Article

Protective Effect of Salvianolic Acid B in Acetic Acid-Induced Experimental Colitis in a Mouse Model

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Abstract: In its prominent experimental studies salvianolic acid B (Sal B) is novel because of its well-defined, common physiological effects, which include anti-inflammatory, anti-depressant, cardioprotective, DNA protective, neuroprotective and hepatoprotective activity in experimental animals. Initially, Sal B was studied for its anti-inflammatory properties, used as a remedy for a wide range of disease conditions, but its specific efficacy on inflammatory bowel disease is still unclear. The aim of this current study was to understand the therapeutic potential of Sal B in an acetic acid (AA)—triggered experimental mouse colitis model. Colitis was triggered by intrarectal injection of 5% AA, and then laboratory animals were given Sal B (10, 20 and 40 µg/kg) for seven days. The ulcerated colonic mucosa was assessed by clinical experiment, macroscopical, biological and histopathological analysis. The results showed depleted SOD, CAT, GSH levels and consequential elevated MPO and MDA levels and aberrant crypt foci and mast cells were seen in the AA-induced colonic mucosa of experimental animals. The data obtained from this study demonstrate that a dose of 40 µg/kg showed an efficacious anti-ulcer effect against AA-induced experimental colitis. Based on its antioxidant efficacy, Sal B may therefore be useful as a therapeutic approach for ulcerative colitis.

Keywords: ulcerative colitis; salvianolic acid B; acetic acid; myeloperoxidase; ACF; mast cells

1. Introduction

Ulcerative colitis (UC) is a widespread significant complication of irritable bowel syndrome and chronic gastrointestinal tract immunological disorder that causes inexorable...
ulceration, tenesmus, diarrhea, bloody stools, inflammation in the intestine and colon and mucosal tissue damage [1–3]. Inflammatory bowel disease (IBD) affected about 2.5 million people in the United States in 2020. By 2030, it is expected that the number of people living with IBD in the United States will have increased to about 3.5 million [4,5]. Ulcerative colitis-related malignancy has some characteristics, including abdominal pain, anemia, diarrhea, slightly raised plaques, and villous adenomas that become malignant [6,7]. However, the etiology of IBD is not clearly understood. The current therapeutic approaches are known for being ineffective at both inducing injury recovery and stopping it from continuing to spread. The inflamed mucosa produces reactive oxygen species (ROS), which may play a role in the observed increased risk of colorectal cancer [8]. The body’s antioxidant defense mechanisms protect tissues from ROS under normal conditions. The over-expression of free radicals acts as a severe contributor to the pathological events of ulcerative colitis [9]. Under physiological circumstances, a balanced condition between free radicals and antioxidants is maintained, which is disrupted during pathological conditions [10,11]. Excess levels of ROS lead to oxidative stress and, thus damage to DNA due to variations among innate exogenous antioxidants. Sal B (Figure 1) is a bioactive phenolic compound extracted from Salvia miltiorrhiza roots; these compounds mostly have antioxidant and free radical scavenging activity [12]. Sal B’s biological properties were documented to include anti-inflammatory, cardioprotective [13], neuroprotective [14], osteoporosis protective [15] and hepatoprotective effects [16]. Our current research aimed to assess the therapeutic potential of Sal B against AA-induced experimental colitis in a mouse model.

Figure 1. The molecular structure of Sal B.

2. Materials and Methods

2.1. Chemicals

Salvianolic acid B was obtained from Sigma-Aldrich Chemical Company (St. Louis, MO, USA). Acetic acid, ketamine, phosphate buffer saline, hydrogen peroxide, ethylene diamine tetraacetic acid (EDTA), nitroblue tetrazolium (NBT), ethanol (EtOH), hexadecyltrimethylammonium bromide (CTAB), methylene blue and all other chemicals were obtained from Sigma-Aldrich Chemical Company (Bangalore, India) and of analytical or reagent grade.

2.2. Animals and Ethical Approval

Male Swiss Albino mice weighed 20–25 g approximately; (8–9 weeks old) were purchased from Sri Venkateshwara Enterprises (Bangalore, India). Animals were maintained in a sterile environment (25°C, 12 hr daylight/dark period). Mice had free access to feed pellets and were able to drink water whenever they desired. The Ethics Commission of Periyar University approved this study (approval number: 1085/ac/07/CPCSEA/PUAE/EC/March-2014/06).
2.3. Induction of Experimental Colitis in Mice

The acute UC model was induced by the administration of 5% AA [17]. Following 24 h fasting, animals were gently anesthetized with ketamine and xylazine (80/10 mg/kg) intraperitoneally. A 3.5 F catheter was subsequently inserted into the stomach with the tip placed 4 cm from the rectum. One mL of AA (5% v/v) with 0.9% saline was used to induce colitis in the colon’s lumen along with a sustained supine Trendelenburg position for 30 s to avoid any outflow of the intra-colonic AA dose. Control mice in this study were treated by a similar protocol but given only 0.9% saline.

2.4. Animals

Animals were randomly divided into five groups (n = 6): (1) Control (0.9% saline); (2) AA-treated mice (50 µL/kg body weight per day); (3) AA + Sal B 10 µg/kg; (4) AA + Sal B 20 µg/kg; (5) AA + Sal B 40 µg/kg body weight per day (Figure 2).

![Experimental design](image)

**Figure 2.** Experimental design. Mice in the AA and Sal B group received an intrarectal dose of 1 mL of 5% AA, whereas the normal control group animals were given 1 mL of sterile saline. On day 8 blood and colonic tissues were collected.

2.5. Macroscopic Damage Score

The colon was dissected, opened longitudinally as well as the injury score was determined by specific characteristics previously described [18]:

- 0 = No damage
- 1 = localized hyperemia but no ulcers
- 2 = linear ulcers with no significant inflammation
- 3 = linear ulcers with inflammation at one site
- 4 = two or more sites of ulceration & inflammation
- 5 = two or more sites of ulceration with major inflammation >1 cm along the length of the colon.

For calculation of values for morphological scores, individual scores given by the predicted values were averaged and the mean of the scores were determined and are given as mean ± SEM.

2.6. Histopathological Evaluation of Colitis

For histopathological examination, the colon was separated from the proximal rectum, removed from adherent tissues, and washed in PBS. The colonic tissues were fixed in 10% formalin, dehydrated with a sequential increase of EtOH concentration, fixed within paraffin, and then sectioned. Sections were fixed on slides, cleaned, hydrated, and stained with hematoxylin at a thickness of 5 µm. The histopathological scoring was evaluated as per the earlier reported method [19], as follows: (0) no abnormality; (i) mild but unequivocal increase; (ii) squatty leukocyte percolation; (iii) neutrophils in the epithelium, hardening of the colon parietes, (iv) excess of neutrophils in part of the crypt, lack of goblet cells,
elevated vascular density, unequivocal erosion ulcer or granulation tissue and hardening of the colon parietes.

2.7. Biochemical Measurements

The animals were sacrificed by cervical dislocation under anesthesia ketamine/xyazine (80/10 mg/kg) administered intraperitoneally, and the distal colon sections were collected, stored at −80 °C until further examination. 1 to 2 g of distal colon tissues was collected for further investigation. The colon sections were washed, cut into smaller pieces, and homogenized in 10 mM of ice-cold Tris-HCl buffer (pH 7.1) to obtain tissue homogenate. The samples were centrifuged at 4000 rpm for 15 min at 4 °C in a cooling centrifuge (REMI micro refrigerated centrifuge, Chennai, India), and the colonic tissue homogenate was utilized for biochemical measurements [20,21].

2.8. Measurement of CAT Activity in Colon Tissues

Catalase (CAT) activity in intestinal colonic tissue supernatants was measured using a spectrophotometric technique developed by Aebi [22]. 20 µL of the homogenate was taken and mixed with 2 mL of PBS (50 mM, pH 7.0) including 10 mM H_2O_2. Catalase activity is defined as the quantity of enzyme needed to degrade 1 nmol of H_2O_2 per minute at 25 °C and pH 7.0. CAT activity was measured by using a UV-Visible spectrophotometer (Shimadzu UV 1800, Torrance, CA, USA) at 240 nm. CAT activity was expressed as U/mg protein.

2.9. Measurement of SOD Activity in Colon Tissues

The hyperactivity of SOD was assessed by quantifying its ability to restrain the photochemical reduction of nitro blue tetrazolium (NBT). Superoxide dismutase activity was measured in colonic tissues following the procedures given in [23] with small modifications. The photochemical diminution of riboflavin produces O_2− that deoxidizes the NBT to generate formazan salt, which is measured at an optical density of 560 nm. The conversion of superoxide radical (O_2−) to peroxide inhibits the reduction of NBT. In a dark room, 20 µL of supernatant, 0.5 mL of 100 mM PBS, 0.2 mL of 200 mM EDTA, and 0.1 mL of 15 mM L-methionine, 200 µL of NBT, 250 µL of riboflavin were added and this sample was exposed to a fluorescent lamp (15 W) for 15 min. The SOD activity was measured by using a UV-visible spectrophotometer (Shimadzu UV 1800) at 560 nm. SOD activity was expressed as U/mg protein.

2.10. Measurement of GSH Activity in Colon Tissues

The reduced GSH concentration was measured in colon tissues [24]. 500 µL tissue homogenate was added to 400 µL ice-cold H_2O and 100 µL of 60% trichloroacetic acid. All test tubes were centrifuged at 4000 rpm at 4 °C for 15 min. The optical density of the supernatant was observed by using a UV-Visible spectrophotometer at 412 nm. For quantitative assessment of GSH (10–100) in nmoles/mg protein, a standardized graph was prepared.

2.11. Measurement of LPO Activity in Colon Tissues

Malondialdehyde (MDA) production was detected as TBARS and evaluated using a previously reported method [25]. In brief, 200 µL of supernatant was added to 200 µL of 15% trichloroacetic acid; this reaction solution was kept in an ice bath for 10 min and centrifuged at 4000 rpm for 10 min. 200 µL of supernatant was mixed in 200 µL of thiobarbituric acid. This reaction solution was kept in a boiling water bath for 5 min and instantly kept in an ice bath for 5 min. The optical density of the supernatant was observed by using a UV-Visible spectrophotometer at 532 nm. The amount of MDA was shown as nmoles of MDA/mg of protein.
2.12. Measurement of MPO Activity in Colon Tissues

Myeloperoxidase hyperactivity was estimated in colonic tissues [26]. In brief, the supernatant was added to a solution containing 0.5% CTAB dissolved in 50 mM PBS (pH 6.5) further, incubated for 10 s in a freezing bath. The supernatant was thawed twice and then centrifuged at 5000 rpm for 10 min. 0.2 mL of the supernatant was added to 2.8 mL of 50 mM PBS containing o-dianisidine dihydrochloride and H₂O₂ (0.005%), and the color developed was measured at 460 nm by using a UV-Visible spectrophotometer. The myeloperoxidase hyperactivity was calculated according to the following formula:

\[
\text{MPO activity (mU/mg)} = 1000 \times \frac{X}{\text{weight of the colon tissue taken (mg)}}
\]

where \(X = 10 \times \) difference in absorption per minute/volume of tissue homogenate taken in the final reaction.

2.13. Determination of Aberrant Crypt Foci (ACF)

The ACF count was measured using a previously reported technique [27]. The intact colon was excised longitudinally and washed thoroughly in a saline solution prior to fixation process. Then the colon tissue was fixed in 5% buffered formaldehyde. To detect the ACF, the colon was stained with 0.22% methylene blue for 3–4 min. Mucosal ACF was enumerated by using a light microscope (40×). The aberrant crypts were spotted from normal crypts by its enlarged size, the clogged and deeply stained perceptive zones.

2.14. Mast Cell Staining on Colon of Experimental Animals

Histopathological examination of mast cells was examined in colon tissues [28]. The 5 µm thick colon tissues were de-waxed in xylene and re-hydrated by decreasing EtOH concentration in d.H₂O (distilled water). These sections were then stained with toluidine blue for 3 min, washed with double d.H₂O, dehydrated in enhancing concentrations of alcohol from beginning to end series, and then fixed with dibutyl phthalate in xylene (DPX). A high-power objective field (40×) was selected for counting mast cells in 10 alternative fields/slides.

2.15. Statistical Analysis

All data were presented as mean ± standard deviation. Statistical analysis of the data was performed by one-way analysis of variance (ANOVA) followed by Tukey’s multiple range tests for post-hoc analysis using the Sigmastat Version 3.5 software (Systat Software Inc., San Jose, CA, USA). The significance level was set at \(p < 0.05\).

3. Results

3.1. Effect of Sal B on AA-Induced Colitis

By comparing the colitis group to the untreated control group, physiological investigations revealed significant changes in colon length, body weight, fecal blood, and general health. Compared with group 1 (Control), the % of body weight drastically decreased in the AA-induced colitis group 2 mice. The minimal disease activity state was elevated in the AA-induced colitis group. However, the supplementation of Sal B ameliorated the body weight, and statistically, major differences were obtained starting on day 5. The length of the colon is considered being inversely proportional to the severity of AA-induced experimental colitis. The consequential shrinking of the colon length (Figure 3A) and macroscopic score (Figure 3B) was observed in the AA-induced mice, along with marked infiltration of inflammatory cells, extensive eradication of mucosal layer, loss of crypts, and submucosal edema, and these morphological changes suppressed following treatment with 40 µg/kg of Sal B (Figure 3A–C).
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Figure 3. Therapeutic effect of Sal B on the body weight changes of mice with colitis. (A) The mice were given only saline (a. control group), saline containing 5% AA (b. colitis group 2), AA + 10 μg/kg Sal B (c. group 3), AA + 20 μg/kg Sal B (d. group 4), AA + 40 μg/kg Sal B (e. group 5), (A) Representative morphological changes of colon tissues. In each group, colon samples were collected from mice. Interestingly, Sal B decreased the degree of ulcer index and colon shortening in a dose-dependent manner. (B) Macroscopic assessment of colons from AA-induced experimental colitis. (C) Effect of daily treatment with Sal B changes in body weight in experimental colitis. AA-acetic acid; Sal B-Salvianolic acid B, * p <0.05, ** p < 0.01, *** p < 0.001 as compared with control.

3.2. Sal B Decreases Microscopic Colon Damage in AA-Induced Experimental Colitis

The histopathological character of the colonic tissue was evaluated by hematoxylin and eosin (H&E) staining. Typical histopathological results are shown in Figure 4. The control mice showed clear colonic mucosa without edema, erosion, or ulceration (Figure 4A). The anatomy of the colon wall in the colitis model mice (group 2) had a characteristic morphology with crypts, edema, thickness and erosion of the colonic mucosa, numerous goblet cells, and a minimal number of lamina propria mononuclear cells. Compared with group 1 (Control), microscopic histopathological scores were considerably elevated in mice from the AA-induced colitis (group 2) (Figure 4B). Mild histopathological changes occurred in group 3 and group 4 (Figure 4C,D). After the treatment of Sal B a considerable decrease in the histopathological count, intact colonic mucosa structure, milder congestion and edema and less inflammatory cell infiltration were observed in group 5 mice as compared with the control group 1 (Figure 4E).
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Figure 4. Histopathological changes in colons from mice stained with H&E. (A) Control mice receiving saline only, showing normal histological structures. (B) 5% AA-induced colon with ulcerative colitis, Group 2 showed marked neutrophil infiltration, thickening of colonic mucosa, severe destruction of epithelial structure with ulceration and infiltration of inflammatory cells compared with control (group 1). (C, D) Mice were administered with AA, Sal B (10, 20 µg/kg respectively), Group 3 and 4 showed mild histological colonic mucosa changes (C, D). Group 5 showed the intact colonic mucosa structure, milder congestion and edema and less inflammatory cell infiltration (E).

3.3. Effect of Sal B on CAT Activity in Colon Tissues

Catalase acts as an enzyme essential for the detoxification of H$_2$O$_2$ during the reaction mediated by SOD. In the current study, the catalase activity was drastically decreased in AA-induced mice colonic tissue (group 2) (Figure 5A). Catalase frames the energetic cellular defense that involved carcinogen deactivation of ecosystem factors. In this catalase, enzymatic activity was increased in the treatment of Sal B in the experimental animals. Mice treated with 10, 20 µg/kg of Sal B showed a considerable enhancement of CAT activity when compared with colitis group 2. Treatment with Sal B (40 µg/kg) significantly increased the colonic tissue CAT activity in group 5 when compared with colitis group 2 (Table 1).
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![Catalase activity](image)

**Figure 5.** Effect of Sal B on enzymatic activities in mice colon tissue with AA-induced experimental colitis (n=6). (A) Catalase activity, * p < 0.05, ** p < 0.01, *** p < 0.001 as compared with control, (B) Superoxide dismutase activity and (C) GSH level. The data are represented as mean ± SD and evaluated using one-way ANOVA, followed by Tukey’s multiple range tests for post hoc analysis. # p < 0.05, ## p < 0.01, ### p < 0.001 as compared with control.

Table 1. Effect of Sal B on colon catalase, SOD, and GSH concentration.

| Parameters/Groups         | Colon Catalase Activity (μmol H$_2$O$_2$/min/mg Protein) | Colon SOD Activity (U/mg Protein) | Colon GSH Activity (nmol/mg Protein) |
|---------------------------|----------------------------------------------------------|----------------------------------|--------------------------------------|
| Control (Saline)          | 9.15 ± 0.489                                             | 8.74 ± 0.53                      | 10.74 ± 0.72                         |
| Colitis (AA)              | 2.72 ± 0.213                                             | 2.31 ± 0.19                      | 3.36 ± 0.23                          |
| Colitis + 10 μg Sal B     | 6.47 ± 0.331                                             | 7.78 ± 0.45                      | 4.99 ± 0.39                          |
| Colitis + 20 μg Sal B     | 5.99 ± 0.361                                             | 6.61 ± 0.35                      | 5.41 ± 0.43                          |
| Colitis + 40 μg Sal B     | 7.22 ± 0.443                                             | 6.78 ± 0.39                      | 7.74 ± 0.56                          |

Data are expressed as mean ± SEM, n = 6. Statistical analysis was performed using One-Way ANOVA followed by Tukey-Kramer multiple comparisons test (p < 0.05).

3.4. Effect of Sal B on SOD Activity in Colon Tissues

The reduced SOD activity in the colon with inflammation could lead to a decrease in abdominal immunity versus oxidative stress, which is directly linked to colon injury in ulcerative colitis. The control group 1 mice showed significantly increased colonic SOD activity while comparing to the AA-induced colitis group 2 mice. Experimental animals treated with 10, 20 μg/kg of Sal B showed a significant increase of SOD activity when compared with colitis group 2. Table 1 represented supplementation of Sal B (40 μg/kg) ameliorates the colonic tissue SOD activity when compared with colitis group 2 mice (Figure 5B).
3.5. Effect of Sal B on GSH Activity in Colon Tissues

Glutathione is a tripeptide thiol molecule and is a reducing agent, being the foremost low molecular weight thiol found in mammals. The effectiveness of Sal B on AA-induced mice in the GSH content was found to significantly improved. In our study, Sal B has considerably reduced the level of GSH content in group 2 when compared with group 1. When compared to control mice, AA treatment resulted in a significant decrease in colonic GSH levels. Experimental animals treated with 10, 20 µg/kg of Sal B showed a moderate increase of GSH activity when compared with colitis group 2. GSH levels were considerably lower in group 5 mice following supplementation with 40 µg/kg of Sal B (Figure 5C) compared to group 2 mice (Table 1).

3.6. Effect of Sal B on MDA Levels in Colon Tissues

Sal B inhibits lipid peroxidation caused by AA-induction in mice. Compared with group 1, the AA-induced colitis group 2 mice, and leads to a significant elevation of MDA levels. MDA is a known biomarker that plays a key role in tissue damage. AA-induced colitis group 2 shows an elevated level of MDA when compared with untreated control group 1. Mice treated with 10, 20 µg/kg of Sal B indicated a mild decline of MDA level when compared with colitis group 2. When compared to AA-induced experimental colitis mice, supplementation of Sal B with higher dosages (40 µg/kg) significantly decreases MDA levels in group 5 (Figure 6A).

![Figure 6. Effect of Sal B on LPO and MPO activity in mice’s colon tissue with AA-induced experimental colitis (n = 6). (A) Effect of Sal B on lipid peroxidation * p < 0.05, ** p < 0.01, *** p < 0.001 as compared with control, (B) Effect of Sal B on MPO activity. The data are represented as mean ± SD and evaluated using one-way ANOVA, followed by Tukey’s multiple range tests for post hoc analysis. # p < 0.05, ## p < 0.01, ### p < 0.001 as compared with control.](image)

3.7. Effect of Sal B on Myeloperoxidase Activity in Colon Tissues

The inflammatory state of the colonic mucosa causes an increase in MPO enzyme expression, which leads to increased oxidative stress. In this study, colonic myeloperoxidase activity was considerably elevated in the AA-induced colitis mice when compared to group 1. Mice treated with 10, 20 µg/kg of Sal B showed a moderate decrease of myeloperoxidase activity when compared with colitis group 2. In addition, Table 2 indicates mice treated with Sal B with higher doses (40 µg/kg) in group 5 showed significantly decreased MPO activity (Figure 6B).
Table 2. Effect of Sal B on colon MDA levels, and MPO activity.

| Parameters/Groups | Colon MDA Levels (nmol/mg Protein) | Colon MPO Activity (U/mg Tissue) |
|-------------------|-----------------------------------|---------------------------------|
| Control           | 18.02 ± 1.87                      | 1.25 ± 0.14                    |
| Colitis           | 45.29 ± 2.34                      | 18.42 ± 0.98                   |
| Colitis + 10 µg Sal B | 23.34 ± 1.83                      | 6.43 ± 0.45                    |
| Colitis + 20 µg Sal B | 15.21 ± 1.24                      | 9.28 ± 0.72                    |
| Colitis + 40 µg Sal B | 10.44 ± 0.91                      | 4.58 ± 0.36                    |

Data are expressed as mean ± SEM, n = 6. Statistical analysis was performed using One-Way ANOVA followed by Tukey-Kramer multiple comparisons test (p < 0.05). Significantly different vs. normal control (p ≤ 0.05).

3.8. Methylene Blue Staining of Aberrant Crypt Foci (ACF) of Experimental Mice

The untreated colonic mucosa of colonic tissue-stained crypts was investigated with methylene blue staining to detect aberrant crypts. Their enormous size, oval form, and thick epithelial lining along with a more massively colossal per crypt zone characterized from normal crypts AA-induced colonic crypts, as shown by their gloomy staining (Figure 7A–E). In the AA-induced mice, group 2 showed a high number of ACF compared with the control group (normal crypt) (Figure 7A,B).

Figure 7. Morphological observations of aberrant crypt foci. (A) The control group showed typical crypt appearance, (B) AA-induced colitis group showed an increased number of crypts per foci, (C,D) Group 3 and 4 showed only a few crypts, (E) Group 5 showed normal crypt-like control.

There are moderate changes in colon tissues treated with 10, 20 µg/kg of Sal B were shown in groups 3 and 4. In our study confirmed, 75% reduction of ACF prevalence in mice administrated with 40 µg/kg of Sal B (group 5), while there was no ACF prevalence in control mice (group 1) as clearly presented in Figure 7E.

3.9. Mast Cell Count of Control and Experimental Mice

The accumulation of many inflammatory cells, mast cells, and eosinophils in the gut mucosa is a well-known morphologic feature of ulcerative colitis. The crypt epithelium was depleted from goblet cells and also observed the ulcer crypt. In addition, inflammatory cell infiltration in lamina propria such as lymphocytes and plasma cells was observed
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Figure 8. Photomicrographs of mast cell count. (A) The control group showed normal architecture lack of mast cells. (B) Ulcerated colonic crypts showed a considerably increased concentration of mast cells. (C, D) showed a moderate concentration of mast cells. Additionally, 40 µg/kg of Sal B treated Group 5 mice showed a significantly reduced concentration of mast cells (E).

4. Discussion

Ulcerative colitis (UC) is a form of inflammatory bowel disease defined as a long chronic inflammatory process, as well as dysbiosis. Diet is an important aspect that influences human health and has a lot of potential in the diagnosis and treatment of UC [29]. This study described that chronic inflammation is connected with an overproduction of unpaired free radical species and has a primary role in numerous pathological conditions such as melanoma, sarcoma, coronary artery diseases, myocardial infarction, obesity, and inflammatory bowel disease [30–32]. Flavonoids have been shown in vivo and in vitro to prevent cells from oxidative stress and to protect the epithelial mucosal layer of the intestines [33]. Antioxidant foods such as blueberries, artichokes, bell peppers, cherries, pecans, squashes, strawberries, red cabbage, raspberry, and tomatoes are recommended as supplementation for individuals with chronic UC and to prevent recurrence [34,35]. The
The current study evaluated the medicinal effects of Sal B treatment in chemically-induced experimental colitis in a mice model that resembles the UC state in human trials. The experimental mice model revealed typical UC indications such as diarrhea, cramps, bloody stools and mucosal ulceration, stimulated increased mucous blood flow and shortened colons \[1,36\].

Sal B, an antioxidant, was helpful in preventing and treating AA-induced colitis in mice, as well as modifying the degree and severity of colitis \[37\]. Oral administration of Sal B in AA-induced colitis mice showed that Sal B improved body weight and enhanced stool stability while reducing the rate of bloody stools, and Sal B protected AA-treated mice from mucosa erosion, colon shortening, inflammatory cell infiltration, submucosal edema, loss and disruption of crypt and villi. Significant increases in body weight, microscopic and macroscopic scores, bleeding in the rectum, histopathological modifications, and shearing of epithelial cells were seen depending on the severity of inflammation in ulcerative colitis \[38\].

The antioxidant system depends on both SOD and GSH are important components. SOD is an antioxidant enzyme that transforms superoxide radicals to H\(_2\)O\(_2\), which is then converted into molecular oxygen and H\(_2\)O by catalase. GSH is an intracellular free radical scavenger \[39,40\]. GSH and GR contribute to detoxifying environmental carcinogenic substances including reactive oxygen species and hydroxyl radicals by combining toxins with GSH and ultimately preventing tissues and organs, toward toxicity induced by carcinogens GSH and GR primarily contribute to the detoxification of environmental carcinogens \[41–43\]. In the current study, SOD and GSH content were reduced in ulcerative colitis caused by acetic acid; however, Sal B therapy was able to reverse the decrease. Sal B showed intrinsic antioxidant properties that could improve intestinal antioxidant status, which is in accordance with previous findings. Oxidative stress is caused by increased oxidant levels and/or a depleted antioxidant system, and it has been related to the development of ulcerative colitis. Lipid peroxidation develops progressively in cellular and subcellular membranes during oxidative stress, causing irreversible damage to cellular function. The lipid peroxidation product MDA was significantly elevated in AA-induced colitis, whereas Sal B administration significantly decreased MDA levels in the colon of AA-induced mice \[37\]. MPO, as a sign of inflammation, might facilitate oxidative tissue damage, whereas Sal B therapy dramatically reduced MPO activity in AA-treated colonic mice.

Mitochondria maintain cellular energy engendered through oxidative phosphorylation \[44\]. The small mitochondrial genes are indispensable for the enzyme involved in the cellular biosynthetic and oxidative function. ROS is a major cause of UC. Cellular macromolecules such as proteins, DNA, and lipids are damaged by oxidative stress. ROS are involved in the development of colitis \[45,46\]. Several researchers have suggested that intra-rectal administration of AA-generated reactive oxygen species leads to modifications in oxidative marker levels, including GSH, MDA and MPO \[47\]. The LPO level is indirectly quantified by the amount of MDA production, which leads to damage to the mitochondrial dysfunction \[47,48\]. However, administration of Sal B decreases the MDA level in experimental colitis. Sal B blocks neutrophil infiltration, increasing the production of cytokines, and oxidative burst into ulcerated tissues. In addition to immune cells, macrophages, antigen-presenting cells and intestinal epithelial cells, cytokines are considered key signals in the gastrointestinal immune system and are directly implicated in the secretion of several cytokines that control the inflammatory response in ulcerative colitis \[49–51\].

Most of the ACF was identified by investigating the sequential early development of erosions in mice colon following acetic acid induction. ACF was characterized by mildly enlarged or small crypts and collections of abnormally appearing colonic crypts that degrade from the surrounding epithelium \[52–54\]. The crypt is concise, as well as similar to a condensed luminal opening. ACF measures resemblance among the tumors regarding the apparent morphology of the colon, pathologic classification, and high crypt multiplicity. The present study results correlated with previous reports, AA-induced mice showed an
elevated number of colonic crypts. Also, the administration of Sal B significantly reduces the number of crypt formations shown in Figure 8.

Mast cells regulate important gastrointestinal activities including blood flow, epithelial cell functions, secretary functions and endothelial permeability, mucus secretion, upregulate membrane expression, GI tract motility, defensins release, neuroimmune interaction, and cancer development [55,56]. Mast cells, as well as tumor cell interactions in their biophysical environment, are essential for cell proliferation and rate of survival. Mast cells are well-known as important effector cells in the gut immune system [57]. Mast cells can be triggered by a range of stimuli, causing them to degranulate and generate a number of biologically active mediators of inflammation, tryptase, hexosaminidase, chemokines, and cytokines, among others [58,59]. An elevation in mast cell levels in the gastrointestinal tract can lead to mast cell-mediated epithelial tissue injury because many mast cell mediators are pro-inflammatory and can trigger gastrointestinal immunological disorders [55,56,60]. This study mainly focused on the induction of AA that increased inflammatory mast cells creating a microenvironment that leads to a progression of colitis.

The role of systemic inflammation in the genesis and recurrence of cancer is a key research topic in tumor cells. At various stages of UC progression and colitis-associated tumorigenesis, ROS and oxidative stress are key components in the initiation, development, and progression. Ulcerative colitis was initiated by an increase in MDA and MPO levels, as well as mitochondrial dysfunction (Figure 9). Within the intestinal mucosa, activated neutrophils can create ROS, increasing oxidative stress and playing a role in the formation of UC [61]. Excessive generation of ROS in mucosal cells during experimental colitis can trigger a series of inflammatory mechanisms that damage gastrointestinal epithelial cells, directly or indirectly, disrupting the intestinal mucosal barrier stability [62].

Figure 9. The mechanism of action of Sal B in AA-induced ulcerative colitis.

5. Conclusions

The major finding of this study is the protective effectiveness of Sal B on local as well as systemic pathological changes induced by acetic acid in colon tissue. The biolog-
ical effects of Sal B effectively decrease oxidative stress, elevated antioxidant properties. Symptoms include body weight loss, colon tissue shortening, diarrhea, and increased colon weight/length ratio, and a reduced disease activity index (DAI) score. Damage to colon tissues was reversed, and biochemical indices such as SOD, CAT, and GSH were improved. Therapeutic approaches that stabilized MPO and MDA levels suggest that Sal B may reduce the risk of chronic conditions. Finally, Sal B considerably inhibited the incidence of ACF, the alternative effect of preneoplastic ulceration and decline of mast cell count in the AA-induced experimental colitis model used. Our findings suggest that Sal B might have good prospects for future development in the clinical treatment for ulcerative colitis and cancer prevention.

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