

# Curcumin use in ulcerative colitis: is it ready for prime time? A systematic review and meta-analysis of clinical trials

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## Abstract

**Background** Curcumin, an active ingredient of the Indian herb turmeric (*Curcuma longa*), has shown promising anti-inflammatory properties. Studies of its potential benefits in treating patients with ulcerative colitis (UC) are limited. We performed a systematic review and meta-analysis of human randomized placebo controlled trials to evaluate the efficacy of adjunctive therapy with curcumin in treating patients with UC.

**Methods** We conducted a search of several databases (from January 2000 to September 2018). A random-effects model was used for analysis. We assessed heterogeneity between study-specific estimates using the Cochran Q statistical test, 95% prediction interval (PI) and  $I^2$  statistics. The outcomes assessed were the pooled odds of clinical response and remission as well as the endoscopic response.

**Results** A total of 7 studies with 380 patients (curcumin n=188; placebo n=190) were included in the final analysis. The pooled odds ratio for clinical remission with curcumin use was 2.9 (95%CI 1.5-5.5,  $I^2=45$ ,  $P=0.002$ ), clinical response was 2.6 (95%CI 1.5-4.5,  $I^2=74\%$ ,  $P=0.001$ ), and endoscopic response/remission was 2.3 (95%CI 1.2-4.6,  $I^2=35.5\%$ ,  $P=0.01$ ).

**Conclusions** Based on our study, combined mesalamine and curcumin therapy was associated with roughly threefold better odds of a clinical response compared to placebo, with minimal side effects. This response was statistically significant, albeit with heterogeneity, probably due to the different severity scoring indices, curcumin dosages and routes of drug delivery used.

**Keywords** Ulcerative colitis, curcumin, meta-analysis

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## Introduction

Ulcerative colitis (UC) is a chronic inflammatory condition that presents with watery and/or hemorrhagic diarrhea associated with rectal urgency. Symptoms can be debilitating and severely affect a person's quality of life. At diagnosis, 30-50% of patients have disease confined to the rectum or the sigmoid colon (distal colitis), 20-30% have left-sided colitis and about 20% have pancolitis [1]. Of the patients with distal colitis, 25-50% progress to more extensive forms of the disease over time [2].

In early 2019, the American Gastroenterology Association (AGA) released clinical guidelines for the management of mild to moderate UC. The recommendation is to start standard dose mesalamine (2-3 g/day) or diazo-bonded 5-amino-salicylic acid (5-ASA), rather than low dose mesalamine, sulfasalazine or no treatment, in patients with extensive mild-moderate UC. The addition of rectal mesalamine to oral 5-ASA is recommended for patients with extensive or left-sided mild-to-moderate UC [3].

There have been reports regarding the efficacy of curcumin, a natural phenol found in the large-leafed herb

*Curcuma longa L.* (common names turmeric, Indian saffron) in the treatment of various diseases, such as hyperlipidemia, diabetes mellitus and non-alcoholic steatohepatitis, as well as UC [4-6]. However, the AGA made no recommendations on its use in mild-to-moderate UC patients already on a 5-ASA agent. The reason for this was stated to be a “knowledge gap”, probably from the lack of large randomized placebo controlled studies (RCTs) evaluating the efficacy of curcumin and its side-effect profile. We therefore aimed at filling this “knowledge gap” by performing a systematic review and meta-analysis of the current evidence in order to evaluate the role of combination curcumin therapy in patients with UC.

## Materials and methods

### Search strategy

We conducted a comprehensive search of several databases and conference proceedings, including PubMed, EMBASE, Google Scholar, SCOPUS and Web of Science databases, for publications from January 2000 to September 2018. We followed the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines [7], using a predefined protocol to identify studies reporting on the use of curcumin in UC. An experienced medical librarian using inputs from the study authors helped with the literature search.

Key words used in the literature search included a combination of “curcumin”, “turmeric”, “inflammatory bowel disease”, and “ulcerative colitis”. The search was restricted to studies in human subjects published in the English language in peer-reviewed journals. Two authors (BPM, SC) independently reviewed the title and abstract of studies identified in the primary search and excluded studies that did not address the research question, based on pre-specified exclusion and inclusion criteria. The full text of the remaining articles was reviewed to determine whether it contained relevant information. Any discrepancy in article selection was resolved by consensus, and in discussion with a coauthor. The bibliographic sections of the selected articles, as well as the systematic and narrative articles on the topic were manually searched for additional relevant articles.

### Study selection

In this meta-analysis, we included clinical trials that evaluated the clinical outcomes of curcumin in UC. Studies were included as long as they provided data needed for the analysis, irrespectively of the sample size, inpatient/outpatient setting and geography. Only RCTs reporting the efficacy of curcumin in UC were included in this meta-analysis. Exclusion criteria comprised: 1) case reports and case series; and 2) studies not published in English. In the event of multiple publications from the same

cohort and/or overlapping cohorts, data from the most recent and/or most appropriate comprehensive report were retained.

### Data abstraction and quality assessment

Data on study-related outcomes in the individual studies were abstracted onto a standardized form by at least 2 authors (SC, OCC), and 2 authors (BPM, SC) did the quality scoring independently. The Jadad scale for RCTs was used to assess the quality of studies, the details of which are provided in Supplementary Table 1 [8].

### Definitions

The response of UC to treatment was assessed using the following indices: Clinical Activity Index (CAI); Simple Clinical Colitis Activity Index (SCCAI); and Disease Activity Index (DAI).

The CAI indexing system comprises 7 items: stool frequency (0-3); blood in stool (0-4); general well-being (0-3); abdominal discomfort (0-3); fever (0-3); extraintestinal manifestations (0-9); and laboratory findings (erythrocyte sedimentation rate and hemoglobin) (0-4) [9].

The SCCAI system comprises 6 items: bowel frequency during the day (0-3); bowel frequency at night (1-2); urgency of defecation (1-3); blood in stool (1-3); general well being (0-4); and extra-colonic features (1 per manifestation) [10].

The DAI, or Mayo score, first developed in 1987, calculates a score between 0 and 12 and includes assessment of stool frequency, rectal bleeding, findings of flexible proctosigmoidoscopy and physician’s global assessment of disease activity [11]. The Mayo endoscopic score has been classified into the following 4 categories: 0, normal or inactive disease; 1, mild disease with erythema, decreased vascular patterns and mild friability; 2, moderate disease with marked erythema, absence of vascular patterns, friability and erosions; and 3, severe disease with spontaneous bleeding and ulceration [12].

Outcomes assessed in the analysis were as follows:

1. Pooled rate of clinical remission, defined as: CAI score  $\leq 4$ ; Ulcerative Colitis Disease Activity Index (UCDAI)  $\leq 2$  or  $< 3$ , SCCAI  $\leq 2$ .
2. Pooled rate of clinical response, defined as: decrease in UCDAI by  $\geq 3$ ; decrease in partial Mayo score by  $\geq 3$ ; and decrease in SCCAI score by  $\geq 3$  points.
3. Pooled rate of endoscopic response and remission, defined as: drop in Mayo score  $\geq 1$  to a score of 0 or 1 for remission and any  $\geq 1$  in Mayo sub-score as response as well as a partial Mayo score  $\leq 1$ .
4. Safety profile, including adverse events.

### Statistical analysis

We used meta-analysis techniques to calculate the pooled estimates in each case, using a random-effects model and

following the methods suggested by DerSimonian and Laird [13]. When the incidence of an outcome was 0 in a study, a continuity correction of 0.5 was added to the number of incident cases before statistical analysis [14]. We assessed heterogeneity between study-specific estimates using the Cochran Q statistical test for heterogeneity, 95% prediction interval (PI), which deals with the dispersion of the effects [15-17], and the  $I^2$  statistics [18,19]. In this, values of <30%, 30-60%, 61-75%, and >75% were suggestive of low, moderate, substantial, and considerable heterogeneity, respectively [20]. Publication bias was ascertained, qualitatively by visual inspection of a funnel plot and quantitatively by the Egger test [21]. When publication bias was present, further statistics using the fail-safe N test and Duval and Tweedie's "Trim and Fill" test was used to ascertain the impact of the bias [22]. Three levels of impact were reported, based on the concordance between the reported results and the actual estimate if there were no bias. The impact was reported as minimal if both versions were estimated to be same, modest if effect size changed substantially but the final finding would still remain the same, and severe if the basic final conclusion of the analysis was threatened by the bias [23]. Predictive factors for the outcomes were assessed by meta-regression methods. All analyses were performed using Comprehensive Meta-Analysis (CMA) software, version 3 (BioStat, Englewood, NJ).

## Results

### Search results and population characteristics

From an initial total of 119 studies, 101 records were screened and 74 full-length articles were assessed. Seven studies (380 patients) were included in the final analysis [24-30]: 188 patients were treated with curcumin as an adjunct to mesalamine and 192 patients were in the control group, receiving placebo with mesalamine. One study [25] reported clinical outcomes using the CAI, 2 studies [29,26] used the UCDAI, 2 studies [30,24] used the Mayo/partial Mayo score, and 2 [27,28] used the SCCAI. The schematic diagram of study selection is illustrated in Supplementary Fig. 1.

There were 174 males and 128 females. Two studies did not report the patients' sex. Mean age ranged from 32.7±8.9 years

to 45.2±15.8 years. The basic population characteristics are described in Supplementary Table 2. Four studies [29,26,28,27] reported the extent of colitis: left sided colitis (52 patients), pancolitis (24 patients), and proctitis (28 patients). In 6 studies [24-28,30], oral curcumin was used, whereas in 1 study [29] the route of administration was rectal. In the study by Masoodi *et al* [28], the actual number of patients who achieved an overall final clinical response was not reported. As the study was otherwise of high quality, the authors decided to include it in the analysis. The primary author of the study was contacted, but it was not possible to obtain the missing information and the most appropriate data were extracted. The potential influence of this study on outcomes, if any, was evaluated using a sensitivity analysis.

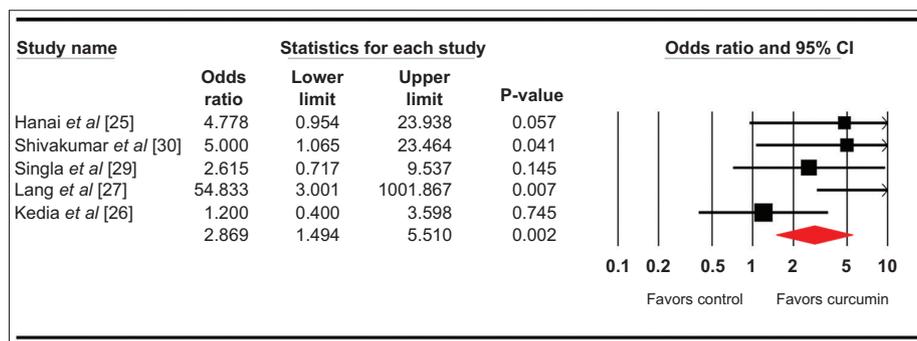
### Characteristics and quality of included studies

All included studies were RCTs. Two studies were published in abstract form [24,30] and the rest as full manuscripts. The detailed assessment of study quality is given in Supplementary Table 1. Overall, all studies were considered to be of high quality based on the Jadad scale. There were no low-quality studies.

### Meta-analysis outcomes

The pooled odds ratio for clinical remission (5 studies) [25-27,29,30] with curcumin use was 2.9 (95% confidence interval [CI] 1.5-5.5, 95% prediction interval [PI] 0.5-33,  $I^2=45$ ,  $P=0.002$ ) (forest plot Fig. 1), while for clinical response with curcumin (5 studies) [24,26-29] it was 2.6 (95%CI 1.5-4.5, 95%PI 0-88,  $I^2=74$ %,  $P=0.001$ ) (forest plot Fig. 2). The pooled odds ratio for endoscopic response/remission (5 studies) [24,26,27,29,30] was 2.3 (95%CI 1.2-4.6, 95%PI 0-14,  $I^2=35.5$ %,  $P=0.01$ ) (forest plot Fig. 3).

With regards to safety and adverse events, Lang *et al* [27] reported 3 serious adverse events resulting in withdrawal of the subjects from the study. Two patients reported worsening UC symptoms and 1 patient reported abdominal pain from a peptic ulcer present prior to initiation of the study medication. Four patients reported mild adverse events, such as nausea, transient increase in stool frequency and abdominal bloating.



**Figure 1** Forest plot. Clinical remission  
CI, confidence interval

Nine adverse events were reported in 7 patients by Hanai *et al* [25], including sensation of abdominal distension, nausea, transient hypertension, transient increase in the number of stools and elevated serum guanosine triphosphate level. Masoodi *et al* [28] reported a total of 8 adverse events, including flatulence, dyspepsia, headache, increased appetite, nausea and yellow stools. There were no serious adverse events.

Meta-regression analysis was done based on the curcumin dosage used. The lowest dose used was 100 mg and the maximum 10000 mg. No significant predictive effect was noted with curcumin dosage on the calculated outcomes (Random effects Knapp-Hartung 2-sided P-value=0.54, 0.34, 0.66 for clinical remission, clinical response and endoscopic response, respectively).

### Validation of meta-analysis results

#### Sensitivity analysis

To assess whether any one study had a dominant effect on the meta-analysis, we excluded one study at a time and analyzed its effect on the main summary estimate. On this analysis, no single study significantly affected the outcome or the heterogeneity. Thus, removing the study by Masoodi *et al* [28], would not have changed our findings.

### Heterogeneity

We assessed the dispersion of the calculated rates using the PI and  $I^2$  percentage values. The PI gives an idea of the range of the dispersion and  $I^2$  tells us what proportion of the dispersion is true versus chance [17]. The pooled rates of primary outcomes had wide prediction intervals with heterogeneity.

### Publication bias

A publication bias analysis was not done, as the total number of studies included in the analysis was less than 10.

### Discussion

Our study demonstrates that adjunctive use of curcumin with mesalamine yields a superior clinical and endoscopic response in the treatment of UC compared to placebo and mesalamine. This study is the first meta-analysis to report on the use of curcumin as an adjunct to mesalamine in the treatment of UC, and it is the most comprehensive review to date of all human trials evaluating the use of adjunctive curcumin therapy in treating UC.

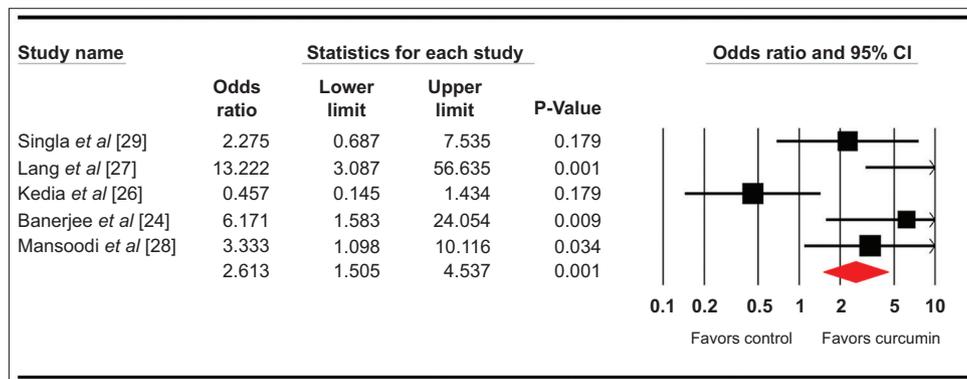


Figure 2 Forest plot. Clinical response  
CI, confidence interval

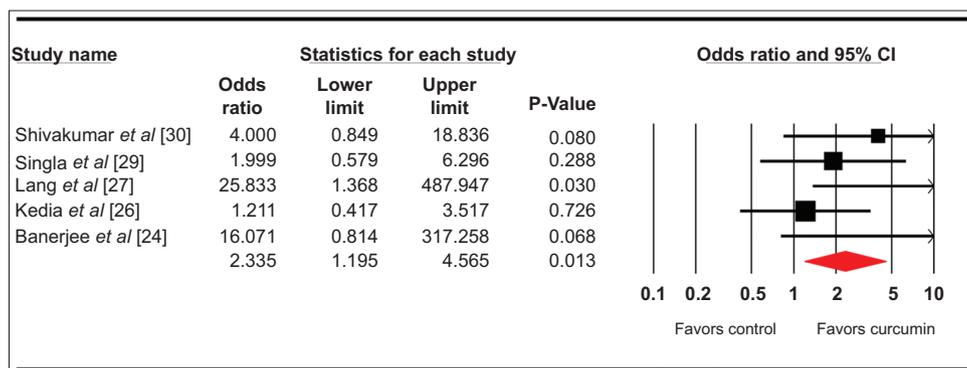


Figure 3 Forest plot. Endoscopic response  
CI, confidence interval

Based on our meta-analysis of clinical trials, the odds of a clinical and endoscopic response to curcumin as an adjunct to mesalamine, compared to placebo with mesalamine, were approximately threefold. A total of 21 adverse events were reported, as mentioned in the Results section, all of which were of mild degree.

In experimental models, curcumin has been shown to prevent colitis induced by tri-nitro-benzene sulfonic acid and dextran sodium sulfate. Its postulated mechanism of action is suppression of nuclear factor  $\kappa$ -light chain enhancer in B-lymphocytes, along with favorable expression of Th1 and Th2 cytokines. Curcumin has also been reported to have anti-interleukin-1 and anti-tumor necrosis factor  $\alpha$  properties, which makes it an attractive naturopathic treatment option for inflammatory diseases such as UC [31,32].

The results of this study are on par with the reported outcomes on the use of curcumin in UC published in the literature [28,25,27,33]. One negative study, that by Kedia *et al* [26], reported no significant differences in the rates of clinical remission, clinical response, mucosal healing, and treatment failure between curcumin and placebo at 8 weeks of treatment. Discrepancies in the drug dosage, drug delivery and duration of treatment are some of the postulated reasons for this outlier. Curcuminoids are lipophilic molecules and their absorption in the gastrointestinal tract can be low and variable [34]. Currently, we do not know whether the therapeutic effects of curcumin depend on its absorption and systemic bioavailability, or are more the result of a topical action on the intestinal mucosa. Clearly, there exists a significant “knowledge gap” regarding the use of curcumin in human beings.

The strengths of this review are as follows: systematic literature search with well-defined inclusion criteria, careful exclusion of redundant studies, inclusion of good quality studies with detailed extraction of data, rigorous evaluation of study quality, and statistics to establish and/or refute the validity of the results of our meta-analysis. We report the prediction intervals, thereby making our comparative pooled estimates applicable to the real population.

There were limitations to this study, most of which are inherent to any meta-analysis. The included studies were not entirely representative of the general population and community practice, with most being performed in tertiary-care referral centers. Heterogeneity was noted, most probably due to differences in the severity scoring indices used, the dosages of curcumin and the route of drug delivery. We wanted to analyze all the published literature on curcumin use in UC and included studies with active, quiescent as well as mild-to-moderate disease activity. We were not able to analyze our results based on the presence of comorbidities and were not able to assess the predictors of treatment success and/or failure. Nevertheless, our study is the best available estimate in the literature thus far with respect to the use of curcumin as an adjunct to mesalamine in the treatment of UC.

In conclusion, our meta-analysis demonstrates that curcumin, when combined with mesalamine, yields a superior clinical and/or endoscopic response in UC, albeit

with heterogeneity. Given the minimal adverse events, we recommend that curcumin be considered as an adjunct to mesalamine in the treatment of UC.

## Acknowledgment

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### Summary Box

#### What is already known:

- American Gastroenterology Association guidelines recommend starting standard dose mesalamine or diazo-bonded 5-amino-salicylic acid (5-ASA) in patients with extensive mild-to-moderate ulcerative colitis (UC)
- Rectal mesalamine is recommended in addition to oral 5-ASA in patients with extensive or left-sided mild-to-moderate UC
- No recommendation was made on the use of curcumin in mild-to-moderate UC patients

#### What the new findings are:

- Based on this meta-analysis of randomized controlled trials, combination therapy of curcumin with mesalamine in patients with mild-to-moderate UC yields a superior clinical and endoscopic response
- Odds ratio (OR) for clinical remission with curcumin was 2.9 (95% confidence interval [CI] 1.5-5.5),  $P=0.002$ ; OR for a clinical response with curcumin was 2.6 (95%CI 1.5-4.5),  $P=0.001$ ; OR for an endoscopic response and/or remission with curcumin was 2.3 (95%CI 1.2-4.6),  $P=0.01$

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**Supplementary Table 2** Study and patient characteristics

Author	Hanai	Shivakumar	Singla	Lang	Kedia	Banerjee	Masoodi
Year, country, study type	2006, Japan, RCT	2011, India, RCT	2013, India, RCT	2015, Israel, RCT	2017, India, RCT	2017, India, RCT	2018, Iran, RCT
Dx Type	UC	UC	Distal UC	UC	UC	UC	UC
Severity	Quiescent	Active	Mild-Mod	Mild-Mod	Mild-Mod	Mild-Mod	Mild-Mod
Measurement of response	Clinical Remission - CAI $\leq 4$ ; Endoscopic - NR	Clinical - Improvement in fecal calpro; Endoscopic response - histology score 1 point decrease	Clinical Response - Decrease in UCDAI by $\geq 3$ , C. Remission UCDAI $< 3$ ; Endoscopic response - histology score 1 point decrease	Clinical Response - Decrease in SCCAI score by $\geq 3$ points, C. Remission SCCAI $\leq 2$ ; E. Remission - Mayo score drop $\geq 1$ to a score of 0 or 1, E. Response - any $\geq 1$ in Mayo subscore	Clinical - Decrease in UCDAI by $\geq 3$ , Remission UCDAI $\leq 2$ ; Endoscopic remission - endoscopic score of 0/1	Clinical - Decrease in partial Mayo score by $\geq 3$ ; Endoscopic Remission - partial Mayo score $\leq 1$	Clinical - improvement in SCCAI score; Endoscopic - NR
Age (mean)							
Curcumin	45.2 $\pm$ 15.8		32.7 $\pm$ 8.9	40.4 $\pm$ 12.7	36 $\pm$ 12		38.21 $\pm$ 16.37
Placebo	39.7 $\pm$ 14.2		35.5 $\pm$ 13.8	41.4 $\pm$ 13.9	34 $\pm$ 7		36.04 $\pm$ 11.78
Disease duration							
Curcumin	98.6 $\pm$ 74.2		60 (36-96)	85.2 $\pm$ 72	45.96 $\pm$ 48		37.75 $\pm$ 56.03
Placebo	93.5 $\pm$ 74.2		33 (12-72)	60 $\pm$ 49.2	43.68 $\pm$ 31.08		33.11 $\pm$ 34.57
Disease involvement							
Curcumin							
Left Sided			12	11	17		12
Pancolitis				5	7		12
Proctitis			11	10	3		4
Placebo							
Left Sided			14	15	21		16
Pancolitis				3	6		8
Proctitis			8	6	2		4
Sex							
Curcumin							
Male	23		12	17	16		15
Female	22		11	9	13		13
Placebo							
Male	26		11	16	25		13
Female	18		11	8	8		15
Intervention							
Curcumin	45	18	23	26	29	19	28
Placebo	44	18	22	24	33	23	28

(Contd...)

**Supplementary Table 2 (Continued)**

Author	Hanai	Shivakumar	Singla	Lang	Kedia	Banerjee	Masoodi
Response to Rx							
Clinical remission							
Curcumin	43	15	10	14	9		
Placebo	36	9	5	0	9		
Response to Rx							
Clinical response							
Curcumin			13	17	6	12	16
Placebo			8	3	12	5	8
Response to Rx							
Endoscopic							
Curcumin		15	12	10/22	10	5	
Placebo		10	8	0/16	10	0	
Total Curcumin							
Dose	2 g	10 g	140 mg	3 g	450 mg	100 mg	240 mg
Route	Oral	Oral	Rectal	Oral	Oral	Oral	Oral
Mesalamine dose	1.5-3 g/day		1.6 g/day				3 g/day
Follow up	24	8	8	4	8	12	4
Adverse events	9	NR	0	3	0	NR	NR
Past Rx							
Curcumin							
5-ASA			18	22	29		24
Steroids			12		0		4
AZT			2		2		7
ASA+Immunomodulator				4			
Anti-TNF							2
Placebo							
5-ASA			16	19	33		23
Steroids			1		0		4
AZT			1		2		10
ASA+Immunomodulator				5			
Anti-TNF							2

RCT, randomized controlled trial; UC, ulcerative colitis; CAI, clinical activity index; SCCAI, simple clinical colitis activity index; DAI, disease activity index; AGA, American gastroenterology association; 5-ASA, 5-aminosalicylic acid; TNF, tumor necrosis factor