−) and red (+), $b^*$ is the axis between blue (−) and yellow (+), $h^*$ indicate Hue angle and $C^*$ indicate chroma parameters.

The inclusion of binding agents within the same variety of cactus pear had no significant effect on $a^*$ and $b^*$ values, so the characteristic color of the fruits was preserved by the peel color itself. Both GM and GL tablets exhibited negative hue ($h^*$) values; according to McLellan, the addition of 180° to a negative value allows us to achieve a positive one [60], and with this adjustment it was possible to determine the yellowish tonalities for the GM tablets and reddish or pink ones for the RM and RL samples. The chroma ($C^*$) value estimates the quantitative attribute of colorfulness, with higher chroma values indicating higher color intensity of samples as perceived by humans [31]. The green cactus pear formulations (GM and GL) presented higher $C^*$ values, followed by RL and RM, so red formulations had less intense tonalities. All samples preserved the color characteristic of the variety used, and attractive colors were observed (Figure 2); therefore, it is not necessary to add artificial colors to make them attractive to the consumer. Besides, cactus pear pigments have been widely used by the food industry as an economical and stable source of pigments [61].

3.5. Antioxidant Properties

The antioxidant properties of the formulations are shown in Table 3. Despite reports that red cactus pear has higher total phenolic content (TPC) in the pulp than in the peel and seeds [62], the results in terms of phenolic compounds were higher ($p < 0.05$) in GL, followed by GM. The TPC in the red cactus pear formulations (RM and RL) did not differ significantly; however, Cejudo et al. reported that yellow varieties of whole cactus pear fruit have higher contents of phenolic compounds in the pericarp and epicarp than in the pulp [63]. Regarding antioxidant activity by DPPH, the radical inhibition of the red formulations was higher (24%) when compared to the green tablets (20%). Differences in antioxidant activity between green and red cactus pear residues were also found by Ramirez-Moreno et al., where seeds from green cactus pear showed greater antioxidant activity than the red variety [64]. These differences in antioxidant capacity could be linked to the antioxidants of lipophilic nature associated with different cactus pear varieties, for instance, lipophilic compounds such as tocopherols may increase the antioxidant activity measured by DPPH [33]. The antioxidant activity by ABTS was similar ($p > 0.05$) among all the tablets. Finally, the RM formulation exhibited higher FRAP values ($p < 0.05$). Although studies on the antioxidant properties of tablets with added fruit wastes are limited, a study on effervescent tablets made with tangerine peel reported a 141 mg GAE/100 g total phenolic content and 27% DPPH inhibition percentage [65], which are lower and slightly higher, respectively, than the cactus pear residue tablets. These results are interesting and are in accord with the study by Bensandon et al.; cactus pear by-products have the potential to be used in the development of food products because their high dietary fiber and antioxidant content can prevent the damage caused by
free radicals [53]. The measures of antioxidant status help to determine why a diet high in antioxidants is often more beneficial, since dietary antioxidants have the ability to inhibit the formation of both reactive oxygen species (ROS) and reactive nitrogen species (RNS), which can adversely affect normal cellular processes and physiological functions. It is well known that excessive oxidative stress could affect proteins, lipids, or genetic material, leading to decreased immune competence and increased inflammatory response, all linked to several diseases such as cancer, cardiovascular disease, obesity and diabetes [66].

Table 3. The antioxidant properties of the cactus pear residue tablet formulations.

|                | GM          | GL          | RM          | RL          |
|----------------|-------------|-------------|-------------|-------------|
| TPC (mg GAE/100 g) | 718 ± 11 b  | 734 ± 8 a   | 649 ± 12 c  | 651 ± 8 c   |
| DPPH (% Inhibition) | 20 ± 1 b    | 20 ± 1 b    | 24 ± 2 a    | 24 ± 1 a    |
| ABTS (µmol TE/100 g) | 4628 ± 412 a | 4637 ± 242 a | 4365 ± 392 a | 4610 ± 326 a |
| FRAP (µmol Fe²⁺/100 g) | 17 ± 0 b   | 16 ± 1 c    | 19 ± 0 a    | 17 ± 0 b    |

Mean values ± standard deviation (n = 3). * Superscript letters indicate significant differences (p < 0.05) between tablet formulations. GM: green cactus pear residue with microcrystalline cellulose; GL: green cactus pear residue with lactose; RM: red cactus pear residue with microcrystalline cellulose; RL: red cactus pear residue with lactose.

Gallic acid (for total phenolic content (TPC)), Trolox (for DPPH and ABTS), and FeSO₄ (for ferric reducing/antioxidant power (FRAP)) were used as reference standards.

On the other hand, studies have been carried out regarding the safety of the consumption of cactus pear fruit; for example, the presence of pesticide residues (malathion, chlorpyrifos, permethrin, diazinon, dimethoate, spinosad, and abamectin) and heavy metals (copper, chromium, arsenic, cadmium, lead, and selenium) has been demonstrated in levels less than the maximum limits of toxic residues established by the North American Free Trade Agreement [67,68], indicating that these phytotoxic elements do not trigger toxicological health risks.

4. Conclusions

Based on the results obtained, cactus pear residue is proposed as an interesting nutritional supplement for the market due to its good quality parameters, fiber functional properties, acceptable color for consumers, and high antioxidant activity, providing probable benefits to consumers. The GL formulation presented better performance as a possible dietary supplement.

Further research on phytotoxicity is needed since there are no existing reports about that, as well as, in vivo model studies in order to evaluate the health effects of a dietary supplement developed from cactus pear by-products.

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Conflicts of Interest: The authors declare no conflict of interest.

Strengths and Limitations: The strengths of the present study consist of the use of an agricultural by-product, through its use in a novel product, which may have beneficial effects on human health and which also represents an economic benefit for the producers of the fruit. On the other hand, among the limitations, the cactus pear is a seasonal fruit, so the residues must be collected and stored during the harvest season. It is also necessary to subject the product to further in vitro tests in order to evaluate any positive or negative effects on human health.
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