Original Article

Protective Effect of Dizocilpine (MK-801) On TNBS-Induced Experimental Colitis in Mice

Ehsan Motaghi, Valiollah Hajhashemi, Parvin Mahzouni and Mohsen Minaiyan

Department of Pharmacology and Isfahan Pharmaceutical Sciences Research Center, School of Pharmacy and Pharmaceutical Sciences, Isfahan University of Medical Sciences, Isfahan, Iran.

Abstract

Ulcerative colitis is chronic and recurrent disease of the gastrointestinal tract with uncertain etiology and incomplete treatment options. N-methyl-d-aspartate (NMDA) receptor suppression has shown anti-inflammatory effects \textit{in-vitro} and \textit{in-vivo}. The aim of present study was to evaluate the role of dizocilpine (MK-801), a noncompetitive NMDA receptor antagonist, on TNBS (trinitrobenzene sulfonic acid)-induced murine model of colitis. Dizocilpine (0.1, 1 and 5 mg/kg) was given to mice intraperitoneally from 24 h before induction of colitis and daily thereafter for 4 days. Dexamethasone (1 mg/kg) was used as the reference drug. Colitis was induced by intracolonic administration of TNBS/Ethanol (50/50 v/v, 40mg/kg). Animals were sacrificed 5 days after colitis induction and distal colons were examined macroscopically and microscopically. The colonic tissue level of pro-inflammatory cytokines including interleukin 1β (IL-1β), interleukin 6 (IL-6), and tumor necrosis factor-α (TNF-α) were assessed by ELISA. Myeloperoxidase (MPO) level was also measured in colon. Dizocilpine, particularly with intermediate dose of 1mg/kg significantly improved animal’s weight loss as well as macroscopic and microscopic signs of colitis, reduced colonic levels of IL-1β, IL-6, TNF-α and MPO activity. Hence, dizocilpine has significant protective effects in TNBS-induced colitis and NMDA suppression may be a new and effective therapeutic strategy in ulcerative colitis via decreasing in pro-inflammatory cytokine production.

Keywords: Dizocilpine; MK-801; Ulcerative colitis; Mice; NMDA.

Introduction

Ulcerative colitis (UC) represents a chronic recurrent inflammation of large intestine. Although exact etiology of UC is unclear, there are circumstantial evidences to link UC to the immune systems imbalance and genetic and environmental factors (1). UC occurs in the gut of susceptible individuals with genetic innate immune defects. Although, there is no conclusive evidence for involvement of a specific bacteria in this disorder, the interaction between the antigen-presenting cells and some bacteria in the colon leads to an excessive activation of local T lymphocytes and subsequently overproduction of pro-inflammatory cytokines such as tumor necrosis factor-α (TNF-α), interleukin-1β (IL-1β), and interleukin 6 (IL-6) (2). The cytokines activate T and B lymphocytes, neutrophils, eosinophils, and basophils and increase T-lymphocytes...
viability. T-lymphocytes accumulation via increased viability finally causes colon inflammation (2, 3).

Despite the great deal of attention given to this disease during the past years, its pharmacological treatment is not completely established yet. The high frequency of adverse effects and treatment failure and disease relapse, encourage scientists to search new treatments with better outcomes (4).

The N-methyl-D-aspartate (NMDA) receptors are ionotropic glutamate receptors that could be found mainly in nervous system and are involved in controlling of pain perception, synaptic plasticity, and memory function (5). However, recent studies have identified this receptors in several types of immune competent cells such as thymocytes, lymphocytes, and neutrophils (6). Therefore, it has been assumed that NMDA antagonists have the potential to be used in treatment of different inflammatory disorders. Blockade of the NMDA receptor channel with memantine, a potent NMDA antagonist, significantly inhibits the early postoperative IL-6 inflammatory response in human (8). Dizocilpine, a highly selective noncompetitive antagonist on NMDA receptors, is an anticonvulsant drug that shares several pharmacological properties with phencyclidine and ketamine (9).

Many therapeutic potentials have been demonstrated for dizocilpine such as the treatment of traumatic brain injury (10), and neurodegenerative disorders like Alzheimer’s (11) and Huntington’s disease (12). Moreover, it has been established that dizocilpine may has beneficial effects in inflammatory disorders including allergic lung inflammation (13), osteoarthritis (14), and autoimmune encephalomyelitis (15).

Although dizocilpine has a low therapeutic index, it may clarify the probable role of NMDA receptors in UC pathology and may represent a future therapeutic intervention for inflammatory bowel disease (IBD). Hence, the current study was accomplished to evaluate the protective effects of dizocilpine on TNBS induced UC in mice.

Experimental

Chemicals
Dizocilpine, hexadecyl trimethylammonium bromide (HTAB), O-dianisidine dihydrochloride, and TNBS were procured from Sigma-Aldrich (Saint Louis, Missouri, USA). Dexamethasone was also purchased from Raha Pharmaceutical Company (Isfahan, Iran).

The ELISA (enzyme-linked immunosorbent assay) kits for mouse TNF-α, IL-1β and IL-6 were bought from Boster company (Pleasanton, CA, USA). Diethyl ether oxide and formalin solution (35%) were supplied by Merck (Darmstadt, Germany). L-glutamic acid was purchased from MP biomedical company (Netherland).

Animals
Swiss albino male mice (25 - 30 g) were used in the experiments. The animal room was maintained at 22 °C – 23 °C and 12:12 h light/dark cycle. The mice were fasted 16-18 hours prior to TNBS instillation. Thereafter, the animals were fed with standard pelleted chow and tap water ad libitum. All experiments were performed after receiving approval from the Ethics Committee of the Isfahan University of Medical Sciences, Isfahan, Iran.

Experimental groups
The mice were randomly distributed into following groups: Sham (normal mice treated with normal saline) and TNBS-induced colitis groups treated with vehicle (colitis control) or with dizocilpine (0.1, 1 and 5 mg/kg, i.p.), L-glutamic acid (2 g/kg, p.o.) or dexamethasone (1 mg/kg, i.p.). L-glutamic acid was used as a NMDA receptor agonist (16). Colitis was induced in mice by a single intra-rectal instillation of 0.1 mL TNBS/Ethanol (50/50 v/v, 40mg/kg).

Sham group received intra-colonic normal saline (0.9%) in an equivalent volume. Each group composed of 6 mice. The animals were treated once daily starting 24 h before induction of colitis and continued for 4 days until the animals were sacrificed at the day fifth (17).
Effect of dizocilpine (MK801-) on colitis

Animal weight assessment
The animals were weighted each day just before the time of drug administration, during the experiments.

Colitis induction
The animals were fasted for 16-18 h before the TNBS enema. Acute colitis was induced by intracolonic instillation of TNBS with a 3 cm length tube under ether anesthesia. The animals were then kept in head down position for approximately 90 seconds to prevent the leakage of TNBS solution. Afterward, food and water were ad libitum (18).

Macroscopic assessment of colon inflammation
The mice were euthanized by ether, 24 h after final treatment. The distal colon was dissected, opened longitudinally, and then macroscopically scored and its length (cm) and weight (mg) were measured.

Macroscopic scoring of colonic ulceration and inflammation was done according to Wallace et al. (19). Then, the colon was placed on a nonabsorbent white tissue paper and a digital imaging was recorded by a well-adjusted Canon camera (Powershot G9, 12 megapixel, Japan) to determine erosion area. Fiji-win 32 software was used for analyzing the images to measure erosion area (20).

Then, the colon sample was cut in tree pieces. One piece was used for colon myeloperoxidase (MPO) activity measurement and other pieces for hematoxylin and eosin (H&E) staining and ELISA evaluation respectively.

Measurement of colon MPO activity
The segment specified for MPO activity assessment was immediately frozen in liquid nitrogen and stored at -70 ºC until used.

Seven microliters of the sample supernatant fluid were added to phosphate-buffered saline (PBS) containing 0.167 mg/mL O-dianisidine dihydrochloride and 0.0005% hydrogen peroxide. Difference in absorbance at 450 nm was recorded at 30 min intervals for 90 second on a plate reader.

MPO activity was reported as units/100 mg of tissue. One unit of MPO activity was expressed as 1 micromole of peroxide degraded per minute at 25 ºC.

Histopathological evaluation of colitis
Colon samples were fixed in formalin buffered with PBS (10%) and processed routinely and embedded in paraffin. They cut into sections 4 μm-thick using a microtome. The slides were deparaffinized in xylene and counterstained with H&E using standard techniques and scored as described previously (22).

Cytokine measurements
The tissue concentration of pro-inflammatory cytokines including TNF-α, IL-1β and IL-6 were assessed using a commercially obtained ELISA kit (Boster company, Pleasanton, CA, USA) according to manufacturer’s protocols.

An antibody specific for each murine cytokine was pre-coated onto the microplate supplied in the kits. Briefly, standards, samples and a biotinylated secondary goat antibody were pipetted into the microplate pre-coated with antibodies specific for each cytokine and incubated at 37 ºC for 60 min.

After being thrice washed with PBS, plate incubated with the avidin-biotin-peroxidase complex at 37 ºC for 30 min. Following 5 times PBS washing, 100 μL of prepared tetramethylbenzidine (TMB) was added to wells. Then, absorbance was read at 450 nm after addition of TMB stop solution.

Data analysis
Parametric data have been shown as mean ± SEM and analyzed by one-way ANOVA, followed by post hoc Tukey test. Nonparametric data have been displayed as median (range) and analyzed using Kruskal-Wallis test and
subsequently Mann–Whitney U test. A value of $P < 0.05$ was delineated as statistically significant. Statistical analyses were performed with GraphPad Prism (GraphPad software Inc, California, USA).

**Results**

**Effect of dizocilpine on body weight**

Induction of colitis reduced body weight both in colitis control and L-glutamic acid groups. Dizocilpine (0.1, 1 and 5 mg/kg, i.p.) treatment significantly attenuated the weight loss as compared to colitis control. Treatment with dexamethasone reduced body weight loss in mice with colitis too. Nevertheless, L-glutamic acid had no apparent effect on animal’s body weight. Moreover, the animal’s weight increased in Sham group relative to colitis control group (Figure 1).

**Effect of dizocilpine on macroscopic damage of colitis**

TNBS induced severe lesions were associated with edema, hyperemia of the mucosa with large areas of erosion in distal colon while Sham group showed an intact mucosa (Figure 2).

![Figure 1](image1.png)

*Figure 1.* Weight change in mice with colitis treated with dizocilpine (0.1, 1 and 5 mg/kg, i.p.), L-glutamic acid (2 g/kg, p.o.) or dexamethasone (1 mg/kg, i.p.). Animals were treated 24 h prior to induction of colitis and continued daily for 4 days. Data are presented as mean ± SEM. n = 6 per group. * P < 0.05, ** P < 0.01, *** P < 0.001 vs colitis control.

![Figure 2](image2.png)

*Figure 2.* Photos of the distal colon appearance at day 5 after treatment of TNBS-induced colitis in mice. (A) Normal colon in Sham group; (B) Colitis control treated with normal saline; (C and D) Colitis treated with dizocilpine Dizocilpine has repeated and one of them should be deleted. (1 and 5 mg/kg, i.p.) (E) L-glutamic acid (2 g/kg, p.o.) and (F) dexamethasone (1 mg/kg, i.p.).
Effect of dizocilpine (MK801-) on colitis

As it is expected, colon damage score (Table 1 and Figure 2), colon weight (Figure 3) and erosion area (Figure 4) increased in colitis control group in comparison with Sham group (P < 0.001), and dexamethasone as reference drug was effective in reducing the tissue injuries (P < 0.001). Dizocilpine reduced colon weight (P < 0.05) and damaging score (P < 0.01) at doses of 1 and 5 mg/kg respectively. However, the lowest dose of dizocilpine (0.1 mg/kg) had no significant impact on both parameters. Percentage of erosion area of 3 cm of distal colon decreased in all dizocilpine treatment groups compared with colitis control group. Additionally, the protective effect of dizocilpine at two doses of 1 and 5 mg/kg was significantly more prominent than the dose of 0.1 mg/kg. L-glutamic acid however, had no significant effect on macroscopic variables of colon inflammation even with the high dose of 2 g/kg compared to control (Figures 3, 4). As it is expected, colon damage score (Table 1 and Figure 2), weight (Figure 3) and erosion area (Figure 4) increased in colitis control group in comparison with Sham group (P < 0.001), and dexamethasone as reference drug was effective in reducing the tissue injuries (P < 0.001). Dizocilpine reduced colon weight (P < 0.05) and damaging score (P < 0.01) at doses of 1 and 5 mg/kg respectively. However, the lowest dose of dizocilpine (0.1 mg/kg) had no significant impact on both parameters. Percentage of erosion area of 3 cm of distal colon decreased in all dizocilpine treatment groups compared with colitis control group. Additionally, the protective effect of dizocilpine at two doses of 1 and 5 mg/kg was significantly more prominent than the dose of 0.1 mg/kg. L-glutamic acid however, had no significant effect on macroscopic variables of colon inflammation even with the high dose of 2 g/kg compared to control (Figures 3, 4).

Table 1. Effect of dizocilpine (0.1, 1, and 5 mg/kg, i.p.) on macroscopic colonic damage score and pathologic parameters of TNBS- induced colitis in mice.

| Groups                | Macroscopic colitis score (0-10) | Microscopic score |
|-----------------------|----------------------------------|-------------------|
|                       | Inflammation severity (0–3)       | Inflammation extent (0–3) | Crypt damage (0–4) | Microscopic colitis index |
| Sham                  | 0 (0-0)**                         | 0 (0-0)**          | 0 (0-0)**          | 0±0**                      |
| Colitis control       | 6 (4-8)                           | 3 (2-3)            | 3 (2-3)            | 8.33±0.61                   |
| Dizocilpine 0.1 mg/kg | 5 (4-6)                           | 3 (0-3)            | 2.5 (1-3)          | 5.83± 0.79                  |
| Dizocilpine 1 mg/kg   | 3 (2-4)**                         | 2 (1-2)**          | 1.5 (1-3)          | 4.33± 0.76**                |
| Dizocilpine 5 mg/kg   | 2 (0-4)**                         | 2 (1-2)**          | 1.5 (1-2)**        | 4± 0.45**                   |
| L-glutamic acid 2 g/kg| 5.5 (4-7)                         | 3 (1-4)            | 2.5 (1-3)          | 8±0.93                      |
| Dexamethasone 1 mg/kg | 4 (2-6)**                         | 0.5 (0-1)**        | 0 (0-1)'           | 1.83±0.75''                 |

The values are presented as median (range) for scoring parameters. Microscopic colitis index is presented as mean ± SEM. N = 6 for each group, *P < 0.05, **P < 0.01, ***P < 0.001 compared to colitis control (n = 6).
tissues showed loss of goblet cells, submucosal edema, hemorrhage, neutrophils infiltration in the mucosa, inflammation extending to the muscularis mucosae and submucosa, damaged crypts, and destruction of the epithelium (Figure 5B). There was no sign of healing in colonic tissue in L-glutamic acid group compared to colitis control group (Figure 5C). The histologic sections of the dizocilpine (1 and 5 mg/kg) and dexamethasone treatment groups showed ameliorating of histopathologic scores including restoration, progressive re-epithelialization, improvement of the glandular structure, and reduction in edema and congestion in comparison with those of the colitis control group ($P < 0.05$) (Figure 5 D, E, F, G and Table 1).

**Effect of dizocilpine on MPO activity**

The colitis control group displayed a significant increase in the level of MPO activity as compared with Sham group ($P < 0.001$) (Figure 6). By using dizocilpine, a favorable effect was detectable on MPO activity level in all doses but the magnitude of effect was significant only at 5 mg/kg dose. However, no statistically significant difference was observed between L-glutamic acid group and colitis control group. Moreover, dexamethasone reduced the enzyme activity in significant magnitude ($P < 0.001$).

**Effects of dizocilpine on the colonic level of pro-inflammatory cytokines**

In contrary to Sham group that showed an undetectable amount of IL-1$\beta$ by immunoassay,
Effect of dizocilpine (MK-801) on colitis

Changes in MPO activity of distal colon of mice with colitis treated with dizocilpine (0.1, 1 and 5 mg/kg, i.p.), L-glutamic acid (2 g/kg, p.o.) or dexamethasone (1 mg/kg, i.p.). Sham and colitis control groups received an equal volume of normal saline (i.p.). Animals were treated 24 h prior to induction of colitis and continued daily for 4 days. Data are presented as mean ± SEM. n = 6 per group. * P < 0.05, ** P < 0.01, *** P < 0.001 vs colitis control.

In the same manner, IL-6 level was significantly increased in the colitis control group compared to the Sham group (P < 0.001). IL-6 levels at doses of 0.1 and 1 mg/kg of dizocilpine tend to be declined compared to colitis control group. However, the difference was only significant at the greatest dose (5 mg/kg) of dizocilpine (P < 0.05) (Figure 8).

Discussion

Figure 6. Changes in MPO activity of distal colon of mice with colitis treated with dizocilpine (0.1, 1 and 5 mg/kg, i.p.), L-glutamic acid (2 g/kg, p.o.) or dexamethasone (1 mg/kg, i.p.). Sham and colitis control groups received an equal volume of normal saline (i.p.). Animals were treated 24 h prior to induction of colitis and continued daily for 4 days. Data are presented as mean ± SEM. n = 6 per group. * P < 0.05, ** P < 0.01 and *** P < 0.001 vs colitis control.

Figure 7. Change in colon IL-1β levels of mice with colitis treated with dizocilpine (0.1, 1 and 5 mg/kg, i.p.), L-glutamic acid (2 g/kg, p.o.) or dexamethasone (1 mg/kg, i.p.). Sham and colitis control groups received an equal volume of normal saline (i.p.). Animals were treated 24 h prior to induction of colitis and continued daily for 4 days. Data are presented as mean ± SEM. n = 6 per group. * P < 0.05, ** P < 0.01 vs colitis control. ## P < 0.05 compared to dizocilpine 0.1 mg/kg.

There are reports suggesting that peripheral NMDA receptors play a key role in peripheral cytokines and MPO (25) occurred in this model. Ulcer, and necrosis (23). Besides overproduction of nitric oxide (24), pro-inflammatory ulcer, and necrosis (23). Besides overproduction of nitric oxide (24), pro-inflammatory cytokines compared to colitis control group. However, the difference was only significant at the greatest dose (5 mg/kg) of dizocilpine (P < 0.05) (Figure 8)

Figure 8. Change in colon IL-6 levels of mice with colitis treated with dizocilpine (0.1, 1 and 5 mg/kg, i.p.), L-glutamic acid (2 g/kg, p.o.) or dexamethasone (1 mg/kg, i.p.). Sham and colitis control groups received an equal volume of normal saline (i.p.). Animals were treated 24 h prior to induction of colitis and continued daily for 4 days. Data are presented as mean ± SEM. n = 6 per group. * P < 0.05, ** P < 0.01, *** P < 0.001 vs colitis control.

Figure 9. Change in colon TNF-α levels of mice with colitis treated with dizocilpine (0.1, 1 and 5 mg/kg, i.p.), L-glutamic acid (2 g/kg, p.o.) or dexamethasone (1 mg/kg, i.p.). Sham and colitis control groups received an equal volume of normal saline (i.p.). Animals were treated 24 h prior to induction of colitis and continued daily for 4 days. Data are presented as mean ± SEM. n = 6 per group. * P < 0.01 and *** P < 0.001 vs colitis control.

colitis control group expressed a higher colonic level of IL-1β and the anti-inflammatory effects of dizocilpine (1 mg/kg) was confirmed by a significant decrease in IL-1β levels in comparison to the colitis control (P < 0.05). Moreover, it was revealed that a significant difference was evident between 0.1 and 1 mg/kg dizocilpine groups (P < 0.05) (Figure 7).
Figure 9 is depicting a significant increase in the colonic TNF-α in the colitis control group. Dizocilpine at all doses was effective to reduce the level of TNF-α compared to Sham group. No significant difference was found for any of the investigated cytokines between L-glutamic acid group and colitis control group. As expected, dexamethasone treatment reduced all investigated cytokines compared to colitis control group (at least $P < 0.05$).

Discussion
This study described anti-inflammatory effects of dizocilpine, an uncompetitive NMDA antagonist, on colitis severity. Our study showed that NMDA receptor antagonists could have potential benefit in the management and treatment of UC because our study demonstrated that dizocilpine decreased gross and microscopic parameters of colitis after a four-day treatment.

TNBS induces a chronic colitis resembling human IBD with immunological background and the symptoms of abdominal pain, anorexia, weight loss, anemia, colonic edema, erythema, ulcer, and necrosis (23). Besides overproduction of nitric oxide (24), pro-inflammatory cytokines and MPO (25) occurred in this model.

There are reports suggesting that peripheral NMDA receptors play a key role in peripheral inflammation responses such as T cell development, edema formation, and muscle inflammatory reactions (6, 26). NMDA receptors are frequently detected within the colon (27). So, we suggested that an additional role for NMDA receptors in colitis pathogenesis would be probable.

Present study showed that dizocilpine, an open NMDA ion channel antagonist, improves colon inflammation parameters including severe weight loss, increased release of pro-inflammatory cytokines and infiltration of inflammatory cells, and distorted mucosal structure in colitic mice. It was found that two greater tested doses were nearly effective to reduce most colitis markers; however, there was no clear relationship between the dose and related response. By considering the wide range of dizocilpine dosage applied in this study, it is concluded that a ceiling effect could be considered for NMDA implementation in colitis prevention and/or therapy.

The pathogenesis of UC remains unclear, but overproduction of pro-inflammatory cytokines, such as IL-1β, IL-6, and TNF-α are believed to play a critical role in inflammation formation (28).

There are several evidences that TNF-α plays a critical role in the pathogenesis of UC especially in TNBS model. TNF-α also induces the release of other cytokines (29). Indeed previous researches on experimental colitis have shown a positive correlation of TNF-α, IL-1β, and IL-6 levels in animal with colitis and the degree of colonic inflammation. So, it is plausible that by inhibition of synthesis or release of these cytokines, dizocilpine could prevent or cause a protection against colitis.

Results of the present study are in agreement with the previous works in which dizocilpine and other NMDA antagonists such as memantine and ketamine have shown efficacy in preventing inflammatory processes (7, 29). Dizocilpine significantly inhibited IL-1β induced gene expression of cytokines and enzymes involved in cartilage damage and inflammation (14). On the other hand, dizocilpine blocked NMDA receptor-mediated production of nitric oxide and subsequently pro-inflammatory cytokines (30). In another work, memantine attenuated CNS inflammation following endotoxin exposure in rat (31). Ketamine decreased TNF-α release from macrophage in response to bacterial lipopolysaccharides (32). One work also indicated that ketamine decreases in-vitro TNF-α secretion into whole blood after exposure to bacterial endotoxins (33).

Other possible mechanism of anti-colic effects of dizocilpine is via interaction with nitric oxide synthase enzyme by decreasing in calcium entry through NMDA receptor-operated channels (34). This interaction can lead to decrease in level of nitric oxide (NO) related oxidative stress and tissue damage partially mediated by peroxynitrite, a strong and relatively long-lived oxidant that is produced by reaction between NO and superoxide (35).

NMDA receptors are involved in regulation of intracellular calcium and ROS levels in
neutrophils (6). Intraperitoneal administration of dizocilpine decreased neutrophil activation indicated by MPO activity. This result is in accordance with previous reports about effects of NMDA antagonist on neutrophils function (6).

As noted in introduction, the main etiology of UC is interaction among antigen presenting cells, colon flora, and T lymphocytes. However, how much glutamate and NMDA receptors may be involved in this interaction, remains to be cleared.

It is worthy to be mentioned that oral administration of L-glutamic acid, a NMDA receptor agonist was not able to elevate or change the colitis markers; likely because of special circumstances such as no requirement for endogenous glutamate for formation of inflammation due to enough activation of NMDA receptors by exogenous glutamate, down regulation of NMDA receptors due to exogenous glutamate administration, limited glutamate delivery to colon due to rapid intestinal absorption and high hepatic clearance.

**Conclusion**

Taken together, dizocilpine showed a protective and anti-inflammatory effect on all features of experimental colitis. It is indicated that NMDA suppression may be a new and effective therapy target in UC patients. However, further investigations are necessary to evaluate whether a similar efficacy can be achieved by other NMDA antagonists and in how much, this mechanism could be effective in clinical setting.

**Acknowledgment**

Financial support for this study was received from the School of Pharmacy and Pharmaceutical Sciences, Isfahan University of Medical Sciences, Isfahan, Iran [grant number 394789].

**References**

(1) Xavier R and Podolsky D. Unravelling the pathogenesis of inflammatory bowel disease. *Nature* (2007) 448: 427-34.
(2) Mudter J and Neurath MF. IL-6 signaling in inflammatory bowel disease: Pathophysiological role and clinical relevance. *Inflam. Bowel Dis.* (2007) 13: 1016-23.
(3) Bochner BS, Luscinaskas FW, Gimbrone M, Newman W, Sterbinsky S, Dersc-Anthony CP, Klunk D and Schleimer RP. Adhesion of human basophils, eosinophils, and neutrophils to interleukin 1-activated human vascular endothelial cells: contributions of endothelial cell adhesion molecules. *J. Exp. Med.* (1991) 173: 1553-57.
(4) Kornbluth A and Sachar DB. Ulcerative colitis practice guidelines in adults (update): American College of Gastroenterology, Practice Parameters Committee. *Am. J. Gastroenterol.* (2004) 99: 1371-85.
(5) McKenna JE and Melzack R. Blocking NMDA receptors in the hippocampal dentate gyrus with AP5 produces analgesia in the formalin pain test. *Exp. Neurol.* (2001) 172: 92-9.
(6) Boldyrev A, Bryushkova E and Vladychenskaya E. NMDA receptors in immune competent cells. *Biochemistry (Moscow).* (2012) 77: 128-34.
(7) SS, Mydel P, Hellward A, Jonsson IM and Bokarewa MI. The N-methyl-d-aspartic acid receptor antagonist memantine ameliorates and delays the development of arthritis by enhancing regulatory T cells. *Neurosignals* (2011) 20: 61-71.
(8) Dale O, Somogyi AA, Li Y, Sullivan T and Shavit Y. Does intraoperative ketamine attenuate inflammatory reactivity following surgery? A systematic review and meta-analysis. *Anesth. Analg.* (2011) 112: 934-43.
(9) Mozayani A, Schrode P, Carter J and Danielson TJ. A multiple drug fatality involving MK-801 (dizocilpine), a mimic of phencyclidine. *Forensic. Sci. Int.* (2003) 133: 113-17.
(10) Imen M, Omay B, Uzunkol A, Erdem T, Sabanci PA, Karasu A, Albayrak SB, Sencer A, Hepgul K and Kaya M. Effect of magnesium, MK-801 and combination of magnesium and MK-801 on blood–brain barrier permeability and brain edema after experimental traumatic diffuse brain injury. *Neuro. Res.* (2009) 31: 977-81.
(11) Kato T. Memantine: a therapeutic drug for Alzheimers disease and the comparison with MK-801. *Nihon Yakurigaku Zasshi. Folia. Pharmacol. Jpn.* (2004) 124: 145-51.
(12) Hussain N, Flumerfelt B and Rajakumar N. Glutamatergic regulation of long-term grafts of fetal lateral ganglionic eminence in a rat model of Huntington’s disease. *Neurobiol. Dis.* (2004) 15: 648-53.
(13) Hamasato EK, de Oliveira APL, Lino-dos-Santos-Franco A, Ribeiro A, de Paula VF, Peron JPS, Damazo AS, Tavares-de-Lima W and Palermo-Neto J. Effects of MK-801 and amphetamine treatments on allergic lung inflammatory response in mice. *Int. Immunopharmacol.* (2013) 16: 436-43.
(14) Piepoli T, Mennuni L, Zerbi S, Lanza M, Rovati L
and Caselli G. Glutamate signaling in chondrocytes and the potential involvement of NMDA receptors in cell proliferation and inflammatory gene expression. Osteoarthritis Cartilage (2009) 17: 1076-83.

(15) Šulkowska G, Dąbrowska-Bouta B, Chalimoniuk M and Struzyska L. Effects of antagonists of glutamate receptors on pro-inflammatory cytokines in the brain cortex of rats subjected to experimental autoimmune encephalomyelitis. J. Neuroimmunol. (2013) 261: 67-76.

(16) Oba M, Koguchi S and Nishiyama K. A concise diasteroselective synthesis of (2S, 3S, 4R)-3, 4-dihydroxyglutamic acid. Tetrahedron Lett. (2001) 42: 5901-02.

(17) Motaghi E, Hajhashemi V, Mahzouzi P and Minaiyan M. The effect of memantine on trinitrobenzene sulfonic acid-induced ulcerative colitis in mice. Eur. J. Pharmacol. (2016) 793: 28-34.

(18) Motavallian-Naeini A, Andalib S, Rabbani M, Mahzouzi P, Afsharipour M and Minaiyan M. Validation and optimization of experimental colitis induction in rats using 2, 4, 6-trinitrobenzene sulfonic acid. Res. Pharm. Sci. (2012) 7: 159-69.

(19) Wallace JL, Mac Naughton W, Morris G and Beck P. Inhibition of leukotriene synthesis markedly accelerates healing in a rat model of inflammatory bowel disease. Gastroenterology (1989) 96: 29-36.

(20) Minaiyan M, Sajadi S-E, Naderi N and Taheri D. Anti-inflammatory effect of Kelussia odoratissima mosaffa. hydroalcoholic extract on acetic acid-induced acute colitis in rats. J. Rep. Pharm. Sci. (2014) 3: 28-35.

(21) Kim JJ, Shahjib MS, Manocha MM and Khan WI. Investigating intestinal inflammation in DSS-induced model of IBD. J. Vis. Exp. (2012) e3678.

(22) Ishiguro K, Ando T, Maeda O, Hasegawa M, Kadomatsu K, Ohmiya N, Niwa Y, Xavier R and Goto H. Paeonol attenuates TNBS-induced colitis by inhibiting NF-kB and STAT1 transactivation. Toxicol. Appl. Pharmacol. (2006) 217: 35-42.

(23) Cronin CC and Shanahan F. Understanding symptoms and signs in inflammatory bowel disease. Inflammatory Bowel Diseases. Springer (2005) 253-67.

(24) Tun X, Yasukawa K and Yamada K-i. Involvement of nitric oxide with activation of Toll-like receptor 4 signaling in mice with dextran sodium sulfate-induced colitis. Free Radic. Biol. Med. (2014) 74: 108-17.

(25) Allhagamhammad MH, Lemberg DA, Day AS, Tan L-Z, Ooi CY, Krishnan U, Gupta N, Munday JS and Leach ST. Advancing nutritional therapy: A novel polymeric formulation attenuates intestinal inflammation in a murine colitis model and suppresses pro-inflammatory cytokine production in ex-vivo cultured inflamed colonic biopsies. Clin. Nutr. (2016) 36: 497-505.

(26) Ro JY. Contribution of peripheral NMDA receptors in craniofacial muscle nociception and edema formation. Brain Res. (2003) 979: 78-84.

(27) Zhou Q, Caudle RM, Price DD, Del Valle-Pinero AY and Verne GN. Selective up-regulation of NMDA-NR1 receptor expression in myenteric plexus after TNBS induced colitis in rats. Mol. Pain (2006) 2: 3-10.

(28) Menegat JSB, Lira-Junior R, Siqueira MA, Brito F, Carvalho AT, Fischer RG and Figueredo CM. Cytokine expression in gingival and intestinal tissues of patients with periodontitis and inflammatory bowel disease: An exploratory study. Arch. Oral. Biol. (2016) 66: 141-6.

(29) Loix S, De Kock M and Henin P. The anti-inflammatory effects of ketamine: state of the art. Acta Anaesthesiol. Belg. (2011) 62: 47-58.

(30) Vladychenskaya E, Tyulina O, Urano S and Boldyrev A. Rat lymphocytes express NMDA receptors that take part in regulation of cytokine production. Cell Biochem. Funct. (2011) 29: 527-33.

(31) Rosi S, Vazdarjanova A, Ramirez-Amaya V, Worley P, Barnes C and Wens G. Memantine protects against LPS-induced neuroinflammation, restores behaviorally-induced gene expression and spatial learning in the rat. Neurosciences (2006) 142: 1303-15.

(32) Spentzas T, Shapley RK, Aguirre CA, Meals E, Lazar L, Rayburn MS, Walker BS and English BK. Ketamine inhibits tumor necrosis factor secretion by RAW264. 7 murine macrophages stimulated with antibiotic-exposed strains of community-associated, methicillin-resistant Staphylococcus aureus. BMC Immunol. (2011) 12: 11.

(33) Kawasaki T, Ogata M, Kawasaki C, Ogata Ji, Inoue Y and Shigematsu A. Ketamine suppresses proinflammatory cytokine production in human whole blood in-vitro. Anesth. Analg. (1999) 89: 665-9.

(34) Bhardwaj A, Northington FJ, Carhuapoma JR, Falck JR, Harder DR, Traystman RJ and Koehler RC. P-450 epoxygenase and NO synthase inhibitors reduce cerebral blood flow response to N-methyl-D-aspartate. Am. J. Physiol. Heart. Circ. Physiol. (2000) 279: 1616-24.

(35) Conners W, Whitebeck C, Chicester P, Legget R, Johnson A, Kogan B, Levin R and Mannikarottu A. L-NAME, a nitric oxide synthase inhibitor, diminishes oxidative damage in urinary bladder partial outlet obstruction. Am. J. Physiol. Ren. Physiol. (2006) 290: 357-63.

This article is available online at http://www.ijpr.ir