Esculentoside A could attenuate apoptosis and inflammation in TNBS-induced ulcerative colitis via inhibiting the nuclear translocation of NF-κB

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Background: Esculentoside A (EsA) has had a remarkable curative effect on a variety of experimental acute and chronic inflammatory and autoimmune diseases. However, the role of EsA in the pathological process of ulcerative colitis (UC) is still unknown.

Methods: Rat colonic smooth muscle cells (SMCs) were identified by immunofluorescence. The effect of EsA and/or lipopolysaccharide (LPS) on the viability, proliferation, and apoptosis of SMCs was explored via 3-(4,5-dimethylthiazolyl-2)-2,5-diphenyltetrazolium bromide (MTT), 5-ethynyl-2'-deoxyuridine (EdU) staining, flow cytometry, and TdT-mediated dUTP nick end labeling (TUNEL) staining, respectively. The changes of apoptosis-related proteins were performed via western blotting. The expression and nuclear translocation of NF-κB were detected via western blotting, immunohistochemistry (IHC), and immunofluorescence staining, respectively. Enzyme-linked immunosorbent assay (ELISA) was used to detect the expression of IL-6 and TNF-α.

Results: The EsA treatment greatly up-regulated the viability of LPS-suppressed SMCs. The LPS-induced cell apoptosis was significantly reversed by EsA treatment, which was achieved via down-regulating Bax and cleaved caspase-3 expression and up-regulating Bcl-2 expression. In addition, LPS-induced IL-6, TNF-α expression and NF-κB activation were also largely decreased when treated with EsA. In vivo, the TNBS-induced colon injury including crypt destruction and crypt deformation, disorder, epithelial cell remains or complete destruction, and inflammatory cell infiltration was recovered by EsA treatment. The secretion of IL-6 and TNF-α in the serum of the model group was also down-regulated by EsA treatment. The expression of Bax, cleaved caspase-3, and Bcl-2 showed similar trends as those observed in the in vitro experiments.

Conclusions: Our data provides supportive evidence that EsA can relieve the symptoms of UC and be used as a drug candidate for the treatment of UC.

Keywords: Esculentoside A (EsA); colitis; p65; Bax; cleaved caspase-3

Submitted Apr 29, 2022. Accepted for publication Jun 28, 2022.
doi: 10.21037/atm-22-2675

View this article at: https://dx.doi.org/10.21037/atm-22-2675
Introduction

Ulcerative colitis (UC) (1), as a kind of recurrent chronic nonspecific inflammatory disease, is characterized by repeated episodes of stools containing mucous and blood, abdominal pain, and diarrhea (2). Due to its complicated etiology, difficult cure, high recurrence rate, poor prognosis, and increased risk of colon cancer, UC is considered a refractory disease (3,4). Currently, the clinical treatment of UC mainly involves glucocorticoids, 5-aminosalicylic acid, sulfasalazine, mesalazine, and so on, which achieve their therapeutic effect via reducing the intestinal inflammation, promoting intestinal mucosal repair, and improving the intestinal mucosal barrier (5-7). However, these drug treatments cannot effectively inhibit the development of UC, especially in the prevention of recurrence, and long-term use of these drugs often leads to significant side effects (8). For example, non-steroidal anti-inflammatory drugs (NSAIDs) have a great impact on the gastrointestinal tract, and long-term use of glucocorticoids often leads to immune suppression and adrenal cortical suppression (9,10). Considering that multiple cellular signaling pathways, especially NF-κB pathway, oxidative stress, activation of pyrin domain-containing protein 3 (NLRP3) inflammasome and autophagy all take part in the pathogenesis of UC (11), searching safe and effective drugs that can improve inflammatory response and inhibit NF-κB pathway are useful for provide new choice for UC treatment.

To date, a large number of studies have been conducted to seek naturally-sourced drugs for the treatment of UC in order to replace or supplement the available drug options, and lots of naturally-sourced drugs are discovered to exert beneficial influence on UC treatment (12,13). Traditional Chinese medicine (TCM) has the characteristics of multipotency, bidirectional regulation, and fewer adverse reactions (14). It is of great significance to discover new anti-inflammatory drugs derived from natural products with less adverse effects to treat with UC. Esculentoside A (EsA), as the main active ingredient extracted from *Euphorbiae sapotin*, has a wide range of pharmacological effects such as anti-inflammatory, diuretic, and immunological regulation (3,4). Ma et al. showed that EsA can play an inhibitory role in nephritis via anti-inflammatory and pro-apoptotic mechanisms (15). It also exerts an inhibitory effect on various respiratory inflammatory conditions by inhibiting the inflammatory cytokines and signaling pathway-related proteins including the PPAR-γ, NF-κB, and ERK signal pathways and the Nrf-2 pathway (16,17). A study has also demonstrated that EsA suppresses neuroinflammation in the hippocampus of model mice via decreasing the phosphorylation levels of ERK, JNK, and p38 (18). Although EsA exerts an anti-inflammatory effect in various diseases, its function in UC is still unknown.

Often, UC is accompanied by diarrhea, abdominal pain, and abnormal intestinal motor function (3). The biological dysfunction of intestinal smooth muscle cells (SMCs) often leads to abnormal intestinal motility, eventually resulting in abnormal gastrointestinal motility (19). Although the SMCs play the most important role in the progression of UC, most researchers have focused on the intestinal dynamics of SMCs in UC, and studies regarding the apoptosis and inflammation of UC have mainly focused on the intestinal epithelial cells (20). Therefore, we aimed to explore the effects of EsA on the inflammation and apoptosis of SMCs.

In this study, we used lipopolysaccharide (LPS) to treat SMCs to establish the inflammatory cells and 2,4,6-trinitrobenzene sulfonic acid (TNBS) to construct a rat UC model. We then studied the effect of EsA on apoptosis and inflammation in vitro and in vivo, with the aim of providing evidence that EsA could be used as an effective drug for the treatment of UC. To the best of our knowledge, this is the first study concerning the possible beneficial effect of EsA on UC. The findings of our research may offer experimental evidence for further understanding the suppressive effect of EsA on inflammation in UC. We present the following article in accordance with the ARRIVE reporting checklist (available at https://atm.amegroups.com/article/view/10.21037/atm-22-2675/rc).

Methods

Animals

A total of 32 male Wistar rats (180–200 g) were obtained from the Zhejiang Vital River Laboratory Animal Technology Co., Ltd. (Jiaxing, China). We purchased EsA from Shanghai Yuanye Biological Co., Ltd. (Shanghai, China). The animals were raised in plastic cages, the relative humidity of the room was maintained at 50–60%, and the temperature at 20–23 °C under 12-hour light/dark cycles. All rats were given free access to water and food. The animal experiments were approved by the Institutional Animal Ethics Committee of Guilin Medical University (No. GLMC202203270), in compliance with Guide for the Care and Use of Laboratory Animals, 8th edition. A protocol was prepared before the study without registration. In this study, the 32 rats were randomly divided into four
groups, as follows: (I) sham group (n=8; normal saline); (II) model group (n=8; normal saline + TNBS); (III) model + negative control (NC) group [n=8; normal saline + TNBS + double-distilled H₂O (ddH₂O)]; and (IV) model + EsA group (n=8, normal saline + TNBS + EsA).

**TNBS-induced UC**

Rats were anesthetized by intraperitoneal injection of 10% chloral hydrate (0.3 mL/kg) after fasting for 24 hours with free access to water. We induced UC in the rats by gentle and slow intrarectal administration of TNBS (Sigma-Aldrich, St. Louis, MO, USA; 100 mg/kg dissolved in 0.25 mL of 50% ethanol) using a flexible gastric lavage needle inserted 8 cm into the colon. Then, the rats were kept head down for 1 minute and lain on their backs in the cage. The Sham model group was administered with 0.25 mL saline as a control. In the model + EsA group, the rats were intraperitoneally injected with EsA (20 mg/kg, Shanghai Yuanye Biological Co., Ltd.) immediately after the UC model was induced and the rats were sacrificed after 10 days of continuous injection of EsA (21). In the model + NC group, the model rats were administered with an equal volume of saline as a NC.

**Isolation of colonic SMCs**

Rats were euthanized by CO₂ inhalation, abdominal incisions were made vertically, about 10 cm of the colons were quickly removed starting from 2 cm distal to the anus and placed in normal saline, enema irrigation was repeatedly applied, and the serum was obtained for the subsequent measurements. Then, the colons were placed in HEPES-buffered Ringer solution (G-clone, Beijing, China) containing 300 U/mL streptomycin (Sigma-Aldrich, USA) and soaked for 15 minutes. The mucosal and serosa layers were gently scraped from the muscle layer by mechanical erosion until the remaining tissue layer was transparent under the stereomicroscope. The tissue layer was cut up and homogenized, added to digestive fluid (4 mL) containing the 0.1% type II collagenase (Worthington Biochemical Corp., Lakewood, NJ, USA) and 0.01% soybean trypsin inhibitor (Worthington, USA), then incubated at 30 °C for 20 minutes, centrifuged, and the supernatant was discarded. The precipitates were again digested by digestive fluid (6 mL) at 30 °C for 30 minutes. The partially digested tissues were washed using collagenase-free medium and collected via centrifuging at 1,000 r/min for 5 minutes. After that, the partially digested tissues were resuspended in Dulbecco’s modified Eagle medium (DMEM) and the SMCs were collected via filtration using 100 mesh cell sieves. The SMCs were plated at a concentration of 1x10⁵ cells/mL and cultured with DMEM (Gibco, Grand Island, NY, USA) medium containing streptomycin (200 μg/mL), penicillin (Sigma-Aldrich, 200 U/mL), amphotericin B (Sigma-Aldrich, 2.5 μg/mL), gentamycin (Sigma-Aldrich, 100 μg/mL), and 10% fetal bovine serum (FBS) (HyClone, Logan, UT, USA) at 37 °C with 5% CO₂. The cell viability was detected using the Trypan blue assay (Thermo Fisher Scientific, Waltham, MA, USA; 0.4%, w/v). The cells were cultured for 24 hours, then the medium was replaced and the non-adherent cells were discarded. After that, the DMEM medium was replaced every 3–4 days as needed until reaching about 90% confluence.

**Cell treatment**

In this study, the in vitro experiments were divided into 3 parts. In the first part, the SMCs cells were pretreated with EsA (0, 4.5, 9, and 18 μM; Topscience, Shanghai, China) for 1 hour and then treated with LPS (Sigma-Aldrich, 1 μg/mL) for 24 hours. In the second part, the SMCs cells divided into four groups: (I) control group; (II) EsA (9 μM) group (cells were treated with EsA for 24 h); (III) LPS group (1 μg/mL 24 h); and (IV) LPS (1 μg/mL) + EsA (9 μM) group (cells were co-treated with LPS and EsA for 24 h). In the third part, the SMCs cells divided into four groups: (I) control; (II) LPS (1 μg/mL) group (cells were treated with LPS for 24 h); (II) LPS (1 μg/mL) + NC group (cells were pretreated with LPS for 24 h and then treated with solvent control); and (IV) LPS (1 μg/mL) + EsA (9 μM) group (cells were pretreated with LPS for 24 h and then treated with EsA for 2 h).

**Immunofluorescence staining**

Generally, SMCs were fixed, permeabilized, blocked, and then incubated with anti-α-actin [Cell Signaling Technology (CST), Danvers, MA, USA; 6487, 1:100 dilution] or anti-p65 (CST, 69994, 1:400 dilution) antibodies at 4 °C overnight. Then, cells were stained with FITC-conjugated immunoglobulin G (IgG; 1:100 dilution) and the nucleus were stained with 4’,6-diamidono-2-phenylindole (DAPI). Lastly, the pictures were captured using a fluorescence microscope (Olympus, Tokyo, Japan) at 400x magnification.
3-(4,5-dimethylthiazolyl-2)-2,5-diphenyltetrazolium bromide (MTT) assay

The total of 3×10^3 SMCs were seeded into 96-well plates and treated as described previously. The cell viability was tested using a MTT assay kit (Sangon Biotech Co., Ltd.) at 24 hours after drug treatment, according to the manufacturer instructions. Lastly, the MTT in cells were dissolved with 150 μL dimethyl sulfoxide (DMSO; Sigma-Aldrich, USA) and the absorbance at 490 nm was measured using a microplate reader (BMG LABTECH, Offenburg, Germany).

EdU staining

The treated SMCs cells were labeled with 50 μM 5-ethynyl-2′-deoxyuridine (EdU) (Thermo Fisher Scientific, USA) for 2 hours according to the protocol, and the cells nuclei were stained with DAPI. The cells were observed using a fluorescence microscope (Olympus, Japan) with the scale bar set at 50 μm.

Cell apoptosis assay

After treatment as previously described, SMCs were collected and re-suspended with 600 μL binding buffer containing 1% propidium iodide and Annexin V-FITC (Thermo Fisher Scientific, USA), then the cells suspension was incubated away from light for 20 minutes. A flow cytometer (Beckman Coulter, Miami, FL, USA) was used to analyze cell apoptosis.

Nuclear and cytoplasmic separation

Nuclear and cytoplasmic separation were carried out using Nuclear and Cytoplasmic Extraction Reagents (CER; Thermo Fisher Scientific, USA). We used CER I to resuspend the cells and incubate on ice for 10 minutes, and then CER II was added to the tube and incubated on ice for 1 minute. Then, the supernatant (cytoplasmic extract) was obtained by centrifuging the tube at maximum speed and the insoluble (nuclear pellets) fractions were extracted with the nuclear extraction reagent (NER) on ice and stored at −80 °C until use.

Western blotting

Briefly, SMCs isolated from the colon were solubilized in radioimmunoprecipitation assay (RIPA) lysate (Beyotime, Nanjing, China) containing protease and phosphatase inhibitors. The concentrations of protein were determined with the bicinchoninic acid (BCA) kit (Beyotime). The proteins were fractionated by sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE) and the separated proteins were transferred to polyvinylidene fluoride (PVDF) membranes. The blots were incubated with the primary antibodies against Bax (CST, 14796, 1:1,000), Bcl-2 (Abcam, ab194583, 1:1,000), cleaved caspase-3 (Invitrogen, Carlsbad, CA, USA; PA5-114687, 1:1,000), p65 (CST, 69994, 1:1,000), GAPDH (CST, 5174, 1:1,000), β-actin (CST, 4970, 1:1,000), and Lamin B1 (CST, 13435, 1:1,000) at 4 °C for 12 hours, and then incubated with secondary antibody conjugated with horseradish peroxidase (HRP) for 2 hours. Then, the protein bands were visualized by ECL-Plus reagent (Thermo Fisher Scientific, USA) using the Gel Imaging System.

Enzyme-linked immunosorbent assay (ELISA)

The inflammatory cytokines IL-6 and TNF-α in the supernatant and serum were detected via ELISA. All the detection reagents were obtained from R&D Systems (Minneapolis, MN, USA).

Hematoxylin and eosin (H&E) staining and histological score

The colonic tissues were collected and fixed with 4% paraformaldehyde immediately after the rats were sacrificed. The 4-μm colonic tissues sections were stained with H&E solution (H8070, Solarbio, Beijing, China) according to the standard procedures. The images were captured using a microscope (Nikon, Japan) at 200x and 400x magnification, respectively. The TNBS-induced colitis model was assessed using histological scoring system. Briefly, each colon was scored in consideration of (I) the extent of inflammation; (II) the severity of inflammation; and (III) crypt damage. The total score of histology was between 0 and 10. The specific scoring criteria were as described by Xiao and Fattahi et al. (22,23).

TdT-mediated dUTP nick end labeling (TUNEL) staining

Briefly, the colonic tissues on slides (4 μm) were colored using TUNEL probes (Beyotime, Nanjing, China), and the cell nuclei were stained with DAPI. A fluorescence
microscope (Olympus, Tokyo, Japan) was used to capture the images at 400× magnification.

**Immunohistochemistry (IHC)**

Briefly, the colonic tissues on slides (4 μm) were treated according to the instructions. The sections were incubated with a p65 antibody (CST; 1:100) at 4 °C overnight, and then incubated with a secondary antibody at room temperature for 2 hours. Immuno-reactivity was visualized by using the 3,3-diaminobenzidine (DAB) chromogen. The sections were stained with hematoxylin and examined under a microscope (Olympus, Japan).

**Statistical analysis**

The software GraphPad Prism 7.0 (GraphPad Software Inc., San Diego, CA, USA) was used to process all data, and mean ± standard deviation was used to evaluate the data. One-way analysis of variance (ANOVA) was used to compare the multiple groups to evaluate the statistical significance. A P value <0.05 was considered as statistically significant.

**Results**

The effect of LPS on apoptosis and proliferation in colonic SMCs were partially reversed by EsA

We selected EsA due to it having the highest content of saponins in the Euphorbiae family and the structure is shown in Figure 1A. It has a remarkable curative effect in inflammatory diseases (24). However, the role of EsA in the pathological process of colitis is still unknown. To study the effect of EsA on colitis, the SMCs were isolated from rat colons and identified via immunofluorescence staining. The data showed that the isolated cells expressed α-actin, which is a marker of SMC cells (Figure 1B), indicating that the SMC cells had been isolated from the rat colons successfully. To explore the role of EsA on colitis, the SMCs were pretreated with EsA and then treated with LPS to induce the colitis model in vitro. As the results demonstrated that EsA could effectively improve cell viability which reduced by LPS in a dose-dependent manner, especially for 24 and 48 hours (Figure 1C). Moreover, the SMC cells were treated with LPS or EsA alone or in combination to explore the cell proliferation and apoptosis. The data showed that co-treatment with EsA and LPS could effectively increase the cell viability reduced by LPS (Figure 1D). Furthermore, co-treatment with EsA and LPS could effectively decrease the cell apoptosis increased by LPS in SMCs (Figure 1E,1F). The EdU staining showed the opposite results in cell proliferation (Figure 1G). All the data indicated that the apoptosis and proliferation of SMCs induced by LPS were greatly reversed by EsA.

The apoptosis and inflammation induced by LPS were largely attenuated by EsA via inhibiting the nuclear translocation of NF-κB in vitro

To further explore the effect of EsA in apoptosis and inflammation induced by LPS, the cell apoptosis and inflammatory factors including IL-6 and TNF-α were detected. As the results showed that the cell apoptosis induced by LPS were partly reduced by EsA (Figure 2A,2B). In addition, the expression of Bax and cleaved caspase-3 which was increased by LPS were partly decreased by EsA; however, the expression of Bcl-2 showed the opposite results (Figure 2C). Moreover, the secretion of IL-6 and TNF-α promoted by LPS were largely inhibited by EsA (Figure 2D,2E). Furthermore, the western blot and immunofluorescence staining showed that LPS could activate NF-κB and promote its nuclear translocation, and that EsA could inhibit the nuclear translocation of NF-κB induced by LPS (Figure 2F,2G).

EsA treatment attenuated TNBS-induced UC in rats

The histological data showed that muscularis propria, mucosa, submucosal, and serosa in the colon of the control group were normal in morphology, without inflammatory cells (Figure 3A). Conversely, the TNBS group and TNBS + NC group displayed disrupted colonic architecture, and infiltration of inflammatory cells with depletion of crypt and thickened muscle layer. Nevertheless, the EsA treatment group reduced the severity of TNBS-induced intestinal damage (Figure 3A,3B). Furthermore, the inflammatory cytokines TNF-α and IL-6 in serum of TNBS-induced UC rats were markedly increased when compared with the control group. Moreover, EsA notably inhibited the secretion of TNF-α and IL-6 in TNBS induced UC rats (Figure 3C,3D).

EsA treatment reduced TNBS-induced apoptosis and NF-κB p65 expression in colon tissue

The apoptosis and apoptosis-related proteins in colon
Figure 1 EsA attenuated the effect of LPS on apoptosis and proliferation in vitro. (A) The structure of EsA. (B) The expression of α-actin in SMC cells isolated from rat colon were detected via immunofluorescence staining. (C,D) The effect of LPS and EsA on cell viability were detected by MTT assay. (E,F) The effect of LPS and EsA on cell apoptosis were observed via flow cytometry. (G) The role of LPS and EsA in cell proliferation were detected via EdU staining. ***P<0.001 vs. Control; *P<0.05 vs. LPS; **P<0.01 vs. LPS; ***P<0.001 vs. LPS. DAPI, 4',6-diamidino-2-phenylindole; OD, optical density; LPS, lipopolysaccharide; EsA, esculentoside A; EdU, 5-ethynyl-2'-deoxyuridine; SMC, smooth muscle cells; MTT, 3-(4,5-dimethylthiazolyl-2)-2,5-diphenyltetrazolium bromide.
Figure 2 EsA attenuated cell apoptosis, inflammation and the NF-κB nuclear translocation in vitro. Cells were pretreated with LPS for 24 h and then treated with EsA for 2 h. (A,B) The cell apoptosis was detected via flow cytometry. (C) Western blotting was used to detect the expression of proteins. (D,E) The secretion of TNF-α and IL-6 in the supernatant were detected via ELISA assay. (F,G) The nuclear translocation of NF-κB were detected via western blotting and immunofluorescence staining, respectively. ***P<0.001 vs. Control; ##P<0.01 vs. LPS. LPS, lipopolysaccharide; NC, negative control; EsA, esculentoside A; TNF-α, tumor necrosis factor-α; IL-6, interleukin-6; DAPI, 4’,6-diamidino-2-phenylindole; ELISA, enzyme-linked immunosorbent assay; NF-κB, nuclear factor-κB.
tissues were detected via TUNEL staining and western blotting, respectively. The results demonstrated that apoptosis was increased in the TNBS group when compared with the control group, and EsA administration could reduce the apoptosis in the colon tissues of TNBS-induced UC rats (Figure 4A). Besides, the pro-apoptotic proteins including Bax and cleaved caspase-3 increased by TNBS were decreased by EsA in UC rats (Figure 4B), and the expression of anti-apoptotic protein Bcl-2 showed the opposite results (Figure 4B). Moreover, the expression of NF-κB p65 were detected via immunostaining on colonic tissues. The data showed that the expression of NF-κB p65 was up-regulated in the TNBS group when compared with the control group. However, EsA administration resulted in the reduction of the expression of NF-κB p65 compared with the model + NC group (Figure 4C).

Discussion

The bowel disease UC is a chronic, non-specific, inflammatory disease occurring in the colon and rectum with an extended duration and high recurrence rate (3). To date, studies have shown that the occurrence of UC is mainly caused by environmental deterioration, genetics, immunity, and other factors, especially the abnormal immune response of the body (3,4). Moreover, long-term unhealed UC can increase the risk of colon cancer. Inflammatory response-resulted DNA damage, oxidative stress, microsatellite and chromosomal instabilities can contribute to the UC-related carcinogenesis (25). It is of great significance to seek anti-inflammatory drugs for the treatment of UC. In this study, firstly, we used EsA to treat the LPS-induced SMC cell inflammatory damage
and demonstrated that EsA can significantly alleviate the SMC cell inflammatory injury. The symptoms and histological changes of the TNBS/ethanol model were similar to those of human UC, which could simulate the chronic recurrence process of UC, and could be used to evaluate the therapeutic effect of new treatment methods on human UC (26,27). The TNBS-induced UC model has the advantages of simple operation, good reproducibility, short

Figure 4 EsA treatment reduced TNBS-induced apoptosis and NF-κB p65 expression in colon tissue. (A) The apoptosis in colonic tissue were detected by TUNEL staining. (B) The expression of proteins in colonic tissue were detected by western blotting. (C) The NF-κB p65 expression in colonic tissue were observed via immunostaining. NC, negative control; EsA, esculentoside A; TUNEL, TdT-mediated dUTP nick end labeling; DAPI, 4',6-diamidino-2-phenylindole; TNBS, 2,4,6-trinitrobenzene sulfonic acid; NF-κB, nuclear factor-κB.
induction time, and long duration of inflammation, which
can reflect the complex process of the transformation from
acute inflammation to chronic inflammation, and create
conditions for the study of inflammation chronization and
the exploration of drug therapy (28). Therefore, consistent
with the earlier study (29), we also constructed rat UC
model via TNBS/ethanol method to explore the therapeutic
effect of EsA on UC. After treatment with 20 mg/kg EsA
for 10 days, the mucosal inflammatory cell infiltration was
significantly decreased, ulcers were healed, necrotic mucosal
epithelium was repaired, crypt abscess was alleviated or disappeared, and histological score was significantly
reduced, indicating that EsA had a good therapeutic effect
on TNBS-induced UC.

Reports have demonstrated that the imbalance between
pro-inflammatory cytokines and anti-inflammatory
cytokines disrupts the relief of inflammation in the colon
which further aggravates the pathogenesis of UC (30).
Among these cytokines, TNF-α and IL-6 are considered key
pro-inflammatory cytokines that cause intestinal mucosal
damage and lead to inflammation (31). Notably, TNF-α
could stimulate monocyte macrophages and neutrophils to
synthesize IL-6, IL-8, and other cytokines (32). Meanwhile,
IL-6, as a pro-inflammatory cytokine with a wide range of
effects, is involved in the inflammatory process and can
regulate the immune response (33,34). The excessive
release of IL-6 often leads to the inflammatory lesions in
UC. Therefore, its level could reflect the condition change
of UC and be used as a biological indicator of efficacy and
prognosis. Ju et al. (21) discovered that EsA could reduce
LPS-caused TNF-α, IL-1 and IL-6 production of peritoneal
macrophages in mice. Liu et al. (35) reported that EsA
inhibited breast cancer progression via suppressing IL-6/
signal transducer and activator of transcription 3 (STAT3)
pathway. In this study, we also showed that the levels of
TNF-α and IL-6 were effectively suppressed after treatment
with EsA in LPS-treated SMCs and TNBS-induced UC
rat model, indicating that EsA could significantly improve
the inflammatory response of mice with UC and the
inflammatory response of SMCs induced by LPS.

As a transcription factor regulates the expression of
inflammatory factors such as IL-6 and TNF-α, NF-κB
is closely related to the pathogenesis and progression of
various types of enteritis (36). The main component of NF-
κB signaling pathway, p65/p50 dimer, is in a suppressed
state when it combines with IκB in cytoplasm (37). Under
the stimulus of external signals, IκB is phosphorylated and
degraded, and the p65/p50 dimer is released and migrated
to the nucleus, thus activating the NF-κB signaling
pathway (17). Earlier literature reported that EsA treatment
inhibited the activation of NF-κB caused by LPS in kidney
tissue of mice (38). Moreover, EsA was discovered to
participated in NF-κB nuclear translocation in experimental
acute liver injury of mice (17). Here, we also showed that
LPS-promoted p65 nucleation was significantly suppressed
when treated with EsA, highlighting that EsA played its
protective role in UC via regulating NF-κB pathway. In the
in vivo experiments, we demonstrated that the up-regulated
p65 expression, which was mainly found in the muscularis
externa section in the UC group, was also greatly decreased
under the treatment of EsA, further confirming that EsA
might treat UC by regulating the NF-κB pathway.

The imbalance of Bcl-2 (an anti-apoptotic protein) and
Bax (a pro-apoptotic protein) often leads to changes in the
apoptotic ability of cells (39,40). The caspase protein family
is involved in the initiation and regulation of apoptosis,
and caspase-3 is a key enzyme in the activation of various
apoptosis-stimulating factors (1). In this study, we showed
that EsA could decrease the apoptosis rate of LPS-treated
SMCs, suppress the Bax protein level, and increase the
protein levels of Bcl-2 and caspase-3, indicating that EsA
could greatly inhibit the apoptosis of LPS-induced SMCs.
Besides, the similar trends were observed in the in vivo
experiments, further confirming that EsA could significantly
reduce the apoptosis in UC.

Intestinal mucosal immune is an important part of
human immune system (41). There are diffuse lymphoid
tissue, isolated lymphoid nodules, collective lymphoid
nodules, lymphocytes, macrophages and plasma cells in
intestinal digestive tract, which work together to form
intestinal mucosal immune system (42). Previous literature
reported that the defect or damage of intestinal mucosal
immune might be associated with UC (43). Considering
that EsA had excellent anti-inflammatory activity, we think
that EsA might also have a beneficial effect on inflammation
in intestinal mucosal immunity. In the future, we will
further explore the possible beneficial influence of EsA in
intestinal mucosal immune.

**Conclusions**

Our study showed that EsA can inhibit the inflammatory
response and cell apoptosis by regulating the NF-κB
signaling pathway thereby inhibiting inflammatory factors

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*Ann Transl Med* 2022;10(14):771 | [https://dx.doi.org/10.21037/atm-22-2675](https://dx.doi.org/10.21037/atm-22-2675)
TNF-α and IL-6 expressions in LPS-induced inflammatory cells and the rat UC model, which provided evidence that EsA might be used as a novel therapeutic drug for UC treatment. Due to the pathogenesis of UC is very complex, which involves in multiple inflammatory-related factors, intestinal cell apoptosis and autophagy, as well as intestinal mucosal immune (44), how EsA modulates the nuclear translocation of NF-κB is still unclear, more animal and clinical experiments are still needed to further explore the efficacy and safety of EsA on UC treatment in the future.

**Acknowledgments**

**Funding:** The study was funded by the National Natural Science Foundation of China (No. 81960106).

**Footnote**

**Reporting Checklist:** The authors have completed the ARRIVE reporting checklist. Available at [https://atm.amegroups.com/article/view/10.21037/atm-22-2675/rc](https://atm.amegroups.com/article/view/10.21037/atm-22-2675/rc)

**Data Sharing Statement:** Available at [https://atm.amegroups.com/article/view/10.21037/atm-22-2675/dss](https://atm.amegroups.com/article/view/10.21037/atm-22-2675/dss)

**Conflicts of Interest:** All authors have completed the ICMJE uniform disclosure form (available at [https://atm.amegroups.com/article/view/10.21037/atm-22-2675/coif](https://atm.amegroups.com/article/view/10.21037/atm-22-2675/coif)). The authors have no conflicts of interest to declare.

**Ethical Statement:** The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. The animal experiments were approved by the Institutional Animal Ethics Committee of Guilin Medical University (No. GLMC202203270), in compliance with Guide for the Care and Use of Laboratory Animals, 8th edition.

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(English Language Editor: J. Jones)

Cite this article as: Liu Y, Wei W, Liang S, Fang H, Cao J. Esculentoside A could attenuate apoptosis and inflammation in TNBS-induced ulcerative colitis via inhibiting the nuclear translocation of NF-κB. Ann Transl Med 2022;10(14):771. doi: 10.21037/atm-22-2675