Original article

Effect of supplementing fava bean (Vicia faba L.) on ulcerative colitis and colonic mucosal DNA content in rats fed a high-sucrose diet

Areej A. Eskandrani
Chemistry Department, College of Science, Taibah University, Medina 30002, Saudi Arabia

A R T I C L E   I N F O

Article history:
Received 14 November 2020
Revised 3 March 2021
Accepted 7 March 2021
Available online 19 March 2021

Keywords:
Acetic acid
Fava bean
Sucrose
Ulcerative colitis

A B S T R A C T

Ulcerative colitis (UC) is an inflammatory bowel disease with high morbidity. Acetic acid-induced damage of colonic mucosa in rats is a commonly used experimental animal model of UC. This research aimed to explore for the first time the ameliorative effect of dietary supplementation with fava bean on the incidence of UC in rats fed with sucrose containing diet. Rats were divided into five groups as follows: G1, control healthy rats; G2, colitic rats; G3, colitic rats fed diets containing 30% sucrose, G4, healthy rats fed diets containing 30% sucrose and G5, colitic rats fed diets containing 30% sucrose supplemented with dried ground fava bean. Colonic injury and inflammation were evaluated through a disturbance of oxidative biomarkers, a significant increase in inflammatory biomarkers and inflammatory cytokines, and histological abnormalities in colonic tissues accompanied by colonic mucosal DNA damage. Colitic rats fed on sucrose containing diet demonstrated additional histological, biochemical, and DNA alterations in colonic mucosa of rats. Dietary supplementation with dried ground fava bean significantly corrected the impaired oxidative and inflammatory biomarker levels and modulated histological features and DNA alterations. Finally, fava bean attenuated the oxidative damage and colonic injury induced by acetic acid, which confirmed its high anti-oxidant and anti-inflammatory properties.

© 2021 The Author(s). Published by Elsevier B.V. on behalf of King Saud University. This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/by-nc-nd/4.0/).

1. Introduction

Inflammatory intestinal disease is a chronic gastrointestinal tract disease (Rohr et al., 2018). It has two main profiles: epithelial ulceration and ulcerative colitis (UC). Various etiological factors may predispose to UC, such as genetic, environmental and immunological factors (Jiao et al., 2018). It affects the quality of a patient’s life and increases the incidence of colon cancer (Abraham et al., 2017).

UC only includes the rectum and colon. Although its aetiology is not fully understood, UC has been linked to lowered antioxidant capacity and raised production of free radicals, which are responsible for lipid peroxidation (LPO) and may lessen the ability of intracellular anti-oxidants, thus ultimately leading to prominent colonic inflammation. In numerous studies, UC pathogenesis indicated an important relationship between excessive inflammation and oxidative stress. Improving LPO and free scavenging of radicals afford a valuable, protective, and therapeutic treatment for UC (Scioli et al., 2014). Various risk factors are associated with UC, including long duration of the disease, low energy activity, complicated primary sclerosing cholangitis or stenotic disease and insufficient pharmacological therapy (Abdin, 2013).

Approximately 50% of patients with UC undergo removal of part of their colons because of failure of conservative medicinal treatments. In addition, these patients are not sufficiently strong to maintain long-term remission (Santos et al., 2020). However, treatments are linked to serious adverse side effects, such as abdominal cramps, diarrhoea, fever and increased blood pressure. New therapeutic selections with lower toxicity and slight side effects need to be developed (Nguyen et al., 2020).

Colonic inflammation has been widely used in numerous chemically induced animal models to elucidate its primary pathophysiological mechanisms and to assess the potential of chemicals that induce UC, such as acetic acid (AA) (Dothel et al., 2013). Colitis induced by AA is a common model of experimental colitis used to examine inflammatory bowel disease. This model is generated via intra-rectal injection of AA, to induce inflammation and ulcer-
ation in the rat rectum and colon. Oxidative destruction in this situation is thought to be a pathogenic factor, and this damage is also observed in the colon's gross morphology (Cagin et al., 2016).

It has been demonstrated that dietary factors are responsible for the pathogenesis of different conditions, including colon cancer and celiac disease; thus, the diet presumably involves in etiology of inflammatory bowel disease (Costello et al., 2020). Diets rich in fibre, such as those rich in cereals, fruits and vegetables have a positive effect on health, due to their effect on reducing the incidence of different types of diseases as it increase the volume of fecal bulk, reducing the time of intestinal transit, cholesterol and glycaemic levels, also fibres was able to trap harmful substances (mutagenic and carcinogenic agents), stimulating the proliferation of the intestinal flora. (Dhingra et al., 2012).

Legumes are an important source of nutrients and occupy an important part in traditional diets in many countries. In addition, they can be eaten in processed or raw forms (Singh et al., 2013). They are rich in dietary fibre, protein, starch, fatty acids, vitamins, trace minerals and phenolic anti-oxidants (Alcázar-Valle et al., 2020).

The fava bean (Vicia faba L.) FB, commonly referred as broad bean, belongs to the Fabaceae family. In the Mediterranean, it is considered a chief winter crop, whereas in other regions of Europe and South America it is considered a spring crop. Fava beans, belong to the Fabaceae family. In the Mediterranean, it is considered a chief winter crop, whereas in other regions of Europe and South America it is considered a spring crop. Fava beans consist of 12% fibre, 22.4%–36% protein, 1.2%–4% lipids and 57.8%–61% carbohydrates (Bouchenak and Lamri-Senhadji, 2013). They are considered an appropriate food for diabetic patients and can protect against heart disease and reduce blood glucose levels (Lonnie et al., 2020). Moreover, they are rich in I-3.4 dehydroxyphenylalanine (I-DOPA), which is a precursor to neurotransmitter, catecholamine (Ramya and Thaakur, 2007). This study aimed to examine for the first time the effect of dietary supplementation of fava beans in experimental UC rats fed on high-sucrose diet.

2. Materials and methods

2.1. Materials

Fava beans were purchased from local markets in Jeddah, Makkah Province, washed, let to dry, and then ground to a fine powder before use.

2.2. Induction of ulcerative colitis

Colitis was induced in colon tissues via an intracolonical single administration of 4% AA (2 mL) through a soft paediatric lubricated catheter under a low dose of ether anaesthesia. Rats were kept in a horizontal position after AA administration for 2 min, to prevent the draining of AA. The same procedure was performed in control animals using equal volumes of saline solution; treatments were started after colitis induction (Shanmugam et al., 2020).

2.3. Experimental design

Fifty male albino rats weighing 150–175 g were housed at the King Fahd Medical Research Center Animal Facility Breeding Colony were allowed free access to basal diet (the basal diet was prepared according to AOAC (2000)) containing 10% corn seed oil, 10% casein, 70% corn starch; 4% salt mixture, 5% cellulose, 1% vitamin mixture, 1%, and water ad libitum. The Ethical Committee of King Fahd Medical Research Center, Jeddah, KSA. approved the study.

Rats were randomly divided into five groups (10 animals/group) as detailed below. Group 1: a healthy control that received normal saline (2 mL) once intrarectally and was fed the basal diet for 21 days; Group 2: a colitic group that received the AA intraperitoneally as a single administration and was fed the basal diet for 21 days; Group 3: colitic rats fed the basal diet supplemented with 30% sucrose (sucrose was replaced with corn starch, to keep the concentrations of other nutrients and carbohydrates constant) for 21 days; Group 4: healthy rats that were fed the basal diet supplemented with 30% sucrose, with dietary administration of dried ground fava seeds as a 252 g/kg diet according to León-Espinosa et al. (2016) (dried ground fava seeds were replaced with casein to keep the concentration of protein in the diet constant) for 21 days; and Group 5: colitic rats that were fed the 30% sucrose containing basal diet as well as dried ground fava seeds (252 g/kg) for 21 days.

At the end of the experimental regimen, the final body weight and the food intake were recorded. Rats were sacrificed, and serum was separated from the obtained blood samples by centrifugation. Serum samples were stored at – 20°C for later use in biochemical analyses and a tissue specimen was obtained from the distal colon and stored in saline solution for DNA analysis and histological investigation.

2.4. Assessments of colitis

2.4.1. Calculation colon/body weight ratio

Eight-cm-long part from the rear of the colon was removed and longitudinally opened. Colon was emptied of faecal materials by spraying 0.9% saline blotted dried and weighed. The oedema index was determined from ratio of colon segment to body weight (Distrutti et al., 2009).

2.4.2. Estimation of anti-oxidant and inflammatory biomarkers in colon homogenates

The collected colorectal tissue was homogenised in 10 mM Tris-HCl buffer (pH 7.1). The obtained colon homogenate was used to measure anti-oxidant enzyme levels, including the following: non-protein sulphydryl (NP-SH) and total glutathione sulphydryl (T-GSH), according to Sedlak and Lindsay (1968), and catalase (CAT) activity and nitric oxide (NO) using commercial kits (Biowision, USA). Protein carbonyl contents (PCOs) in colon homogenates were determined spectrophotometrically according to Levine et al. (1990) based on the reaction of the carbonyl group with 2,4-dinitrophenylhydrazine. Colonic myeloperoxidase (MPO) activity was determined according to Krawisz et al. (1984), whereas colonic alkaline phosphatase (ALP) activity was measured using disodium p-nitrophenol phosphate as the substrate (Dorai and Bachhawat, 1977).

2.4.3. Determination of colonic mucosal cytokines

Inflammatory mediators including IL-6, IL-β and TNF-α and prostaglandin E2 (PGE2), were determined in colon tissues by ELISA based assays in accordance with the (Kamiya Biomedical Co. CA. USA) supplier’s instructions.

2.4.4. Determination of colon mucosal DNA

Colonic DNA was isolated from colon mucosal tissue following the guidelines of manufacturer (QIAGEN, USA). About 25 mg of colon tissues were minced into smaller pieces to facilitate complete lysis. To the tissue lysate, 180 μL of ATL buffer, 20 μL of proteinase K were added and incubated at 56 °C. The sample was vortexed during incubation, to homogenize it, or placed in a shaking water bath and then vortexed for 20 s. Next, to this sample mixture, 200 μL of AL buffer and 200 μL of absolute was added, and vortexed to mix thoroughly. Finally, DNA purity and content were measured and further analysed by agarose gel electrophoresis.
2.4.5. Macroscopic assessment

Several parameters were assessed for microscopic scoring. Each of the studied parameters including mucosal and submucosal inflammatory cell infiltration, mucosal oedema, and necrosis of the mucosa was allotted a grade (+++, severe; ++, moderate; +, mild; and –, no change) based on the shift in its severity in relation to respective parameter in control.

2.4.6. Histopathological investigations

Following excision, colon tissue was fixed in 10% formalin. Tissue samples were embedded into blocks of paraffin wax and cut into thin sections. They were then stained with haematoxylin and eosin H&E and examined under light microscope.

2.5. Statistical analysis

Data were analysed by (SPSS program for windows, version 20) (SPSS Inc., Chicago, IL, USA). The difference between groups was made using One Way ANOVA (LSD) test. The data were presented as means ± SD. The p < 0.01 was considered as statistically significant.

3. Results

3.1. Anthropometric parameters and food intake

Data on the effects of treatments on anthropometric parameters and food intake in control and colitic rats are presented in Table 1. Food intake in colitic rats was not affected by the presence of colitis, as it was comparable to that observed in control rats. However, food intake in normal and colitic rats fed a diet with 30% sucrose-supplementation was significantly (p < 0.01) augmented compared with normal diet fed control and colitic rats without supplementation. The final body weights of normal rats that fed on a 30% sucrose supplemented diet was significantly elevated (p < 0.01) compared those in control rats. In contrast, body weights of colitic rats were significantly decreased (p < 0.01) compared with control rats. However, the administration of a 30%-sucrose-supplemented diet to colitic rats led to a significant increase (p < 0.01) in their body weights compared with colitic rats. The body weights of colitic rats fed a sucrose-supplemented diet were further increased after the supplementation of their diets with FB, compared with colitic rats fed a diet with or without supplementation with sucrose. The colon-to-body weight ratio was significantly (p < 0.01) higher in normal rats fed on a diet supplemented with 30%-sucrose. Similarly, matched to control rats, colitic rats fed with or without the sucrose-supplemented diet had significantly (p < 0.01) elevated colon to body weight ration. Conversely, colitic rats fed a diet added with both sucrose and FB had a significantly (p < 0.01) lower colon-to-body weight ratio.

3.2. Oxidative stress markers

The effects of different treatments on oxidative stress markers in colon tissues are listed in Table 2. Compared with the control animals, T-GSH, NP-SH and CAT levels were significantly decreased (p < 0.01), and NO, PCO, MPO and ALP levels were significantly increased (p < 0.01) in colitic rats. Feeding of a 30%-sucrose-supplemented diet to colitic rats further downregulated GSH, NP-SH and CAT levels and further augmented the NO, PCO, MPO and ALP levels significantly compared with colitic rats that were fed a normal diet. Conversely, supplementation of the diet of colitic rats with FB together with 30% sucrose led to a significant reversal (p < 0.01) in the levels of the studied oxidative stress markers compared with those detected in colitic rats fed on a diet with or without supplementation with 30% sucrose. Compared with control, normal healthy rats fed on a sucrose-supplemented diet showed no significant change in any of the studied oxidative stress markers.

3.3. Inflammatory markers

The effects of various treatments on inflammatory indices are presented in Table 3. Colitic rats had significantly elevated levels of IL-6, TNF-α, PGE2 and IL-1β (p < 0.01) compared with control rats. Colitic rats fed on a diet containing 30%-sucrose exhibited a further increase (p < 0.01) in these markers matched to those in colitic rats that were fed a normal diet. In contrast, colitic rats fed on a diet supplemented with sucrose as well as FB exhibited significantly decreased levels (p < 0.01) of all the studied inflammatory markers compared with colitic rats fed on diet with or without sucrose supplementation. Conversely, the levels of studied inflammatory markers were comparable between control and normal rats that were given a 30%-sucrose containing diet.

3.4. Histopathological analysis

The scores of the histological analysis of colonic mucosa are illustrated in Table 4. A significant necrosis of colonic mucosa, infiltration of inflammatory cells in mucosa and submucosa and submucosal oedema was noted in colitic rats as well as in colitic rats fed a 30%-sucrose-supplemented diet. Necrosis of the mucosa and submucosal inflammatory cell infiltration were more severe in colitic rats that were fed a normal diet compared with colitic rats that were fed a 30%-sucrose containing diet. In contrast, submucosal oedema was more pronounced in colitic rats that were fed a sucrose containing diet compared with colitic rats fed with a diet without 30% sucrose supplementation. Moreover, rats fed with a diet mixed with sucrose as well as FB exhibited a significant improvement in necrosis of mucosa, oedema and mucosal inflammatory cell infiltration. No significant changes were noticed in normal healthy rats fed a 30%-sucrose containing diet compared with control rats (Table 4).

Table 1

|                      | Control | 30% sucrose | Colitic | Colitic + 30% sucrose | Colitic + 30% sucrose + FB |
|----------------------|---------|-------------|---------|-----------------------|-----------------------------|
| Food intake (g)      | 260 ± 18.7 | 285 ± 19.9*  | 264 ± 17.9 | 280 ± 19.5*  | 282 ± 18.4*                |
| Final body weight (g) | 310 ± 21.2 | 350 ± 19.4*  | 250 ± 17.8* | 266 ± 19.9*  | 280 ± 20.4*                |
| Colon/body weight ratio | 5.1 ± 0.94  | 6.2 ± 0.96*  | 6.8 ± 0.99* | 7.1 ± 0.81*  | 5.8 ± 0.91*                |

FB, fava beans; *compared with the control; †compared with the colitic group; ‡compared with the colitic + 30% sucrose group; §compared with the 30% sucrose, colitic + 30% sucrose and colitic + 30% sucrose + FB groups. The data are shown as the mean ± SD (N = 10); p < 0.01.
supplemented with FB.

and oedema in submucosa were noted in colitic rats given a diet necrosis, submucosal and mucosal inflammatory cell infiltration that were given a normal diet exhibited more severe mucosal tology of colonic mucosa and absence of submucosal oedema and with 30% sucrose compared with control rats, in which normal his-

A significant improvement in mucosal supplementation. Submucosal oedema was more pronounced in pared with colitic rats that were provided with a diet with sucrose

3.5. Histology of mucosal tissue

The histological observations performed via H&E staining of mucosal tissues of rats collected after the different treatments are provided in Fig. 1. A significant mucosal necrosis accompanied by submucosal and mucosal infiltration of inflammatory cells and oedema in submucosa were observed in both colitic rats fed on a normal diet and in colitic rats fed with a normal diet supplemented with 30% sucrose compared with control rats, in which normal histology of colonic mucosa and absence of submucosal oedema and inflammatory cell infiltration were observed. In turn, colitic rats that were given a normal diet exhibited more severe mucosal necrosis and submucosal infiltration of inflammatory cells compared with colitic rats that were provided with a diet with sucrose supplementation. Submucosal oedema was more pronounced in colitic rats fed a sucrose diet than it was in those without the sucrose diet. In contrast, a significant improvement in mucosal necrosis, submucosal and mucosal inflammatory cell infiltration and oedema in submucosa were noted in colitic rats given a diet supplemented with FB.

3.6. DNA damage

The effects of the different treatments on mucosal DNA fragmentation are presented in Fig. 2. Relatively, 55% and 68% fragmentation of mucosal DNA was noticed in colitic rats fed a normal diet and in colitic rats given a normal diet supplemented with 30% sucrose, respectively, compared with the 1% fragmentation of colonic DNA detected in control rats. Colitic rats fed on a diet supplemented with 30% sucrose and FB showed a marked reduction in colonic DNA fragmentation, as only 42% of DNA was found to be fragmented, indicating a significant protection of FB against DNA damage. Rats that were fed a 30% sucrose diet exhibited 7% DNA fragmentation, which was comparable to the 1% fragmentation of colonic DNA found in control rats.

4. Discussion

Acetic-acid-induced UC is a widely used animal model for ulcerative colitis (Ashry et al., 2016). It is similar to human UC in terms of histopathological characteristics, pathogenesis and the inflammation, rendering it to be a useful tool for anti-colitic drug screening (Randhawa et al., 2014).

In this study, AA elicited an inflammatory response in colon tissues, as evident from the reduction of anti-oxidant biomarkers (TSH, NP-SH and CAT) and the increase of NO, PCO, inflammatory cytokines and inflammatory markers. Here, a high-sucrose diet was administered and the results of the G2 colitic group and the G3 colitic group were compared. We found that this diet generated multiple effects on the occurrence of colon inflammation caused by a single dose of AA. Because of the hydrolysis of sucrose into monosaccharides, a rat feeding study reported that sucrose increased the proliferation of colon cells to a greater extent than fructose and glucose did shortly after its ingestion (Wang et al., 2009). A study performed by Chen et al. (2015) confirmed that sucrose acts as a co-initiator or promoter of colon tumour formation. Pettan-Brewer et al. (2011) demonstrated that sucrose augments inflammation markers and acts as a proinflammatory agent. Epidemiological studies presented important correlations between inflammation of the colon and high intake of sucrose.
Different studies have recently demonstrated that increased consumption of sucrose correlated with a raised colon cancer risk (Rawla et al., 2019). In an animal study sucrose elevated the rate of mutations in the colon, and also changed oxidative stress by glycoxidation and changed the colonic environment, leading to a variation in pH and short-chain fatty acids, which are fermentation products, a central supply of colonic epithelium energy (Gu et al., 2019).

The results indicated that UC negatively affected the final body weight; this may be attributed to an impairment in the absorption of water in the inflamed mucosa, contributing to diarrhoea in this animal model in cases of human inflammation of the intestines (Bitiren et al., 2020). The weight/length ratio was also used to determine the weight gain caused by oedema or chronic inflammation shortening, which are indices of the severity of colitis (Ağış et al., 2015). The increase in body weight in sucrose-supplemented groups may have led to an increase in palatability because of the effect of sucrose. In addition, a body weight increase was detected in G5 (colitic rats + 30% sucrose supplemented with FB), which was due to the effect of FB seeds on palatability and their high protein and carbohydrate content (Basheer-Salimia et al., 2014).

The increased colon to body weight ratio is considered a sensitive and reliable index for assessing severity of ulcers, oedema and increased levels of inflammatory cytokines, such as TNF-α and IL-6 (Palla et al., 2016). In this study, AA-treated rats exhibited a significant increase in the weight ratio of the colon to the body weight. Oxidative stress is a driving force in the initiation and pathogenesis of IBD. AA causes vascular dilation and the accumulation of white blood cells (WBCs), and elevates flow of blood, resulting in higher production of reactive oxygen species. Bitiren et al. (2020) demonstrated the role of free radicals in the etiology of mucosal injury. AA reduced the content of anti-oxidant biomarkers (T-SH, NP-SH, CAT, NO and PCO). In contrast, the absence of significant differences between the G1 and G4 control groups appears to indicate the role of AA in augmented oxidative stress in lieu of sucrose.

Earlier studies reported that sucrose causes colon, liver or plasma oxidative stress (Van Hecke et al., 2019). Several studies proved that intra-rectal AA administration decreases the defensive anti-oxidant system, including GSH and catalase activity (Krishnan et al., 2014). ROS produce several inflammatory cytokines that exacerbate tissue damage in different tissues. Free radicals cause severe cell damage due to lipid peroxidation (Jena et al., 2012). GSH is an important anti-oxidant intracellular enzyme that is expressed in the gut of mammals, and plays crucial role tissue repair because of its ability to inhibit ROS. Inflammation causes decrease in GSH, and the colon mucosa is severely degraded. GSH, therefore, plays pivotal role in intestinal cell protection and abrogating inflammation (Zińczuk et al., 2020). GSH or NP-SH peptide sulphhydryl groups are widely used in all biological systems and...
act as an enzyme-free anti-oxidant. GSH due to its sulfhydryl group directly negates ROS and protects oxidative related injuries and it can also indirectly participate in ROS detoxification as a coenzyme or cofactor (Bai et al., 2019).

Nitric oxide and nitric oxide synthase NOS are potential colitis facilitators. The inflammatory reactions observed during colitis, such as enhanced interstitial oedema and intestinal wall thickening, are linked to inflammatory molecules, such as NO (Gillberg et al., 2012). Clinically, colonic tissues of patients with colitis release ROS and NO, which lead to increased membrane peroxidation and DNA attacks (Venkatachalam et al., 2020). Our results indicate that dietary supplementation with dried fava beans can restore the oxidative status of colitic rats in colon homogenates. Ganesan and Xu (2017) indicated that the anti-proliferative and anti-oxidant potentials of legumes were correlated with the occurrence of different classes of phenolic compounds.

Fava bean pods also contain up to 100 mg/kg of caffeoyl-L-malic acid. Similarly, earlier studies (Xu and Chang, 2010) have demonstrated that pro-antioxyanidins are important phenolic compounds present in fava beans. Pro-antioxyanidins can, however, alleviate oxidative stress by scavenging free radicals and altering signalling pathways (Puiggròs et al., 2014).

In human, animal and culture studies, which have collectively demonstrated the potential of PCs to prevent or treat oxidative-stress-related diseases, the anti-oxidant efficacy of PCs has been verified (Casassa et al., 2013). MPO is a neutrophilic enzyme; its activity therefore, correlates to neutrophil population in the inflamed tissue. MPO activity measurement was, therefore, considered to be a sensitive test for the detection of acute intestinal inflammation (Molodecky et al., 2012). The increased activity of MPO is considered a neutrophil infiltration index and indicates the formation of potent cytotoxic oxidants, such as H2O2, hydrochloric acid and chloride ions (Costello et al., 2020). The disturbed redox system can also be reflected in the increased levels of the protein carbonyls PCO and LPO (Garcia et al., 2019). Acute administration of AA through the rectal route increased the MPO and PCO activities, which were reduced after fava bean dietary supplementation in colitic rats. The modelling effect of fava beans on inflammatory markers was linked to the high content of powerful phenolic compounds and other bioactive compounds that could protect the mucosal tissue by increasing its immunity (Esmaillzadeh and Azadbakht, 2012). In this work, AA was adversely affected by inflammatory reaction after colonic injury, leading to a massive intracellular acidification that caused immense epithelial damage (Hall et al., 2020). Cytokines mediate immune responses after AA administration and manage the expansion of colonic mucosal damage (Pearl et al., 2013).

UC was related to an impaired immune response followed by the release of inflammatory molecule, IL-1β, which is expressed as inactive pro-IL-1β and is activated under caspase-1 control (Opipari and Franchi, 2015). Of note, Lazaridis et al. (2017) reported that, in the inflamed colonic mucosa, IL-1β was obviously enhanced and correlated with the severity of the disease. TNF-α is an important pro-inflammatory cytokine that in early inflammation is secreted from lymphocytes and macrophages. It was reported that inflammatory bowel disease plays an integral role in the pathogenesis of UC (Sheng et al., 2020). Earlier human studies (Dai et al., 2020; Zhang et al., 2020) indicated that TNF-α and IL-6 were increased in inflammatory bowel conditions and suggested that these factors play an integral role in their pathogenesis. In this study, the obtained histological findings and oedema formation caused by the elevated cytokines and epithelial cell necrosis were as reported previously by clinical studies (Tahan et al., 2011).

The results of elevated colonic PGE2 levels in colitic rats obtained here were consistent with those of Yu et al. (2019). Furthermore, many pro-inflammatory cytokines induce apoptosis in intestinal epithelia during the intestinal inflammation process. In colonic tissues, pro-inflammatory cytokines are involved in T-cell growth and B-cell proliferation (Santos et al., 2020). However, Bai et al. (2020) and Cibor et al. (2020) reported that IL-6 serum levels were significantly high and proportional to the intensity of colonic mucosal injury.

Fava bean is rich in pro-antioxyanidins, which have antioxidant and anti-inflammatoryary properties. Several studies have demonstrated that pro-antioxyanidins can have a beneficial effect on endothelium injury, platelet aggregation and inflammation caused by systemic disease pathogenesis (Huang et al., 2014). In this study, the histological examinations confirmed the biochemical measurements, i.e., that AA caused marked mucosal necrosis and inflammatory cell infiltration. The study reported by Pastrelo et al. (2017) demonstrated that acetic acid-induced colitis leads to different lesions in the colonic mucosa and necrosis associated with neutrophils. Evidence suggests that peroxides and hydroxyl groups destabilize cell-membrane, cause damage to DNA and cell death due lipid peroxidation (Krzyżek-Korpacka et al., 2020).

Regarding the results of colonic mucosal DNA, the supplementation of the diet with sucrose led to changes in the colonic mucosal DNA. According to Guercio et al. (2018) dietary sucrose played a role of promoter or co-initiator of colon tumour formation. Genotoxic and mutagenic potential of sucrose in colonic tissues was attributed to oxidative stress induced DNA damage or derailed DNA repair mechanism (Van Hecke et al., 2019). Notably, Yu et al. (2018) concluded that a diet rich in sucrose doe dependently contributed to increased mutation rate in rat colon directly or indirectly, and lowers DNA levels. In addition, Heredia-Garcia et al. (2019) found that sucrose promoted DNA damage in the colon, indicating pro-carcinogenic nature of sucrose. In contrast, fava beans have anti-mutagenic effects because of their high phenolic and flavonoid content, thus exerting a protective effect against DNA damage (Siah et al., 2012).

5. Conclusions

Increased levels of inflammatory mediators in the rat colon indicated that acetic acid triggered an inflammatory state in colon tissues. In addition, high-sucrose diets increase colonic inflammation and induce histological and mucosal DNA alterations. Dietary supplementation with dried ground fava bean significantly corrected the impaired oxidative and inflammatory biomarker levels and modulated histological features and DNA alterations. Finally, fava bean attenuated the oxidative damage and colonic injury induced by acetic acid, which confirmed its high anti-oxidant and anti-incediary properties.

Funding

This research did not receive any specific grant from funding agencies in the public, commercial or not-for profit sectors.

Declaration of Competing Interest

None.

References

Abdin, A.A., 2013. Targeting sphingosine kinase 1 (SphK1) and apoptosis by colon-specific delivery formula of resveratrol in treatment of experimental ulcerative colitis in rats. Eur. J. Pharmacol. 718, 145–153.
Abraham, B.P., Ahmed, T., Ali, T., 2017. Inflammatory bowel disease: pathophysiology and current therapeutic approaches. Gastrointest. Pharmacol. 239, 115–146.

A.A. Eskandrani Saudi Journal of Biological Sciences 28 (2021) 3497–3504
Carnitine is Efficacious in Ulcerative Colitis Through its Action on the Immune Function and Microvasculature. Clin. Transl. Gastroenterol. 5, 55.

Sedlak, J., Lindsay, R.H., 1968. Estimation of total, protein-bound, and nonprotein sulfhydryl groups in tissue with Ellman’s reagent. Anal. Biochem. 25, 192–205.

Shanmugam, S., Thangaraj, P., dos Santos Lima, B., Trindade, G.G.G., Narain, N., Mata de Oliveira, E., Silva, A., Santin, J.R., Broering, M.F., Serafini, M.R., Quintans-Júnior, L.J., de Souza, Antunes, A., 2020. Protective effects of flavonoid composition rich P. subpeltata Ortega on indomethacin induced experimental ulcerative colitis in rat models of inflammatory bowel diseases. J. Ethnopharmacol. 248, 112350.

Sheng, N., Ma, Z., Zhou, Y., Xu, J., Gao, Y., Fu, X.Y., 2020. Cholesterol 25-hydroxylase protects against experimental colitis in mice by modulating epithelial gut barrier function. Sci. Rep. 10, 14246.

Van Hecke, T., De Vrieze, J., Boon, N., De Vos, W.H., Vossen, E., De Smet, S., 2019. Combined Consumption of Beef-Based Cooked Mince and Sucrose Stimulates Oxidative Stress, Cardiac Hypertrophy, and Colon Outgrowth of Desulfovibrionaceae in Rats. Mol. Nutr. Food Res. 63, e1800962.

Venkatachalam, K., Vinayagam, R., Arokia Vijaya Anand, M., Isa, N.M., 2020. Ponnaiyan R Biochemical and molecular aspects of 1,2-dimethylhydrazine (DMH)-induced colon carcinogenesis: a review. Toxicol. Res. (Camb) 9, 2-18.

Wang, B., Bohe, G., LaPres, J.J., Bourquin, L.D., 2009. High sucrose diets promote intestinal epithelial cell proliferation and tumorigenesis in APC (Min) mice by increasing insulin and IGF-1 levels. Nutr. Cancer 61, 81–93.

Xu, B., Chang, S.K.C., 2010. Phenolic substance characterization and chemical and cell-based antioxidant activities of 11 lentils grown in the Northern United States. J. Agric. Food Chem. 58, 1509–1517.

Yu, Y., Cheng, D., Parfrey, P., Liu, G., Savas, S., 2018. Two functional indel polymorphisms in the promoter region of the Brahma gene (BRM) and disease risk and progression-free survival in colorectal cancer. PLoS One 13, e0198873.

Yu, Y., Wong, F.H., Park, H.H., Baek, S.Y., Kim, Y.J., Lee, S., Kim, Y.L., Seo, K.W., Kang, K.S., 2019. Preconditioning with interleukin-1 beta and interferon-gamma enhances the efficacy of human umbilical cord blood-derived mesenchymal stem cells-based therapy via enhancing prostaglandin E2 secretion and indoleamine 2,3-dioxygenase activity in dextran sulfate sodium-induced colitis. Tissue Eng. Regen. Med. 13, 1792–1804.

Zhang, T., Jiang, J., Liu, J., Xu, L., Duan, S., Sun, L., Zhao, W., Qian, F., 2020. MK2 Is Required for Neutrophil-Derived ROS Production and Inflammatory Bowel Disease. Front. Med. (Lausanne) 7, 207.

Zin’czuk, J., Maciejczyk, M., Zaręba, K., Przybylska, A., Dymicka-Piekarska, V., Kamińska, J., Kober-Lenkiewicz, O., Matowicka-Karna, J., Kędra, B., Zalewska, A., Guzińska-Ustymowicz, K., 2020. Pro-Oxidant Enzymes, Redox Balance and Oxidative Damage to Proteins, Lipids and DNA in Colorectal Cancer Tissue. Is Oxidative Stress Dependent on Tumour Budding and Inflammatory Infiltration?. Cancers (Basel) 12, 1636.70.