Yacon (Smallanthus Sonchifolius)-Based Product Increases Fecal Short-Chain Fatty Acids Concentration and Up-Regulates T-Bet Expression in the Colon of BALB/c Mice During Colorectal Carcinogenesis

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Abstract

Colorectal Cancer (CRC) is the third most diagnosed type of cancer worldwide. Prebiotics containing Fructooligosaccharides (FOS) and inulin have been reported to improve CRC in experimental models. We hypothesized whether consumption of the yacon-based product (PBY-Smallanthus sonchifolius concentrate), as a source of FOS and inulin, could mitigate colonic pre-neoplastic lesion development in mice by enhancing fecal Short-Chain Fatty Acid (SCFA) production besides modulating intestinal immune responses. Therefore, we investigate the effects of PBY consumption on anatomical and fecal characteristics, serum biomarkers, fecal SCFA concentration, intestinal lymphocytes population, expression of transcription factors of the adaptive immune response and Aberrant Crypt Foci (ACF) count in the colon of mice chemically induced to pre-neoplastic lesions. Male BALB/c mice were injected intraperitoneally with 1,2-dimethylhydrazine (20mg/kg body weight/week) for 8 weeks. Thereafter, mice were fed either control (AIN-93M) or PBY diet (AIN-93M supplemented with PBY, 6.0% FOS +Inulin) for 8 weeks and then euthanized. PBY was not successful in reducing ACF counts; however, it improved fecal SCFA concentration (formic, acetic, propionic, butyric and valeric), reduced fecal pH and increased humidity and viscosity of feces. Although there were no significant changes in the intestinal lymphocytes population (CD4, CD8, Natural Killer, Treg and Th17), PBY consumption modulates the immune response in the colon reducing the expression of FOXP3 and increasing RORγt and T-bet, which would contribute to activation and proliferation of CD8 T lymphocytes and better CRC prognosis. Therefore, we suggest that PBY might improve intestinal health during the early stages of colorectal carcinogenesis, especially by modulating SCFA production and colonic adaptive immune response.

Keywords: Colorectal cancer; Immunomodulation; Prebiotic; Short-chain fatty acid; Transcription factor; Yacon

Introduction

Colorectal Cancer (CRC) is the third most diagnosed type of cancer in the world [1]. CRC etiology cross talks with genetic factors, eating habits, immune response and intestinal microbiota composition [2-4]. The tumor-infiltrating lymphocyte profile strongly associates with CRC prognosis [5,6]. CD8 T lymphocytes and Natural Killer (NK) cells, for instance, have been related to disease delay and better prognosis [7-9]. Furthermore, the Th1 immune response may contribute to remission and better CRC prognosis [10,11]. On the other hand, Th17 cells have been associated with worse CRC prognosis and disease progression [12], although it has already been associated with a better prognosis [9]. Regulatory T cells (Treg) also evidence a dual role [13,14], sometimes suppressing the anti-tumor immune response [15], sometimes attenuating the inflammatory immune response [16].

Evidence suggests that prebiotic food consumption induces the growth of probiotic bacteria [17], which improves the anti-inflammatory activity [18] and favors the modulation of cells in the gut-associated lymphoid tissue [19,20]. Yacon is a known source of Fructooligosaccharides (FOS) and inulin and therefore, considered a prebiotic food. This tuberous root is originated from the Andes, but it can easily adapt to different climatic regions, altitudes and soils [21]. For this reason, yacon has been cultivated in several countries such as United States, Japan, Italy, Brazil, Argentina, Bolivia, Czech Republic,
Yacon consumption might modulate the intestinal microbiota by increasing the number of bifidobacteria and lactobacilli [28,29], which directly inhibit the growth of pathogenic bacteria [30,31]. Studies have shown that yacon increase fecal Short-Chain Fatty Acids (SCFA) concentration [20,26,29,32,33], improve intestinal transit [28,34], control satiety [20], contribute to weight loss [27] and reduce glycemia [24], triglycerides and cholesterol [35,36]. Yacon also modulates the intestinal inflammatory response by increasing fecal secretory immunoglobulin A (sIgA) and serum interleukin (IL)-10 and IL-4 [33,37,38], besides preventing pre-neoplastic lesions during colorectal carcinogenesis in animals [26,39]. Recently, our research group verified the role of the yacon-based product (PBY- *Smallanthus sonchifolius*) concentrate on intestinal immunity. PBY consumption in the colon of BALB/c mice and mitigated the inflammatory profile by down regulating the expression of RORγt [20].

Yacon exhibits antioxidant properties owing to its high polyphenol content, such as chlorogenic acid, caffeic acid, coumaric acid and protocatechuic acid [25,40]. Its content of fructans, mainly FOS and inulin, favor SCFA production by the intestinal microbiota, especially butyrate, which has been reported to be closely linked to neoplastic cells apoptosis [41], Histone Deacetylase (HDAC) inhibition [42,43] and cell cycle phase, thus impairing colorectal carcinogenesis and tumor progression [44].

The benefits of yacon are directly related to the modulation of intestinal microbiota [29] and intestinal immune response [20,38]. Nevertheless, the immune response profile induced by yacon during colorectal carcinogenesis has still not been elucidated. In this study, we use a mouse model that allows *in situ* evaluation of pre-neoplastic lesions, known as aberrant crypt foci (ACF) [26,39], as well as we assess biomarkers similar to used in human studies. Therefore, this model supports the evaluating of the effects of prebiotics on colorectal carcinogenesis [45]. We used yacon concentrate (PBY) that provides higher concentrations of FOS and inulin, besides, be lesser perishable than fresh yacon. As recommended by Paula and co-workers [46], PBY was added to the animals’ diet to supply 6% FOS + inulin and hence modulate intestinal function. Thus, we hypothesized whether PBY consumption, as a source of FOS and inulin, could mitigate ACF development in mice by enhancing fecal SCFA concentration besides modulating intestinal immune responses. Moreover, the effects of PBY consumption were explored on anatomical and fecal characteristics, serum biomarkers, intestinal lymphocyte populations, and expression of transcription factors of the adaptive immune response in the colon of BALB/c mice chemically induced to pre-neoplastic lesions.

### Methods and Materials

#### Animals and experimental design

Thirty-six nine-week-old male BALB/c mice were obtained from the Central Biotechnology (Health and Biology Science Center) of the Universidade Federal de Viçosa, Brazil. The animals were housed in the Experimental Nutrition lab in a temperature-controlled room (22±2°C) with a 12-hours light/dark cycle and *ad libitum* access to water and food. The animal protocol was approved by the Ethics Committee on Animal Experimentation from the Universidade Federal de Viçosa, Brazil, under the process number 30/2016 and performed according to the Guide for the Care and Use of Laboratory Animals, National Academy of Sciences (US). Upon arrival, mice were induced to pre-neoplastic colorectal lesions by intraperitoneal injection of 1,2-Dimethylhydrazine (DMH) (Sigma, Saint Louis, USA) (20mg/kg body weight) once a week during 8 weeks [45]. DMH was dissolved in 0.9% saline solution containing 1mM EDTA and 10mM sodium citrate, at pH 8 [47]. Then, the animals were randomly assigned to two experimental groups: The group receiving control diet (DC; n=17) or the group receiving PBY-supplemented diet (DY; n=19) (Table 1). The sample size was calculated according to the aberrant crypt data reported by de Moura and co-workers [34], to achieve a statistical power of 90%. Diets were offered *ad libitum* for 8 weeks. Subsequently, mice were anesthetized using 3% isoflurane and blood was collected from the retro-orbital sinus. Mice were euthanized by cervical dislocation and tissue and feces were harvested for analysis.

#### Table 1: AIN-93M composition in the control and PBY diets.

**Note:** †Centesimal composition and digestible content of carbohydrate, inulin, and FOS on PBY in 100g of product: Fructose: 15.25g; Glucose: 8.59g; Sucrose: 6.35g; FOS: 12.81g; Inulin: 4.16g; Total carbohydrate: 45.49g; Fibers: 1.99g; Humidity: 36.82g; Ashes: 3.39g; Lipids: 0.21g; Protein: 2.89g; AIN: American Institute of Nutrition; FOS: Fructooligosaccharides; PBY: yacon-based product.

#### Yacon-Based Product (PBY) and experimental diet

Yacon was purchased from a local market in Viçosa-Minas Gerais, Brazil. The PBY was processed according to the methodology proposed by Rodrigues and co-workers [23], which is currently going...
through patent request PI 1106621-0. The chemical composition of PBY (carbohydrates, proteins, fats, fiber, ash, and humidity) was determined according to the AOAC methodology (AOAC, 1997). FOS and inulin contents were determined by High-Performance Liquid Chromatography (HPLC) with a BIO-RAD brand HPX-87p column (lead stationary phase) using purified water for the mobile phase.

The experimental purified diets were based on the AIN93-M diet, as recommended by the American Institute of Nutrition [49]. The PBY-supplemented diet contained 6.0% FOS + Inulin from PBY (Table 1), as suggested by Paula and co-workers [46]. Casein, sucrose, dehydrinized starch, starch, and fiber adjustments were made to control and PBY diets to obtain similar amounts of carbohydrates, lipids, proteins, fibers, and calories. The diets were made in a pellet format and stored at -20°C for a maximum period of thirty days before consumption.

The calculation of the human equivalent amount of dietary PBY consumed by male BALB/c mice was performed according to the body surface area normalization method as previously describe [50]. PBY supplementation was 353.56g PBY/kg of diet. In our study, the average daily consumption was 6g per mouse. This is equivalent to 2,121g of PBY (corresponding to 272mg of FOS and 88mg of inulin) daily for an adult mouse of 45g, which approximately corresponds to 47g PBY/kg body mass/day. Taking into account that the average human adult weight is 70kg, it is thus converted to 229g per day for humans. It is worth noting that the equivalent dose calculation according to Reagan-Shaw and co-workers [50] was only used as a comparative method, with no intention to extrapolate the results for human intake, since the dose-response to nutrient intake may vary according to the experimental model used or to the individual participant in a study.

**Body weight and dietary intake**

In order to evaluate the weight loss/gain of the animals, individual body weight was weekly recorded using a digital weighing scale. Dietary intake was determined according to the difference between the diet offered and wasted. Quantification was done every 3 to 5 days during the whole dietary intervention on a digital weighing scale. Data represent the diet consumption per cage (7 to 10 animals/cage). Diet consumption (g) was corrected by humidity (fresh diet weight/g/diet weight (g) in the cage per day).

**Fecal characteristics**

Fresh feces were harvested and stored at -80°C for pH determination. For each animal, an aliquot of feces was diluted in distilled water (1:10), homogenized, and the pH was measured on a digital pH meter (Hx 7.8mm, Bio-rad®, Rio de Janeiro, Brazil) at 32°C with acidified water (0.005 M H2SO4) as eluent at a flow rate of 0.6mL/minute.

**Fecal SCFA quantification**

SCFA quantification was assessed according to the method used by Smiricky-Tjardes and co-workers [54], with some modifications. Fifty mg of frozen feces were weighed and thoroughly vortexed with deionized water (950μL). While incubated on ice for 30 minutes, the samples were homogenized for 2 minutes every 5 minutes. Samples were centrifuged (10,000 x g, 30 minutes, 4°C) three times and the supernatants were collected. The final supernatant from each sample was filtered through a 0.45μm membrane and transferred to vials. SCFA were measured by High-Performance Liquid Chromatography - HPLC (Shimadzu®) on an Aminex HPX 87H column (300 x 7,8mm, Bio-rad®, Rio de Janeiro, Brazil) at 32°C with acidified water (0.005 M H2SO4) as eluent at a flow rate of 0.6mL/minute. The products were detected and quantified by an ultraviolet detector (model SPD-20A VP) at 210nm. Standard curves for formic, acetic, propionic, isobutyric, butyric, isovaleric, valeric, isocaproic, and caproic acids (SUPELCO®) were performed. Results are expressed as μmol SCFA/g feces.

**Aberrant Crypt Foci (ACF) counts**

After removal, the large intestine was washed in PBS solution, opened along the mesenteric margin, placed in paraffin plates with the mucous facing the top of the plate, and fixed in Carson’s formalin [55], for 24 hours. Following fixation, flat colons were equally divided into three segments (proximal, medium, and distal) and stained with 0.1% methylene blue for 2 minutes to quantify ACF under a BX-60 light microscope (Olympus, Tokyo, Japan) with a magnification of 200X. ACF were counted across the mucosal surface of the large intestine by three independent observers. The ACF categorization was based on the number of aberrant crypts per focus: foci with fewer than 3.

**Anatomical characteristics**

After euthanasia, the organs (liver, spleen, small intestine, cecum, colon, and abdominal adipose tissue) were excised, washed in Phosphate-Buffered Saline (PBS) (NaCl: 0.85%, Na2HPO4: 0.023%, NaHPO4,2H2O: 0.15%, pH 7.2) and weighed using a semi-analytical weighing scale. The hepatosomatic index was obtained by dividing the liver weight by the animal’s body weight. The colon length was obtained by measuring from the end portion of the cecum till the end portion of the rectum on a flat surface using a millimeter ruler.
or equal to three crypts (ACF ≤3) and foci with more than three crypts (ACF >3) [56].

Histopathological score

Paraffin-embedded colonic tissues were sectioned at 5-μm thickness and subjected to Hematoxylin-Eosin (H&E) staining. Each colonic section was scored as previously described by Kang and co-workers [57]. The scoring system was based on three independent parameters: severity of inflammation (0-3), depth of injury (0-3) and crypt damage (0-4). The sum of such scores provides a total histopathological score with 0 being a normal tissue and 10 being the most extensive/severe disease symptoms. The assessment of histopathological score was performed using a BX-60® light microscope (Olympus, Tokyo, Japan) at 200x magnification. Three sections per animal that covered the length of 500μm of the proximal colon were stained and evaluated in a blinded manner.

Determination of leukocytes by immunophenotyping

Leukocytes were quantified and characterized in the colon mucosa as previously described [58], with some modifications. The colon was removed and washed in ice-cold PBS, cut into small fragments and incubated in a cell culture medium, DMEM, pH 7.2 (Sigma-Aldrich ™) for 90 minutes at 37°C. The suspension was centrifuged three times at 42 g for 5 minutes to collect the supernatant, and lastly at 543 g for 10 minutes. After the last centrifugation, the remaining pellet was then resuspended with PBS buffer (100μL, pH 7.2). Cell viability was assessed with Trypan blue exclusion and cells were counted in a Neubauer chamber. The obtained leukocytes were incubated with the following antibodies, according to the manufacturer’s instructions: Anti-CD4 (PeCy5), anti-CD25 FITC-conjugated, anti-CD196 (anti-CCR6) PE-conjugated, anti-CD49b (anti-PanNK) APC-conjugated, anti-CD8 PECD7-conjugated (Biolegend, San Diego, CA, USA). Leukocytes (1x10⁶ events) were acquired (FACS Verse™ and BD FAC Suite software; BD Biosciences Phar Mingen San Jose, CA, USA) according to size (forward scatter) and granularity (side scatter). One or two stains were used to identify TCD4 lymphocytes (CD4⁺), TCD8 lymphocytes (CD8⁺), regulatory T cells (CD4⁺CD25⁺), Th17 lymphocytes (CD4⁺CD169⁺) and Natural Killer cells (CD49b⁺). Results are expressed as means ± Standard Error of the Mean (SEM) of the percentage of each cell subpopulation specifically stained within a selected gated.

Real-time PCR

Total RNA was extracted from the whole colon using Trizol reagent (Invitrogen™, Carlsbad, CA, USA) according to the manufacturer’s instructions. cDNA was synthesized using 5μg of RNA through a reverse transcriptase reaction using a specific kit (GoScript™ Reverse Transcription System, Promega, Madison, WI, USA). Random primer and RNase free water were added to the sample and heated for 5 minutes at 70°C. Subsequently, a mixture containing the reverse transcriptase, dNTPs and ribonuclease inhibitor was added to the sample and heated for 60 minutes at 37°C. Quantitative mRNA analysis was performed by Real-time PCR on the Applied Biosystems® 7500 Real-Time PCR System. Sybr green (Ludwig, Biotech). Primer sense, primer antisense (400nm/reação), D Nas free water and the sample (250ng cDNA/μl) were added at each reaction. Standard PCR conditions were used during the reading (7500 software V2.3, Applied Biosystems®). The sequences of murine primers (Integrated DNA Technologies®) were as follows: FOXP3, sense: 5’-AGG AGC CGC AAG CTA AAA GC- 3’, antisense: 5’-TGC CTG CGT GCC CAC TGT-3’; RORyt, sense: 5’-GGA CGT CTG CCA GAA TGA CC-3’, antisense: 5’-CAG GGT TCG AAA CAG CAC CTC-3’; Tbet, sense: 5’-AGC AAC AGC GGC GAA TGT T-3’, antisense: 5’-GGG TGG ACA TAT AAG CGG TTC TCC TGC-3’, GAPDH, sense: 5’-TCA ACA GCT CCC ACT CCT CTA CCA-3’, antisense: 5’-ACC CTG TTG CTG ATC TAG CGG GC-3’. mRNA values were calculated according to the constitutive GAPDH gene on the basis of the ΔΔCt algorithm.

Statistical analysis

Results are expressed as means ± SEM. Data were analyzed using Graph Pad Prism version 6.0. Mean values were firstly tested by Kolmogorov-Smirnov test and the groups with normal distribution were tested using unpaired Student’s t-test. Groups that did not show normal distribution were tested using Mann-Whitney test. Statistical differences were considered when the p<0.05 (*) or p<0.001 (**).

Results

PBY-supplemented diet does not alter diet intake or body weight

There was no difference in the weekly weight gain of the animals either during pre-neoplastic lesion induction (1st to 8th week) or dietary intervention (9th to 16th week) (Figure 1A). In addition, there was no change in the average dietary intake between the groups during dietary treatment (Figure 1B).

Figure 1: Weight gain and food intake in BALB/c mice induced to colonic pre-neoplastic lesions and submitted to 8 weeks of dietary treatment. (A) Average weight gain of animals over the weeks. The data are expressed as means ± SEM (n=17 (DC); n=19 (DY)). (B) Mean food intake (g/mice/day) of PBY diet or control diet consumed during the period of dietary intervention (8 weeks). The data are expressed as means ± SEM of each experimental group (means referring to the group/cage consumption pool (4 pools/week). Statistical differences between groups were analyzed by the unpaired Student’s t-test or Mann Whitney, (*) p<0.05, (**) p<0.001. DC: Dimethylhydrazine-induced mice treated with control diet; DY: Dimethylhydrazine-induced mice treated with PBY diet; PBY: yacon-based product.

PBY-supplemented diet reduces fecal pH and increases fecal humidity

The physical-chemical evaluation of feces contributes to determining the direct and indirect effects of prebiotics on the intestine.
In this study, fecal pH was lower in the animals treated with PBY-supplemented diet (p<0.05) (Figure 2A). Fecal humidity was higher in the group receiving the PBY-supplemented diet when compared to control (p<0.001) (Figure 2B). Qualitative analysis through the fecal score shown 70% of PBY-treated animals in score 2 (viscous non-diarrheic stools) and 30% in score 3 (aqueous feces characteristic of diarrhea). Animals fed control diet were 100% in score 2.

**PBY-supplemented diet increases bowel and adipose tissue weight**

PBY-supplemented diet increased colon (p<0.05), cecum (p<0.001), small intestine (p<0.05) and abdominal adipose tissue weight (p<0.05) when compared to controls (Figure 3B-E). There was no significant change in colon length, liver and spleen weight or hepatosomatic index between the groups (Figure 3 A-H).

**PBY-supplemented diet increases fecal SCFA concentrations**

The intestinal bacterial activity was assessed through fecal SCFA concentrations. PBY-treated animals showed increased formic (p<0.001), acetic (p<0.05), propionic (p<0.05), butyric (p<0.001), and valeric (p<0.05) acid concentrations when compared to controls (Figure 5A-C, E and G). There was no difference in isobutyric, isovaleric, isocaproic, and caproic acid concentrations between the groups (Figure 5).

**PBY-supplemented diet does not change ACF counts**

ACF was assessed to confirm pre-neoplastic lesions formation in DMH-induced mice and to detect the effectiveness of PBY diet in reducing its formation. In both groups, a greater number of ACF categorized as ≤3 were observed concerning ACF >3 (Figure 6). In ACF ≤3 categorization, ACF were more abundant in the proximal colon, followed by medial and distal colon (Figure 6). There was no significant difference in total ACF count or categorized (ACF ≤3 and ACF >3) per colon segment between the groups (Figure 6).

**PBY-supplemented diet does not alter the colonic histopathological score**

In both groups, DMH was able to promote colonic histopathological changes at crypt level (Figure 7B), inflammation (Figure 7C),...
depth of damage (Figure 7D), and histopathological score (Figure 7A). Both groups showed crypt damages. Damage was considered when the crypt light does not reach the intestinal lumen, besides the oval aspect. The control group mostly presented crypts with 2/3 of damage (Figure 7E). PBY-treated animals showed crypts with 1/3 of damage. Inflammatory infiltrates were found in both groups and were minimally expressive (Figure 7E). Damage was mostly restricted to the mucosa.

PBY-supplemented diet does not alter the percentage of colonic lymphocytes

Immunophenotyping of colonic immune cells was performed to evaluate the intestinal lymphocyte population. There were no significant changes in the percentage of CD4+ and CD8+ T cells, Treg cells (CD4+CD25+), Th17 cells (CD4+CCR6+), NK cells, and CD4/CD8 ratio between the groups (Figure 8). Although not significant, there was a reduction on the percentage of CD4+ T lymphocytes (30.4%), and increments in CD8+ T cells (37.7%), CD8+/CD4+ ratio (156.9%) and Treg cells (84.8%) in the colon of PBY-treated animals when compared to controls.

PBY-supplemented diet increases the expression of RORγt and T-bet in the colon

Transcription factors of the adaptive immune response were evaluated to verify the pattern of the intestinal immune response in the colon. PBY-treated animals showed down-regulation of FOXP3 (p<0.05) and up-regulation of RORγt (p<0.05) and T-bet (p<0.05) when compared to controls (Figure 9). GATA-3 expression, although up-regulated in the PBY-treated groups (p <0.05), should not be considered biologically relevant due to the very low expression in both groups (DC: 0.023; DY: 0.046).

Discussion

Yacon has been associated with the reduction of pre-neoplastic lesions in experimental CRC models [26,39], especially by its high content of FOS and inulin. The benefits of yacon are directly related to the growth of probiotic bacteria and their metabolite production, such as SCFA [59].
Despite the promising role of yacon on the immune system modulation, as indicated by increased IgA production [33,37,38], reduction of pathogenic bacteria [28], increased fecal SCFA, enhanced Treg cells, and down-regulation of RORγt in healthy BALB/c mice [20], it is still unknown the adaptive immune response induced by yacon consumption during colorectal carcinogenesis. Thus, this study aimed at investigating the effects of PBY consumption on ACF improvement in DMH-induced mice as well as to assessing the immunological mechanisms behind this process.

We observed reduced fecal pH and increased humidity and viscosity of feces in PBY-treated mice. Yacon-supplemented diet has already been shown to reduce fecal pH in rodents [26] and improve fecal consistency and intestinal transit in humans [28]. In obese individuals, yacon consumption also improves satiety and decreases body weight [27]. However, in our study, dietary intake and body weight remained unchanged between the groups during the dietary intervention, probably because such effects should be more pronounced in overweight conditions. The increased weight of colon, cecum, and small intestine verified in PBY-treated mice are supposedly due to the trophic effect of yacon on the intestinal mucosa. Yacon is demonstrated to increase the number and depth of intestinal crypts [31,60] and enhance SCFA production, like butyric acid, by intestinal microbiota metabolism. This SCFA can be used as an energetic substrate for colonicocytes [61], thus favoring the trophic effect on the intestine.

Metabolic function biomarkers are essential to assess health, especially during carcinogenesis. In this study, the PBY diet did not alter biomarker concentrations of lipid metabolism and hepatic and renal function. However, it was verified increased albumin and alkaline phosphatase in PBY-treated animals. Higher albumin levels might be interesting for carcinogenesis-induced animals, as albumin levels decrease during inflammation, which is considered a negative reagent of the acute-phase inflammatory response [62]. Also, albumin levels might have increased owing to higher fecal SCFA concentration in PBY-treated mice, once albumin is responsible to carry fatty acids in the bloodstream [63]. The increase in alkaline phosphatase can be attributed to its role in osteogenesis [64], as yacon promotes an increase in intestinal calcium absorption and bone deposition [60].

Fecal SCFA concentration reflects bacterial activity in the colon, an important indicating of intestinal health. PBY-supplemented diet promoted increased formic, acetic, propionic, butyric, and valeric acid concentrations on feces. Such an increase may be attributed to the probiotic role of yacon, which contains high concentrations of FOS and inulin [65], stimulating probiotic bacteria growth such as lactobacilli and bifidobacteria [28,29]. The use of yacon flour has already been shown to increase propionic and butyric acid concentrations in rodents feces [26]. Besides the trophic effect on the intestinal mucosa, butyrate has also apoptotic action on cancer cells [66]. SCFA, mainly butyric acid, and to a lesser extent propionic acid, are supposed to act as histone deacetylase (HDAC) inhibitors [42,43]. HDAC inhibitors can trigger histone hyperacetylation, which allows the opening of chromatin and, therefore, the gene transcription activation. This process promotes the expression of silenced genes related to apoptosis and cell cycle arrest, contributing to the suppression of colon cancer cell growth [41,44]. The SCFA-induced cell growth blockade is associated with increased expression of the p21 cell cycle inhibitor and down regulation of cyclin B1 in colon cancer cells [42]. In addition, the role of the SFCA- SFCA receptor axis in the suppression of
bacterial invasion, chronic inflammation and inflammation-associated intestinal carcinogenesis has recently been demonstrated [67].

ACF is considered to be pre-neoplastic lesions or a previous event to CRC development [56]. In the present study, DMH-induced colorectal tumorigenesis in male BALB/c mice resulted in a greater amount of ACF in the proximal colon, followed by the medial colon, and in lesser number in the distal colon. Most of crypts foci was containing 3 aberrant crypts utmost. ACF induction also compromised the structure of intestinal mucosa, thus affecting the architecture of intestinal crypts, which usually leads to the recruitment of inflammatory infiltrates. PYB-supplemented diet did not alter intestinal ACF counts. Although an improvement in the crypt structure could be noticed, the histopathological score remained unchanged. Some authors have shown that DMH-induced animals fed with yacon-containing diets presented a reduction of ACF count [26,39]. Yacon flour containing 20.4% FOS, added to the diet at 5%, offered for 13 weeks after DMH-induction, significantly reduced the lesions in rats [39]. Similarly, a yacon flour-supplemented diet (7.5% FOS) offered for 8 weeks to DMH-induced rats (25mg/kg body weight) showed more than 40% reduction in ACF count [26]. The lack of changes in ACF counts in our study might be justified by the lesser percentage of FOS and inulin (6%) in PYB-supplemented diet, yacon processing type, or the short period of dietary intervention (8 weeks), which only started after ACF induction.

The profile of immune cells in the colon has been studied as a prognostic factor in CRC. Regarding CD4+ and CD8+ T lymphocytes, Treg cells (CD4+CD25+), Th17 cells (CD4+CCR6+), NK cells and CD4+CD8 ratio, no significant differences between the groups were noticed in our study. Nevertheless, PYB-treated animals had an increase in CD8+/CD4+ T cells ratio (156.9%), which denotes a higher prevalence of CD8+ T over CD4+ T cells. CD8+ T lymphocytes are strongly related to the anti-tumor role in CRC [9]. There was also an increase in Treg cells (84.8%) that might be attributed to the modulatory role of PYB, recently described by our research group [20]. However, the role of Treg cells is still controversial in CRC, as they are related to either tumor progression [68-70] or better CRC prognosis [16,71,72]. Treg cells play a regulatory role in the immune response, secreting cytokines and immune-modulating factors [73], able to suppress the effector function of both cytotoxic T lymphocytes and NK cells, which act eliminating tumor cells [74]. Furthermore, Treg cells may not only mediate the immunosuppression of immune responses to the tumor [15], but also attenuate the inflammatory immune response, especially when triggered by intestinal bacteria [16]. This fact denotes the strong immunosuppressive role of Treg cells during intestinal dysbiosis, usually seen in the CRC. Similarly to Treg cells, Th17 lymphocytes might have a dual role in CRC [9]. Although related to tumor progression owing to its main cytokine profile (IL-17A, IL-17F, IL-21, IL-22 and, IL-6) [12], Th17 lymphocytes may also favor neutrophils recruitment through IL-8 secretion, and drive highly cytotoxic CCR5CCR6CD8+ T cells into the tumor tissue, through CCL5 and CCL20 release [9].

Concomitantly, we investigate what could be the main immune pathways induced by PYB diet during colon carcinogenesis. Thus, the expression of transcription factors key to the adaptive immune response were assessed. The PYB-supplemented diet was able to down-regulate FOXP3 expression and up-regulated both RORγt and T-bet expression in the colon. FOXP3 is the key transcription factor for Treg cell differentiation. The differentiation of Th17 lymphocytes, in turn, requires the RORγt transcription factor [75], which is expressed in CD4+ T cells in an opposite manner of FOXP3 expression. High amounts of transforming growth factor-β (TGF-β) and pro-inflammatory cytokines, such as IL-6, IL-21, and IL-23 promote RORγt expression, whereas a low amount of pro-inflammatory cytokines and high amount of TGF-β induce FOXP3 expression [75]. Intestinal FOXP3 down-regulation in PYB-treated animals could be contributing to reduce Treg cells, which may be beneficial since such cells are being involved in anti-tumor immune response suppression [15]. Interestingly, cancer cells may also express FOXP3 [76], thus generating a confounding factor in the overall tissue analysis. Somehow, FOXP3 down-regulation may indicate a positive effect of PYB supplementation; however, it is necessary to differentiate FOXP3+ cancer cells from FOXP3- lymphocyte populations in intestine tissue for better comprehension of this data. Regarding Th17 cells, we did not observe changes between the groups; nevertheless, RORγt up-regulation might suggest a possible stimulation of theTh17 profile in the immune response.

T-bet transcription factor initiates the development of the Th1 cell line, which secretes interferon-gamma (IFN-γ), the main cytokine of this lineage. It also produces IL-2 and TNF-α [77]. In contrast, GATA-3 is the specific transcription factor for Th2 cell line development, which secretes the key cytokine IL-4 in addition to IL-5, IL-6, IL-9, and IL-13 [77-79]. It is widely recognized that IFN-γ-producing Th1 cells, as well as CD8 T lymphocytes, play an important role in inhibition and death of tumor cells [80]. Moreover, Th1 cells produce IL-2, which is essential for cytotoxic T lymphocyte proliferation [81]. Th1 cells have already been shown to be associated with better CRC prognosis [10,11]. Therefore, increased T-bet expression in PYB-treated animals could be contributing to impairing tumor growth and, thus, to better disease prognosis.

In this study, we confirmed our hypothesis that PYB may increase fecal SCFA concentration and modulate the intestinal immune response against colorectal carcinogenesis, verified by FOXP3 down-regulation, and RORγt and T-bet up-regulation. Despite our findings, and knowing the benefits of SCFA (butyric and propionic acids) in neoplastic cell apoptosis, as well as the role of Th1 cells (T-bet expression) in CRC control, it was not possible to verify changes in ACF counts with PYB consumption. Therefore, certain limitations in this study should be considered, for instance, the amount of FOS and inulin in PYB-supplemented diet, and the short period of dietary intervention. Besides, the restricted number of samples per analysis that required the entire colon, each one (ACF counts, immunophenotyping, and Real-time PCR) may have contributed to the absence of statistical significance, especially concerning immunophenotyping analysis. We have also considered that the human equivalent amount of dietary PYB (229g PYB per day for a 70kg adult) might be difficult to be achieved at once. Therefore, we suggest that PYB dosage could be spread throughout the day, thus favoring the inclusion of small portions in the usual diet, for example, in juices, shakes or bakery products. Interestingly, Santana and co-workers (de Souza Lima Sant’Anna et al., 2015) reported that in humans a lower amount of PYB (52g PYB twice daily, containing 10g FOS + inulin), mixed with orange juice for 30 days was effective in improving intestinal constipation and microbiota modulation. This fact suggests that prebiotic consumption long term may be more effective for a healthy gut.
Conclusion

In summary, PBY consumption did not reduce colonic ACF count; however, it promoted the reduction of fecal pH, and increased humidity and viscosity of feces, improving fecal characteristics. PBY also increased fecal SCFA concentration (formic, acetate, propionic, butyric, and valeric acids), thus indicating probiotic activity stimulity. Besides, the increase in SCFA, mainly butyric acid, can contribute to the activation of tumor suppressor genes. PBY consumption also down-regulated FOXP3 expression and up-regulated RORγt and T-bet expression in the colon, which would contribute to activation and proliferation of CD8 T lymphocytes and better CRC prognosis. Thus, we might infer that during the early stages of colorectal carcinogenesis in mice, PBY consumption contributes to driving the anti-tumor immune response. Therefore, we suggest that PBY, as a functional food, might improve intestinal health during the early stages of colorectal carcinogenesis, especially by modulating both SCFA production and colonic adaptive immune response. Studies concerning cell cycle arrest genes and apoptosis could contribute to the comprehension of this research.

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Conflict of Interest

The authors declare no conflict of interest.

Ethical Approval

All the trials and experimental protocols were approved by the Ethics Committee on Animal Experimentation of Universidade Federal de Viçosa, Brazil, under process number 30/2016, and performed according to the Guide for the Care and Use of Laboratory Animals, National Academy of Sciences (US).

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