

Cite this: *Food Funct.*, 2026, **17**, 3291

# Prune (dried plum) consumption does not reduce colonic tumor formation but drives beneficial changes in the gut microbiome of rats

Minghong Jiang,<sup>a</sup> Andres Gomez,<sup>b</sup> Davis M. Seelig<sup>c</sup> and Daniel D. Gallaher<sup>a\*</sup>

Previous research has highlighted the potential benefits of prune consumption, including a changed gut microbiome composition and a reduction in colon cancer risk factors. This study investigated whether prune consumption reduced colon tumor development and led to positive changes in the large intestinal microbiome in a chemically induced colon cancer model in rats. Male Wistar rats were fed one of three diets: 5% (by energy) prune, 10% prune, or a prune-free control. Rats were fed the diets for 32 weeks. Rats received weekly injections of 1,2-dimethylhydrazine for 15 weeks to induce colon tumorigenesis. Colonic tumor number or size did not differ among the diet groups. However, there was a trend toward fewer small intestinal tumors in the 10% prune diet group ( $p < 0.1$ ). Groups fed prune had heavier cecum tissue, indicating greater large intestinal fermentation. The prune diets increased taxonomic richness and altered bacterial species composition. Specifically, prune consumption was associated with increased abundance of *Methanospaera* genus and taxa from the Lachnospiraceae family, such as *Blautia* and *Coproccoccus*. Prune diets also increased total cecal SCFA amount, notably butyrate. However, 24 hour fecal excretion of *p*-cresol, indole, and total bile acids did not differ significantly among the groups. While prune consumption did not show a significant reduction in colonic tumor formation, potential benefits were noted in a trend towards reducing small intestine tumors, increasing large intestinal fermentation and SCFA production, and increasing microbial richness, suggesting prune consumption may provide other health benefits.

Received 8th August 2025,  
Accepted 27th January 2026

DOI: 10.1039/d5fo03398e

rsc.li/food-function

## 1. Introduction

Colorectal cancer (CRC) is the third leading cause of cancer-related mortality in the United States and the second globally.<sup>1,2</sup> The incidence of CRC has increased rapidly in countries that have adopted a Westernized diet. The Western diet, characterized by the consumption of large amounts of red and processed meat, refined grains, and sugar, coupled with low intake of whole grains, fruits, and vegetables has been associated with dysbiosis of the gut microbiome and inflammation in the gut, thus contributing to the development of CRC.<sup>3</sup> Due to advancements in treatment and screening, there has been a 57% decline in overall CRC mortality in the U.S. from 1970 to 2020. However, there is a concerning recent trend of an upward trajectory in CRC cases in individuals under 50 years old, accompanied by a significant shift towards advanced disease stages.

Fruit intake has been inversely associated with CRC risk, but most epidemiological studies report only total fruit intake rather than specific types, making it difficult to assess the chemopreventive potential of individual fruits.<sup>4,5</sup> Prunes, recognized as a functional food, offer various health benefits, including alleviating constipation, lowering cholesterol, and improving bone density. Derived from the dried fruit of *Prunus domestica*, prunes are rich in dietary fiber, sorbitol, and phytochemicals such as polyphenols and carotenoids.<sup>6</sup> However, compared to more commonly studied fruits like citrus and apples, prunes (dried plums) are less frequently consumed and even less studied, leaving their chemopreventive effects largely unknown.<sup>5</sup>

Yet there is evidence that prune consumption may reduce colon cancer risk. In carcinogen-treated rats, feeding a plum beverage reduced inflammation and colonic pre-neoplastic lesions known as aberrant crypt foci (ACF), which have been commonly used as an early biomarker of colon carcinogenesis. Polyphenols extracted from prunes have been shown to exhibit strong antioxidant and anti-inflammatory properties,<sup>7,8</sup> reducing two established hallmarks of cancer.<sup>9</sup> In cancer cell lines, prune extracts or isolated polyphenols induced apoptosis and suppressed cell proliferation,<sup>10,11</sup> indicating potential chemopreventive effects against colonic tumor formation. However,

<sup>a</sup>Department of Food Science and Nutrition, University of Minnesota, St. Paul, MN, 55108, USA. E-mail: dgallaher@umn.edu

<sup>b</sup>Department of Animal Science, University of Minnesota, St. Paul, MN, 55108, USA

<sup>c</sup>Department of Clinical Pathology, Veterinary Clinical Sciences, University of Minnesota, St. Paul, MN, 55108, USA



feeding a prune powder in another study did not show a reduction in ACF, although changes in other colorectal cancer risk factors, including reduced total and secondary bile acids, were observed.<sup>12,13</sup> Due to these inconsistent findings, it is uncertain whether prunes or prune products will reduce the risk of colon cancer.

There is also growing interest in the role of the gut microbiome in the development of CRC. CRC patients have lower microbial diversity and different fecal microbial population patterns compared to healthy individuals.<sup>14</sup> Prunes exert beneficial effects on the gut ecosystem, likely due to its being a rich source of fiber, with 40–80% of the fiber being pectin, and a high concentration of insoluble fibers, polyphenols, and sorbitol.<sup>6</sup> These nutrients undergo fermentation by the gut microbiota in the large intestine, leading to increased production of short chain fatty acids (SCFAs), including butyrate, which is an important energy source for the colonic epithelium. Butyrate has also been found to reduce CRC risk.<sup>15,16</sup> Recent research involving prune supplementation in human studies revealed significant shifts in microbial composition in prune consumption groups.<sup>17,18</sup> Studies found that prune consumption increased bacterial populations typically regarded as beneficial, such as *Bifidobacterium* and *Lactobacillus*, while reducing harmful bacteria such as *Escherichia coli* and *Clostridium perfringens* (*C. perfringens*).<sup>19,20</sup> The increased abundance of the Lachnospiraceae family was correlated with a reduced plasma concentration of the inflammatory cytokine IL-1 $\beta$ , suggesting a positive effect of prune consumption in modulating the gut microbiome and reducing systemic inflammation.<sup>17</sup>

These observations support the chemopreventive potential of prune consumption. Therefore, we hypothesized that diets containing prunes would reduce dimethylhydrazine-induced colonic tumors in rats. Furthermore, given the role of the gut microbiome in modulating colon cancer risk,<sup>21</sup> we analyzed bacterial abundance in the large intestine as well as determined bacterial metabolites implicated in colon cancer risk, including short chain fatty acids, bile acids, and markers of inflammation.

## 2. Materials and methods

### 2.1. Animals and diet

Eighty-one male Wistar rats, initially weighing between 100 and 170 g, were purchased from Envigo (Indianapolis, IN). The rats were housed in pairs and maintained under a 12 hour light/dark cycle, with *ad libitum* access to food and water. All animal handling procedures were conducted in accordance with the guidelines set by the National Institutes of Health and met the institutional standards for animal care. The experimental protocol was approved by the University of Minnesota Animal Care and Use Committee (protocol #2012-38704A).

The prune product (hereafter referred to as prune) was provided by Sunsweet Growers Inc. (Yuba City, CA). It was pre-

pared by blending 75% prune concentrate with 25–30% prune paste by weight. The prune concentrate was produced by removing pulp and pits, followed by vacuum evaporation until reaching 68° Brix, while the prune paste was made from steamed plums adjusted to 22–25% moisture. The prune used in this study had a moisture content of approximately 29.8%. The primary compositional difference between the prune and whole prunes was the fiber content. The prune contained more soluble fiber (6.4 g vs. 2.1 g per 100 g) and less insoluble fiber (1.1 g vs. 4.1 g per 100 g) compared with whole prune but was otherwise similar. The macronutrient composition of the prune, whole prunes, prune juice, and fresh plums is shown in Table S1.

All diets were formulated to provide equal macronutrient and total energy levels (Table 1). The pectin concentrations in the 5% and 10% prune diets were 0.28% and 0.56%, respectively, while sorbitol concentrations were 1.15% and 2.30%, respectively. The basal diet contained cellulose as the sole fiber source.

The prune was incorporated into the AIN-93G diet<sup>21</sup> at concentrations providing 5% and 10% of total energy. The prune was thoroughly mixed with other dry ingredients and soybean oil using a commercial mixer to achieve a uniform texture. The diets were fed as a powder and stored at 4 °C.

### 2.2. Experimental design

Rats were randomly assigned to one of the three diets. After a two-week adaptation period to their respective diets, the rats were administered a weekly subcutaneous injection of dimethylhydrazine (20 mg per kg body weight) for 15 consecutive weeks. Following the completion of all injections, the rats were placed in metabolic cages for 24 hours to collect urine and fecal samples, which were subsequently stored at –80 °C. The rats continued to be fed their respective diets for an additional

**Table 1** Composition of experimental diets

	Basal	5% prune	10% prune
<b>Ingredients (g kg<sup>-1</sup>)</b>			
Dried prune puree	0	76	154
Sucrose	100	73	66
Maltodextrin	132	132	132
Cornstarch	368	368	368
Casein	200	198	196
Soybean oil	70	70	70
Cellulose	80	73	66
AIN-93G mineral mix	35	35	35
AIN-93G vitamin mix	10	10	10
L-Cystine	3	3	3
Choline bitartrate	2.5	2.5	2.5
TBHQ	0.014	0.014	0.014
<b>Macronutrient composition (% of total energy)</b>			
Carbohydrate (% of energy)	62.6	62.6	62.6
Fat (% of energy)	16.5	16.4	16.4
Protein (% of energy)	20.9	20.9	20.9
Prune pectin and sorbitol (% of energy)	0	1.4	2.8
Total energy (kcal g <sup>-1</sup> )	3.8	3.7	3.5



15 weeks until the end of the feeding trial. Throughout the study, food intake and body weight were recorded weekly.

At the end of the 32 week period, the rats were fasted overnight and then anesthetized the next morning using isoflurane. Blood was collected through cardiocentesis and the liver, epididymal fat pad, and cecum were excised, rinsed of adhering blood, and weighed. Cecum contents were collected, and the empty cecum was weighed. Tissues were rapidly frozen in liquid nitrogen before being stored at  $-80\text{ }^{\circ}\text{C}$ . Serum was separated and stored at  $-20\text{ }^{\circ}\text{C}$  for subsequent analysis.

The colon was dissected from the anus to the cecum and opened longitudinally, and colon pellets were collected and snap-frozen in liquid nitrogen before storage at  $-80\text{ }^{\circ}\text{C}$ . Tumors were counted and tumor volume calculated using the formula  $V = \text{length} \times \text{width} \times \text{diameter} \times \pi/6$ , where length is the longest dimension, width is the middle dimension, perpendicular to length, and diameter is the smallest dimension, perpendicular to both length and width.<sup>22</sup> Following tumor excision, each tumor with a segment of normal colonic interstitial tissue was fixed in 10% neutral buffered formalin for 72 hours and then transferred to 70% ethanol for histopathological analysis. The remaining colon tissue was placed on an ice-cold metal tray, and the mucosa was carefully scraped off using a glass slide. The resulting mucosal scraps were snap-frozen in liquid nitrogen and stored at  $-80\text{ }^{\circ}\text{C}$  for subsequent measurement of inflammatory markers.

### 2.3. Tumor histopathologic analysis

The formalin-fixed colon tissue underwent a standard processing procedure, involving dehydration through a series of graded alcohols, followed by embedding in paraffin wax. Subsequently,  $4\text{ }\mu\text{m}$  tissue sections were cut and stained with hematoxylin and eosin. The stained tissue sections were examined by a board-certified veterinary pathologist according to established consensus guidelines on the pathology of rodent models of intestinal cancer.<sup>23,24</sup> The pathologist was blinded to treatment groups during the assessment of the proliferative lesions. Lesions were categorized as hyperplasia, adenoma, intramucosal adenocarcinoma, or adenocarcinoma, based on specific morphological criteria and cellular characteristics observed in the tissue sections.<sup>23,25</sup>

### 2.4. Analysis of fecal *p*-cresol and indole

Fecal pellets collected during housing in the metabolic cages were freeze-dried and ground into a fine powder. Five mL of methanol and  $100\text{ }\mu\text{L}$  of internal standard ( $300\text{ }\mu\text{g mL}^{-1}$  4-ethylphenol in ethanol) were added to duplicate samples of  $0.5\text{ g}$  of fecal powder, and the samples were mixed and refluxed at  $100\text{ }^{\circ}\text{C}$  for 15 minutes. After cooling, the samples were centrifuged at  $2000g$  for 5 minutes, and the supernatant was carefully decanted into a new tube. The extracted samples were then resuspended in  $5\text{ mL}$  of methanol, refluxed again, and centrifuged as previously described. The sample was further washed with  $0.5\text{ mL}$  of methanol and centrifuged once more. The pooled supernatants were dried under nitrogen gas and reconstituted with  $0.3\text{ mL}$  of methanol and filtered through a  $0.45\text{ }\mu\text{m}$  syringe

filter. Quantification of *p*-cresol and indole was performed using HPLC with fluorescence detection, following a modified version of the method described by King *et al.*<sup>26</sup> Separation was achieved using a C18 column with a flow rate of  $1\text{ mL min}^{-1}$ . The mobile phase consisted of a 7:3 water/acetonitrile solution adjusted to pH 3.2 with glacial acetic acid.

### 2.5. Analysis of total fecal bile acids and cecal SCFAs

Analysis of total bile acids was conducted on ground fecal samples collected during housing in the metabolic cages. The bile acids were extracted from dried fecal powder and partially purified using solid-phase extraction columns, as described by Lockett and Gallaher.<sup>27</sup> Bile acids were assayed enzymatically using  $3\alpha$ -hydroxysteroid dehydrogenase.<sup>28</sup> SCFAs were extracted from cecal contents and quantified using gas chromatography, as described by Carlson *et al.*<sup>29</sup>

### 2.6. Determination of inflammatory markers

Colon mucosal scrapings were homogenized in  $1\text{ mL}$  of phosphate-buffered saline solution (pH 7.4) containing protease and phosphatase inhibitors (Roche, Switzerland). After centrifugation at  $10\text{ }000g$  at  $4\text{ }^{\circ}\text{C}$  for 5 minutes, supernatants were diluted 1:3 with PBS, proteins separated on a 10% SDS-PAGE gel, and the bands transferred to nitrocellulose membranes.

The membranes were blocked with 5% non-fat dry milk solution in TBST (0.1% Tween-20) and incubated with primary antibodies overnight at  $4\text{ }^{\circ}\text{C}$ . The membranes were incubated with secondary antibodies for an hour at room temperature. Protein bands were visualized using Super-Signal West Atto Ultimate Sensitivity Chemiluminescent Substrate (Thermo Fisher Scientific) and scanned using the Licor Odyssey imaging system.

The primary antibodies used in this study included NF $\kappa$ B, p-NF $\kappa$ B p65 (Ser536),  $\beta$ -catenin, and  $\beta$ -actin (Cell Signaling Technology, Beverly, MA). Serum calprotectin levels were determined using an ELISA assay following the manufacturer's guidelines (S100A8/S100A9, Immundiagnostik AG, Bensheim, Germany).

### 2.7. DNA extraction and 16S rRNA gene sequencing

The colonic pellets were mechanically disrupted by bead beating for 3 minutes at  $6\text{ m s}^{-1}$ , and microbial DNA was extracted using the QIAamp Fast DNA Stool Mini Kit (QIAGEN, Germantown, USA), following the manufacturer's instructions. The 16S rRNA bacterial gene (V4 region) was sequenced at the University of Minnesota Genomics Center (UMGC), on the Illumina MiSeq sequencing platform, using an optimized method for amplicon-based library preparation.<sup>30</sup>

### 2.8. Processing of sequencing data

Amplicon sequence variants (ASVs) were identified from raw Illumina sequences using a combination of Cutadapt, BBDMap, and the DADA2 plugin within QIIME 2.<sup>31–33</sup> Briefly, adapter sequences, barcodes, primers, and low-quality sequences were eliminated from the pair-end reads using Cutadapt with default parameters. Subsequently, the pair-end reads were merged using the BBDMap merger function, and singleton



sequences were discarded. The resulting quality-filtered data were then imported into QIIME2 (version 2022.8) and processed through the DADA2 pipeline, where sequences underwent demultiplexing, quality filtering, and denoising. Taxonomic analysis of the ASVs was performed using the Greengenes 13\_8 database.<sup>34</sup> Following preprocessing, approximately 70% of the reads were retained, yielding an average of 17 000 reads per sample (range: 3315 to 34 582).

The ASV table generated from these steps was then imported into R Studio for further analysis. Bray–Curtis beta-diversity and Shannon alpha-diversity were calculated using the Vegan package in R.<sup>35</sup> Additionally, indicator species analysis was used to identify ASVs associated with each dietary group by calculating an indicator value (IndVal), the product of an ASV's frequency and abundance within a group. IndVal ranges from 0 to 1, with 1 indicating the strongest association within a specific group.<sup>36</sup> IndVal analysis was conducted at both the genus and species levels and significance was assessed using the Kruskal–Wallis test.<sup>36</sup>

### 2.9. Statistical analysis

Statistically significant differences among the groups were inspected using the analysis of variance (ANOVA) test for numerical variables using Statistical Analysis Software (SAS, version 9.4), with final body weight used as a covariate for liver and fat pad weight analyses. Given the unequal variance observed in tumor data, according to the Shapiro–Wilk test, the data were ranked and subsequently assessed using the ANOVA test within the SAS software. Duncan's multiple range test was used to identify statistically significant differences among groups. Comparisons of tumor incidence among groups were conducted using the chi-square test.

Microbial community differences were evaluated using PERMANOVA, which tests significant differences in group centroids, and ANOSIM (Analysis of Similarities), which assesses whether between-group dissimilarities exceed within-group dissimilarities. Both were performed with 999 permutations on Bray–Curtis distances using the vegan R package.<sup>35</sup> Alpha diversity indices, richness, and individual taxa relative abundances were tested for normality using the Shapiro–Wilk test. Based on the results, group differences were assessed using one-way ANOVA for normally distributed data or the Kruskal–Wallis test for non-normal data. Following this, pairwise comparisons were performed using Tukey's HSD *post hoc* test or Dunn's test to identify specific group differences.

Associations between individual taxa and SCFAs were determined by calculating Spearman's correlation coefficients. The top five taxa strongly correlated with SCFAs underwent further analysis using a multiple regression model to explore the association between each taxon, serving as a predictor, and each individual SCFA. The multiple regression model handles multicollinearity of independent variables, providing a more reliable estimate of the association between taxa and SCFAs. Taxa exhibiting a non-significant association with specific SCFAs were excluded from the multiple regression model. Before conducting the multiple regression analysis, normality

and multicollinearity assumptions were thoroughly assessed, which included checking normality using Q–Q plots and evaluating multicollinearity through a correlation plot of all independent variables. These steps were essential to ensure the reliability and validity of the subsequent statistical analyses.

## 3. Results

### 3.1. Tissue weight, energy intake, and weight gain

Throughout the study duration, the weekly and final body weights of rats showed no significant differences among the groups, indicating that prune consumption did not influence body weight. Similarly, energy intake remained equivalent across all groups throughout the study period. Differences were observed in liver and epididymal fat pad weights among the groups. The group consuming the 5% prune diet exhibited significantly lower liver and epididymal fat pad weights compared to the basal group ( $p = 0.05$  and  $p = 0.04$ , respectively, Table 2), despite no significant differences in overall body weight. Empty cecum weight, an indicator of large intestinal fermentation, was significantly greater in the prune diet groups ( $p = 0.01$ ). Furthermore, the cecal content weight was greater in the 5% prune diet group compared to the basal group ( $p = 0.04$ ), indicating greater large intestinal contents due to prune consumption.

### 3.2. Tumor number and volume

The physical appearance of the colon segments varied, as shown in Fig. 1. The healthy (*i.e.*, uninvolved) colon segments exhibited a smooth and uniform appearance, lacking any visible irregularities or protrusions. Adenomas, early-stage tumor growths, were identifiable by their smooth and round polyp-like formations along the inner lining of the colon. Adenocarcinoma, indicative of advanced and malignant growth, presented as irregular masses with an uneven surface. Additionally, the colon segment lining displayed a pale appear-

**Table 2** Body weight, energy intake, and organ weights

	Diet		
	Basal	5% prune	10% prune
Final body weight (g)	613 ± 18	575 ± 18	608 ± 18
Energy intake (kcal d <sup>-1</sup> )	89.56 ± 3.47	91.15 ± 3.32	89.28 ± 3.52
Liver weight (g)	16.0 ± 0.81*	14.8 ± 0.69 <sup>†</sup>	15.40 ± 0.65* <sup>†</sup>
Epididymal fat pad weight (g)	7.10 ± 0.49*	5.60 ± 0.37 <sup>†</sup>	6.60 ± 0.43* <sup>†</sup>
Cecum weight (g)	0.58 ± 0.02*	0.80 ± 0.08 <sup>†</sup>	0.75 ± 0.03 <sup>†</sup>
Cecum content weight (g)	2.13 ± 0.16*	4.71 ± 1.23 <sup>†</sup>	3.22 ± 0.14* <sup>†</sup>
Fecal weight (g per 24 h)	1.66 ± 0.31	1.49 ± 0.11	1.71 ± 0.20

Values are mean ± SE;  $n = 29$  (basal), 25 (5% prune), and 21 (10% prune); values not sharing a common superscript are statistically different, ( $P < 0.05$ ). Liver and fat pad weights were analyzed using body weight as a covariate, as body weight was a significant covariate for both ( $P < 0.0001$  and  $P = 0.002$ , respectively).



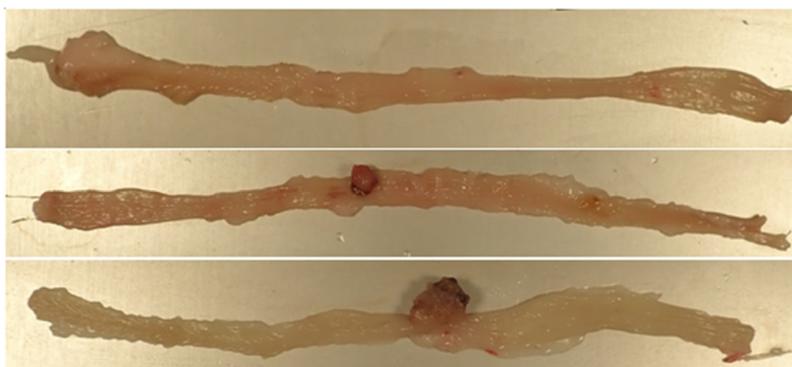


Fig. 1 Top, normal colon; middle, colon with adenoma; bottom, colon with adenocarcinoma.

ance, potentially signaling pathological changes or inflammation within the tissue.

No significant differences were observed in the count or volume of non-cancerous lesions, colon adenocarcinoma, total colonic tumors, or total GI tumors across the studied groups. However, both the 5% and 10% prune diets displayed a trend towards fewer small intestine tumors ( $p = 0.1$ ) (Table 3). Moreover, the 10% prune group also had a tendency toward a lower volume of small intestine tumors compared to the basal diet group ( $p < 0.1$ ). Finally, the incidence of tumors, defined as the percentage of rats with tumors, did not significantly differ among the experimental groups.

Table 3 Tumor count, volume, and incidence

	Diet group		
	Basal	5% prune	10% prune
<b>Count</b>			
Small intestine adenocarcinomas	44.2 ± 3.6*	37.1 ± 2.6 <sup>†</sup>	36.3 ± 2.7 <sup>†</sup>
Colonic non-cancerous lesions	40.3 ± 3.8	39.9 ± 4.5	38.1 ± 3.9
Colonic adenomas	40.3 ± 3.1	41.8 ± 3.6	36.1 ± 2.8
Colonic adenocarcinomas	39.3 ± 3.8	37.5 ± 4.8	41.8 ± 4.3
Total colonic tumors	39.3 ± 3.8	38.9 ± 4.9	40.3 ± 3.8
Total GI tumors	44.3 ± 3.8	35.4 ± 4.7	37.9 ± 4.6
<b>Volume (mm<sup>3</sup>)</b>			
Small intestine adenocarcinoma	38.9 ± 3.3*	33.4 ± 3.0* <sup>†</sup>	31.9 ± 2.7 <sup>†</sup>
Colonic non-cancerous lesions	39.9 ± 3.8	39.5 ± 4.4	38.9 ± 4.2
Colonic adenoma	35.5 ± 3.0	38.0 ± 3.7	31.5 ± 2.8
Colonic adenocarcinoma	34.0 ± 4.1	33.5 ± 4.1	37.6 ± 4.4
Total colonic tumor	34.5 ± 4.0	34.7 ± 3.9	35.9 ± 4.7
Total GI tumor	36.1 ± 4.2	33.0 ± 3.8	35.1 ± 4.2
<b>Incidence (%)</b>			
Small intestine tumor	31	16	17
Colonic adenoma	24	28	13
Colonic adenocarcinoma	76	72	87
Colonic tumor	86	80	92
Total GI tumor	90	92	84

Tumor count and volume are presented as ranked means ± SE;  $n = 29$  (basal), 25 (5% prune), and 24 (10% prune). Tumor incidence is expressed as the percentage of animals with tumors in each diet group. Values not sharing a common superscript indicate a trend toward significance ( $P < 0.1$ ) based on the chi-square test.

### 3.3. Fecal and cecal metabolites

Analysis of cecal content SCFAs revealed significant increases in butyrate ( $P < 1 \times 10^{-4}$ ) and total SCFA amount ( $p = 0.01$ ) per cecum in both the 5% prune and 10% prune diets (Fig. 2). Butyrate increased linearly with the dietary concentration of prune ( $R^2 = 0.98$ ). Animals fed the 5% prune diet had the highest amount of cecal propionate ( $p = 0.03$ ) and the branched short chain fatty acid isovalerate ( $p = 0.049$ ) compared to other dietary groups. Additionally, there was a trend towards higher acetate and valerate amounts ( $p = 0.1$ ) in the 5% prune diet. Across all diet groups, acetate was present in the largest amount among all SCFAs.

In contrast, the analysis of total fecal bile acids did not reveal any statistically significant differences among the groups ( $p = 0.44$ ) (Table 4). There were no significant differences among the groups in the fecal concentration of *p*-cresol or indole (Table 4). These findings suggest that while prune intake influenced SCFA levels, particularly butyrate and propionate, it did not significantly impact fecal bile acids or metabolites of aromatic amino acids in this study.

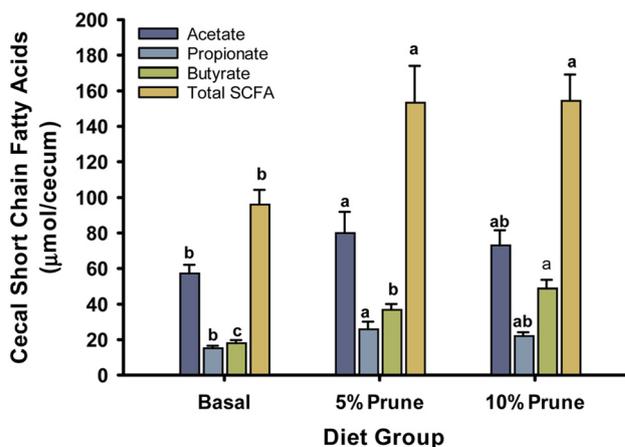


Fig. 2 Amount of short chain fatty acids in cecal contents. Values are mean ± SE. Values not sharing a common letter are statistically different,  $P < 0.05$ .



**Table 4** Daily excretion of fecal bacterial metabolites

	Diet		
	Basal	5% prune	10% prune
<i>p</i> -Cresol ( $\mu\text{g d}^{-1}$ )	7.94 $\pm$ 1.59	6.24 $\pm$ 1.02	7.99 $\pm$ 1.23
Indole ( $\mu\text{g d}^{-1}$ )	0.53 $\pm$ 0.21	0.27 $\pm$ 0.08	0.62 $\pm$ 0.20
Total bile acids ( $\mu\text{mol d}^{-1}$ )	63.97 $\pm$ 12.06	50.21 $\pm$ 9.18	59.60 $\pm$ 10.84

Values are expressed as mean  $\pm$  SE. Sample sizes were as follows: *p*-cresol and indole,  $n = 16$  (basal), 13 (5% prune), and 18 (10% prune); bile acids,  $n = 11$  (basal), 14 (5% prune), and 12 (10% prune).

### 3.4. Inflammatory markers

The inflammatory markers NF $\kappa$ B, p-NF $\kappa$ B, the p-NF $\kappa$ B/NF $\kappa$ B ratio, and  $\beta$ -catenin were analyzed in non-involved colon tissue by western blotting. There were no statistically significant differences among the diets for any marker (Fig. 3A and B). Serum calprotectin, a well-known inflammatory marker utilized for diagnosing inflammatory bowel diseases,<sup>37</sup> did not differ among the dietary groups (Fig. 3C).

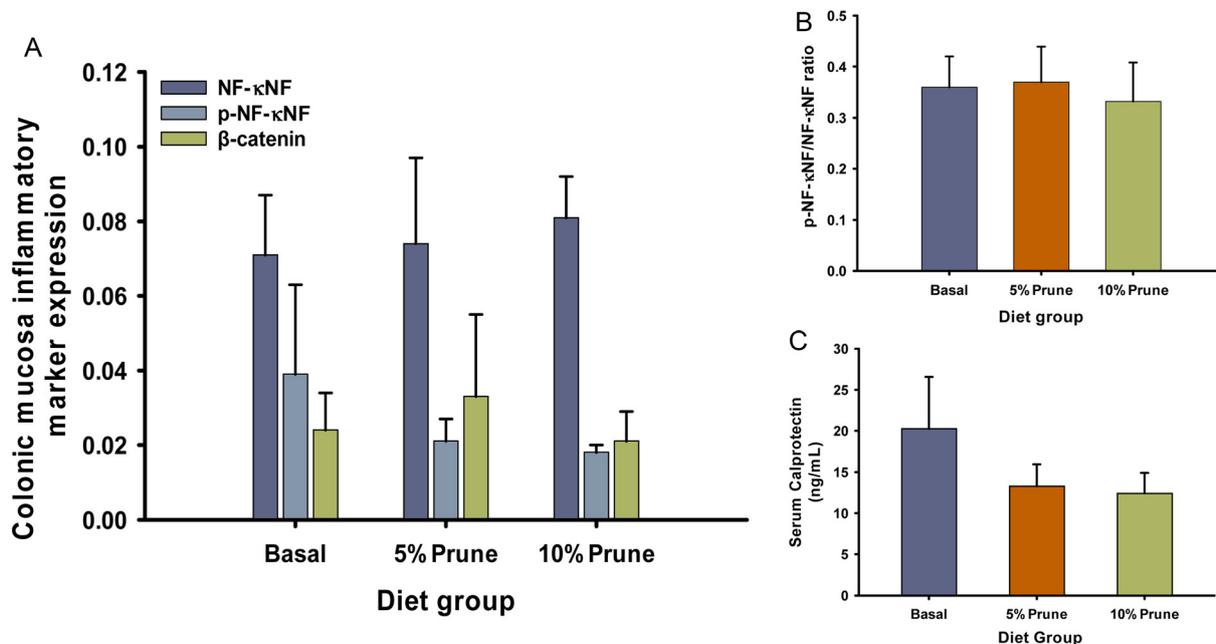
### 3.5. Changes in the gut microbiome

The Bray–Curtis dissimilarity index shows a clear separation in microbiome composition between the prune-containing diets and the basal diet in both cecum and colon contents (Fig. 4A and B). PERMANOVA analysis confirmed significant group differences (colon:  $R^2 = 0.16$ ,  $p = 0.001$ ), indicating that 16% of the variation in the colonic microbiota was explained by

diet. ANOSIM analysis further supported a moderate separation ( $R = 0.48$ ,  $p = 0.001$ ). In contrast, the cecal microbiome showed a smaller dietary effect ( $R^2 = 0.10$ ,  $p = 0.001$ ), suggesting that prune consumption may have had a slightly more pronounced impact on the colonic microbial community structure. Alpha diversity was assessed using the Shannon index and richness. The Shannon index was significantly higher in both prune groups compared with the basal group (colon:  $p = 1.9 \times 10^{-5}$ ; cecum:  $p = 0.024$ ; Fig. 4C and D), indicating greater microbial diversity and evenness of species. Similarly, richness increased in the colon ( $p = 0.0025$ ; Fig. 4E), reflecting an overall greater bacterial diversity with prune supplementation.

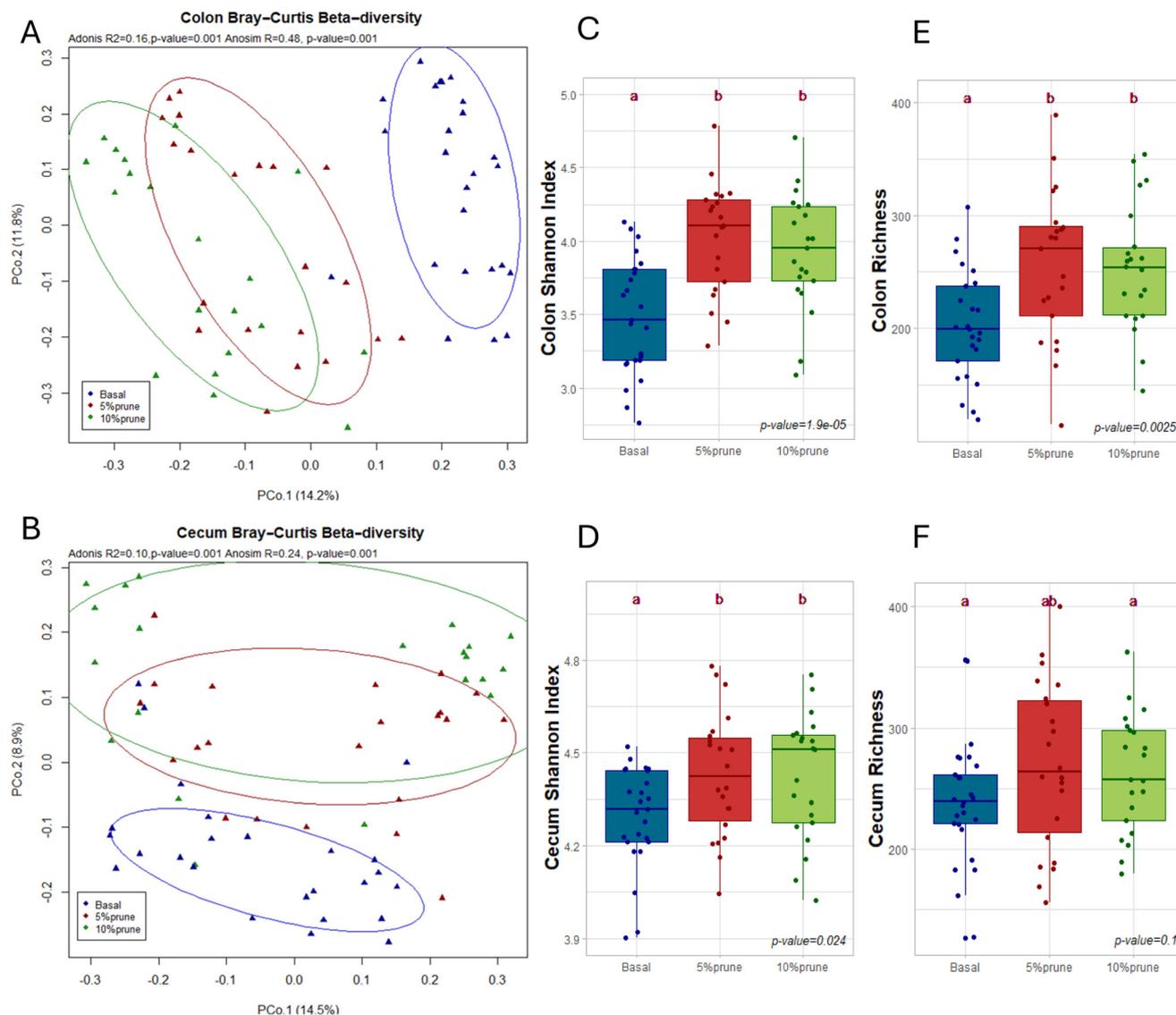
Indicator species analysis identified taxa differentiating the dietary groups at both genus and species levels (Fig. 5A and B). Prune consumption increased the relative abundance of *Prevotella*, *Paraprevotella*, *Coprococcus*, *Coprobacillus*, *Methanosphaera*, and *Collinsella*, while reducing *Clostridium*, *Ruminococcus*, and *Bifidobacterium*. At the species level, *Lactobacillus salivarius*, *Ruminococcus flavefaciens*, *Ruminococcus bromii*, *Coprococcus eutactus* and *C. perfringens* decreased with prune intake, whereas *Collinsella aerofaciens* increased. *Blautia* was enriched only in the 5% prune group, suggesting a nonlinear response to prune in the diet. At the family level, Coriobacteriaceae, Lachnospiraceae, and Barnesiellaceae were more abundant in prune-fed rats, with the largest increase observed for Barnesiellaceae in the 5% prune group (Fig. 5C).

Associations between microbial taxa and cecal SCFAs are summarized in Table 5. *Methanosphaera* showed consistent positive correlations with all SCFAs, including acetate, buty-



**Fig. 3** (A) Protein expression of NF- $\kappa$ B ( $P = 0.92$ ), p-NF- $\kappa$ B ( $P = 0.57$ ), and  $\beta$ -catenin ( $P = 0.84$ ) in colonic mucosa. (B) Ratio of p-NF- $\kappa$ B to NF- $\kappa$ B ( $P = 0.93$ ). (C) Serum calprotectin concentration ( $P = 0.81$ ). Values are presented as mean  $\pm$  SE. No significant differences were observed among diet groups.





**Fig. 4** (A and B) Beta-diversity analysis of colonic and cecal samples, respectively. Each dot represents a single microbiome, color-coded by diet group. Results from Adonis and ANOSIM tests are shown with the corresponding  $R/R^2$  values and  $P$ -values. (C and D) Shannon diversity index of colonic and cecal samples, respectively, with  $P$ -values. (E and F) Taxonomic richness in colonic and cecal samples, respectively, expressed as the number of observed taxa per sample, with  $P$ -values.

rate, valerate, and total SCFAs, suggesting a broad role in SCFA metabolism. *Prevotella* exhibited the strongest correlation with butyrate, while *Coriobacteriaceae*, *Blautia*, and *Paraprevotella* were positively associated with both butyrate and propionate. Multiple regression analysis further identified *Methanosphaera* as a key predictor for SCFA levels and confirmed significant associations between *Coriobacteriaceae* and *Prevotella* with butyrate levels (Table 6).

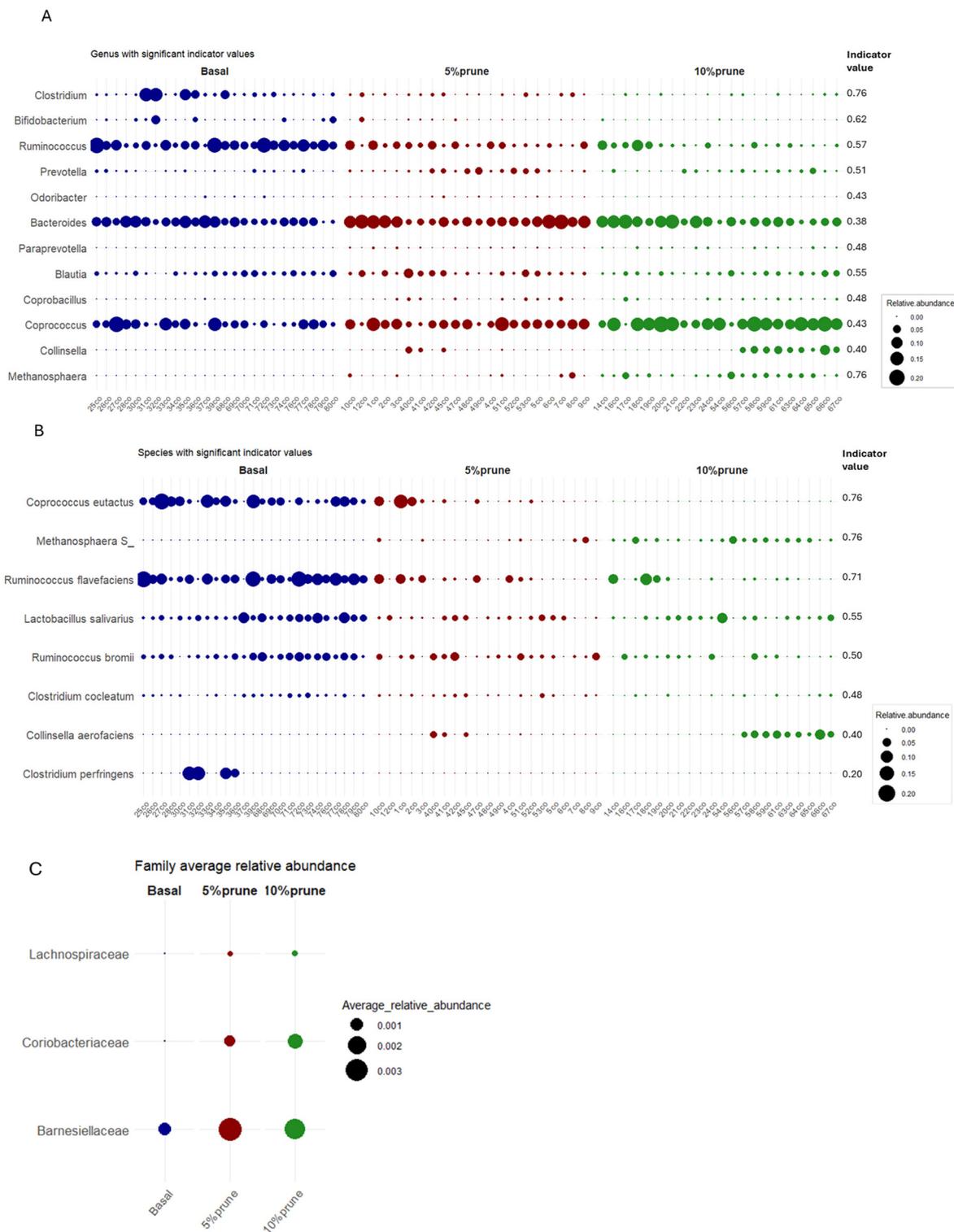
## 4. Discussion

In the present study, we incorporated a prune product (prune) that was substantially similar to whole prunes into the diets of

carcinogen-treated rats to evaluate its effect on tumor formation. Our hypothesis was that the prune, as a food high in fermentable fibers, sorbitol, and polyphenolics, would decrease tumor development. However, prune consumption did not reduce colonic tumors. We also carried out both microbiome and metabolite analyses of large intestinal contents to assess changes in the microbial composition and bacterial metabolites. Prune feeding resulted in substantial changes to the microbiome but did not affect daily fecal bacterial metabolite excretion.

Previous cohort studies have reported that high dietary fiber intake, particularly from cereals and fruits, is associated with reduced colorectal adenomas and distal colon cancer.<sup>38</sup> However, in the present study, all three diets were matched for





**Fig. 5** Bubble plots indicating the relative abundance of a taxon in an individual animal sample, color-coded by diet group. Sample sizes:  $n = 25$  (basal), 20 (5% prune), and 21 (10% prune). Indicator values are displayed for relevant taxa. (A) Genus-level bubble plot with indicator values. (B) Species-level bubble plot with indicator values. (C) Family-level bubble plot showing average relative abundance.

total fiber content, which was primarily cellulose in all three diets (80, 73, and 66 g kg<sup>-1</sup> in the basal, 5%, and 10% prune diets, respectively), a fiber that is highly resistant to fermenta-

tion. Pectin, a highly fermentable soluble fiber, and sorbitol, a highly fermentable sugar alcohol, were present only in the prune diets. Consequently, this study represents a comparison



**Table 5** Spearman's correlation  $\rho$  and  $P$ -value of top five taxa to each short chain fatty acid (SCFA)

	Acetate	Butyrate	Propionate	Valerate	Total SCFAs
<i>g_Methanosphaera</i>	0.42 ( $5.8 \times 10^{-4}$ )	0.53 ( $1.2 \times 10^{-5}$ )	0.32 ( $1.0 \times 10^{-2}$ )	0.44 ( $3.1 \times 10^{-4}$ )	0.52 ( $1.2 \times 10^{-5}$ )
<i>g_Blautia</i>	0.37 ( $3.0 \times 10^{-3}$ )	0.50 ( $4.1 \times 10^{-5}$ )			0.45 ( $2.0 \times 10^{-4}$ )
<i>g_Ruminococcus</i>	0.37 ( $3.2 \times 10^{-6}$ )			0.31 ( $1.4 \times 10^{-2}$ )	
<i>f_Barnesiellaceae</i>			0.33 ( $7.8 \times 10^{-3}$ )	0.28 ( $2.9 \times 10^{-2}$ )	0.37 ( $3.2 \times 10^{-3}$ )
<i>g_Christensenella</i>	0.33 ( $8.4 \times 10^{-3}$ )				
<i>f_Victivallaceae</i>	0.34 ( $6.5 \times 10^{-3}$ )				
<i>f_Coriobacteriaceae</i>		0.66 ( $6.7 \times 10^{-9}$ )	0.42 ( $7.1 \times 10^{-4}$ )	0.25 ( $5.0 \times 10^{-2}$ )	0.41 ( $1.0 \times 10^{-3}$ )
<i>g_Prevotella</i>		0.71 ( $1.2 \times 10^{-10}$ )	0.41 ( $9.3 \times 10^{-4}$ )	0.35 ( $5.7 \times 10^{-3}$ )	0.48 ( $9.5 \times 10^{-5}$ )
<i>g_Paraprevotella</i>		0.46 ( $1.7 \times 10^{-4}$ )	0.30 ( $1.8 \times 10^{-3}$ )		

For each SCFA and the total SCFA, the top five taxa with the strongest correlations are listed. Values are presented as Spearman's correlation coefficients ( $\rho$ ) along with the corresponding  $p$ -values in parentheses.  $\rho = 0.1$ – $0.4$ , weak positive correlation;  $0.4$ – $0.7$ , moderate positive correlation;  $>0.7$ , strong positive correlation.

**Table 6** Standardized multiple regression estimates ( $\beta$ ) and  $p$ -values for microbial taxa significantly associated with each SCFA

	Acetate	Butyrate	Propionate	Valerate	Total SCFAs
<i>g_Methanosphaera</i>	0.47 ( $2.7 \times 10^{-6}$ )		0.73 ( $1.5 \times 10^{-8}$ )	0.55 ( $3.1 \times 10^{-7}$ )	0.63 ( $9.8 \times 10^{-8}$ )
<i>g_Ruminococcus</i>	0.54 ( $3.2 \times 10^{-6}$ )			0.42 ( $1.6 \times 10^{-5}$ )	
<i>f_Coriobacteriaceae</i>		0.45 ( $7.1 \times 10^{-3}$ )			
<i>g_Prevotella</i>		0.33 ( $9.8 \times 10^{-3}$ )			
Adjusted $R^2$	0.57 ( $4.2 \times 10^{-10}$ )	0.46 ( $1.5 \times 10^{-7}$ )	0.50 ( $2.0 \times 10^{-8}$ )	0.57 ( $3.0 \times 10^{-10}$ )	0.46 ( $1.4 \times 10^{-7}$ )

Values are presented as standardized regression coefficients ( $\beta$ ) with their respective  $p$ -values in parentheses.

not of differences in the dietary concentration of dietary fiber but of the type of fiber – non-fermentable *versus* fermentable.

It has sometimes been assumed that feeding highly fermentable fibers would reduce colon cancer risk, largely based on the belief that short chain fatty acids, particularly butyrate, produced by large intestinal fermentation, are protective against colon cancer. However, feeding isolated pectin to carcinogen-treated rats has resulted in either no difference<sup>39–43</sup> or increases<sup>44–47</sup> in tumorigenesis compared to the cellulose control, with only two studies reporting reduced tumor number.<sup>48,49</sup> Feeding of other types of highly fermentable fiber such guar gum resulted in higher tumor number in carcinogen-treated rats<sup>50</sup> as well. Finally, in carcinogen-treated mice, feeding inulin, a highly fermentable fiber, resulted in greatly increased tumor incidence compared with cellulose feeding.<sup>51</sup> Thus, in most studies, feeding a highly fermentable purified fiber did not decrease tumor development and often increased it. Our finding that prune, a rich source of fermentable materials, in the form of pectin and sorbitol, did not reduce tumor number is consistent with these studies.

Prunes are a rich source of phenolic compounds, primarily chlorogenic and neochlorogenic acids.<sup>52</sup> Certain phenolic compounds, such as ferulic acid<sup>53</sup> and curcumin,<sup>54</sup> reduce tumor incidence in carcinogen-treated rats. Although no reduction in tumor incidence has been found with feeding chlorogenic acid,<sup>55,56</sup> reductions in tumor multiplicity<sup>55</sup> and in colonic precancerous lesions (aberrant crypt foci, ACF)<sup>57</sup> have been reported. Chlorogenic acid upregulates the antioxidant enzyme glutathione S-transferase and suppresses ROS-mediated NF $\kappa$ B, AP-1 and MAPK activation,<sup>58</sup> suggesting a

molecular mechanism for chemoprevention. In our study, however, no increase in NF $\kappa$ B was noted with prune feeding. It may be that the quantity of chlorogenic acid consumed by rats fed prune was too little to reduce carcinogenesis.

A difference in phenolic acid content may explain the difference observed between two previous studies examining prune- or plum-beverage on colonic precancerous lesions in rats. Banerjee *et al.*<sup>12</sup> reported a reduction in ACF in carcinogen-treated rats administered plum beverage, whereas Yang *et al.*<sup>13</sup> found no reduction in ACF with prune powder-containing diets. The two prune treatments differed substantially in their phenolic acid content. The concentration of neochlorogenic and chlorogenic acids in the plum beverage used by Banerjee *et al.* was approximately ten times higher than those in the prune powder used by Yang *et al.* In the latter study, prunes were ground and dried at 120–148 °C to produce a powder. Such a heat treatment would have oxidized much of the phenolic compounds present, as has been shown for red grape pomace dried at temperatures between 100 and 140 °C.<sup>59</sup> In addition, the plum beverage and prune powder differed in their pectin contents. Although neither study disclosed exact pectin concentrations, treatment of the plum beverage with pectinase by Banerjee *et al.* would have markedly reduced its pectin content, whereas the prune powder used by Yang *et al.* retained the full complement of pectin. Thus, the divergent results could be due to the difference in pectin or phenolic acid content of the prune products used in the two studies.

High butyrate concentrations have long been suggested to reduce colon cancer risk, based in part on cell culture studies demonstrating its ability to inhibit histone deacetylase



activity.<sup>60,61</sup> We demonstrated that feeding prune greatly increased total cecal SCFAs and increased butyrate amount linearly with the dietary concentration of prune (total SCFA and butyrate,  $R^2 = 0.44$  and  $0.73$ , respectively). Our results are consistent with earlier studies demonstrating that fermentable fibers significantly increase fecal butyrate and propionate levels.<sup>62,63</sup> As discussed above, these diets are generally not effective in reducing tumorigenesis. It is possible that the failure of high large intestinal butyrate to reduce colon cancer risk in *in vivo* studies could be due to experimental design differences among studies. However, it may be that butyrate produced by fermentation in the large intestine is not chemopreventive. Whether this might be due to insufficient butyrate production, the location in the large intestine where it is produced, or simply that butyrate is not chemopreventive is unclear. Furthermore, animal studies examining the chemopreventive effect of administered butyrate have not clarified the situation, as they are conflicting. For example, oral administration of butyrate to carcinogen-treated rats increased fecal butyrate levels but did not reduce ACF or colonic tumors.<sup>64,65</sup> In contrast, Tian *et al.* reported that oral administration of a SCFA mixture significantly increased cecal SCFAs and reduced colonic tumors in mice given DSS.<sup>66</sup> The cecal butyrate concentration ( $26 \mu\text{mol g}^{-1}$ ) achieved through oral SCFA administration was substantially higher than the concentration observed in our 10% prune diet group ( $14 \mu\text{mol g}^{-1}$ ). It may be that a threshold concentration of butyrate must be achieved to exert protective effects in the colon, a threshold that is difficult to achieve by dietary means. The concept of a threshold is supported by the finding that low doses of butyrate (0.5 and 1.0 mM) stimulated growth of HCT116 colon carcinoma cells, while higher doses (2.0 and 5.0 mM) had an inhibitory effect on growth.<sup>61</sup> Thus, whether increases in large intestinal butyrate produced through dietary means can be chemopreventive remains an open question.

Although pectin feeding has rarely been found to reduce large intestinal tumors, it has been shown to decrease small intestinal tumors in DMH-treated rats in two previous studies.<sup>39,67</sup> This observation is consistent with our finding of a trend toward fewer small intestinal tumors in both prune diet groups. Prune pectin is composed primarily of rhamnogalacturonans and type I arabinogalactans. Oral administration of rhamnogalacturonans has been reported to significantly increase the expression of MUC1 and the number of mucin-producing goblet cells in the colons of DSS-treated mice.<sup>68</sup> Increased MUC1 expression attenuated inflammatory responses by dampening NF- $\kappa$ B activation.<sup>69</sup> Thus, the role of pectin in tumorigenesis appears to be complex. It has shown potential colon tumor-promoting effects, yet it has demonstrated protective effects against small intestinal tumor development. Understanding the mechanisms underlying these contrasting effects and identifying the physiological conditions under which pectin exerts protective *versus* promotive influences on intestinal tumorigenesis would be most helpful in clarifying the role of fermentable dietary fiber in carcinogenesis.

Prunes contain fiber, polyphenols, and sorbitol, all of which are known to be fermentable in the large intestine. Empty cecum weight, a well-established marker of fermentation,<sup>70</sup> was significantly increased in both prune-supplemented diets compared to the fiber (cellulose)-matched basal diet, confirming the greater degree of fermentation with these diets. A higher cecal content mass accompanied this greater cecal fermentation. However, daily fecal weight did not differ among the groups, indicating that the non-fermentable cellulose fiber was primarily responsible for influencing fecal mass.

Another noteworthy finding was the decrease in fat pad weight in animals fed 5% prune, and a trend towards a decrease in the 10% prune group. Since fat pad weight is highly correlated with whole body adiposity,<sup>71</sup> this indicates that long-term feeding of prune reduces body fat. This is most likely due to the small intestinal viscosity produced by the pectin in the prune, as viscosity appears to be the characteristic of dietary fiber responsible for reduction in body fat.<sup>72</sup>

Analysis of the large intestinal microbiome indicated that prune feeding promotes greater fermentation and bacterial richness relative to the basal diet. At the genus level, prune feeding led to increases in *Prevotella*, *Paraprevotella*, *Collinsella*, *Blautia*, and *Coprococcus*. Simpson *et al.* reported that greater abundance of *Blautia* was observed in the  $50 \text{ g d}^{-1}$  prune-fed group rather than the  $100 \text{ g d}^{-1}$  group in post-menopausal women. They also reported an inverse correlation between *Blautia* and total urinary phenolics, suggesting *Blautia* may be involved in metabolism of polyphenols.<sup>17</sup>

*g\_Blautia*, *g\_Prevotella*, *g\_Paraprevotella*, and *f\_Coriobacteriaceae* abundance was positively correlated with cecal butyrate amounts. While *g\_Prevotella* and *g\_Blautia* are not classified as primary butyrate producers, they may contribute indirectly to butyrate production through cross-feeding mechanisms, by providing precursor substrates such as succinate or acetate.<sup>73,74</sup> In addition, *g\_Prevotella* and *g\_Paraprevotella* are known to ferment dietary glycans and plant-derived polysaccharides, which are abundant in prunes.<sup>75</sup> *f\_Coriobacteriaceae* have also been identified as butyrate producers, based on the presence of the gene for butyrate kinase (*buk*), a marker gene for butyrate production.<sup>76</sup> In mice fed either a carbohydrate fraction or a polyphenol fraction from prunes, only the carbohydrate fraction increased the abundance of *f\_Coriobacteriaceae*.<sup>77</sup> These findings suggest that prune feeding enhances butyrate production through fiber fermentation, not polyphenol metabolism.

We found that prune feeding reduces the abundance of the probiotic genus *Bifidobacterium*. A similar reduction of *Bifidobacterium* was observed in mice fed a 25% prune diet.<sup>77</sup> However, findings from human studies are inconsistent. Lever *et al.*<sup>19</sup> reported a trend towards an increase in *Bifidobacteria* abundance in humans fed prunes for 9 weeks, whereas Simpson *et al.*<sup>17,18</sup> found no difference in abundance in humans consuming prunes for 12 months.

A novel finding from our microbiome analysis was the greater abundance of *g\_Methanosphaera* in the prune diets



relative to the basal diet. This abundance was positively correlated with all SCFAs, showing the strongest correlation with butyrate ( $R = 0.53$ ). *Methanosphaera* is an archaeal genus characterized by its ability to utilize hydrogen to reduce various organic compounds to methane.<sup>78</sup> Methane's oxidation byproduct, formaldehyde, is considered a genotoxic compound. The relationship between methanogens and colon cancer risk has become an area of growing interest. For instance, one study reported a greater abundance of methanogens in colon biopsy samples from CRC patients.<sup>79</sup> Similarly, a multi-cohort study found increasing abundance of Methanobacteriota in fecal samples from healthy controls to patients with adenomas to patients with CRC.<sup>80</sup> The role of diet, particularly a diet high in fermentable constituents, in influencing abundance of methanogens in the large intestine, and its relation to chemoprevention would seem to warrant further study.

Epidemiological studies report both elevated primary and secondary bile acid concentrations in colon cancer patients, particularly cholic acid, deoxycholic acid, and lithocholic acid.<sup>81–83</sup> Tinker *et al.* reported that in humans consuming a prune supplement for 4 weeks, total fecal bile acid concentration showed a trend toward reduction, with a significant reduction in secondary bile acids, relative to a grape juice control group.<sup>84</sup> Similarly, Yang *et al.* found that rats fed a 5% prune powder diet had significantly reduced total fecal bile acid concentrations compared to both basal and fiber-matched control groups, suggesting that the type of fiber in prunes contributes to this effect.<sup>13</sup> In the present study, fecal bile acid concentration did not differ among the diets. Since the fecal weight did not differ among the diets, no doubt due to non-fermentable cellulose being the major dietary fiber in all diets, this suggests that prune did not alter the bile acid pool or bile acid reabsorption from the intestinal tract.

Dietary fiber and undigested protein represent the major macronutrients entering the large intestine. Fermentation of fiber (saccharolytic fermentation) produces metabolites considered neutral or beneficial, such as short chain fatty acids and fermentation gases, whereas protein fermentation (proteolytic fermentation) produces metabolites that may be detrimental, such as branch chain fatty acids and derivatives of aromatic amino acids, such as phenols and indoles. For example, proteolytic byproducts such as *p*-cresol and phenol are often genotoxic.<sup>85</sup> However, in some cases, proteolytic fermentation products may be beneficial, as indole enhances epithelial barrier function, decreases pathogen colonization, and attenuates NF- $\kappa$ B activation in HCT-8 cells.<sup>86</sup> Since increasing saccharolytic fermentation has been found to reduce the products of proteolytic fermentation,<sup>87</sup> we determined the fecal excretion of *p*-cresol and indole. Neither prune-containing diet reduced these products of protein fermentation, suggesting that prune supplementation did not reduce proteolytic fermentation.

Colonic inflammation is well established to increase colon cancer risk.<sup>3</sup> NF- $\kappa$ B, a pro-inflammatory transcription factor regulating the synthesis of inflammatory cytokines, was determined to assess colon tissue inflammation. Banerjee *et al.* previously reported reduced colonic mucosal NF- $\kappa$ B levels in rats

fed a plum beverage.<sup>12</sup> However, prune feeding in the present study had no effect on NF- $\kappa$ B. The unchanged NF- $\kappa$ B levels observed here may be due to the absence of any dietary component that would induce intestinal inflammation (*e.g.*, high fat or high sugar), resulting in a very low baseline level of colonic inflammation. Furthermore, we observed no reduction in serum calprotectin, an established marker of inflammation. Whether prune consumption can ameliorate intestinal inflammation should therefore be examined under conditions where some degree of inflammatory stress is induced.

Finally, we examined  $\beta$ -catenin, a key regulator in the Wnt signaling pathway that controls cell proliferation and differentiation. Activation of  $\beta$ -catenin, the central regulator of the canonical Wnt pathway, results in enhanced transcription of factors required for normal intestinal function. However, hyperactivation of Wnt/ $\beta$ -catenin signaling is associated with progression to colon cancer. Previous studies have shown that fruit-derived polyphenols, such as stilbenes, pterostilbene and resveratrol, can reduce  $\beta$ -catenin expression.<sup>88</sup> However, in our study, no significant differences in  $\beta$ -catenin protein expression were observed among dietary groups, suggesting that prune consumption did not modulate the Wnt/ $\beta$ -catenin signaling pathway.

## 5. Conclusion

Our study found that prune consumption does not significantly induce or reduce tumor formation in the colon, although a trend towards reducing small intestine tumors was observed. Prune consumption, however, induced significant shifts in the gut microbiome, particularly enriching bacteria from families such as Prevotellaceae and Lachnospiraceae, while reducing the abundance of potentially pathogenic species such as *C. perfringens*. The strong correlation observed between cecal butyrate levels and specific bacterial genera highlights the butyrogenic effect of prune consumption. However, our findings raise the question of whether the amount of SCFAs, particularly butyrate, that can be produced from dietary sources containing fermentable components at levels achievable in the human diet would be sufficient to reduce colon cancer risk. Therefore, understanding whether there is a threshold concentration of butyrate necessary for colon cancer prevention will be crucial for developing effective dietary strategies. Moreover, future studies should evaluate the potential chemoprotective effects of prunes under conditions of pre-existing intestinal inflammation, as it may be that a chemopreventive effect of prunes will only become apparent in the presence of an inflamed colon. This information will be valuable for developing therapeutic strategies for patients with pre-existing colonic inflammation.

## Author contributions

Conceptualization: D. D. G. and A. G.; experiments and formal analysis: M. J. and D. M. S.; methodology and data curation:



M. J., A. G., and D. D. G.; funding acquisition: D. D. G. and A. G.; draft preparation: M. J. and A. G.; finalizing the manuscript: D. D. G.; project administration and supervision: D. D. G. All authors have read and agreed to the published version of the manuscript.

## Conflicts of interest

D. D. G. is a member of the Nutrition Advisory Panel of the California Prune Board. No other authors have a conflict of interest.

## Abbreviations

ACF	Aberrant crypt foci
AIN-93G	American Institute of Nutrition-93G
ASV	Amplicon sequence variant
CRC	Colorectal cancer
DSS	Dextran sulfate sodium
IndVal	Indicator value
NF- $\kappa$ B	Nuclear factor kappa B
SCFA	Short chain fatty acid

## Data availability

Microbiome-related data files may be accessed at <https://hdl.handle.net/11299/166578>.

Supplementary information (SI) describing the composition of the dried prune product, as well as the composition of dried prunes, prune juice, and fresh plums, is available. See DOI: <https://doi.org/10.1039/d5fo03398e>.

## Acknowledgements

This work was supported by funding from the California Prune Board to D. D. G. and A. G.

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