

# Can vitamin D induce remission in patients with inflammatory bowel disease?

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## Abstract

**Background** The prevalence of vitamin D (VD) deficiency in adults with inflammatory bowel disease (IBD) has been reported to be very high. In adults with ulcerative colitis (UC) it has been estimated to be around 45-50%, and in the case of Crohn's disease (CD) it ranges from 35-100%. This systematic review aimed to address the effects of VD in inducing remission in patients with CD and UC.

**Methods** PubMed, Embase, and Cochrane were searched, and PRISMA guidelines were followed. Nine studies with active disease subjects were evaluated; 5 of them investigated patients with UC, 2 studied patients with CD, while 2 studies included patients with both diseases.

**Results** The studies show that oral daily doses from 1000 IU can raise VD levels above 20 ng/mL in patients with active CD or UC. Doses close to 50,000 IU per week are sufficient to raise VD levels above 40 ng/mL. Moreover, daily doses from 2000 IU can be related to improvements in the clinical scores of the disease and improvement in patients' quality of life.

**Conclusion** It is challenging to define the best doses of VD as a therapeutic adjunct to induce remission in IBD patients, as there are biases in the inclusion of studies, since the route of administration of the VD can vary, as well as the doses and the time of use.

**Keywords** Vitamin D, cholecalciferol, ulcerative colitis, Crohn's disease, inflammatory bowel disease

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## Introduction

Ulcerative colitis (UC) and Crohn's disease (CD) are the primary elements of inflammatory bowel disease (IBD). These conditions may be designated as chronic relapsing idiopathic inflammatory processes of the gastrointestinal tract, whose etiology is not entirely understood. Genetic and environmental factors are related to the disruption of the intestinal epithelial

barrier, leading to increased permeability and an enhanced uptake of microorganisms that triggers immune system activation. The resulting imbalance may be associated with infections, oxidative stress, high fat and sugar diet, food additives, medications, a decrease in the intake of fibers, and reduced levels of vitamin D (VD) [1-3].

The world is experiencing an increasing number of patients with IBD, considered as a global burden. Some authors have estimated that more than 1 million people in the United States and 2.5 million in Europe are suffering from this disease. This alarming rise is mainly observed in developed countries, and there is also an increase in the newly industrialized population. People with IBD are affected by abdominal pain symptoms, diarrhea, bleeding, bowel obstruction, extra-intestinal manifestations, and a reduced quality of life (QoL) and capacity for work [4-6].

VD and its active form 1,25-dihydroxyvitamin D is defined as a fat-soluble hormone synthesized in skin exposed to sunlight or obtained from the diet. When produced by plants and fungus, it is named ergocalciferol (VD<sub>2</sub>), and when produced in the skin, it is called cholecalciferol (VD<sub>3</sub>): 1,25-dihydroxyvitamin D<sub>3</sub> 1,25(OH)<sub>2</sub>D<sub>3</sub>. The active form, 1,25(OH)<sub>2</sub>D<sub>3</sub>, is associated with biological effects [7,8].

Besides being associated with the regulation of mineral and bone homeostasis, it modifies adaptive and innate immune

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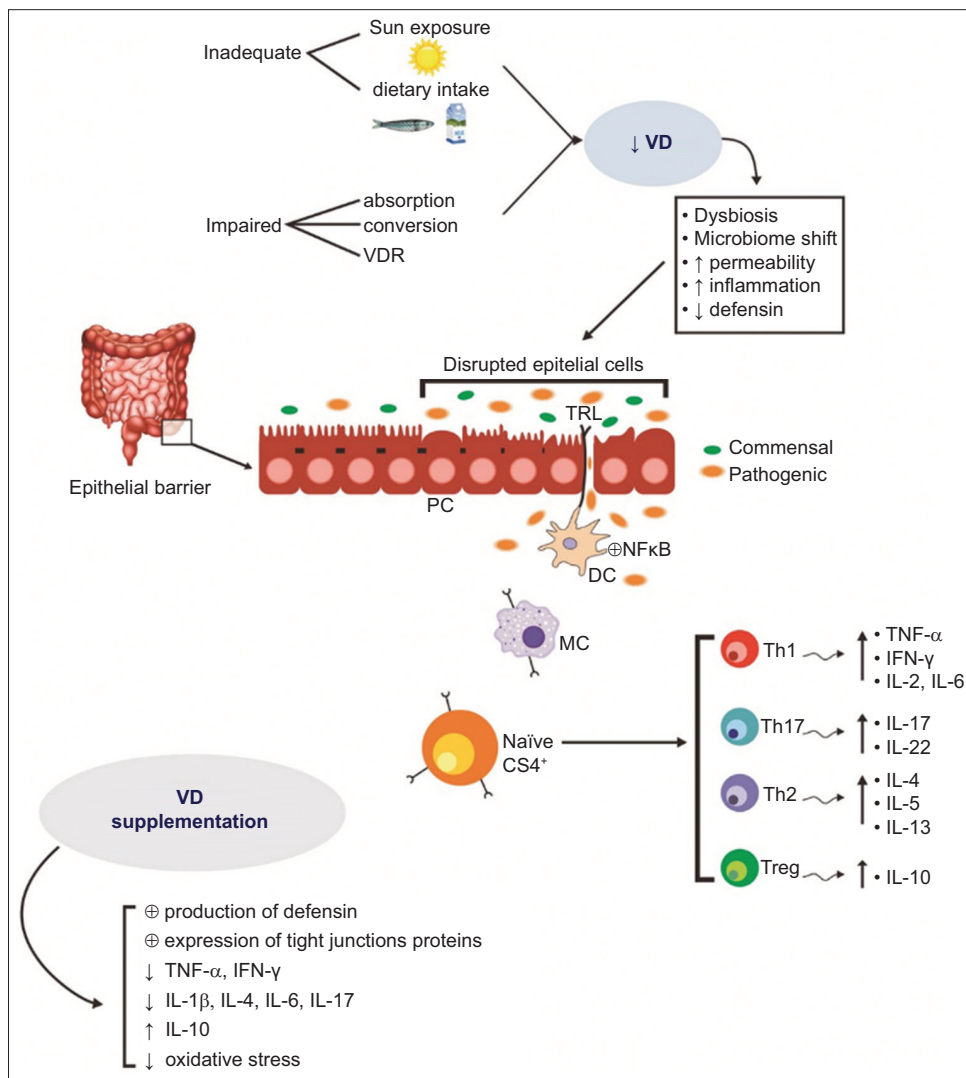
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responses. For these reasons, it has been related to several aspects of inflammation pathways. VD receptors in the thymus and peripheral T-cells are associated with the release of biomarkers of inflammation, such as interleukin (IL)-2 and IL-10 in dendritic cells and macrophages [9,10]. VD is related to the upregulation of T regulatory cells, IL-4, IL-10, transforming growing factor- $\beta$ , and increased expression of proteins related to cell adhesion, proliferation and migration. On the other hand, it is associated with the downregulation of the expression of tumor necrosis factor (TNF)- $\alpha$ , interferon (IFN)- $\gamma$ , IL-6, IL-12, the *ATG16L1* gene, and lysozyme by Paneth cells. It also reduces bacterial translocation to extra-intestinal tissues, inhibits inflammatory cell infiltrates and reduces colorectal tumors in C57BL/6J and NOD2<sup>-/-</sup> mice [2,11]. Intestinal epithelium and mucosal immune cells express the VD receptor. Some studies have shown that this vitamin may regulate the proteins of tight junctions and reduce intestinal apoptosis, thus protecting the gut barrier [12-14]. Fig. 1 summarizes the effects of VD in IBD.

The prevalence of VD deficiency in adults with UC has been estimated at 45-50% and 35-100% in CD. This deficiency may be attributed to decreased sunlight exposure, low VD intake, altered enterohepatic circulation, and augmented loss due to enteropathy. In addition, low serum levels of 25-hydroxyvitamin have been reported as a biomarker for disease activity and as a predictive factor for poor clinical outcomes [15,16].

It has not been established whether VD deficiency leads to IBD or *vice versa*. As shown by Zullow *et al* [17] in their study of 255 subjects with IBD, most of them had low VD levels, UC was associated with a lower risk of VD insufficiency, and in CD patients VD sufficiency was associated with reduced disease activity and improved QoL.

Although VD has become popular and is considered to aid in the treatment of patients with IBD, few clinical trials have investigated the use of this vitamin in patients with active UC and CD. More than 20 studies evaluated the use of VD



**Figure 1** Sources of vitamin D (VD) and its effects on the altered epithelial barrier of patients with inflammatory bowel disease. VD induces immune cells to adopt an anti-inflammatory pattern  
 IL, interleukin; IFN, interferon; PC, Paneth cell; TH1, T helper 1; TH2, T helper 2; TH17, T helper 17; Treg, regulatory T cells; TNF- $\alpha$ , tumor necrosis factor- $\alpha$ ; VDR, vitamin D receptor

in patients with IBD, but only 9 included patients with active disease. Therefore, systematic reviews and meta-analyses that consider patients at different stages of the disease may be at risk of significant bias. Since many studies have shown that VD profoundly influences the inflammatory pathways and may help prevent or remission of IBD, this systematic review aimed to evaluate the effects of this vitamin in UC and CD patients in relapse.

## Materials and methods

### Focused question

The focused question used for this review was, “Can vitamin D assist in the remission of IBD patients?”

### Databases

We searched the MEDLINE/PubMed (National Library of Medicine, National Institutes of Health), Cochrane and EMBASE databases for relevant articles addressing the focused question.

As different terms are used to refer to IBD and VD, the following combinations of MeSH terms were used: Inflammatory Bowel Disease or Ulcerative Colitis or Crohn’s Disease and vitamin D or calcidiol or calcitriol or 25 hydroxycholecalciferol or cholecalciferol. From the resulting list of studies that included these combinations of terms, we selected the articles that investigated the effects of VD in relapsing IBD patients. Two authors evaluated the screening of the studies. This search was not limited to any specific period. We followed the PRISMA (preferred reporting items for a systematic review and meta-analysis) guidelines [18].

### Language

Only studies in English were selected.

### Inclusion criteria

Clinical trials associating the use of VD in patients with active UC and CD were included. We included both full and not full texts available in the consulted databases.

### Exclusion criteria

We excluded studies not in English, reviews, editorials, case reports, studies not involving humans (in animals or *in vitro* studies), studies with patients in remission, and pediatric cohorts (patients aged less than 18 years).

### Eligibility criteria

Our review’s eligibility criteria followed the PICO (population, intervention, comparison, and outcome) format for randomized controlled trials (RCT).

### Data extraction

Two independent judges performed the search for identifying RCT in the databases. The articles’ abstracts were evaluated, and full- and no full-text articles were retrieved to support decision-making.

### Outcomes

The outcomes considered were simple clinical colitis activity index (SCCAI), CD activity index (CAAI), IBD questionnaire (IBDQ), Mayo score, short IBDQ (SIBDQ), C-reactive protein (CRP), erythrocyte sedimentation rate (ESR), fecal calprotectin (FC), and QoL. Remission was defined as resolution of visible inflammation and ulceration on endoscopy, a reduction in the Disease Activity Index, and the patient showing cessation of diarrhea, bloating, and pain [19-21].

### Quality assessment

The evaluation of the risk of bias regarding the selection, detection, and reporting bias of each RCT was evaluated according to the Cochrane Handbook for systematic reviews of interventions. Other risks of bias in the selection of patients, classification of interventions, missing data, and measurement of outcomes were also evaluated.

## Results

Fig. 2 shows the selection of the studies. Table 1 summarizes the included RCTs, and Table 2 shows a description of the main biases. All the patients of the included studies maintained the standard drug therapy for IBD during intervention with VD. Nine studies (3 from Iran, 1 from India, 1 from Brazil, 1 from China, 1 from Australia, and 2 from the United States) with active disease subjects were included in this systematic review. Five of them were based on patients with UC, 2 with CD, and 2 studies included patients with both diseases. In total, the studies included 387 patients with UC and 124 with CD. The exact number of men and women could not be calculated because 4 studies did not specify the patients’ sex.

Although we only included studies with active disease patients, there was a wide variation in the doses administered (daily oral doses of 1000-50,000 IU, 150,000/3 months, or a single injectable dose of 300,000 IU). The treatment duration ranged from 1 day for 300,000 IU [22,23], to

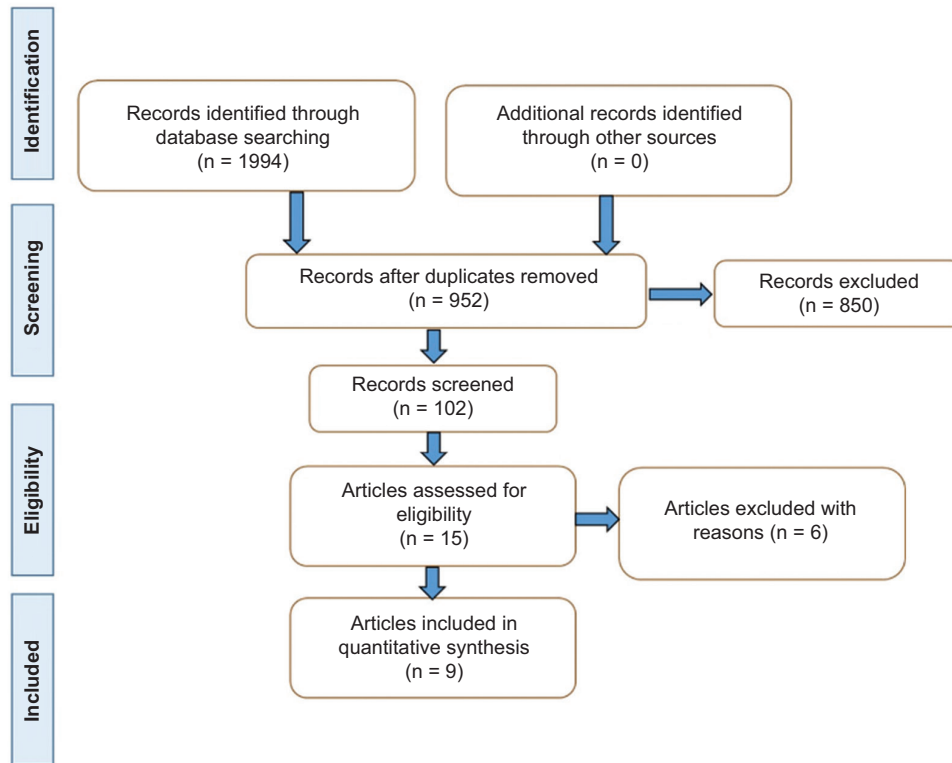


Figure 2 Flow diagram showing the selection of the studies - PRISMA guidelines – [17]

8 days for 60,000 IU [24], and 8 weeks to 1 year for the other doses [25-30].

All studies showed significant improvement in serum VD levels after supplementation (Table 1). The studies of Emani *et al* [22] and Sharif *et al* [23], who used a single injectable dose of 300,000 IU, reported a mean level of 40.8±5.2 ng/mL. Ahamed *et al* [24] achieved similar values (40.83 ng/mL) with a daily dose of 60,000 IU/8 days. Even higher values of 46.14±12.7 and 45±19 ng/mL were obtained by Baffuto *et al* [28] and Yang *et al* [29], respectively (although half of the patients included in this last study, who received 5000 IU, did not reach values greater than 40 ng/mL). The lowest initial (10.58±3.82 in CD and 10.68±4.11 in UC) and final (23.04±9.66 for CD and 28.09±11.6 for UC) VD values were seen in the study of Tan *et al* [30]. Karimi *et al* [27] observed the lowest percentage increases in serum VD levels: the group that received 2000 IU presented 21.83±9.69 at baseline, and 28.99±8.69 after 12 weeks, while the group that received 1000 IU presented 24.37±8.14 initially and 28.75±11.90 ng/mL after 12 weeks. In the study of Bafutto *et al* [28], the authors compared supplementation with 2000 (G1), 10,000 (G2), and 50,000 IU (G3)/week and concluded that the highest dose is the best for VD replacement (G1 19.5 ± 5.1 at baseline and 26±6.7 at the end of the study, P=0.07; G2: 19.1±4.1/26±5.8, P=0.04; and G3: 19.5±6.4/46.4±12.7, P<0.001). This study, and the trial performed by Ahamed *et al* [24] that used 60,000 IU daily for 8 days, showed a higher percentage of augmenting serum levels of VD.

Only 2 studies [24,29] reported remission after VD treatment. Four studies reported an improvement in IBD

scores (UCDAI, SCCAI, CDAI) [24,25,27,29] and 4 reported improvements in QoL (IBDQ) [26-29].

One study showed that VD supplementation could suppress the levels of visfatin (3.1±1.4 vs. 1.4±2.4 ng/mL; P=0.03) and vascular endothelial growth factor (VEGF) (368±151 at baseline and 262±112 at the end of the study) in patients with low (insufficiency) of VD [22], suggesting VD might be helpful in reducing proangiogenic factors in UC patients. Nevertheless, the results of this study did not show a significant improvement in visfatin and VEGF at the end of the intervention.

Three studies [24,26,30] observed a significant reduction in CRP (with doses that varied from 2000 IU to 150,000 IU) in UC and CD patients, respectively, whereas others did not see any changes in this marker [23,25].

Only 3 studies investigated the levels of FC, and different results were found. The study of Garg *et al* [25] did not observe changes in this marker, using 5000 IU of VD daily for 3 months, but the sample had only 10 patients and that may have affected the results. On the other hand, the trial of Ahamed *et al* [24] showed significant reductions in FC with the use of 6000 IU daily for 8 days in UC patients. Bafutto *et al* [28] observed a reduction in FC in mild-severe CD patients who received 50,000 IU of VD (but not in those that received 2000 or 10,000 IU).

Only one [23] of the included studies investigated the levels of IFN-γ, TNF-α, and IL-12p70 and observed significant reductions (respectively, P<0.001, P=0.001, and P=0.001) after an injection of 300,000 IU in UC patients.

Also, only one study [27] investigated the antioxidant/oxidant status of UC patients treated with 1000 or 2000 IU of VD and

**Table 1** Clinical trials that investigated the effect of vitamin D in patients with ulcerative colitis and Crohn's disease

Type of study	Population	Intervention / Comparison	Main outcomes	Remission	AE	Reference
Studies with ulcerative colitis patients						
Double-blind randomized controlled trial with a parallel design and 1:1 ratio of 2 groups (Iran)	90 UC patients not in the relapse phase (26-47y; ? 27) (using sulfasalazine, mesalamine or azathioprine)	300,000 IU of VD (single intramuscular injection) or 1 mL saline as placebo	Baseline VD = 33.3±7.0; final = 40.8±5.2 (P<0.05). There were no significant differences in visfatin and VEGF levels between the 2 groups following supplementation	NR	No AE	Emani <i>et al</i> [22]
Single-center double-blind randomized parallel placebo-controlled trial (India)	60 patients with active UC (UCDAI ≥3) (29-37y; 36) (using mesalamine)	60,000 IU of VD daily/8d (nano liquid formulation) or a similar placebo	Baseline VD levels: 15.4 ng/mL, and at the end of 4 w: 40.83 ng/mL (P<0.05). The 3-point reduction in UCDAI score was seen more frequently in the VD group than placebo: 16/30 (53.3%) vs. 4/30 (13.3%), P=0.002. VD levels showed significant negative correlation with baseline and post-treatment UCDAI scores (P = -0.0.274, P=0.034; P = -0.455, P≤0.001). Increased VD levels showed a significant correlation with a decrease in ESR (ρ = -0.256, P=0.049), CRP (ρ = -0.598, P≤0.001), calprotectin (ρ = -0.368, P=0.004), and histology score (ρ = -0.608, P≤0.001). The mean VD of the entire group was 15.4 at baseline and 40.83 mg/dL in VD group (P≤0.001)	Yes	Transient vomiting and dyspepsia	Ahamed <i>et al</i> [24]
Double-blind randomized controlled trial (Iran)	90 patients with mild to moderate UC (28-46y; ? 27) (using sulfasalazine, mesalamine or azathioprine)	300,000 IU of VD (single intramuscular injection) or 1 mL saline as placebo	Baseline VD levels: 33.3±7 ng/mL, and after injection in the treated group: 40.8±5.2 ng/mL (P<0.001). Significant decrease on IFN-γ, TNF-α, and IL-12p70 levels. VD levels were significantly higher in the treated group after 3 m (P<0.001)	NR	NR	Sharifi <i>et al</i> [23]
Double blind randomized clinical trial (Iran)	50 patients with mild to moderate UC (22-55y; 23 27) (using mesalamine, corticosteroids or azathioprine)	1000 or 2000IU/d of VD (respectively as low dose or high dose group)/ 12 w	After 12 w, VD levels were significantly higher in the high dose group (from 21.83±9.69 to 28.99±8.69, P<0.001) than in the low dose group (24.37±8.14 to 28.75±11.90, P=0.192). Serum Total Oxidant Status reduced significantly only in the high-dose group (P=0.023). IBDQ-9 mean score was significantly increased in the high dose group (P=0.001), and the SCCAI score was reduced in both groups (P=0.045)	NR	NR	Karimi <i>et al</i> [27]
Prospective double-blinded, randomized trial (EUA)	18 UC subjects (18-56y; 5 13) (using azathioprine, anti TNF or 5-aminosalicylic)	2000 IU or 4000 IU of oral VD daily/90 days	In the group receiving 4000 IU of VD, the increase in quality life scores (SIBDQ) was significant (P=0.017) but not in the 2,000 IU VD group (P=0.87). 4,000 UI of VD increased serum levels more than 2,000 (P<0.001); there was an increase of 16.8±9.15 in the higher dose group and 5.0±3.82 in the lower dose group. The improvement in the Partial Mayo UC was not statistically significant in both groups. CRP levels decreased significantly in both groups	NR	No	Mathur <i>et al</i> [26]

(Contd...)

Table 1 (Continued)

Type of study	Population	Intervention / Comparison	Main outcomes	Remission	AE	Reference
Studies with Crohn's disease patients						
Double-blind, randomized, prospective study (Brazil)	30 patients with mild-severe CD (18-67y; ?♀♂) (using anti TNF)	2000 IU of VD/w/8w (group 1); or 10,000 IU of VD/w/8w (group 2); and 50,000 IU of VD/w/8w (group 3)	Significant augment of VD levels (baseline for G1 was 19.5±5.1ng/mL to 26.0±6.7 at 8w (P=0.07); G2: 19.1±4.1 to 26.0±5.8 (P=0.04) and G3: 19.5±6.4 to 46.4±12.7 (P<0.001)); improvement in IBDQ were observed in all groups (significant improvement was observed in G2 (P=0.04) and G3 (P=0.01). Significant reduction in FC was seen in G3 (P=0.04)	NR	NR	Bafutto <i>et al</i> [28]
Open-labeled prospective clinical trial (EUA)	18 patients with CD (CDAI: 150 - 400 and serum levels of VD ≤40ng/mL) (21-55y; 11♀, 7♂) (using purine analogs or 5-aminosalicylic)	1000IU/d/2w, after which the dose was escalated to 5000IU/d/24 w	The treatment led to a significant increase of serum levels of VD from 16.0±10 to 45±19 ng/mL at the end of the treatment (P<0.001) and decreased the mean CDAI scores from 230±74 to 118±66 (P<0.001). QoL scores also improved with supplementation (P=0.0004)	Yes	No significant AE is directly related to the use of VD	Yang <i>et al</i> [29]
Studies with both ulcerative colitis and Crohn's disease patients						
Prospective, randomized, parallel controlled, pen-label study (China)	74 patients with UC and 71 patients with CD (>18 y; ?♀♂ group) (using 5-aminosalicylic, glucocorticoids or anti TNF)	3 arms/12 m: A (VD3 150,000 IU/every 3 m plus calcium 600 mg oral daily); B (calcium 600 mg oral daily), and C (control)	No significant differences were observed of baseline VD in Arm A between UC and CD patients (10.62±4.11 ng/mL vs. 10.58±3.82ng/mL, P=0.97). After 12 m the levels increased to 28.09±11.60 ng/mL and 23.04±9.66 ng/mL in respectively UC and CD subjects (P=0.001). There were no significant differences in either UC or CD subjects in CRP, ESR, and Mayo score/CDAI among the 3 arms at baseline (p>0.05) and after 12 m	NR	Hyper-phospha-temia hypercal-cemia, and intoxication	Tan <i>et al</i> [30]
A pilot study (Australia)	5 patients with mild-moderately active CD and 5 with UC, (24-67 y; 3♀7♂) (using mesalazine or anti TNF)	1000 to 10,000 IU of VD/d/3 m according to serum levels	VD levels at baseline were 10.8-29.2 ng/mL and at the end ranged from 35.6-50 ng/mL (P<0.001 without differences for UC or CD patients). Fecal calprotectin and biomarkers of inflammation (platelet count, serum albumin 2L, and CRP) did not improve. Clinical disease activity (HBI and SSCAI) significantly reduced over 12 weeks in CD (P=0.019), and a trend was seen in UC (P=0.051)	NR	Hypercalciuria	Garg <i>et al</i> [25]

AE, adverse events; CD, Crohn's disease; CDAI, Crohn's disease activity index; CRP, C-reactive protein; ESR, erythrocyte sedimentation rate; FC, fecal calprotectin; HBI, Harvey-Bradshaw Index; IBDQ, quality of life questionnaire; QoL, quality of life; SSCAI, simple clinical colitis activity index; SIBDQ, short IBD questionnaire; UC, ulcerative colitis; VD, vitamin D; VEGF, vascular endothelial growth factor; TNF, tumor necrosis factor; IFN, interferon; IL, interleukin

observed a reduction only in total antioxidant status ( $P=0.023$ ).

As can be seen in Table 1, 7 of the 9 studies included in this systematic review compared disease activity indexes after using VD (3 studies with UC patients, 2 with CD patients, and 2 studies that included both UC and CD subjects). The single-center, double-blinded study of Ahamed *et al* [24] showed that there was a significant reduction in UCDAI scores after treatment with 60,000 IU of VD daily for 8 days in UC patients compared to the placebo group ( $P=0.001$ ). Karimi *et al* [27] observed a significant reduction in SCCAI score ( $P=0.045$ ) in UC patients that received 1000 or 2000 IU daily for 12 weeks, and a significant increase in IBDQ-9 ( $P=0.001$ ) in those patients that received 2000 IU daily. Moreover, Mathur *et al* [26] used 2000 or 4000 IU daily and observed that UC patients who received the higher dose showed a significant improvement in SIBDQ.

In CD patients, Bafutto *et al* [28] observed significant improvement in IBDQ in the groups that received 2000, 10,000 and 50,000 IU for 8 weeks. Yang *et al* [29] observed a significant reduction in CDAI scores and improved QoL scores in the entire sample that started receiving 1000 daily, with escalation to 5000 IU.

Two studies included both UC and CD patients [25,30]. Garg *et al* [25] observed improvement in clinical disease activity using oral 1000-10,000 IU daily for 3 months. Tan *et al* [30] did not see significant differences in Mayo score/CDAI in patients who received 150,000 IU every 3 months for a year.

In summary, we can see from the results above that patients with UC achieved improvement in IBDQ-9 with oral doses of VD of 2000 IU (daily/3 months) [27] and improvement in SIBDQ with 4000 IU (daily/3 months) [26]. A significant reduction in UCDAI was observed with use of VD 60,000 IU (daily/8 days) [24]; a reduction in SCCAI with the use of VD 1000-2000 IU (daily/3 months) [27]; and a reduction in SSCAI and HBI (Harvey Bradshaw Index) was seen with the use of VD 1000-10,000 IU (daily/3 months) [25]. Unfortunately, these disease activity scores were not evaluated in the trial with a single intramuscular injection [22]. Furthermore, the results show that, for CD patients, there was a significant improvement in IBDQ with the daily use of VD 2000, 10,000, and 50,000 IU for 2 months [28]. QoL scores also improved with the dose of 2000 escalated to 5000 IU [29]. The use of 1000-10,000 IU (daily/3 months) also significantly improved CD patients' clinical disease activity [25].

In the study of Bafutto *et al* [28], the authors suggest that the dose of 50,000 IU per week is the most appropriate for improving the immunomodulation of CD patients. However, the immunomodulation was evaluated only by improving FC, which is a potential bias.

## Discussion

Although there is much variation related to the dose (1000-300,000 IU) and form of administration (oral liquid or capsules, or injectable) of VD and the duration of supplementation (1-12 months), the results in patients with active CD and UC can be promising in inducing remission. However, relevant biases should be considered (Table 2), which

can interfere with the studies' results. Indeed, some studies used only a small number of participants [25,26,29]. The studies of Karimi *et al* [27], Mathur *et al* [26], Bafutto *et al* [28], Yang *et al* [29] and Garg *et al* [25] did not make comparisons with a placebo group. As previously mentioned, the outcomes assessed were also different.

In addition to clinical trials that included CD or UC patients with active disease, other studies investigated the use of VD in patients in remission. The retrospective cohort study performed by Janssen *et al* [31] with 384 patients, intended to determine risk factors for VD deficiency, found a high prevalence of CD or UC (63% and 55%, respectively). Their results also showed that VD deficiency is related to disease activity in CD and to anemia in UC subjects. On the other hand, increasing VD levels correlated with improved CD activity.

Narula *et al* [32] investigated VD's effects in CD patients in remission. In this randomized and double-blind placebo-controlled trial, the authors evaluated the effects of VD use at 10,000 IU daily compared to 1000 IU for one year and observed significantly higher serum levels in the first group ( $P=0.02$ ). Although the relapse rate was not significant in either group, there was an improvement in anxiety and depression scores in both groups treated with vitamin D3.

VD can also affect CD patients as it leads to reduced stimulation of dendritic cells by LPS, explaining the positive effects of VD on CD patients [33]. The open-label pilot study of Wiese *et al* [34] enrolled CD patients with active disease (CDAI score >150) treated with 16 oz of IBD nutrition formula (IBDNF) for 4 months. The 28 subjects were consuming regular and similar medications for CD and were asked to consume 16 oz of IBDNF/day. Nutrition status was assessed, as well as CDAI and IBDQ. The results of this exciting study showed that at the end of 4 months there was a significant increase in the levels of vitamin D (25-OH) in all patients ( $P<0.001$ ). Moreover, the investigators observed a significant reduction in plasma levels of arachidonic acid, augmented eicosapentaenoic and docosahexaenoic acids, and improved QoL in CD subjects.

Jorgensen *et al* [35] also investigated the effects of VD in a randomized and double-blind placebo-controlled trial in 108 patients with CD in remission. Subjects were randomized to intake 1200 IU VD ( $n=46$ ) or placebo ( $n=48$ ) daily for 12 months and the authors observed that the use of the vitamin increased serum levels from a mean of 69 nmol/L to 96 nmol/L after 3 months of treatment ( $P<0.001$ ). This paper's exciting results showed that the relapse rate was lower in the treated patients ( $P=0.06$ ). In the same clinical trial, Bendix-Struve *et al* [36] observed that the CD patients who received VD (1200 IU daily) presented higher serum levels of the vitamin and less release of IL-6 compared to the placebo group ( $P<0.02$ ). Moreover, VD use increased the proliferating stimulated CD4+ T cells ( $P=0.02$ ).

Guzman-Prado *et al* [15] carried out a systematic review of 19 studies involving the use of VD in patients with either CD or UC. They concluded that supplementation with VD is helpful in improving serum levels; additionally, it is associated with improvement in IBD scores and clinical and biochemical parameters. Although this is a very interesting study, the authors included patients at different stages of the disease, which could have led to bias in the results.

**Table 2** Description of the main biases of the clinical trials that investigated the effects of vitamin D on inflammatory bowel diseases

Study	Question focus	Appropriate randomization	Allocation blinded	Double-blind	Losses (<20%)	Prognostics or demographic characteristics	Outcomes	Intention to treat analysis	Sample calculation	Adequate follow up
Ulcerative colitis										
Emani <i>et al</i> [22]	Yes	Yes	Yes	Yes	Yes	Yes (incomplete)	Yes	Yes	Yes	Yes
Ahamed <i>et al</i> [24]	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes
Sharifi <i>et al</i> [23]	Yes	Yes	Yes	Yes	Yes	No	Yes	NR	Yes	Yes
Karimi <i>et al</i> [27]	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	No	Yes
Mathur <i>et al</i> [26]	Yes	Yes	Yes	Yes	Yes	Yes	Yes	No	No	Yes
Crohn's disease										
Bafutto <i>et al</i> [28]	Yes	NR	NR	Yes	No	No	Yes	NR	NR	Yes
Yang <i>et al</i> [29]	Yes	No	No	No	Yes	Yes	Yes	Yes	No	Yes
Crohn's disease and ulcerative colitis										
Tan <i>et al</i> [30]	Yes	Yes	Yes	No	Yes	No	Yes	No	No	NA
Garg <i>et al</i> [25]	Yes	No	No	No	Yes	Yes	Yes	Yes	No	Yes

NR, Not reported; NA, Not applicable

According to the results of the systematic review performed by Gubatan *et al* [16], insufficient levels of VD may increase the disease activity among IBD subjects and may be a marker of disease activity. Low VD levels in these patients may occur as a result of malnutrition and malabsorption in flaring patients, or less sun exposure due to poor general wellbeing. Further, low VD status is linked to a higher risk of future clinical relapse. With these considerations, the authors suggest that VD levels may play a cause-and-effect role in the clinical outcomes.

In our review, 5 studies showed flared patients with VD values at baseline lower than 20 ng/mL [24,25,28-30]. The highest VD levels at baseline were close to 30 mg/mL in the study of Emani *et al* [22]. These findings make us wonder if the cutoff value for VD of 20 ng/mL is sufficient to maintain bone tissue health and maintain or induce remission in IBD patients. Our results, however, indicate that, when patients with low values reach levels above 20 ng/mL, they are already showing improvement in the clinical scores of the disease and in their QoL. Moreover, supplementation with 1000-2000 IU daily is enough to elevate the 20 ng/mL levels in these patients. On the other hand, our findings show that patients with relapse of the disease always present VD levels lower than 40 ng/mL.

We recommend that VD levels be monitored in IBD patients and that supplements are provided for those deficient in this component. Supplementation should raise levels above 20 ng/mL, which can be achieved with daily doses above 1000 mg/dL. Longer-term administration should be monitored to maintain

normal levels of VD and remission of the patients. Another important finding of this review is that the studies show that patients with relapse of the disease never present VD levels above 40 ng/mL. This finding leads us to consider whether patients with IBD should be supplemented with higher doses so their serum levels are not less than 40 ng/mL.

This is the first systematic review that includes only studies performed on relapsed patients to the best of our knowledge. Although our study proposed to use only trials with patients in the same stage of the disease (active disease), we could not eliminate all biases since the doses used and the duration of use of VD was very different. Therefore, we believe it is challenging to define the best doses of this vitamin as a therapeutic adjunct to induce remission in UC and CD patients. Oral daily doses from 1000 IU/12 weeks can raise VD levels above 20 ng/mL in UC or CD patients with active disease, and doses close to 50,000 IU per week are sufficient to raise VD levels above 40 ng/mL. Moreover, doses from 2000 IU per day can be related to improvements in the clinical scores of the disease and improvement in patients' QoL.

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## Summary Box

### What is already known:

- Vitamin D is a modulator of adaptive and innate immune responses
- Vitamin D reduces bacterial translocation to extra-intestinal tissues
- Vitamin D is associated with the downregulation of tumor necrosis factor- $\alpha$ , interleukins -6 and -12, thereby offering benefit to patients with inflammatory bowel disease

### What the new findings are:

- The effects of vitamin D on the remission of inflammatory conditions such as ulcerative colitis and Crohn's disease are still somewhat obscure
- The use of vitamin D in flared patients can interfere with remission
- The dosages of vitamin D can influence the outcomes

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