

# The endogenous cannabinoid system in the gut of patients with inflammatory bowel disease

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Activation of cannabinoid receptors (CBs) by endocannabinoids impacts on a number of gastrointestinal functions. Recent data indicate that CB1 agonists improve 2,4-dinitrobenzene sulfonic acid-induced colitis in mice, thus suggesting a role for the endocannabinoid agonist anandamide (AEA) in protecting the gut against inflammation. We here examined the gut endocannabinoid system in inflammatory bowel disease (IBD) patients, and investigated the *ex vivo* and *in vitro* effects of the non-hydrolysable AEA analog methanandamide (MAEA) on the mucosal proinflammatory response. The content of AEA, but not of 2-arachidonoyl-glycerol and *N*-palmitoylethanolamine, was significantly lower in inflamed than uninfamed IBD mucosa, and this was paralleled by lower activity of the AEA-synthesizing enzyme *N*-acyl-phosphatidylethanolamine-specific phospholipase D and higher activity of the AEA-degrading enzyme fatty acid amide hydrolase. MAEA significantly downregulated interferon- $\gamma$  and tumor necrosis factor- $\alpha$  secretion by both organ culture biopsies and lamina propria mononuclear cells. Although these results are promising, further studies are needed to determine the role of cannabinoid pathways in gut inflammation.

## INTRODUCTION

Inflammatory bowel diseases (IBD), which encompass Crohn's disease and ulcerative colitis, are driven by an excessive immune response in the gut.<sup>1,2</sup> Despite recent therapeutic advances and improved knowledge of the underlying pathogenic mechanisms, patients with IBD are often resistant to treatment, justifying the continued search for new therapeutic approaches.<sup>3</sup> Recently, understanding of the nature and regulation of endogenous cannabinoids in the gut has suggested that the endocannabinoid system might be a therapeutic target in IBD.<sup>4–6</sup>

The cannabinoid system comprises specific G-protein-coupled receptors (CB1 and CB2), a variety of exogenous (marijuana-derived cannabinoids) and endogenous ligands, and a machinery dedicated to endocannabinoid synthesis and degradation.<sup>7</sup> Endocannabinoids include amides and esters of long-chain polyunsaturated fatty acids, among which anandamide (AEA) and 2-arachidonoylglycerol (2-AG) are the main endogenous agonists of CB1 and CB2.<sup>8</sup> CB1 are predominant in brain and vessels

and mediate the psychoactive effects of cannabis, whereas CB2 are normally expressed on immune cells.<sup>9,10</sup> AEA shows higher affinity for CB1 over CB2, whereas 2-AG binds equally well to both CB1 and CB2.<sup>7–9</sup> Other endocannabinoid-like compounds have been described to date, such as *N*-palmitoylethanolamine (PEA) and oleoylethanolamide, which are unable to bind with significant affinity to either CB1 and CB2.<sup>11</sup>

Endocannabinoid signaling is terminated by specific degradation systems involving their cellular uptake by a facilitated transport mechanism and their hydrolysis by fatty acid amide hydrolase (FAAH) for AEA, monoacylglycerol lipase for 2-AG, and *N*-acylethanolamine-hydrolyzing acid amidase for PEA.<sup>7,11</sup> Although AEA synthesis might be because of several metabolic routes, *N*-acyl-phosphatidylethanolamine-specific phospholipase D (NAPE-PLD) is currently considered the major enzyme responsible for AEA production,<sup>12</sup> whereas 2-AG synthesis is mainly mediated by diacylglycerol lipase.<sup>7,13</sup> Several synthetic compounds have been shown to efficiently block either endocannabinoid uptake or degradation, thus providing

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pharmacological tools to enhance the activity of the endocannabinoid system.<sup>14</sup>

Besides their effects on central nervous system and on gastrointestinal motility, the anti-inflammatory properties of *Cannabis sativa* preparations have long been known, and potential roles of endocannabinoids in modulating the immune response have been recently reviewed.<sup>15–17</sup> In particular, AEA downregulates interleukin (IL)-2 transcription and secretion, reduces tumor necrosis factor (TNF)- $\alpha$  and interferon (IFN)- $\gamma$  production, induces lymphocyte apoptosis, and inhibits neutrophil recruitment at the site of inflammation.<sup>17</sup> Inflammatory stimuli may in turn control endocannabinoids by regulating their degradation. Indeed, lipopolysaccharide stimulation increases lymphocyte intracellular content of AEA by downregulating FAAH gene expression.<sup>18</sup>

There is evidence from experimental studies that the endocannabinoid system is involved in intestinal inflammation.<sup>4,5,15,19</sup> In particular, FAAH-deficient mice are protected from 2,4-dinitrobenzene sulfonic acid-induced colitis, whereas a pharmacological blockade of CB1 leads to a worsening of murine colitis.<sup>20</sup> On this basis, we first examined the levels of endocannabinoids and their receptors in IBD inflamed and uninflamed mucosa, and then investigated the *ex vivo* and *in vitro* effects of the synthetic non-hydrolysable AEA analog methanandamide (MAEA) on proinflammatory cytokine production by gut biopsies and lamina propria mononuclear cells (LPMCs) from IBD patients. Furthermore, we explored the action of MAEA on myofibroblasts isolated from Crohn's disease strictures.

## RESULTS

### *In vivo* mucosal endocannabinoid levels

Mucosal levels of AEA, 2-AG, and PEA were assessed by high-performance liquid chromatography (HPLC)-mass spectrometry (**Figure 1a**). Median levels of AEA were significantly lower in IBD inflamed mucosa (in Crohn's disease: 46 pmol g<sup>-1</sup>, range 15–89; in ulcerative colitis: 71 pmol g<sup>-1</sup>, range 36–286,  $P < 0.05$ ) in comparison with IBD uninflamed mucosa (in Crohn's disease: 123 pmol g<sup>-1</sup>, range 61–427; in ulcerative colitis: 122 pmol g<sup>-1</sup>, range 63–675). No significant difference was found in AEA levels between Crohn's disease and ulcerative colitis uninflamed mucosa and normal colon (131 pmol g<sup>-1</sup>, range 21–542). Unlike AEA, 2-AG median levels were comparable in IBD inflamed mucosa (in Crohn's disease: 5.0 nmol g<sup>-1</sup>, range 1.2–8.5; in ulcerative colitis: 6.5 nmol g<sup>-1</sup>, range 2.1–19.5), IBD uninflamed mucosa (in Crohn's disease: 4.6 nmol g<sup>-1</sup>, range 1.0–8.1; in ulcerative colitis: 5.2 nmol g<sup>-1</sup>, range 2.1–11.5) and normal colon (4.0 nmol g<sup>-1</sup>, range 1.5–9.1). PEA median concentration did not significantly differ in IBD inflamed mucosa (in Crohn's disease: 123 pmol g<sup>-1</sup>, range 57–1,127; in ulcerative colitis: 151 pmol g<sup>-1</sup>, range 55–380), IBD uninflamed mucosa (in Crohn's disease: 163 pmol g<sup>-1</sup>, range 54–677; in ulcerative colitis: 310 pmol g<sup>-1</sup>, range 55–1,394), and normal colon (109 pmol g<sup>-1</sup>, range 28–401). Based on these results, we further investigated the activity of AEA-metabolic enzymes. Among the 34 control subjects, no difference was found in the mucosal levels of AEA, 2-AG, and PEA between those who turned out to have

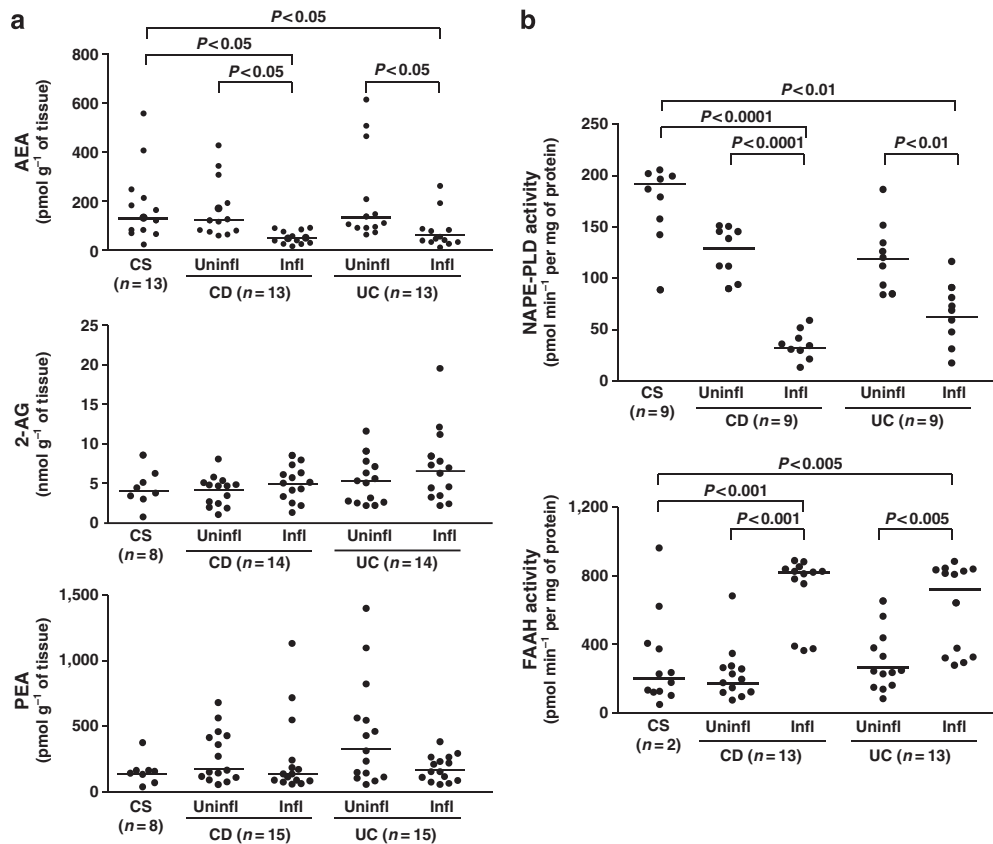
functional diarrhea at the end of their diagnostic work-up and those undergoing colectomy for colon cancer.

### *In vivo* mucosal enzyme activity

The activity of NAPE-PLD and FAAH, which respectively synthesize and degrade AEA, was assessed by radiochromatography (**Figure 1b**). Median NAPE-PLD activity was significantly lower in IBD inflamed mucosa (in Crohn's disease: 32 pmol min<sup>-1</sup> mg<sup>-1</sup>, range 13–58,  $P < 0.0001$ ; in ulcerative colitis: 63 pmol min<sup>-1</sup> mg<sup>-1</sup>, range 17–116,  $P < 0.01$ ) in comparison with IBD uninflamed mucosa (in Crohn's disease: 138 pmol min<sup>-1</sup> mg<sup>-1</sup>, range 89–150; in ulcerative colitis: 120 pmol min<sup>-1</sup> mg<sup>-1</sup>, range 83–186). No significant difference was found in NAPE-PLD activity between IBD uninflamed mucosa and normal colon (193 pmol min<sup>-1</sup> mg<sup>-1</sup>, range 74–208). On the contrary, median FAAH activity was significantly higher in IBD inflamed mucosa (in Crohn's disease: 820 pmol min<sup>-1</sup> mg<sup>-1</sup>, range 361–886,  $P < 0.001$ ; in ulcerative colitis: 711 pmol min<sup>-1</sup> mg<sup>-1</sup>, range 275–881,  $P < 0.005$ ) in comparison with IBD uninflamed mucosa (in Crohn's disease: 184 pmol min<sup>-1</sup> mg<sup>-1</sup>, range 73–680; in ulcerative colitis: 241 pmol min<sup>-1</sup> mg<sup>-1</sup>, range 80–651). No significant difference was found in FAAH activity between IBD uninflamed mucosa and normal colon (median 206 pmol min<sup>-1</sup> mg<sup>-1</sup>, range 98–911). Among the 34 control subjects, no difference was found in NAPE-PLD and FAAH activity between those who turned out to have functional diarrhea at the end of their diagnostic work-up and those undergoing colectomy for colon cancer.

### *In vivo* mucosal CB1 and CB2 expression

CB1 and CB2 expression was investigated through enzyme-linked immunosorbent assay (ELISA; **Figure 2a**), immunoblotting (**Figure 2b**), and immunohistochemistry (**Figure 2c–f**). CB1 concentration, detected by ELISA and expressed as a mean percentage of control subjects, was significantly higher in the inflamed mucosa of 14 IBD patients (in Crohn's disease: mean 262 $\pm$ 10%,  $P < 0.001$ ; in ulcerative colitis: mean 204 $\pm$ 13%,  $P < 0.05$ ) in comparison with the uninflamed mucosa (in Crohn's disease: mean 100 $\pm$ 14%; in ulcerative colitis: mean 100 $\pm$ 14%; **Figure 2a**). These results were confirmed both by immunoblotting experiments, which showed a higher expression of CB1 in inflamed than uninflamed mucosa of both Crohn's disease and ulcerative colitis patients (**Figure 2b**), and immunohistochemistry, which showed increased CB1 expression by both epithelial cells and LPMCs in inflamed IBD mucosa (**Figure 2d**) in comparison with IBD uninflamed mucosa (**Figure 2c**). On the contrary, no significant difference was found in CB2 levels detected by ELISA between inflamed IBD mucosa (in Crohn's disease: mean 106 $\pm$ 10%; in ulcerative colitis: mean 205 $\pm$ 12%) and uninflamed IBD mucosa (in Crohn's disease: mean 100 $\pm$ 19%; in ulcerative colitis: mean 100 $\pm$ 5%; **Figure 2b**). Accordingly, no difference was found in CB2 expression detected both by immunoblotting in inflamed mucosa in comparison with uninflamed mucosa, both in ulcerative colitis and Crohn's disease patients (**Figure 2b**), and immunohistochemistry that showed a similar pattern of CB2 expression in IBD uninflamed (**Figure 2e**)



**Figure 1** *In vivo* mucosal levels of endocannabinoids and anandamide (AEA)-synthesizing and AEA-degrading enzymes. **(a)** Levels of AEA, 2-arachidonoylglycerol (2-AG), and *N*-palmitoylethanolamine (PEA) were measured by high-performance liquid chromatography (HPLC)-tandem mass spectrometry, and **(b)** activity of *N*-acyl-phosphatidylethanolamine-specific phospholipase D (NAPE-PLD) and fatty acid amide hydrolase (FAAH) were assessed by radiochromatography, in the inflamed (Infl) and uninflamed (Uninfl) intestinal mucosa of Crohn's disease (CD) and ulcerative colitis (UC) patients, and in the normal gut of control subjects (CS).

and inflamed mucosa (**Figure 2f**), mostly limited to scattered mononuclear cells in the lamina propria. Among the 34 control subjects, no difference was found in CB1 and CB2 expression between those who turned out to have functional diarrhea at the end of their diagnostic work-up and those undergoing colectomy for colon cancer.

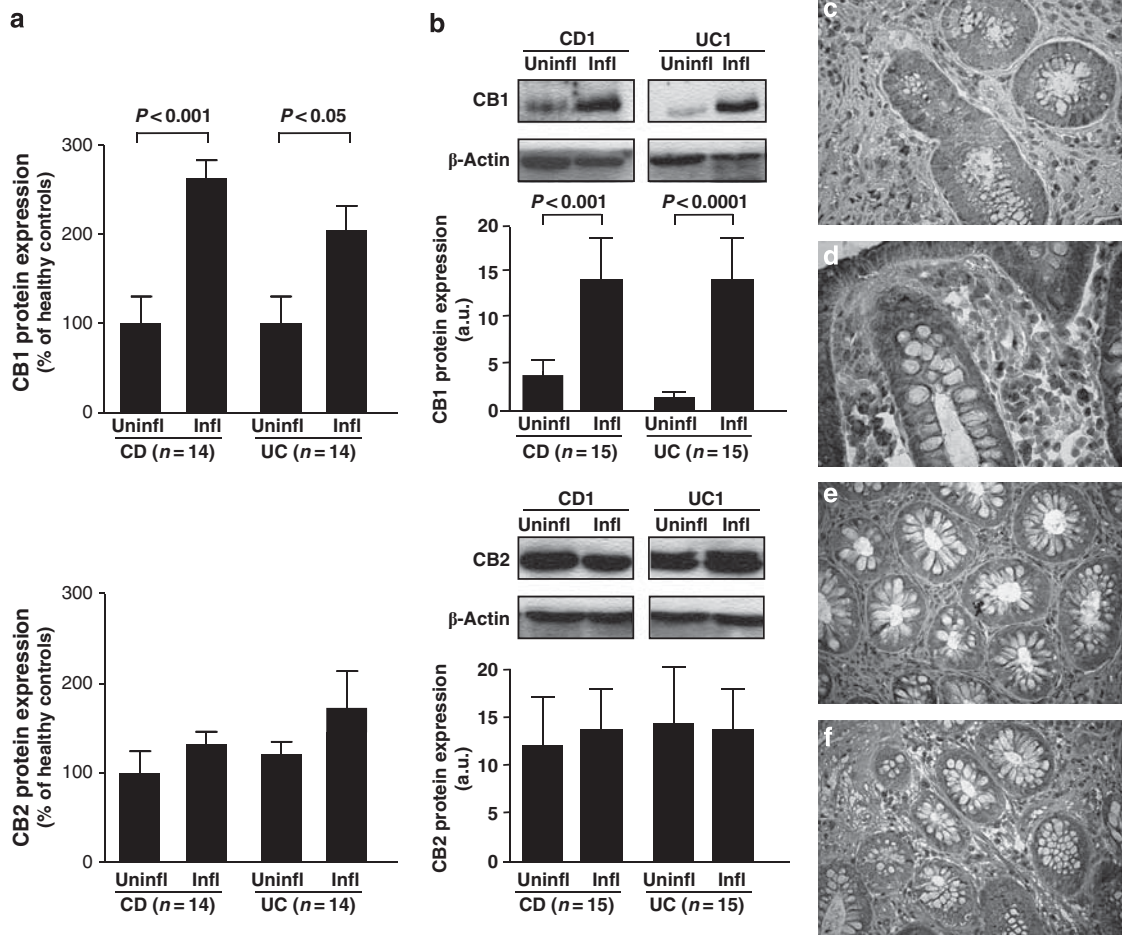
#### ***In vitro* and *ex vivo* effects of MAEA on proinflammatory cytokines**

To investigate the effect of the non-hydrolysable AEA analog MAEA on proinflammatory cytokine production, we measured IFN- $\gamma$  and TNF- $\alpha$  concentration in the supernatants of anti-CD3/CD28-stimulated LPMCs from inflamed areas of 33 IBD patients (18 with Crohn's disease and 15 with ulcerative colitis) after incubation with increasing concentrations of MAEA (**Figure 3a**). As shown in **Table 1a**, both IFN- $\gamma$  and TNF- $\alpha$  production by stimulated IBD LPMCs were significantly inhibited in a concentration-dependent manner by MAEA. It is noteworthy that the maximal effect of MAEA was comparable with that exerted by FK506. No appreciable difference in the efficacy of MAEA between Crohn's disease and ulcerative colitis samples was observed. In addition, we investigated the effect of MAEA on IFN- $\gamma$  and IL-17 production by purified CD3<sup>+</sup> LPMCs from 16 patients with IBD (8 with Crohn's disease and 8 with

ulcerative colitis), and we found that MAEA exerted a concentration-dependent downregulatory effect on IFN- $\gamma$  and IL-17 production by T cells (**Table 1b**). We then explored the influence of MAEA on proinflammatory cytokine production by organ culture biopsies from inflamed mucosa of the same patients (**Figure 3b**). A significant dose-dependent decrease of both IFN- $\gamma$  and TNF- $\alpha$  concentration was observed in the supernatants of IBD inflamed biopsies incubated with increasing concentrations of MAEA, in comparison with biopsies incubated with medium only: IFN- $\gamma$  (mean 344 $\pm$ 65 pg ml<sup>-1</sup> with medium alone vs. 69 $\pm$ 24 pg ml<sup>-1</sup> at 1  $\mu$ M ( $P$ <0.001) and 27 $\pm$ 9 pg ml<sup>-1</sup> at 10  $\mu$ M ( $P$ <0.0001)) and TNF- $\alpha$  (mean 132 $\pm$ 26 pg ml<sup>-1</sup> with medium alone vs. 52 $\pm$ 12 pg ml<sup>-1</sup> at 1  $\mu$ M ( $P$ <0.01) and 23 $\pm$ 9 pg ml<sup>-1</sup> at 10  $\mu$ M ( $P$ <0.001)). No appreciable difference in the efficacy of MAEA between Crohn's disease and ulcerative colitis samples was observed.

#### ***In vitro* effect of MAEA on T-bet and pSTAT4 expression**

We then tested whether MAEA influenced the expression of the transcription factor T-bet and the phosphorylated form of signal transducer and activator of transcription 4 (pSTAT4). Anti-CD3/CD28-stimulated LPMCs from inflamed areas of 21 IBD patients (11 with Crohn's disease and 10 with ulcerative colitis) were cultured in the presence or absence of 10  $\mu$ M MAEA and,



**Figure 2** *In vivo* mucosal expression of type-1 (CB1) and type-2 (CB2) cannabinoid receptors. CB1 and CB2 expression was detected by (a) enzyme-linked immunosorbent assay (ELISA), (b) immunoblotting, and (c–f) immunohistochemistry in the inflamed (Infl) and uninflamed (Uninfl) mucosa of Crohn's disease (CD) and ulcerative colitis (UC) patients. (a) Results are expressed as mean percentage of control subjects  $\pm$  s.d. (b) Each example shown in the upper panel is representative of experiments performed in 15 CD patients and 15 UC patients. Blots were stripped and analyzed for  $\beta$ -actin as an internal loading control. The lower panel shows the densitometry of CB1 and CB2 expression normalized for  $\beta$ -actin. Results are mean  $\pm$  s.d. arbitrary units (a.u.). CB1 expression, which is evident at epithelial and lamina propria mononuclear cell level, is higher in (d) inflamed than in (c) uninflamed inflammatory bowel disease (IBD) mucosa. CB2 expression, which is limited to a few mononuclear cells in the lamina propria, is similar in (e) uninflamed and (f) inflamed IBD mucosa.

after culture, T-bet and pSTAT4 were determined by immunoblotting on cell lysates and respectively normalized for  $\beta$ -actin and total STAT4 (Figure 4a and b). A significant ( $P < 0.001$ ) increase of both T-bet and pSTAT4 expression was observed when LPMCs were stimulated with anti-CD3/CD28 antibodies. This increase was significantly ( $P < 0.001$ ) inhibited by MAEA. No appreciable difference in the effect of MAEA on T-bet and pSTAT4 expression was observed between Crohn's disease and ulcerative colitis samples.

#### ***In vitro* effect of MAEA on Crohn's disease strictured myofibroblasts**

In order to assess the influence of the endocannabinoid system on intestinal fibrogenesis in Crohn's disease, we explored the effect of MAEA on collagen production and migration by myofibroblasts isolated from the submucosa of intestinal strictures of 11 patients with fibrostenosing Crohn's disease. Concentration of total soluble collagen was significantly ( $P < 0.01$ ) lower (mean  $140 \pm 33 \mu\text{g ml}^{-1}$ ) in the supernatant

of myofibroblasts cultured with MAEA in comparison with those cultured with medium alone (mean  $319 \pm 72 \mu\text{g ml}^{-1}$ ; Figure 5a). An *in vitro* wound-healing scratch assay was performed to assess the effects of MAEA on myofibroblast migration, which was measured as a percentage of wound repair. MAEA significantly ( $P < 0.05$ ) increased the migration of Crohn's disease strictured myofibroblasts at 4 h (from mean  $3.9 \pm 1.1$  to  $10.9 \pm 1.4\%$ ), 8 h (from mean  $5.1 \pm 1.0$  to  $15.1 \pm 1.6\%$ ), 12 h (from mean  $8.0 \pm 1.5$  to  $20.3 \pm 3.5\%$ ), 16 h (from mean  $10.8 \pm 2.1$  to  $21.9 \pm 1.3\%$ ), 20 h (from mean  $12.1 \pm 1.4$  to  $24.1 \pm 3.5\%$ ), and 24 h (from mean  $15.2 \pm 1.5$  to  $26.7 \pm 3.2\%$ ) in comparison with myofibroblasts cultured with medium only evaluated at the same time points (Figure 5b).

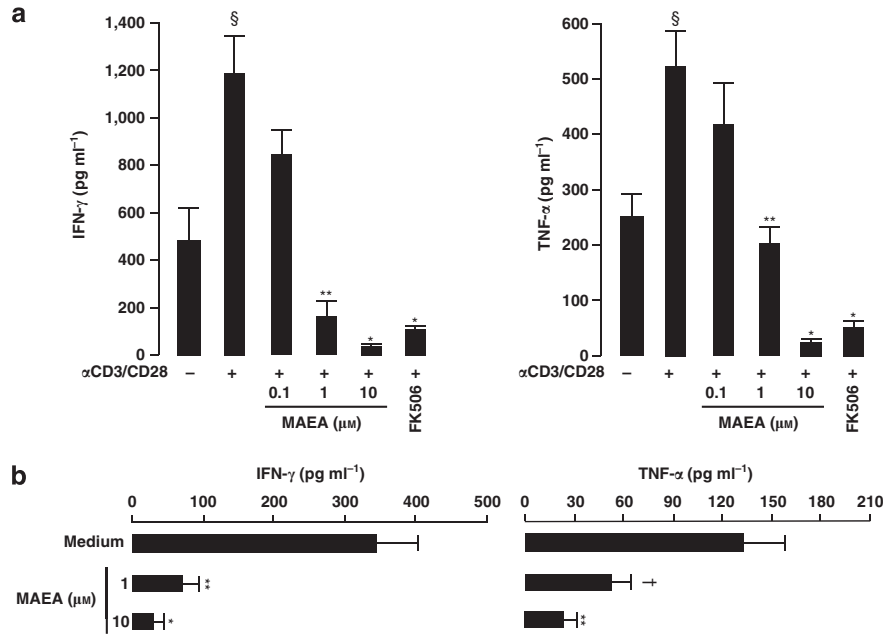
#### **DISCUSSION**

In this study we show that the content of the major endocannabinoid AEA is reduced in IBD inflamed mucosa as a consequence of both defective synthesis and increased degradation. The synthetic AEA analog MAEA exerts anti-inflammatory

*in vitro* and *ex vivo* functions, as shown by the downregulation of proinflammatory cytokines following incubation of IBD LPMCs and biopsies with the synthetic compound. Moreover, MAEA inhibits collagen production and restores migration of myofibroblasts isolated from Crohn's disease strictures.

Endogenous cannabinoid agonists, together with the enzymes and receptors involved in their metabolism and signaling processes, are present in different tissues such as skin, brain, spleen, uterus, kidney, and gut, both in health and disease.<sup>8</sup> There is increasing evidence that the endocannabinoid system plays an

autoprotective role in immunologically mediated disorders.<sup>15,16</sup> Nevertheless, most of the studies on this topic were conducted in animal models, and the way the endocannabinoid system exerts its protective activity in humans awaits further elucidation. We here detected AEA, 2-AG, and PEA levels in gut mucosa of IBD patients, and we found that AEA, but not 2-AG and PEA levels, are significantly reduced in inflamed compared with uninflamed areas, both in ulcerative colitis and Crohn's disease. We then asked the question whether this was secondary to a dysregulated AEA metabolism. On the basis of our results showing lower



**Figure 3** *In vitro* and *ex vivo* effects of methanandamide (MAEA) on proinflammatory cytokines. Interferon (IFN)- $\gamma$  and tumor necrosis factor (TNF)- $\alpha$  levels were detected by enzyme-linked immunosorbent assay (ELISA) (a) in the supernatants of lamina propria mononuclear cells (LPMCs) isolated from inflamed areas of 33 patients with inflammatory bowel disease (18 with Crohn's disease and 15 with ulcerative colitis), cultured for 48 h with anti-CD3/CD28 antibodies ( $\alpha$ CD3/CD28) in the absence or presence of increasing concentrations of MAEA (0.1–10  $\mu$ M) or 10  $\mu$ M FK506, and (b) in the supernatants of biopsy specimens from inflamed areas of the same patients cultured for 24 h in the absence or presence of increasing concentrations of MAEA (1 and 10  $\mu$ M). Results are mean  $\pm$  s.d. (a) <sup>§</sup> $P < 0.001$  vs. medium; \* $P < 0.0001$  vs.  $\alpha$ CD3/CD28; \*\* $P < 0.001$  vs.  $\alpha$ CD3/CD28; (b) \* $P < 0.0001$  vs. medium only; \*\* $P < 0.001$  vs. medium alone; <sup>†</sup> $P < 0.05$  vs. medium alone.

**Table 1a Levels of IFN- $\gamma$  and TNF- $\alpha$  (pg ml<sup>-1</sup>) in the supernatants of anti-CD3/CD28-stimulated ( $\alpha$ CD3/CD28) LPMCs isolated from the inflamed colon of 18 CD and 15 UC patients, and cultured for 48 h with increasing concentrations (0.1, 1, and 10  $\mu$ M) of MAEA or with 10  $\mu$ M FK506**

	IFN- $\gamma$		TNF- $\alpha$	
	CD	UC	CD	UC
Medium	536 (148)	466 (161)	266 (33)	238 (51)
$\alpha$ CD3/CD28	1,289 (173) <sup>§</sup>	1,103 (188) <sup>§</sup>	478 (53) <sup>§</sup>	564 (72) <sup>§</sup>
$\alpha$ CD3/CD28 + 10 $\mu$ M MAEA	52 (7)*	34 (12)*	35 (7)*	19 (4)*
$\alpha$ CD3/CD28 + 1 $\mu$ M MAEA	184 (46)**	156 (32)**	173 (25)**	229 (36)**
$\alpha$ CD3/CD28 + 0.1 $\mu$ M MAEA	935 (129)	777 (107)	432 (79)	402 (64)
$\alpha$ CD3/CD28 + 10 $\mu$ M FK506	144 (39)*	102 (42)*	62 (13)*	42 (9)*

Abbreviations: CD, Crohn's disease; IFN- $\gamma$ , interferon- $\gamma$ ; LPMC, lamina propria mononuclear cell; MAEA, methanandamide; TNF- $\alpha$ , tumor necrosis factor- $\alpha$ ; UC, ulcerative colitis.

Results are mean (s.d.).

<sup>§</sup> $P < 0.001$  vs. medium; \* $P < 0.0001$  vs.  $\alpha$ CD3/CD28; \*\* $P < 0.001$  vs.  $\alpha$ CD3/CD28.

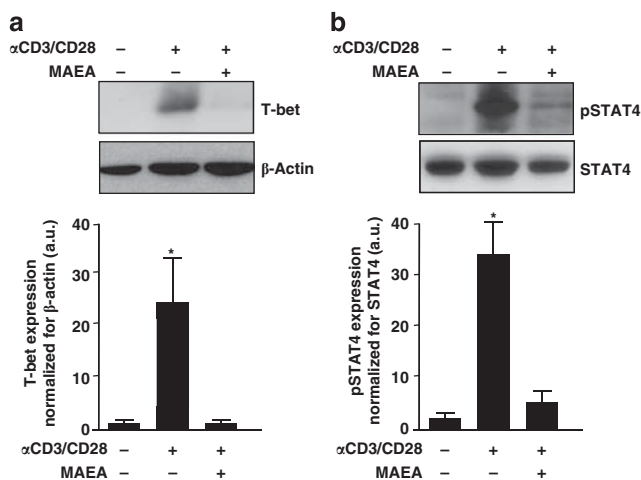
**Table 1b Levels of IFN- $\gamma$  and IL-17 (pg ml<sup>-1</sup>) in the supernatants of anti-CD3/CD28-stimulated ( $\alpha$ CD3/CD28) T cells isolated from the inflamed colon of 8 CD and 8 UC patients, and cultured for 48 h with increasing concentrations (0.1, 1, and 10  $\mu$ M) of MAEA or with 10  $\mu$ M FK506**

	IFN- $\gamma$		IL-17	
	CD	UC	CD	UC
Medium	1,035 (278)	1,175 (299)	1,328 (489)	1,149 (351)
$\alpha$ CD3/CD28	11,098 (2,671) <sup>§</sup>	12,953 (2,938) <sup>§</sup>	8,035 (1,738) <sup>§§</sup>	6,791 (1,272) <sup>§§</sup>
$\alpha$ CD3/CD28 + 10 $\mu$ M MAEA	2,537 (358)*	2,409 (497)*	2,098 (571)*	1,880 (430)*
$\alpha$ CD3/CD28 + 1 $\mu$ M MAEA	4,793 (987)**	5,031 (1,065)**	3,905 (727)**	3,781 (639)**
$\alpha$ CD3/CD28 + 0.1 $\mu$ M MAEA	9,107 (1,876)	9,328 (2,172)	7,692 (1,526)	6,391 (1,034)
$\alpha$ CD3/CD28 + 10 $\mu$ M FK506	1,239 (302)***	1,108 (263)***	1,572 (503)***	1,283 (467)***

Abbreviations: CD, Crohn's disease; IFN- $\gamma$ , interferon- $\gamma$ ; IL-17, interleukin-17; MAEA, methanandamide; UC, ulcerative colitis.

Results are mean (s.d.).

<sup>§</sup> $P < 0.0001$  vs. medium; <sup>§§</sup> $P < 0.001$  vs. medium; \* $P < 0.001$  vs.  $\alpha$ CD3/CD28; \*\* $P < 0.05$  vs.  $\alpha$ CD3/CD28; \*\*\* $P < 0.0001$  vs.  $\alpha$ CD3/CD28.



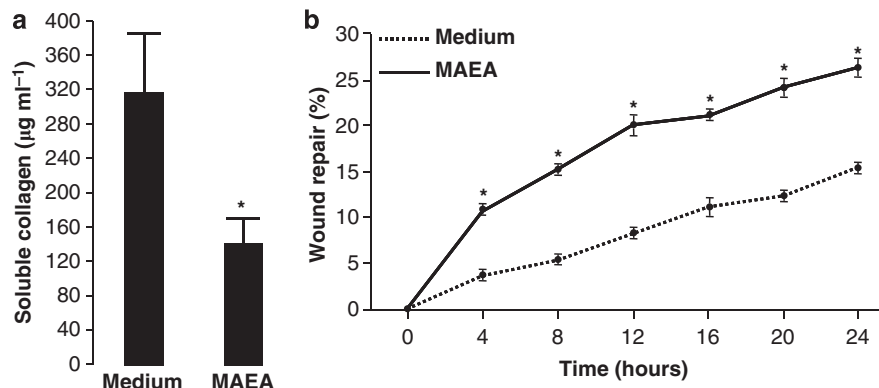
**Figure 4** *In vitro* effect of methanandamide (MAEA) on the expression of T-bet and the phosphorylated form of signal transducer and activator of transcription 4 (pSTAT4) in lamina propria mononuclear cells isolated from inflamed areas of patients with inflammatory bowel disease. (a) T-bet and (b) pSTAT4 were detected by immunoblotting of anti-CD3/CD28 ( $\alpha$ CD3/CD28)-stimulated lamina propria mononuclear cells incubated for 48 h in the absence or presence of 10  $\mu$ M MAEA. In the upper panels, each example is representative of experiments performed in 21 patients (11 with Crohn's disease and 10 with ulcerative colitis). Blots were stripped and analyzed for (a)  $\beta$ -actin or (b) total STAT4 as an internal loading control. The lower panels show (a) densitometry of T-bet normalized for  $\beta$ -actin, and (b) densitometry of pSTAT4 normalized for total STAT4. Results are mean  $\pm$  s.d. arbitrary units (a.u.). \* $P < 0.001$ .

NAPE-PLD activity and higher FAAH activity, we can conclude that reduced AEA levels result from both defective synthesis and increased degradation. FAAH can also degrade 2-AG *in vitro*,<sup>7,13</sup> yet variations of its activity observed in IBD mucosa did not exert any effect on 2-AG levels. Our results are in agreement with the observation of Marquez *et al.*,<sup>21</sup> who found lower NAPE-PLD expression and high AEA levels in inflamed ulcerative colitis when compared with control mucosa, but they diverge from those by D'Argenio *et al.*<sup>22</sup> Moreover, the findings of Marquez *et al.*,<sup>21</sup> who showed the lack of difference in the diacylglycerol lipase/monoacylglycerol lipase ratio between

ulcerative colitis patient and healthy control epithelium, support our data on mucosal 2-AG levels.

We then assessed CB expression in mucosal homogenates, and we found a higher expression of CB1 but not CB2 in inflamed compared with uninflamed areas of both Crohn's disease and ulcerative colitis patients. CB1 stimulation reduces gastrointestinal inflammation in various animal models,<sup>23</sup> and CB2 activation on immune cells downmodulates proinflammatory cytokine production.<sup>17,24–26</sup> CB1 is upregulated in the colon of mice treated with 2,4-dinitrobenzene sulfonic acid,<sup>20</sup> and Börner *et al.*<sup>27</sup> showed that anti-CD3/CD28-stimulation increases CB1, but not CB2, expression on human peripheral blood T cells. Numerous studies have shown a variable gut CB2 expression in animal models of colitis, either because they were performed in different species or because the protocols used diverged considerably.<sup>28</sup> An overexpression of CB2 at the ulcerative margin in active Crohn's disease has been reported,<sup>28</sup> which could be part of a redistribution of CB2 expression during inflammation, thus leaving the overall CB2 levels in the whole inflamed mucosa unchanged.

The endocannabinoid system plays a role in the modulation of several immune processes.<sup>15–17</sup> Cannabinoids reduce the MHC class II expression on the surface of dendritic cells, hence affecting antigen presentation, and inhibit peripheral T-cell activation in response to lipopolysaccharide and anti-CD3 antibodies.<sup>26,29,30</sup> Based on these findings, several studies have focused on the use of cannabinoids for the treatment of immune-mediated diseases, both in experimental models and humans.<sup>29</sup> As the dysregulated proinflammatory response observed in IBD pathogenesis is characterized by a T helper cell type (Th)1/Th17 immune profile,<sup>31</sup> we explored the influence of endocannabinoids on T cell-mediated gut inflammation, by investigating the *in vitro* and *ex vivo* effects of the synthetic non-hydrolysable AEA analog MAEA on proinflammatory cytokine production by anti-CD3/CD28-stimulated LPMCs, CD3<sup>+</sup> LPMCs, and biopsies from inflamed IBD mucosa. MAEA downregulated IFN- $\gamma$  and TNF- $\alpha$  production by LPMCs and biopsies in a concentration-dependent manner. Furthermore, MAEA exerted a direct immunomodulatory effect on T cells,



**Figure 5** *In vitro* effects of methanandamide (MAEA) on myofibroblasts isolated from the submucosa of strictured areas of 11 patients with fibrotic Crohn's disease. Myofibroblasts were cultured with medium alone or with medium containing 0.1 µM MAEA. **(a)** Soluble collagen concentration in myofibroblast culture supernatants. Results are mean±s.d. \* $P < 0.01$ . **(b)** Myofibroblast migration, assessed by *in vitro* wound-healing scratch assay, at 0, 4, 8, 12, 16, 20, and 24 h. Results, which are expressed as percentage of wound repair, are mean±s.d. § $P < 0.05$  vs. medium at 4, 8, 12, 16, 20, and 24 h.

as shown by the dose-dependent inhibition of IFN- $\gamma$  and IL-17 production by CD3<sup>+</sup> LPMCs. These data support a clear role for the endocannabinoid system in downregulating inflammation in IBD.<sup>32</sup> As the activated form of STAT4 (i.e., pSTAT4) and T-bet are essential mediators of naive T-cell polarization toward the Th1 phenotype,<sup>33</sup> we explored the effect of MAEA on these two transcription factors in anti-CD3/CD28-stimulated LPMCs from inflamed areas of IBD patients. MAEA significantly inhibited the anti-CD3/CD28-induced upregulation of both T-bet and pSTAT4, further supporting its role in dampening the Th1 immune response. Moreover, the inhibitory effect of MAEA on TNF- $\alpha$  production by total LPMCs suggests that not only T cells, but also monocytes and macrophages, which are the major source of TNF- $\alpha$ ,<sup>2</sup> are susceptible to cannabinoid action.

Of further interest is the fact that the endocannabinoid system plays a role in antagonizing the fibrogenic process in chronic liver disease.<sup>34</sup> Indeed, CB2 stimulation triggers potent antifibrogenic *in vitro* effects, such as growth inhibition and apoptosis, on hepatic stellate cells and myofibroblasts from patients with active cirrhosis.<sup>34</sup> Additionally, CB2 knockout mice are more sensitive than wild-type littermates to bleomycin-induced dermal fibrosis.<sup>35</sup> As >30% of patients with Crohn's disease develop intestinal fibrosis leading to stricture formation, frequently requiring surgical resection,<sup>36</sup> and myofibroblasts are key cells in the process of wound healing and stricture formation in the gut,<sup>37</sup> we investigated the *in vitro* effects of MAEA on myofibroblasts isolated from intestinal Crohn's disease strictures. We found that MAEA reduced collagen production by strictured Crohn's disease myofibroblasts and increased their migration ability, thus supporting the view that endocannabinoid pathways might be involved in the fibrogenic process in Crohn's disease.

## METHODS

**Patients and tissues.** Endoscopic biopsies or surgical specimens were taken from macroscopically and microscopically inflamed intestinal mucosa or strictured areas of 41 patients affected by Crohn's disease

(Table 2a) and 33 patients affected by ulcerative colitis (Table 2b). Diagnosis of Crohn's disease was ascertained according to the usual clinical criteria,<sup>38</sup> and the site and extent of the disease were confirmed by endoscopy, histology, and enteroclysis in all patients. Diagnosis of ulcerative colitis was ascertained according to the usual clinical criteria,<sup>39</sup> and the site and extent of the disease were confirmed by endoscopy and histology. None of the IBD patients had been ever treated with cyclosporine, methotrexate, infliximab, or adalimumab. Mucosal samples were also collected from the colon of 16 subjects who turned out to have functional diarrhea at the end of their diagnostic work-up (mean age 40.5 years, range 25–63), and from macroscopically and microscopically unaffected colonic areas of 18 patients undergoing colectomy for colon cancer (mean age 47.8 years, range 41–70). Each patient who took part in the study gave informed consent, and ethics committee approval was obtained.

**HPLC-tandem mass spectrometry.** Biopsies were subjected to lipid extraction with chloroform/methanol (2:1, v/v), in the presence of d<sub>8</sub>-AEA, d<sub>8</sub>-2-AG, and d<sub>4</sub>-PEA, as internal standards.<sup>40</sup> The organic phase was dried, pre-purified by chromatography on silica, and then analyzed by HPLC electrospray ionization mass spectrometry using an LCQ Duo quadrupole ion trap mass spectrometry (ThermoFinnigan, San Jose, CA) with a TSP4000 HPLC pump and a TSP AS3000 autosampler. Quantitative analysis was performed by selected ion recording over the respective protonated molecular ions.

**Radiochromatographic method.** The synthesis of [<sup>3</sup>H]AEA through the activity of NAPE-PLD (E.C. 3.1.4.4) was assayed in tissue homogenates (100 µg per test), using 100 µM [<sup>3</sup>H]NArPE,<sup>41</sup> by means of reversed-phase HPLC, as reported.<sup>42</sup> The hydrolysis of [<sup>3</sup>H]AEA by FAAH was assayed in tissue homogenates (50 µg per test) by measuring the release of [<sup>3</sup>H]ethanolamine, as reported.<sup>43</sup> AEA was from Sigma-Aldrich (St Louis, MO), whereas [<sup>3</sup>H]AEA (60 Ci mmol<sup>-1</sup>) and N-[<sup>3</sup>H]arachidonoyl-phosphatidylethanolamine ([<sup>3</sup>H]NArPE, 200 Ci mmol<sup>-1</sup>) were from ARC (St Louis, MO).

**Cell isolation and culture.** LPMCs were isolated as previously described.<sup>44</sup> Briefly, the epithelial layer was removed with 1 mM EDTA (Sigma-Aldrich). After stirring for 1 h at 37°C, the supernatant was removed and the remaining tissue was treated with type 1A collagenase (1 mg ml<sup>-1</sup>; Sigma-Aldrich) for 2 h with stirring at 37°C. The crude cell suspension was allowed to stand for 5 min to permit debris

**Table 2 Clinical features of patients with (a) CD (n=41) and (b) UC (n=33)**

Characteristics and parameters	N	Median (range)
<b>(a)</b>		
Age (years)		35.6 (18–69)
First attack	7	
<i>Intestinal location</i>		
Small bowel and colon	26	
Colon only	15	
<i>Disease behavior</i>		
Fistulizing	7	
Stricturing	11	
Luminal	23	
Duration of disease (months)		57.3 (1–212)
Number of recurrences		1.8 (0–5)
CDAI		203 (69–405)
<i>Treatment</i>		
Mesalazine	16	
Mesalazine + topical steroids	9	
Mesalazine + antibiotics	8	
Mesalazine + azathioprine/ 6-mercaptopurine	3	
Azathioprine/6-mercaptopurine	5	
<b>(b)</b>		
Age (years)		38.4 (19–62)
First attack	6	
<i>Intestinal location</i>		
Pancolitis	19	
Left-sided colitis	14	
Duration of disease (months)		56.9 (1–167)
Number of recurrences		1.8 (0–5)
Clinical Activity Index		6.5 (1–12)
<i>Treatment</i>		
Mesalazine	8	
Mesalazine + topical steroids	11	
Mesalazine + azathioprine/ 6-mercaptopurine	14	

Abbreviations: CD, Crohn's disease; CDAI, Crohn's Disease Activity Index; UC, ulcerative colitis.

sedimentation. Cells from the supernatant were washed twice, resuspended in 1 ml RPMI-1640 medium (Sigma-Aldrich) containing 10% fetal bovine serum, 100 U ml<sup>-1</sup> penicillin, and 100 µg ml<sup>-1</sup> streptomycin, and kept on ice until used. Purified CD3<sup>+</sup> cells were obtained by positive magnetic-activated cell sorting separation using a CD3<sup>+</sup> cell isolation kit (Miltenyi Biotec, Auburn, CA) after removing monocytes by

adherence. The resulting cell preparation contained >95% CD3<sup>+</sup> T cells as assessed by flow cytometry. Cells were not used if viability was <90%. Isolated LPMCs or CD3<sup>+</sup> T cells (2×10<sup>5</sup> cells per well) were stimulated in anti-CD3-coated 96-well plates (BD Biosciences, Oxford, UK) with anti-CD28 antibody (0.5 µg ml<sup>-1</sup>, eBioscience, San Diego, CA), and incubated for 48 h with medium containing MAEA (0.1, 1, 10 µM; Calbiochem, San Diego, CA) or FK506 (10 µM; Sigma-Aldrich). As negative control, 0.1% dimethylsulfoxide (Sigma-Aldrich) was used. After culture, supernatants were snap frozen and stored at -70 °C. Myofibroblasts were isolated from uninfamed mucosa overlying Crohn's disease strictures as previously described.<sup>45</sup> Briefly, after removing the epithelial layer with EDTA treatment, tissue samples were denuded of epithelial cells, and were subsequently cultured at 37 °C in a humidified CO<sub>2</sub> incubator in Dulbecco's modied Eagle's medium (Sigma-Aldrich) supplemented with 20% fetal bovine serum, 1% nonessential amino acids (Invitrogen, Paisley, UK), 100 U ml<sup>-1</sup> penicillin, 100 µg ml<sup>-1</sup> streptomycin, 50 µg ml<sup>-1</sup> gentamycin, and 1 µg ml<sup>-1</sup> amphotericin (Sigma-Aldrich). During culture, numerous cells appeared both in suspension and adherent to the culture dish. The cells in suspension were removed every 24- to 72-h culture period, and the denuded tissue was maintained in culture for up to 6 weeks. Established colonies of myofibroblasts were seeded into 25 cm<sup>2</sup> culture flasks and cultured in Dulbecco's modied Eagle's medium supplemented with 20% fetal bovine serum and antibiotics. At confluence, the cells were passaged using trypsin-EDTA in a 1:2 to 1:3 split ratio. Cells were grown up to at least passage 4 before there were enough to be used in stimulation experiments with MAEA, and were characterized by immunocytochemical staining as previously described.<sup>45</sup>

**Wound-healing scratch assay.** Myofibroblast migration was assessed according to the method of Rodriguez *et al.*<sup>46</sup> as modified by us.<sup>47</sup> Briefly, cells (2×10<sup>5</sup>) were seeded into Nunc cell culture dishes (Nalge Nunc International, Rochester, NY) with 2 mm grids, size 35×10 mm, in 2 ml of Dulbecco's modied Eagle's medium supplemented with 20% fetal bovine serum and antibiotics. The cells were maintained at 37 °C and 5% CO<sub>2</sub> until confluent. Once confluent, each dish of monolayer cells was given a mechanical wound by scoring with a 200 µl pipette tip, parallel to the grid bars along the central grid line. This permits easy viewing of the cells growing back together, and ensures that the 2 mm grid may be used as a reference, so that the wound areas can be measured and compared. Wound placement was checked with an Olympus inverted CK2 microscope (Olympus UK, London, UK). The medium was then removed, and the cells were washed five times with HL-1 serum-free medium (Cambrex Bio Science, Nottingham, UK) supplemented with antibiotics, and then replaced with 1.5 ml HL-1 medium with 0.1% dimethylsulfoxide (Sigma-Aldrich) or MAEA 0.1 µM. Photographs of the cells in each grid along the induced wound were taken at 0, 4, 8, 12, 16, 20, and 24 h, using a digital camera (Olympus Camedia 34–40 zoom, ×20 magnification) attached to a light microscope. The computer program "Image J" was used to measure the area of initial damage (images taken at time 0) and of the remaining damage at subsequent time points. Each grid image was observed separately, and two points per grid at the same position at every time point were measured using imaging software at the same magnification. The percentage of wound repair was then calculated.

**Organ culture.** Biopsy specimens were placed on iron grids in the central well of an organ culture dish and the dishes placed in a tight chamber with 95% O<sub>2</sub>/5% CO<sub>2</sub> at 37 °C.<sup>45</sup> Increasing concentrations of MAEA (1 and 10 µM) were added to the medium containing RPMI-1640 medium supplemented with 10% HL-1 (Cambrex Bio Science), 100 U ml<sup>-1</sup> penicillin, and 100 µg ml<sup>-1</sup> streptomycin. As negative control, 0.1% dimethylsulfoxide (Sigma-Aldrich) was used. After 24 h culture, supernatants were snap frozen and stored at -70 °C.

**Western blotting.** Western blotting was performed according to a modified method described previously.<sup>47</sup> In brief, tissue samples or cells were lysed in ice-cold lysis buffer (10 mM EDTA, 50 mM pH 7.4 Tris-HCl, 150 mM sodium chloride, 1% Triton-X-100, 2 mM phenylmethylsulfonyl



fluoride, 2 mM sodium orthovanadate, 10 mg ml<sup>-1</sup> leupeptin, and 2 mg ml<sup>-1</sup> aprotinin) and the amount of protein was determined by the Bio-Rad Protein assay (Bio-Rad Laboratories, Hemel Hempstead, UK). An equal amount of protein (20 µg) was loaded onto 10% sodium dodecyl sulfate-polyacrylamide gels and blotted onto polyvinylidene fluoride sheets (Amersham Biosciences, Piscataway, NJ) under reducing conditions. Membranes were blocked with 10% nonfat dried milk and 5% bovine serum albumin for 2 h, and then incubated overnight with rabbit anti-CB1 (diluted 1:500, Cayman Chemicals, Ann Arbor, MI), rabbit anti-CB2 (1:300, Cayman Chemicals), and mouse anti-β-actin (1:1,000), mouse anti-STAT4 (1:500), mouse anti-phospho-STAT4 (pSTAT4, 1:500), and mouse anti-T-bet (1:500), all from Santa Cruz Biotechnology (Santa Cruz, CA). Membranes were rinsed and incubated with the appropriate horseradish peroxidase-conjugated secondary antibody (1:2,000) in blocking solution. Detection was performed using West Dura Chemiluminescence System (Pierce, Rockford, IL).

**Immunohistochemistry.** Tissue blocks were fixed in 4% buffered formaldehyde and embedded in paraffin. Blocks were cut into longitudinal 4-mm thick sections that were mounted on glass slides with the positively charged surface and air-dried. After blocking endogenous peroxidase with 0.3% hydrogen peroxide for 8 min, proteolytic digestion was obtained with trypsin 2% in calcium chloride 0.1%, pH 7.8, for 5 min. Sections were then immersed in 0.01 M citrate buffer (pH = 6.0) and microwave-treated for 10 min at 410 W. After washing in phosphate-buffered saline, sections were incubated with an anti-CB1 antibody (Abcam, Cambridge, UK; ab23703; 1:100 dilution) or an anti-CB2 antibody (Abcam; ab45942; 1:100 dilution) overnight at 4°C, and then treated with a secondary biotinylated antibody and peroxidase-conjugated streptavidin (Biogenex, San Ramon, CA). Sections were then dehydrated in ethanol, cleared in xylene, and coverslipped with Eukitt mounting medium (Kindler GmBH, Freiburg, Germany).

**ELISA.** IFN-γ, TNF-α, and IL-17 were measured in culture supernatants using human IFN-γ ELISA, (Endogen, Woburn, MA), and human TNF-α Quantikine ELISA Kit and human IL-17 Quantikine ELISA Kit, both from R&D Systems (Abingdon, UK), according to the manufacturer's instructions. In addition, anti-CB1 (diluted 1:500) and anti-CB2 (diluted 1:300) antibodies were used to determine CB1 and CB2 protein expression by ELISA in mucosal homogenates. The A<sub>405</sub> values in unknown samples (20 µg protein per well) were within the linearity range of a calibration curve drawn with different amounts of homogenates (in the range 0–40 µg protein per well). The results were expressed as mean percentages (±s.d.) of control subjects.

**Collagen assay.** Total soluble collagen was measured using Sircol Collagen Assay kit (Biocolor, Belfast, UK) according to the manufacturer's instructions. The Sircol dye reagent has been formulated to bind specifically to the [Gly-X-Y]<sub>n</sub> helical structure found in collagen types I to XIV. The collagen content in each sample of cell supernatant was obtained as an average of three readings.

**Statistical analysis.** Data were analyzed in the GraphPad Prism statistical PC program (GraphPad Software, San Diego, CA) using the paired *t*-test and the Mann-Whitney *U*-test. A level of *P* < 0.05 was considered statistically significant.

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#### DISCLOSURE

The authors declared no conflict of interest.

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