

Curcumin ameliorates DSS-induced colitis in mice by regulating the Treg/Th17 signaling pathway

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Abstract. Curcumin has a therapeutic effect on ulcerative colitis, but the underlying mechanism has yet to be elucidated. The aim of the present study was to clarify the possible mechanisms. Dextran sulfate sodium-induced colitis mice were treated with curcumin via gavage for 7 days. The effects of curcumin on disease activity index (DAI) and pathological changes of colonic tissue in mice were determined. Interleukin (IL)-6, IL-10, IL-17 and IL-23 expression levels were measured by ELISA. Flow cytometry was used to detect the ratio of mouse spleen regulatory T cells (Treg)/Th17 cells, and western blotting was used to measure the nuclear protein hypoxia inducible factor (HIF)-1 α level. The results demonstrated that curcumin can significantly reduce DAI and spleen index scores and improve mucosal inflammation. Curcumin could also regulate the re-equilibration of Treg/Th17. IL-10 level in the colon was significantly increased, while inflammatory cytokines IL-6, IL-17 and IL-23 were significantly reduced following curcumin treatment. No significant difference in HIF-1 α was observed between the colitis and the curcumin group. It was concluded that oral administration of curcumin can effectively treat experimental colitis by regulating the re-equilibration of Treg/Th17 and that the regulatory mechanism may be closely related to the IL-23/Th17 pathway. The results of the present study provided molecular insight into the mechanism by which curcumin treats ulcerative colitis.

Introduction

The incidence of ulcerative colitis (UC) has recently increased (1). UC is a chronic, non-specific inflammatory disease that affects the colorectal tissue of the body (2). The abnormal immune system in the intestinal mucosa leads to an imbalance in immune cells and inflammatory cytokines, resulting in damage to the intestinal tissues (3,4). Th17 cells have been considered to be closely correlated with inflammatory bowel disease (5), and studies (6,7) have found that there is abnormal mucosal immune response in the intestine of UC, which may be closely related to the imbalance between regulatory T cells (Treg) and Th17 cells (8). Th17 affects innate and adaptive immune responses, and participates in the immunopathological process and prognosis of inflammatory bowel disease (IBD) by releasing interleukin (IL)-17 and other inflammatory cytokines (9). Previous studies have shown that severe Treg/Th17 imbalance in colitis mice has a clear correlation with the occurrence and development of colitis (10,11). The imbalance of Treg/Th17 and cytokines, such as IL-10, IL-17, IL-6 and IL-23 have a crucial role in the development of UC (12,13). The IL-23/Th17 pathway also plays an important role in the activation of Th17 cells and the release of IL-17 and other inflammatory factors (14). The hypoxia inducible factor (HIF)-1 α signaling pathway plays an important role in the development and activation of Th17 cells and other inflammatory factors, such as IL-17. HIF-1 α can also bind to forkhead box P3 (Foxp3), leading to accelerated degradation of Foxp3 and affecting the development and function of Treg (15). Recently, the abnormal Treg/Th17 ratio has become one of the possible targets for IBD therapy (16,17). Curcumin is a non-toxic all-natural compound extracted from turmeric and has many biological activities such as anti-inflammation, anti-infection, anti-coagulation and immune regulation (15,18). It has been reported that curcumin has a good therapeutic effect on experimental colitis in mice and can alleviate intestinal inflammation. It participates in anti-inflammatory, anti-oxidation and other protective effects by reducing the concentration of nitric oxide, myeloperoxidase and tumor necrosis factor (TNF)- α , and inhibiting the activation of NF- κ B. However, its therapeutic mechanism in UC remains

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to be elucidated (8,19,20). Through animal experiments, the therapeutic effect of curcumin on mice with colitis was studied at the histopathological, cytological and molecular levels, with the objective of revealing the possible mechanism of curcumin on mice with colitis and providing clinical treatment ideas. The results of the present study demonstrated that oral administration of curcumin to dextran sulfate sodium (DSS)-treated mice could inhibit the intestinal inflammation, and increase the levels of anti-inflammatory cytokines IL-10 by regulating the balance of Treg/Th17, while reducing the concentration of proinflammatory cytokines IL-23, IL-17, and IL-6. It achieved therapeutic effect through the inhibition of the IL-23/Th17 pathway (but possibly not the HIF-1 α signaling pathway).

Materials and methods

Experimental animal and treatment. SPF grade BALB/c male mice (6-7 weeks old, weight 22-26 g, n=36; Laboratory Animal Center of Southern Medical University, certificate number: SCXK Guangdong, China) were reared in a clean animal room. The temperature was 22-25°C and the relative humidity was ~55%. The environment was a 12-h light/dark cycle. Water was given as required and mice were given free access to food. DSS (average molecular weight 5,000; FUJIFILM Wako Pure Chemical Corporation) was formulated as 5% solution using sterile distilled water. Curcumin was purchased from Guangzhou QiYun Biotechnology Co., Ltd., (purity \geq 98.5%). The mice were fed adaptively for one week and randomly divided into three groups with 12 in each group: Normal group, Colitis control group and Curcumin 100 mg/kg.d treatment group (CUR group). The Normal group was given sterile distilled water for 14 days. The Colitis group and the CUR group were given 5% DSS for 7 days. Once the model was established, they were given distilled water for 7 days. On the 8th day, mice in the Normal group and the Colitis group were given 0.2 ml of 0.5% ethanol per day for 7 days, while mice in the CUR group were given (via gavage) the dose of curcumin 100 mg/kg.d in 0.2 ml of 0.5% ethanol for 7 days. All experiments on animals were approved by the Experimental Animal Ethics Committee of Southern Medical University.

Evaluation of experimental colitis in mice. Mouse daily disease activity index (DAI) was recorded according to the Murthy scoring system (21). The scoring criteria are shown in Table I. DAI = (weight loss rate, stool viscosity, and total score of invisible/visible bloody stools)/3. After the mice were sacrificed, the spleen was dissected and the blood on the spleen surface was washed with physiological saline, blotted, and weighed. Spleen weight (mg)/body weight (g) indicates spleen index (SI) (22).

Histopathology evaluation of mouse colon. Fresh colon tissue (without cecum) of each experimental group was collected and immediately placed in 10% formaldehyde. After fixing at room temperature for 48 h, the tissues were dehydrated using an ascending alcohol series at room temperature: 30% ethanol for 40 min, 50% ethanol for 40 min, 70% ethanol for 30 min, 80% ethanol for 30 min, 90% ethanol for 30 min, 95% ethanol for 30 min, 100% ethanol for 20 min and 100% ethanol for 20 min. Subsequently, the tissue sections

were cut into 5- μ m thick sections and heated at 60°C for 55 min. Then the tissues sections were embedded in paraffin and hematoxylin and eosin (H&E) staining was performed as follows: Hematoxylin for 10 min at room temperature and eosin for 2 min at room temperature. All tissue sections were observed under a BX-51 light microscope (Olympus Corporation; magnification, x200).

Flow cytometry detection of Treg/Th17 cells. The spleen of the mouse was removed from the laminar flow cabinet, and the appropriate amount of spleen tissue was taken. Subsequently, 5 ml mouse lymphocyte separation solution was added to separate the lymphocytes, and the number of lymphocytes was adjusted to 2×10^6 /tube. Samples were prepared according to the manufacturer's protocol of Th17/Treg kit (BD Biosciences). Lymphocytes were stimulated with 50 ng/ml phorbol 12-myristate 13-acetate (PMA; Sigma-Aldrich; Merck KGaA), 1 μ g/ml ionomycin (Sigma-Aldrich; Merck KGaA) and appropriate concentration of Monensin (BD Biosciences) for 5 h. Cells were collected and incubated at room temperature for 30 min with 20 μ l flow antibody in each tube after staining, fixation, membrane breaking and other steps. The main flow antibodies included mouse CD4-PerCP-Cy5.5, IL-17A-PE, and Alexa Fluor 647-FoxP3. A negative control group was also included. Samples were analyzed by flow cytometry using a BD FACSCalibur flow cytometer (BD Biosciences) and FACStation software (version FACS101; BD Biosciences).

ELISA assay for cytokine expression. A total of 50 mg of mouse colon tissue was collected and an appropriate amount of pre-cooled saline was added. Following homogenization with a glass homogenizer on ice, it was centrifuged at low temperature, and the supernatant was collected for quantification of the protein with Nanodrop 2000 (Thermo Fisher Scientific, Inc.). The concentrations of IL-6 (cat. no. DKW12-2060; Dakewe Biotech Co., Ltd.), IL-10 (DKW12-2100; Dakewe Biotech Co., Ltd.), IL-17 (DKW12-2170; Dakewe Biotech Co., Ltd.), and IL-23 (cat. no. M2300; R&D Systems, Inc.) in the intestine of mice were measured in strict accordance with the ELISA kit instructions.

Western blot analysis. A total of 100 mg of colon tissue was taken, and the nuclear protein was extracted using the NE-PER™ Nuclear and Cytoplasmic Extraction Reagents kit (Pierce; Thermo Fisher Scientific, Inc.), protease and phosphatase inhibitors were added. Protein quantification was performed by using Nanodrop 2000 quantitative analyzer (Thermo Fisher Scientific, Inc.) and the BCA method. An appropriate amount of loading buffer was added to adjust to the same concentration. Then 10% protein electrophoresis separation gel and 5% aminated gel were prepared, and 30 μ g nuclear protein was sampled. After the electrical conversion, the PVDF membrane was removed and blocked in 5% skimmed milk powder/TBST solution (0.05% Tween-20) for 1 h at room temperature. The samples were separated by electrophoresis, followed by membrane transfer, blocking, primary and secondary antibodies incubation, ECL imaging (Thermo Fisher Scientific, Inc.), and analysis. The primary antibodies were anti-mouse HIF-1 α (cat. no. 36169S; 1:200; Cell Signaling Technology, Inc.) and Lamin (cat. no. ABP0099; 1:1,500 dilution; Abbkine Scientific Co., Ltd.), and the incubation was at 4°C

Table I. Murthy scoring system.

Score	Weight loss rate	Stool viscosity	Invisible/visible bloody stools
0	(-)	Normal (Granular, shaped)	Normal
1	1-5%		
2	6-10%	Soft (paste, no adhesion to the anus)	Occult blood (+)
3	11-15%		
4	>15%	Diarrhea (water sample, adhesion to the anus)	Bloody stools

overnight. The secondary antibodies for horseradish peroxidase labeled goat anti mouse (cat. no. 79233; 1:1,500 dilution, OriGene Technologies, Inc.) incubated for 1 h at room temperature. Quantity One software (v1-D; Bio-Rad Laboratories, Inc.) was used to calculate the relative grayscale value with Lamin as the reference protein. SPSS v13.0 statistical software (SPSS, Inc.) was used for statistical analysis.

Statistical analysis. The experimental data were all expressed as mean \pm standard deviation. All experimental data were analyzed using SPSS v13.0 statistical software (SPSS, Inc.). One-way analysis of variance was used to analyze the differences between groups. S-N-K (Student-Newman-Keuls, q test) was selected to compare the mean of multiple samples. $P < 0.05$ was considered to indicate a statistically significant difference.

Results

DAI and SI in mice. No animal died during the experiment. The Normal group score was 0. Colitis group and CUR group demonstrated umbelliferous bloody stool, weight loss, and activity decreased in the first 7 d after administration of 5% DSS. As shown in Fig. 1A, compared with the Colitis group, the DAI score was significantly lower in the CUR group on the 10 to 14th day ($P < 0.05$). As shown in Fig. 1B and C, the spleen was significantly enlarged in the Colitis group, while the CUR group spleen decreased significantly and SI decreased compared with the Colitis group ($F = 71.85$, $P < 0.05$).

Mucosal inflammation. As shown in Fig. 2, compared with the Normal group (Fig. 2A-a), the Colitis group (Fig. 2A-b) clearly demonstrated an acute inflammatory response with mucosal erosion, congestion, edema, reduction of crypts and infiltration of inflammatory cells such as neutrophils. Compared with the Colitis group, mucosal inflammatory cell infiltration, erosion and edema in the CUR group (Fig. 2A-c) were significantly improved.

Treg/Th17 cell ratios. As shown in Fig. 3, compared with the Normal group, the ratio of CD4+IL-17+(Th17)/CD4+ lymphocytes was significantly increased in the Colitis group, The Treg/Th17 ratio was the lowest ($P < 0.05$). Compared with the Colitis group, Th17 lymphocytes in the CUR group were significantly decreased, and the Treg/Th17 ratio was significantly increased with the Colitis group ($F = 30.67$, $P < 0.05$).

Alterations to cytokine levels. As shown in Fig. 4, compared with the Normal group, the levels of cytokines IL-6, IL-17,

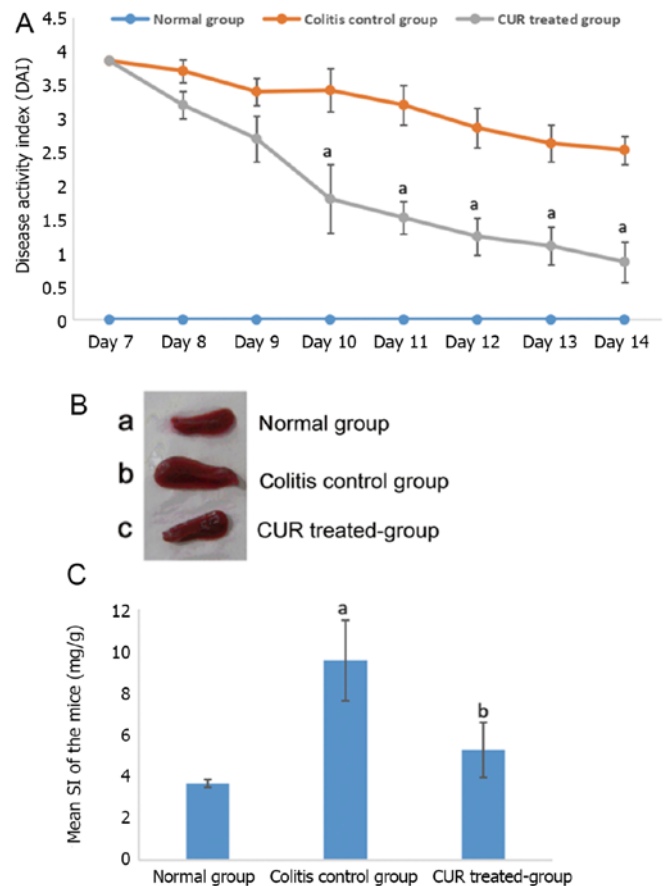


Figure 1. Effects of CUR on disease activity index and mean SI in mice. (A) Disease activity index. (B) Mean SI in the (B-a) normal group, (B-b) colitis control group and (B-c) CUR-treated group. (C) Analysis of mean SI. ^a $P < 0.05$ vs. normal group and ^b $P < 0.05$ vs. colitis control group. CUR, curcumin; SI, spleen index.

and IL-23 in the colon of mice from the Colitis group were significantly increased, while the IL-10 cytokine level was significantly decreased ($F = 67.96$, $P < 0.05$). Compared with the Colitis group, the IL-6, IL-17 and IL-23 in the colon of the CUR group were significantly decreased, while IL-10 was significantly elevated ($F = 21.36$, $P < 0.05$).

Alterations to HIF-1 α protein. As shown in Fig. 5, compared with the Colitis group and the CUR group, the expression of HIF-1 α in the colon tissue of the normal group was significantly decreased ($F = 52.18$, $P < 0.05$). The expression of the nuclear protein HIF-1 α in the colon mucosa in the Colitis group and the CUR group was significantly different ($F = 12.35$, $P < 0.05$).

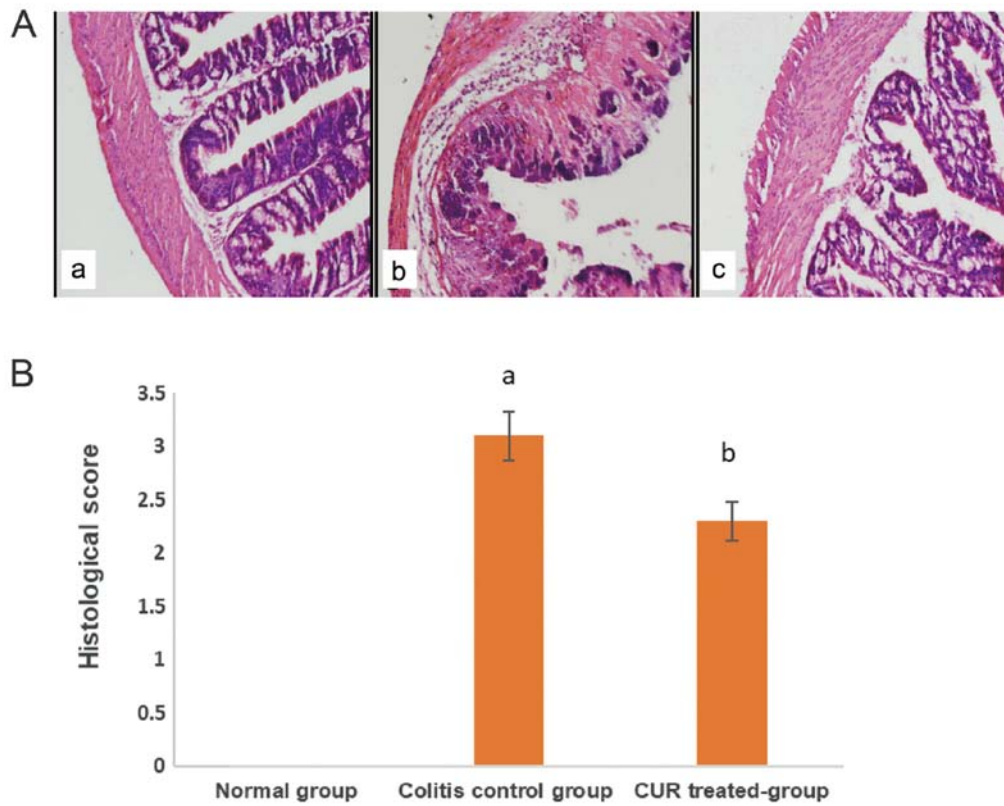


Figure 2. Effects of CUR on histological score of distal colon in mice. (A) Histological examination of colon (magnification, x100) in the (A-a) normal group, (A-b) colitis control group and (A-c) CUR-treated group. (B) Analysis of histological score * $P < 0.05$ vs. normal group and * $P < 0.05$ vs. colitis control group. CUR, curcumin.

Discussion

The present study observed increasing DAI and SI and severe mucosa erosion in colitis mice, which is similar to the symptoms of humans UC. After 7 days of curcumin treatment in experimental colitis mice, DAI and SI were significantly decreased, and colonic mucosal inflammation was significantly improved, suggesting that curcumin has a good therapeutic effect on mice with colitis. Previous studies have achieved similar therapeutic effects on colitis mice (23-25). One of the problems with curcumin in the treatment of colitis is low gastrointestinal absorption, however, IBD is a special case because curcumin does not need to be absorbed through the gastrointestinal tract to trigger a systemic therapeutic effect, but rather serves a therapeutic role through direct intestinal contact. Curcumin can significantly control the symptoms of systemic disease and intestinal inflammation in colitis mice by improving the DAI and intestinal pathology of colitis mice. The effect of curcumin on the spleen of mice attracted our attention, and thus the underlying immune mechanisms that are associated with the therapeutic effect of curcumin were further explored.

It was demonstrated that the CUR group had more significant improvement in DAI and SI compared with the colitis group. The ratio of Th17 cells increased significantly in splenic lymphocytes of colitis mice, suggesting that there was a serious imbalance in Treg/Th17 ratio in colitis. DSS-induced colitis mice experienced a disruption in Treg/Th17 balance, which led to intestinal immune disorders, one of the key factors for the

formation and progression of colitis. It was thus speculated that the spleen serves as an important immune organ and that DSS can induce immunological disorders in colitis mice and cause compensatory enlargement of the spleen. The present study further confirmed that there was a significant Treg/Th17 imbalance in the splenic lymphocytes of mice with colitis, which is consistent with human UC. Th17-related cytokines IL-6, IL-17 and IL-23 in mice with colitis was also increased, while Treg-related cytokine IL-10 was significantly decreased.

It was speculated that there were multiple mechanisms that could mediate the therapeutic effect of curcumin on colitis. First, the imbalance of Treg/Th17 occurs in a variety of inflammatory and autoimmune diseases. The present study confirmed that curcumin primarily regulates Treg/Th17 rebalance by downregulating CD4+IL-17+Th17 cells (26,27). Therefore, it was hypothesized that one of the mechanisms for the therapeutic effect of curcumin in the treatment of IBD is to regulate the re-balance of Treg/Th17. The experiments also revealed that the mechanism of curcumin in regulating Treg/Th17 re-balance may be through the inhibition of the IL-23/Th17 pathway (but possibly not the HIF-1 α signaling pathway) to downregulate of Th17. The present study demonstrated that curcumin can significantly downregulate Th17 cells. In addition, the expression of IL-23 in intestinal tissue was significantly decreased in CUR group. Therefore, it is suggested that the main mechanism of curcumin regulation of Treg/Th17 rebalance is to inhibit the IL-23/Th17 pathway. It has also been shown by previous studies that the IL-23/Th17 pathway is closely related to the development and activation

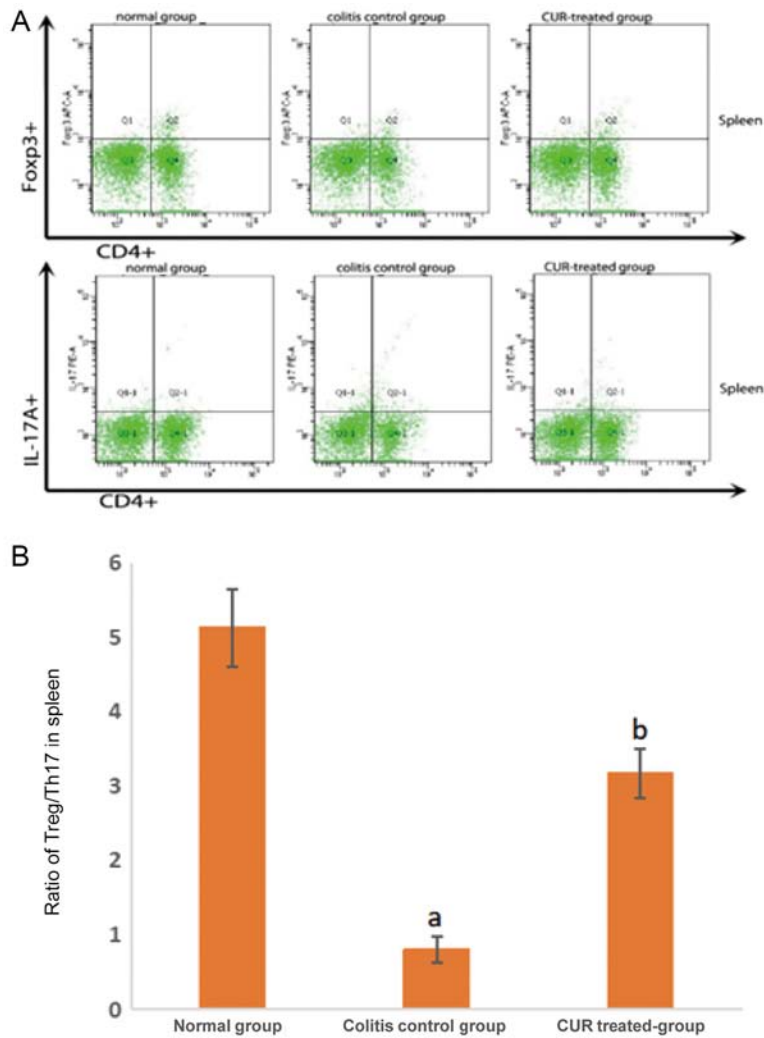


Figure 3. Effects of CUR on Treg/Th17 in the spleen. The ratio of Treg/Th17 in spleen in normal group, colitis control group and CUR-treated group was (A) determined by flow cytometry and (B) quantified. ^aP<0.05 vs. normal group and ^bP<0.05 vs. colitis control group. CUR, curcumin; CD4, cluster of differentiation 4; Fop3, forkhead box P3; IL, interleukin; Treg, regulatory T cells.

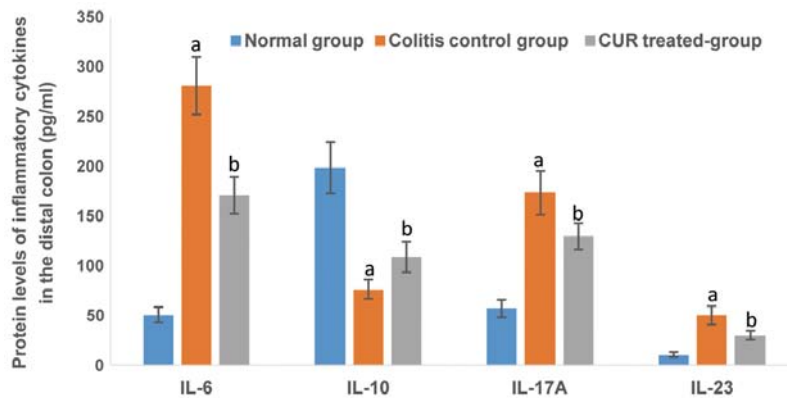


Figure 4. Effects of CUR on cytokine protein levels. Cytokine protein levels in normal group, colitis control group and CUR-treated group. ^aP<0.05 vs. normal group and ^bP<0.05 vs. colitis control group. CUR, curcumin; IL, interleukin.

of Th17 cells, and thus is associated with various diseases such as chronic inflammatory diseases and immune diseases, including rheumatoid arthritis, psoriasis and IBD (28-30). However, studies have suggested that local tissue hypoxia induces HIF-1 α synthesis, promotes its entry into the nucleus

to exert biological functions, upregulates and activates Th17, and simultaneously promotes Fop3 binding (leading to its ubiquitin and degradation) and downregulates and inhibits Treg (31,32). The present study found that curcumin produced no significant downregulation of the colon nuclear

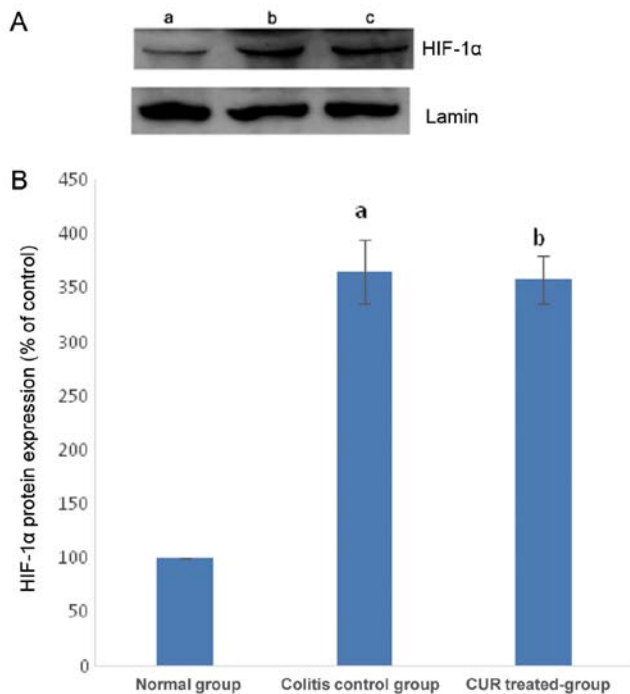


Figure 5. Effects of curcumin on HIF-1 α protein expression. (A) HIF-1 α protein expression in the (a) normal group, (b) colitis control group and (c) CUR-treated group. (B) Protein expression was semi-quantified. ^a $P < 0.05$ vs. normal group and ^b $P > 0.05$ vs. colitis control group. CUR, curcumin; HIF, hypoxia inducible factor.

protein HIF-1 α . Thus, it is hypothesized that CUR may not participate in regulating cell balance by inhibiting the HIF-1 α signaling pathway.

The present study demonstrated that the anti-inflammatory cytokine IL-10 in the CUR treatment group was elevated, and the proinflammatory cytokines IL-6, IL-17, and IL-23 were decreased. Therefore, maintaining the re-equilibration of cytokines serves an important role in curcumin treatment of IBD. IL-10 plays a negative regulatory role in cellular immunity. It can inhibit the antigen presentation and downregulate the transcription and secretion of IL-1 β , IL-6 and TNF- α and other inflammatory factors in T cells and macrophages. Ultimately, it inhibits T cell-mediated immune response and thus improves UC intestinal inflammation (4). Curcumin serves an anti-inflammatory role by elevating the level of IL-10 and reducing pro-inflammatory cytokines. It was speculated that curcumin promotes the proliferation of antigen-specific effector cells by modulating the interaction between immune cells through the regulation of multiple cytokines, and thus improves the local intestinal or systemic inflammatory response and balances immune disorders in mice with colitis (33,34).

Curcumin is an all-natural compound extracted from plants. It has many biological activities including anti-inflammatory, anti-infective and immune-regulating, and it protects the intestinal mucosa and repairs the function of intestinal tissue (35,36). Other similar studies have shown that in the DSS-induced colitis model, curcumin is found to significantly improve intestinal inflammation, repair the intestinal mucosa and inhibit the expression of TNF- α and p38MAPK (23,24).

Curcumin itself has anti-inflammatory and anti-infective effects. By inhibiting production of leukocyte eicosanoid

and related inflammatory cytokines, Treg/Th17 disorder is balanced. Through negative feedback regulation of its own immune and anti-inflammatory system, it further prevented the inflammatory cascade amplification, which serves a crucial role in the recovery of UC. At the same time, curcumin can reduce and inhibit the exudation of neutrophils and macrophages, regulate intestinal immune disorders, reduce intestinal endothelial cell swelling and increase permeability, which further reduces intestinal inflammation (4,37).

Therefore, the present study suggested that the mechanism of curcumin in treating IBD may be through downregulating the proportion of Th17 cells to regulate the re-equilibration of Treg/Th17 and Th17-related cytokines. Curcumin can significantly decrease the DAI and SI of the mice with colitis. It can regulate the re-equilibration of Treg/Th17 by inhibiting the IL-23/Th17 pathway, downregulating IL-6, IL-17, IL-23 and upregulating anti-inflammatory cytokine IL-10. Curcumin, as a good all-natural drug for the treatment of IBD, possesses good prospects in clinical application.

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Availability of data and materials

The datasets used and/or analyzed during the present study are available from the corresponding author on reasonable request.

Authors' contributions

CW designed the study and wrote the manuscript. JW and DL were laboratory personnel; these authors participated in part of the experimental process, guided part of the experimental process design and data analysis, and modified the paper. FX, BW, ML, ZY and HJ performed the histological assessments and evaluations. TL, QT, YL, DZ and ZX performed the flow cytometry and proteomics experiments. LW and JY were responsible for designing the project, guiding the experimental process, revising the paper and providing scientific research funding support. All authors read and approved the final manuscript.

Ethics approval and consent to participate

All experiments on animals were approved by the Experimental Animal Ethics Committee of Southern Medical University, China.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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