The Healing Effect of Grape Seed Oil Enema with or without Sesame Oil in Acetic Acid Induced Ulcerative Colitis of Rats

Fatemeh Hosseinzadeh1,2, Moosa Salehi1,2, Nader Tanideh3,4*, Davood Mehrabani3, Azadeh Sayarifard5, Anahita Sedighi6

ABSTRACT

BACKGROUND
Inflammatory bowel diseases contain two digestive system diseases, ulcerative colitis (UC) and Crohn’s disease with unclear causes. The aim of present study was to investigate the therapeutic effects of administration of the Sesame oil (SO) and grape seed oil (GSO) as enema route in rats suffering from experimental acetic acid induced UC.

METHODS
Eighty male rats were randomly allocated into 8 equal groups as health control (HC1) without any disease treated with 1 ml of normal saline as enema; HC2 received SO; HC3 received GSO; negative control (NC) with induced UC receiving 1 ml of normal saline as enema; and positive control (PC) with induced UC treated by asacol. All treatments were performed identically with 4 mg/kg of medication except for asacol that was 100 mg/kg for 7 days. The weight changes was recorded after seven days. The serum levels of malondialdehyde (MDA), total antioxidant capacity (TAC), interleukin-6, and c-reactive protein (CRP) were measured. Colon macroscopic and microscopic histological changes were also measured at the end of 7th day.

RESULTS
No significant changes were detected in weight in neither groups on day 0 nor at the end of study. No beneficial effects were seen for all treatments regarding healing process and the decrease in inflammation. Between treatment groups, the lowest MDA (7.40±0.98 U/ml), CRP (83.20±10.01 mg/l) and IL-6 levels (130.86±10.70 mU/ml) and highest TAC (1.91±0.43 mmol/l) belonged to GSO group.

CONCLUSION
GSO enema alone can be considered as a treatment of choice for UC due to its antioxidant properties.

KEYWORDS
Grape seed oil; Sesame oil; Ulcerative colitis; Enema
INTRODUCTION

Inflammatory bowel diseases (IBDs) contain two digestive system diseases, ulcerative colitis (UC) and Crohn's disease (CD) with unclear causes. The occurrence of UC can be affected by environmental, genetics, immune factors, and gut microbiota. UC is demonstrated by mucosal inflammation and limited to the colon. A significantly reduced biodiversity in fecal microbial count and diversity in IBD patients compared to those in healthy controls were reported. It has been reported that high intakes of mono- and polyunsaturated fats and vitamin B6 are associated with an increased risk to develop UC.

So it can be suggested that use of any chemical or herbal agents in enema route can be effective in treatment of UC by changing colon microbiome and also by producing local effects. There are several ways to induce UC in experimental animals such as use of acetic acid as acetic acid is inexpensive and available in most laboratories. Grape is a fruiting berry of the deciduous woody vines of the botanical genus Vitis. Grape seed oil (GSO) has many uses ranging from cooking (as a food additive), cosmetics and in controlling several diseases and wound healing potential. Grape seed and GSO are rich in unsaturated fatty acids and antioxidants such as tocopherols, phytosterol, and α- and γ-tocotrienol.

Although oil content and fatty acid composition of grape seed show great variation among various genotypes, but it has been reported that the predominant fatty acids in the GSO were linoleic (65.0%), linolenic (1.5%), oleic (17.0%), and palmitic (8.0%) acids which play significant roles in lipid metabolism. Unlike GSO, sesame oil (SO) is a known oil which showed any therapeutic effects due to its antioxidant, antibacterial and anti-inflammatory agents. There are no reports about the use of GSO with or without SO in the acetic acid induced UC neither in enema route application nor by other routes. Therefore, the aim of the present study was to evaluate the healing effect of GSO alone or in combination with SO in enema route in the acetic acid induced ulcerative colitis.

MATERIALS AND METHODS

The use of experimental animals in this study was approved by Ethical Committee of Shiraz University of Medical Sciences. All efforts were made to minimize animal stress and handling and also the animal welfare was under consideration in all part of the study. Eighty male Sprague Dawley rats (200±20 g) were obtained from the Center of Experimental and Comparative Medicine, Shiraz University of Medical Sciences, Shiraz, Iran. The rats were randomly allocated into 8 equal separated groups as follow and were housed in standard cages under a 12-h light cycle (lights on at 7:00 pm) with an ambient temperature of 22±2°C, and 55% relative humidity.

Healthy control I (HC1) group did not undergo induction of UC and received 1 ml of normal saline by enema route, HC2 group did not undergo UC induction and received 4 mg/kg of SO, HC3 group did not undergo UC induction and received 4 mg/kg GSO. Negative control (NC) group underwent UC induction and received 1 ml of normal saline by enema route, Positive control (PC) group underwent UC induction and received 100 mg/kg of asacol by enema route, SO group underwent UC induction and received 4 mg/kg of SO by enema route, GSO group underwent UC induction and received 4 mg/kg of GSO by enema route, and SO+GSO group that underwent UC induction and received both 4 mg/kg SO and 4 mg/kg GSO by enema route.

Oils used in this study were Zareentalia Grap seeds Oil (Zareentalia Co, Italy) and Samar sesame oil (Samar Co, Iran). These oils were produced from the best quality raw materials and having the standard production certificates. All animals were fasted overnight and their bowels were cleaned before induction of UC to emptying the colon. UC induction was performed using 2 ml of 3% acetic acid transrectally as described before. All treatment were applied in enema route.

The weight changes were recorded before the study (day 0) and at the 7th day (end of the study) by using a digital scale with 0.1 g precision. At the end of the seventh days, the rats were sacrificed in the CO₂ induction box. Then, the distal 8 cm of the colon was removed and dissected by longitudinal incision. We processed, stained and evaluated the colon tissues of rats in each groups as reported before. Indices which presented by Morris et al. as 0-5 degree for macroscopic mucosal injuries, by Murthy et al. as 0-4 degree for crypt injuries, and Onderdonk et al. as 0-3
degrees for level of inflammation were used in this study to evaluate the histopathological changes of the colon tissues.

The serum samples were separated from the heart blood using centrifugation (3000 rpm, 10 min). The malondialdehyde (MDA) level, as the end-products of lipid peroxidation (LPO) and one of the most important indices of oxidative stress, was assessed via the measurement of thiobarbituric acid reactive substances (TBARS) in sera. In addition, the total antioxidant capacity (TAC) in serum was evaluated by colorimetric method using commercial kit (Cayman, USA). The IL-6 content of the sera was measured using rat specific sandwich ELISA kit (Sigma Aldrich, USA). Serum CRP was evaluated by enzyme immunoassay kit (IBL international, Germany).

All of our evaluated variables were biological parameters and have normal distribution in the population. Therefore, we can face with it as parametrical variables. The data are presented as mean and standard deviation. Normality of data were checked by Kolmogorov-Smirnov test. One-way ANOVA and Tukey post hoc test were used for comparison of mean differences in all variables between 8 groups. \( p<0.05 \) was considered statistically significant. Mann-Whitney U test with Bonferroni correction was used for comparing histopathology scores between different groups on different days. SPSS 16.5 software was used to analyze the data.

**RESULTS**

There are no significant differences in weight of rats between different groups in neither day 0 nor 7th day \( (p>0.05) \). However, the highest mean weight belonged to GSO group \( (187.63\pm9.74 \text{ g}) \). The mean and SD of macroscopic and microscopic scores in different groups are presented in Table 1. As shown, NC and then SO+GSO had highest degree of macroscopic lesions which significantly differed from HC group \( (p<0.001 \text{ and } p=0.005, \text{ respectively}) \). The highest microscopic lesions and inflammation severity were seen in NC group which significantly higher than all other 7 groups \( (p<0.05) \). No significant differences were detected in these variables between other 7 groups \( (p>0.05) \).

The mean and SD of oxidant and antioxidant status, inflammatory indices in the serum at the end of the study period (day 7) are presented in Figure 1. Between treatment groups, the lowest MDA \( (7.40\pm0.98 \text{ U/ml}) \), CRP \( (83.20\pm10.01 \text{ mg/l}) \) and IL-6 levels \( (130.86\pm10.70 \text{ mU/ml}) \) and highest TAC \( (1.91\pm0.43 \text{ mmol/l}) \) were belonged to GSO group. This demonstrated the higher efficacy of GSO in treatment of UC and prevention of oxidative stress and inflammation.

**DISCUSSION**

UC as one type of IBD has several treatment methods which are anti-inflammatory and immune modulating drugs such as

| Groups  | Weight (day 0) | Weight (day 7) | Morris\(^{13}\) | Onderdonk\(^{15}\) | Murthy\(^{14}\) |
|---------|---------------|---------------|----------------|----------------|-------------|
| HC\(_1\) | 168.60±9.87\(^a\) | 178.20±10.47\(^a\) | 0.00±0.00\(^b\) | 0.00±0.00\(^a\) | 0.00±0.00\(^a\) |
| HC\(_2\) | 176.00±3.16\(^a\) | 176.00±7.07\(^a\) | 0.60±0.89\(^b\) | 0.60±0.55\(^a\) | 0.60±0.55\(^a\) |
| HC\(_3\) | 174.40±9.40\(^a\) | 171.80±11.34\(^a\) | 1.60±1.82\(^b\) | 0.20±0.45\(^a\) | 0.20±0.45\(^a\) |
| NC     | 177.22±9.90\(^a\) | 167.22±19.29\(^a\) | 3.00±1.32\(^b\) | 2.67±0.87\(^a\) | 2.44±0.53\(^a\) |
| PC     | 165.67±8.04\(^b\) | 166.67±10.39\(^b\) | 0.33±0.52\(^b\) | 0.00±0.00\(^a\) | 0.00±0.00\(^a\) |
| SO     | 172.38±11.20\(^a\) | 172.00±20.20\(^a\) | 1.75±1.58\(^b\) | 0.63±1.19\(^a\) | 0.75±1.39\(^a\) |
| GSO    | 176.63±11.61\(^b\) | 187.63±9.74\(^b\) | 0.88±0.83\(^b\) | 0.88±0.35\(^a\) | 0.88±0.35\(^a\) |
| SO+GSO | 169.29±4.61\(^a\) | 172.86±20.19\(^a\) | 2.43±1.90\(^a\) | 1.00±1.15\(^a\) | 1.29±1.50\(^a\) |

HC\(_1\) group: Neither UC nor treatment; HC\(_2\) group: No UC induction but use of 4 mg/kg sesame oil (SO); HC\(_3\) group: No UC induction but use of 4 mg/kg grapeseed oil (GSO); NC group: UC induction without treatment; PC group: UC induction but treated with 100 mg/kg Asacol; SO group: UC induction but treated with 4 mg/kg SO; GSO group: UC induction but treated with 4 mg/kg GSO; SO+GSO group: UC induction but treated with both 4 mg/kg SO and 4 mg/kg GSO. Different superscript letters in each column indicate significant difference \( (P<0.05) \).
salazosulfapyridine, mesalazine, corticosteroids, azathioprine, 6-mercaptopurine, methotrexate and cyclosporine. However, applying new treatments needs prior animal studies and finding good model of UC induction. Also, several studies reported different animals as experimental models of UC.

In the present study, we evaluated and compared the effects of GSO alone or in combination with SO in the healing of colon tissue suffered from acetic acid induced UC in rat. This performed by macroscopic and microscopic changes comparison, antioxidant status, and inflammatory indices evaluations. No beneficial effects were seen for all treatments in point of weight gain, macroscopic and microscopic healing and decreasing of inflammation severity. But use of GSO alone was decreased oxidative stress and combated with inflammation. Therefore, enema application of GSO can be presented as a drug of choice for treatment of UC.

The major symptoms in the human and animal UC include diarrhea, rectal bleeding and weight loss. It has been reported that use of plants which have antibacterial and antioxidant activities could be prevent the weight loss and also enhanced the rat weight in animal model of UC. It seem that the weight changes in response to any agents may be due to treatment of the health of the colon ulcers and also improving the overall health conditions. However, we could not find significant changes between different groups in neither day 0 nor 7th day. This may be due to the route of administration and also unknown molecular and hormonal pathways which need further clarification.

No macroscopic and microscopic beneficial effects at the same time, having antioxidant and anti-inflammatory properties were detected.
in this study just for GSO. Although, in the previous studies, it had been cleared that SO contain antioxidant factors which combat with inflammation and free radicals,24-26 but we could not find similar properties. There are some reports which expressed that GSO had antioxidant effects but they are not in UC. For instance, it has been reported that pretreatment with GSO exhibited a protective role against diazinon toxicity in rat by the antioxidant role of
its constituents.\textsuperscript{27} Its beneficial antioxidant properties against ethanol,\textsuperscript{28} tetrachloride carbon,\textsuperscript{29} and cyclophosphamide\textsuperscript{30} toxicities and also acrylamide induced lesion\textsuperscript{31} were also reported. All of these beneficial effects plus our findings in UC are related to natural antioxidants which existed in GSO as reported previously.\textsuperscript{32} We conclusively found beneficial effects from the use of GSO alone but not in combination with SO as enema route in induced UC in rat. This mostly related to antioxidative and anti-inflammatory effects and therefore, use of GSO in enema route can be advised in them. However, performing randomized controlled trial to confirm such effects is highly necessary.

\textbf{ACKNOWLEDGMENT}

The present article was extracted from the Ph.D. thesis (ID 7167) written by Dr. Fatemeh Hosseinzadeh and was financially supported by the Shiraz University of Medical Sciences, Shiraz, Iran. Authors wish to thanks Dr. Mahjoob Vahedi, Mr. Omid Koohi-Hosseinabadi and Mr. Mohammad Heidari for their kind cooperation.

\textbf{CONFLICT OF INTEREST}

The authors declare no conflict of interest.

\textbf{REFERENCES}

1. Safarpour AR, Hosseini SV, Mehrabani D. Epidemiology of inflammatory bowel diseases in Iran and Asia; a mini review. \textit{Iran J Med Sci} 2013;38:140.
2. Takhshid MA, Mehrabani D, Ai J, Zarepor M. The healing effect of licorice extract in acetic acid-induced ulcerative colitis in rat model. \textit{Comp Clin Pathol} 2012;21:1139-44.
3. Geerling BJ, Dagnelie PC, Badart-Smook A, Russel MG, Stockbrügger RW, Brummer R-JM. Diet as a risk factor for the development of ulcerative colitis. \textit{Am J Gastroenterol} 2000;95:1008-13.
4. Tanideh N, Afaridi E, Mehrabani D, Azarpira N, Hosseinzadeh M, Amini M, et al. The healing effect of \textit{Berberis vulgaris} in acetic acid-induced ulcerative colitis in rat. \textit{Middle East J Sci Res} 2014;21:1288-94.
5. This P, Lacombe T, Thomas MR. Historical origins and genetic diversity of wine grapes. \textit{Trends Genet} 2006;22:511-9.
6. Shivananda Nayak B, Dan Ramdath D, Marshall JR, Isitor G, Xue S, Shi J. Wound-healing properties of the oils of \textit{Vitis vinifera} and \textit{Vaccinium macrocarpon}. \textit{Phytother Res} 2011;25:1201-8.
7. Zhao L, Yagiz Y, Xu C, Lu J, Chung S, Marshall MR. Muscadine grape seed oil as a novel source of tocotrienols to reduce adipogenesis and adipocyte inflammation. \textit{Food Funct} 2015;6:2293-302.
8. Sabir A, Unver A, Kara Z. The fatty acid and tocopherol constituents of the seed oil extracted from 21 grape varieties (\textit{Vitis} spp.). \textit{J Sci Food Agric} 2012;92:1982-7.
9. Kikalishvili B, Zurabashvili DZ, Zurabashvili ZA, Turabelidze DG, Shanidze LA. Fatty acid of Rkatsiteli grape seed oil, Phellodendron lavellei oil and Amaranthus seeds oil and its comparative byological activity. \textit{Georgian Med News} 2012;212:73-6.
10. Bigdeli Shamloo MB, Nasiri M, Dabirian A, Bakhityari A, Mojab F, Alavi Majd H. The Effects of Topical Sesame (Sesamum indicum) Oil on Pain Severity and Amount of Received Non-Steroid Anti-Inflammatory Drugs in Patients With Upper or Lower Extremities Trauma. \textit{Anesthesiol Pain Med} 2015;5:e25085.
11. Bagheri Nesami M, Shorofi SA, Hashemi-Karoie SZ, Khalilian A. The effects of sesame oil on the prevention of amiodarone-induced phlebitis. \textit{Iran J Nurs Midwifery Res} 2015;20:365-70.
12. Tanideh N, Nematollahi SL, Hosseini SV, Hosseinzadeh M, Mehrabani D, Safarpour A, et al. The healing effect of \textit{Hypericum perforatum} extract on acetic acid-induced ulcerative colitis in rat. \textit{Ann Colorect Res} 2014;2:e25188.
13. Morris GP, Beck PL, Herridge MS, Depew WT, Szewczuk MR, Wallace JL. Hapten-induced model of chronic inflammation and ulceration in the rat colon. \textit{Gastroenterology} 1989;96:795-803.
14. Morthy SNS, Cooper HS, Shim H, Shah RS, Ibrahim SA, Sedergran DJ. Treatment of dextran sulfate sodium-induced murine colitis by intracolonic cyclosporin. \textit{Dig Dis Sci} 1993;38:1722-34.
15. Onderdonk AB, Cisneros RL, Bronson RT. Enhancement of experimental ulcerative colitis by immunization with \textit{Bacteroides}}
Grape seed enema in ulcerative colitis

16 Zal F, Mostafavi-Pour Z, Vessal M. Comparison of the effects of vitamin e and/or quercetin in attenuating chronic cyclosporine a-induced nephrotoxicity in male rats. *Clin Exp Pharmacol Physiol* 2007;34:720-4.

17 Hanauer SB, Present DH. The state of the art in the management of inflammatory bowel disease. *Rev Gastroenterol Disorders* 2003;3:81-92.

18 Tanideh N, Akbari Baseri F, Jamshidzadeh A, Ashraf MJ, Kuhi O, Mehrabani D. The healing effect of strawberry extract on acetic acid-induced ulcerative colitis in rat. *World Appl Sci J* 2014;31:281-8.

19 Mehrabani D, Ziaei M, Hosseini SV, Ghahramani L, Bananzadeh AM, Ashraf MJ, Amini A, Amini M, Tanideh N. The effect of *Calendula officinalis* in therapy of acetic acid induced ulcerative colitis in dog as an animal model. *Iran Red Crescent Med J* 2011;13:884-90.

20 Mehrabani D, Bahrami F, Hosseini SV, Ashraf MJ, Tanideh N, Rezaianzadeh A, Amini M, Amini A. The healing effect of *Teucrium polium* in acetic acid-induced ulcerative colitis in the dog as an animal model. *Middle East J Dig Dis* 2012;4:40.

21 Tanideh N, Bahrani M, Khoshnood-Mansoorkhani MJ, Mehrabani D, A Firoozy D, Koohi-Hosseinabadi O, Iraji A. Evaluating the effect of *Melilotus officinalis* I. aqueous extracts on healing of acetic acid-induced ulcerative colitis in male rats. *Ann Colorec Res* 2016;4:E42856.

22 Khodakarm-Tafti A, Mehrabani D, Homafar L, Farjanikish G. Healing effects of Elaeagnus angustifolia extract in experimentally induced ulcerative colitis in rats. *J Pharmacol Toxicol* 2015;10:29-35.

23 Mehrabani D, Nasibi M, Izadpanah A, Tanideh N, Amini, SV Hosseini M. The comparison of two methods of steroid therapy via appendicostomy and enema in experimental acetic acid induced ulcerative colitis in dog. *World Appl Sci J* 2013;26:1538-45.

24 Hsu DZ, Chiang PJ, Chien SP, Huang BM, Liu MY. Parenteral sesame oil attenuates oxidative stress after endotoxin intoxication in rats. *Toxicology* 2004;196:147-53.

25 Alves N, Valdes S, Silveira C, Duarte-Martino H, Milagro FI, Moreno-Aliaga MJ, et al. Studies on mechanistic role of natural bioactive compounds in the management of obesity an overview. *Open Nutraceut J* 2012;5:193-206.

26 Hsu DZ, Li YH, Chien SP, Liu MY. Effects of sesame oil on oxidative stress and hepatic injury after cecal ligation and puncture in rats. *Shock* 2004;21:466-9.

27 Al-Attar AM. Effect of grapeseed oil on diazinon-induced physiological and histopathological alterations in rats. *Saudi J Biol Sci* 2015;22:284-92.

28 El-Ashmawy IM, Saleh A, Salama OM. Effects of marjoram volatile oil and grape seed extract on ethanol toxicity in male rats. *Basic Clin Pharmacol Toxicol* 2007;101:320-7.

29 Maheswari MU, Rao PGM. Antihepatotoxic effect of grape seed oil in rat. *Indian J Pharmacol* 2005;37:179-82.

30 El-Rahim AH, Hafiz NA. Investigation on the protective effect of Grape seed and Linseed oils against cyclophosphamide induced genotoxicity in mice. *Global Vet* 2009;3:377-82.

31 Hasseeb MM, Al-Hizab FA, Hamouda MA-H. Impacts of grape seed oil supplementation against the acrylamide induced lesions in male genital organs of rats. *Pak Vet J.* 2013;33:282-6.

32 Lutterodt H, Slavin M, Whent M, Turner E, Yu LL. Fatty acid composition, oxidative stability, antioxidant and antiproliferative properties of selected cold-pressed grape seed oils and flours. *Food Chem* 2011;128:391-9.