Exacerbation of Oxazolone Colitis by Infection with the Helminth *Hymenolepis diminuta*

**Involvement of IL-5 and Eosinophils**

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Substantial data show that infection with helminth parasites ameliorates colitis; however, oxazolone-induced colitis is exaggerated in mice infected with the tapeworm, *Hymenolepis diminuta*. We tested the hypothesis that the IL-5 response to helminth infection enhances the severity of oxazolone-induced colitis. Mice were infected with *H. diminuta* and 8 days later were treated with oxazolone ± anti–IL-5 antibodies. Colitis was assessed 72 hours postoxazolone treatment by disease activity scores, myeloperoxidase activity, and histopathology. Other mice received injections of a replication-deficient adenovirus that carried the IL-5 (Ad.IL-5) gene or a control adenovirus (Ad.delete) ± oxazolone. The effect of *H. diminuta* ± oxazolone in CCL11/CCL22 (eotaxin-1 and 2) knockout (KO) mice was determined. Helminth infection and Ad.IL-5 treatment increased IL-5 and eosinophil numbers. In *in vivo* neutralization of IL-5 significantly reduced the severity of colitis in *H. diminuta* ± oxazolone-treated mice, and *H. diminuta* did not exaggerate oxazolone-induced colitis in CCL11/CCL22 KO mice. Mice receiving Ad.IL-5 only had no colitis, while oxazolone-induced colitis was more severe in animals cotreated with Ad.IL-5 (Ad.delete ± oxazolone was not significantly different from oxazolone only). Thus, while there is much to be gleaned about antiinflammatory mechanisms from rodent-helminth model systems, these data illustrate the caveat that infection with helminth parasites as a therapy could be contraindicated in patients with eosinophilia or elevated IL-5 unless coupled to appropriate measures to block IL-5 and/or eosinophil activity. *(Am J Pathol 2010, 177:2850–2859; DOI: 10.2353/ajpath.2010.100537)*

The potential of infection with parasitic helminths to reduce the severity of concomitant disease has been the focus of a cadre of investigators, who, using a variety of model systems, have produced substantial proof-of-concept data that infection with nematode, trematode, or cestode parasites reduces disease in animal models of inflammatory and autoimmune disease. Collectively, these studies have shown that as a consequence of the host response to infection with helminth parasites, there is mobilization of transforming growth factor (TGF-β), interleukin (IL)-10, Foxp3⁺ regulatory T cells, alternatively activated macrophages, and inhibition of IL-17– and interferon (IFN)-γ–driven events: one, or a combination, of these events could block the development of autoimmune and inflammatory disorders. These findings have been complemented by intriguing data suggesting that infection with viable parasitic nematodes could be a treatment for patients with inflammatory bowel disease (IBD) or asthma, whose condition is not managed by conventional therapies.

We have contributed to the concept of ‘helminth therapy,’ noting that while specific helminths may be identified that can be used as a treatment, the information gleaned from analyses of helminth-rodent model systems could also be the basis for the development of novel therapeutic approaches. However, parasitism comes at a cost to the host, and many species take a huge toll on...
human health and well-being. This coupled to the spectrum of iatrogenic disease means that one must be vigilant of the potential for infection with helminth parasites to exaggerate disease, and to provoke deleterious side-effects. In this context, we showed that in contrast to the exaggerate disease, and to provoke deleterious side-effects. In this context, we showed that in contrast to the dinitrobenzene sulfonic acid (DNBS) model of colitis, mice infected with the rat tapeworm *Hymenolepis diminuta* developed significantly more severe colonic inflammation when challenged intrarectally (ir.) with oxazolone. Grossly, DNBS and oxazolone-induced colitis in mice look similar but they are different in etiology and histopathology; one of the more notable differences being an increase in eosinophils in the colon of oxazolone-treated mice. Because a stereotypic human and murine response to infection with helminth parasites is eosinophilia, we hypothesized that the enhancement of oxazolone-induced colitis by infection with *H. diminuta* could be due to IL-5 and consequently eosinophils. To test this postulate, i) *H. diminuta*+oxazolone–treated mice were administered anti–IL-5 antibodies, and ii) mice were treated with adenovirus carrying the murine IL-5 gene ± oxazolone. *In vivo* neutralization of IL-5 blocked the enhanced colitis that develops in *H. diminuta*+oxazolone treated mice and in accordance with these data, overexpression of IL-5 resulted in a more severe disease in oxazolone cotreated mice. Many variables are at play in animal models, and while none fully recapitulate human disease, we present these data as a caveat that individuals with eosinophilia may not be good candidates for ‘helminth therapy,’ unless this is coupled with other therapeutics to block the effects of IL-5 and/or other eosinophil-derived molecules.

**Materials and Methods**

**Mice and Induction of Colitis**

Colitis was induced in anesthetized male BALB/c mice (7–9 weeks old; Charles River Animal suppliers, Quebec, Canada), BALB/c mice engineered to lack CCL11/CCL22 (kindly provided by Dr. M. Rothenberg, University of Cincinnati, Ohio) and BALB/c mice lacking the α chain of the IL-4 receptor (IL-4Rα: kindly provided by Dr. F. Brombacher, University of Cape Town, South Africa) via the intrarectal (ir.) delivery of 3 or 4 mg of oxazolone in 100 μl of 1:1 PBS:ethanol using a 3-cm lubricated polyethylene catheter. Time-matched vehicle-only treated mice served as controls. Mice were humanely sacrificed 72 hours postoxazolone treatment. Some studies using oxazolone use a sensitization and ir. rechallenge protocol. We opted to use a single ir. delivery in accordance with our earlier study and for comparisons with the DNBS model of colitis. Experiments conformed to the Canadian guidelines for animal welfare.

**Adenoviral Administration**

The adenovirus used was engineered to lack the E1 region and thus was replication-deficient and had the murine IL-5 gene inserted into its genome (Ad.IL-5). Replication-deficient adenovirus with no additional gene insertions (ie, Ad.delete) were used as a control. Under light anesthesia, mice received an intraperitoneal injection of either Ad.IL-5 or Ad.delete [10⁸ plaque forming units (pfu) of adenovirus/100 μl sterile PBS: adenovirus kindly provided by Dr. Z. Xing, McMaster University, Hamilton, ON, Canada] and 48 hours later were randomized into those receiving PBS or oxazolone.

**Infection with *H. diminuta* and Treatment with *In Vivo* IL-5 Antibody**

Mice under manual restraint received an oral gavage of 100 μl of sterile PBS containing five viable cysticercoids of *H. diminuta* and were challenged with oxazolone 8 days later. Other mice were administrated a total of 200 μg of anti–IL-5 (TRK5, a rat IgG: Cat# MM550C Pierce Endogen, Rockford, IL) delivered i.p. to lightly anesthetized mice 5 (50 μg AB), 7 (100 μg AB) and 9 (50 μg AB) days postinfection (with oxazolone treatment at 8 days postinfection). Time-matched controls consisted of oxazolone+ anti–IL-5 treatment and mice treated with an irrelevant rat IgG; (Biolegend, San Diego, CA).

**Assessment of Colitis**

**Macroscopic Assessment**

After oxazolone treatment the mice were monitored daily for signs of disease, including diarrhea (ie, wet or faecal-stained anus), weight loss, and behavioral changes (eg, lethargy, ruffled fur). At sacrifice, the entire colon was excised, weighed, and examined for signs of damage/ulceration. A disease activity score (DAS) was determined using a 5 point-scale: >10% drop in body weight (0 or 1); soft/loose stool, empty colon, wet anus (0 or 1); evidence of active bleeding or occult blood (0 or 1); presence of macroscopic ulcers (0 or 1); a maximum score of 5 was given if any animal had to be humanely sacrificed because of reaching a pre-determined end-point in disease severity. Because colitis results in shortening of the colon, colons were divided based on percentage of total length: the distal 10% was discarded, the next 20% was frozen in liquid nitrogen for use in the myeloperoxidase (MPO; an indicator of granulocyte infiltration) and eosinophil peroxidase (EPO) assay, and the next 20% (ie, the most proximal segment) was fixed in formalin for histological analysis.

**Myeloperoxidase (MPO) and Eosinophil Peroxidase (EPO) Determinations**

MPO (a marker of acute inflammation often associated with neutrophil infiltration) and EPO (a specific eosinophil marker) activities were determined in tissue extracts and the human acute promyelocytic leukemia cell line, HL-60 (clone 15) (ATCC), differentiated into eosinophils by culture for 7 days with 0.5 mmol/L butyric acid (Sigma Chemical Co.) and 5 ng/ml granulocyte macrophage col-
ony stimulating factor (GM-CSF: R & D systems)18 via a colorimetric kinetic assay in which H₂O₂ catabolism is measured. One unit of MPO activity in the samples is the amount of enzyme required to degrade 1 μmol/L H₂O₂/minute. The assay was repeated in the presence of 50 mmol/L 3-amin-o-1,2,4-triazole (AMT: Sigma Chemical Co.) to inhibit EPO, whose activity was calculated by subtracting the MPO+AMT value from the MPO value only.13

Histological Assessment

Segments of the colon were immersion-fixed in 10% buffered formalin (72 hours), dehydrated in graded alcohols, cleared in xylene, and embedded in paraffin wax. Sections (3 μm) were collected on coded slides, stained with hematoxylin and eosin and a histological damage score calculated on 12-point scale as previously described.13 Eosinophils were counted in 5 random high power (×40 objective) fields (HPF) of view.

Cytokine Production

Serum was obtained from murine blood samples 24 hours after Ad.IL-5 injection and in additional studies 72 hours after oxazolone and stored at −80°C. Interleukin-5 levels were determined by ELISA. Spleen cells, mesenteric lymph node (MLN) cells, or colon lamina propria lymphocytes (LPLs)19 were isolated and 5 × 10⁶ cells stimulated with concanavalin A (ConA: 2 μg/ml) for 48 hours, cell-free supernatants collected, and levels of IL-4, IL-5, and IL-10 measured. All ELISAs were performed in duplicate using antibody pairs from R&D Systems (Minneapolis, MN) and each assay had a limit of detection of 8 pg/ml.

Eosinophil Counts

Blood smears or cytospin preparations of bone-marrow cells retrieved from murine femurs were stained using the Hema3 differential staining kit (Fisher Scientific, Kalama-zoo, MI) and mononuclear cells, neutrophils, and eosinophils counted.

Epithelial Barrier Function

One million cells of the human colon-derived T84 cell line were seeded onto porous filter supports and cultured as described20 (human cell lines were used for these studies because of the lack of a mouse colon-derived epithelial cell line suitable for use in barrier studies). On reaching confluence (defined as transepithelial resistance (TER) ≥750 Ω/cm²), monolayers were cultured with 1 × 10⁵ HL-60 eosinophils (in the basal compartment of the culture well) ± LPS (Escherichia coli, 100 ng/ml; Sigma Chemical Co.) and epithelial barrier function assessed by measurements of TER and the lumen to basal flux of horseradish peroxidase (HRP) after a 48-hour period: i) TER was measured using a voltmeter and matched electrolytes (Millipore) at the beginning and end of the experiments and is presented as percentage of pretreatment values21; ii) HRP (20 ng type VI; Sigma Chemical Co.) was added to the apical chamber (1 ml vol.) of the well and three 10 μl samples of culture medium were collected from the basolateral chamber, diluted (1:10) in 0.5% HTAB buffer and enzymatic activity determined by spectrophotometric measurement (OD₄₅₀nm) of 3,3′,5,5′-tetramethylbenzidine (TMB) oxidation after the reaction was terminated with 2N H₂SO₄, and HRP determined against a concentration curve.

Statistical Analysis

All data are presented as the mean ± SEM, where n is the total number of mice. Statistical comparisons were performed via one-way analysis of variance, followed by post hoc pair-wise comparisons of the groups with either Student’s t-test or Tukey’s test, as appropriate and P < 0.05 set as a statistically significant difference.

Results

Use of an Anti–IL-5 Antibody Reduces the Exaggeration of Oxazolone-Induced Colitis in Mice Infected with H. diminuta

Direct ir. instillation of oxazolone resulted in an acute and self-resolving colitis: disease severity peaking 1–3 days posttreatment, with significant spontaneous healing by day 7 posttreatment (n = 6–7 mice; data not shown).13 Normal BALB/c animals completely expel H. diminuta, whereas intestinal washings from IL-4R KO mice revealed that these animals failed to spontaneously reject the helminth. Eleven days after infection 12 H. diminuta (length range = 8–12 cm; total biomass = 269.5 mg) were retrieved from the small intestine of 3 IL-4R KO mice (nb. each mouse given 5 cysticercoids), whereas no worms were obtained from time-matched identically infected wild-type BALB/c mice. Oxazolone-induced colitis in IL-4R KO mice was less severe than that observed in wild-type BALB/c mice (DAS = 3.8 ± 0.5 and 1.8 ± 0.5* for wild-type and IL-4R KO mice respectively (n = 7 mice, *P < 0.05), affirming a role for IL-4Rα signaling in the disease process and this is consistent with other studies using this model system.14

Successful infection with H. diminuta in BALB/c mice was confirmed by elevated IL-4 production from ConA-stimulated spleen cells: H. diminuta+oxazolone = 368 ± 144*, H. diminuta+oxazolone+anti–IL-5 = 898 ± 302*, control = 106 ± 19 pg/ml (n = 4; *, P < 0.05 compared to control). Furthermore, IL-5 production by conA-stimulated spleen cells and LPLs was significantly increased in cells from H. diminuta-infected mice ± oxazolone treatment (Figure 1, A and B), and this was paralleled by increased eosinophil production by the bone-marrow (Figure 1C), and eosinophil numbers in the blood (Figure 1D) and colon (Figure 1, E and F). Analysis of circulating and colonic eosinophils revealed no significant differ-
ences between naïve mice and those treated with oxazolone + H. diminuta + anti–IL-5 AB (Figure 1, D and E).

Consistent with earlier findings, oxazolone-induced colitis was characterized by loss of body weight (Figure 2A), shortening of the colon (Figure 2B), increased disease activity scores (Figure 2C), and significant histopathology (Figure 3, A and B). Administration of anti–IL-5 AB did not affect the outcome of oxazolone-induced colitis (n = 4; Figure 2), suggesting that while eosinophils are recruited to the colon14 their overall contribution to oxazolone-induced colitis is small and masked within the natural variation between mice in their response to oxazolone. Oxazolone-induced colitis was more severe in mice previously infected with H. diminuta (Figures 2 and 3)13; however, animals that also received anti–IL-5 AB had less colitis than H. diminuta + oxazolone treated mice and were similar in phenotype to naïve mice. Mice infected with H. diminuta and treated with oxazolone and irrelevant IgG1 had severe colitis (Figures 2 and 3) that was not statistically different from that in H. diminuta + oxazolone–treated mice. These data support an IL-5-eosinophil axis in the exaggeration of oxazolone-induced colitis in H. diminuta–infected mice.

Oxazolone-Induced Colitis in CCL11/CCL22 (Eotaxin 1/2) KO Mice Is Not Exaggerated by Infection with H. diminuta

CCL11/CCL22 KO mice developed colitis in response to topical application of oxazolone that was similar to that observed in wild-type BALB/c mice (Figure 4, A–C). This is consistent with in vivo neutralization of IL-5 having no effect on the outcome of oxazolone-induced colitis in
normal mice (Figure 2, A–C). In contrast to normal BALB/c mice, infection with *H. diminuta* did not exaggerate oxazolone-induced colitis in CCL11/CCL22 KO as gauged by all parameters of disease: disease activity scores, representative histological images and histological damage scores are shown in Figure 4. Indeed, prior infection with *H. diminuta* led to a degree of protection against oxazolone-induced colitis (Figure 4). Enumeration of eosinophils in histological sections of colons from control (n = 7), oxazolone (n = 6), and *H. diminuta*+oxazolone (n = 6)–treated CCL11/CCL22 KO revealed ≤5 eosinophils/HPF, with the majority of HPF containing 0–2 eosinophils. Thus, while eosinophil influx may characterize oxazolone-induced colitis, they are not a prerequisite for disease and significant colitis can develop in the absence of eosinophils as shown by this assessment of CCL11/CCL22 KO mice.

### Ad.IL-5 Enhances Oxazolone-Induced Colitis

Serum levels of IL-5 24 hours after injection of Ad.IL-5 were 488 ± 55* pg/ml compared to controls at 139 ± 21 pg/ml (n = 4; *, P < 0.05), confirming bioactivity of the adenovirus. Similarly, levels of IL-5 were significantly increased in sera from Ad.IL-5 or oxazolone+Ad.IL-5–treated mice at the end of the experiment (Figure 5).

Analysis revealed no statistical differences in the colon of ethanol-treated control mice and mice receiving either Ad.IL-5 or Ad.delete only and this dose of nonreplicating adenovirus did not evoke an increase in blood polymorphonuclear (PMNs) immune cells: control = 29 ± 5% versus Ad.IL5 = 27 ± 5% of blood immune cells. In contrast, mice treated with Ad.IL-5+oxazolone or Ad.delete+oxazolone displayed significant increases in PMNs (n = 4), of 61 ± 11% and 58 ± 9% of blood leukocyte, respectively. In addition, mice treated with Ad.IL-5 ± oxazolone had increased numbers of eosinophils in the colon: control = 9 ± 3, Ad.IL-5 = 28 ± 3*, oxazolone = 19 ± 4, Ad.IL-5+oxazolone = 23 ± 2* and Ad.delete+oxazolone = 18 ± 4* eosinophils/HPF of view (n = 3–4; *, P < 0.05 compared to control). There was no difference between the oxazolone- and Ad.IL-5+oxazolone–treated mice in terms of absolute eosinophil numbers, but this...
gives no indication of the activation status of the recruited cells. These histological observations were supported by significant increases in EPO levels in colonic segments excised from oxazolone/H11006 adenovirus treated mice as compared to naïve mice or mice injected with Ad.IL-5 or Ad.delete only (data not shown).

Mice cotreated with oxazolone/Ad.delete had, on average, slightly more severe disease (likely reflecting a response to the small dose of virus); however, there were no statistical differences in the colitis in Ad.delete/oxazolone compared to oxazolone-treated mice. In contrast, mice that received Ad.IL-5/oxazolone displayed a significant increase in the severity of the colitis (Figure 6, A–C): histological examination revealed greater derangement of colonic architecture, more inflammatory cell infiltrate, and more extensive epithelial erosion and ulceration (Figure 7). Colonic levels of MPO activity were quite variable between experiments, and were statistically significantly increased in tissues from oxazolone (0.9 ± 0.2,* n = 18), oxazolone+Ad.IL-5 (1.8 ± 0.4,*# n = 18) and oxazolone+Ad.delete (0.7 ± 0.2,* n = 9) treated mice compared to those from control mice (0.3 ± 0.05 U/mg tissue, n = 18) (*, P < 0.05 compared to control; #, P < 0.05 compared to oxazolone and oxazolone+Ad.delete).

LPS-Activated Eosinophils Decrease Epithelial Barrier Function

A 48-hour exposure to eosinophils or LPS only had negligible effects on the barrier function of T84 epithelial cell monolayers. However, coculture with the HL-60 eosinophil cell line exposed to 100 ng/ml of E. coli LPS had a significant impact on epithelial permeability as shown by a drop in transepithelial resistance of ~40% and a fourfold increase in the transepithelial flux of HRP (Figure 8, A and B).

Potentially Antiinflammatory Effects of Infection with H. diminuta Do Occur in the Oxazolone Model of Colitis

Consistent with our earlier studies, ConA stimulation of spleen cell or lamina propria lymphocyte cultures from mice infected 11 days previously with H. diminuta ± oxazolone (given at 8 days postinfection) produced more IL-10 than cells from control mice or those treated with oxazolone only (Figure 9).

Discussion

Proinflammatory roles have been shown for IL-5 and eosinophils in the context of inflammation in the airways. In comparison, the significance of increases in eosino-
phils in the blood or colon of patients with IBD, particularly those with ulcerative colitis, is unclear. The correlation of increased numbers of eosinophils in inflamed tissue and the ability of eosinophil-derived molecules (e.g., major basic protein) to cause tissue damage has led to the supposition that these cells contribute to inflammation: a postulate supported by data from animal models of colitis. However, the release of TGFβ from eosinophils can promote tissue restitution. Indeed, recognition of multiple functional phenotypes within a given type of immune cell lineage raises the possibility that subgroups of eosinophils could exert anti-inflammatory effects.

Infection with helminth parasites can reduce the severity of concomitant disease in animal models. Such studies have much to offer in the assessment of pathophysiology and the identification of endogenous mechanisms to down-regulate inflammation. However, the possibility that infection with helminth parasites could exaggerate disease cannot be overlooked, as for example in Citrobacter rodentium-induced enteropathy and oxazolone-induced colitis. Given the commonality of i) increased IL-4 in ulcerative colitis and murine oxazolone-colitis, ii) increased eosinophils in oxazolone-induced colitis and some patients with ulcerative colitis, and iii) the mobilization of IL-5 and eosinophils after infection with helmminth parasites, we hypothesized that IL-5 and eosinophils participated in the worsening of oxazolone-induced colitis in H. diminuta-infected mice. The subsequent analyses support three conclusions: first, the exaggeration of oxazolone-induced colitis observed in H. diminuta–infected mice is inhibited by in vivo neutralization of IL-5; second, increased production of IL-5 worsens the outcome of oxazolone-induced colitis; and, third, despite a more severe disease in H. diminuta/H. diminuta–infected mice compared to control and oxazolone, respectively; m, muscle; l, lumen; arrowhead, friable epithelium; arrow, inflammatory infiltrate; asterisk, ulceration; original magnification, ×200).

Figure 7. Representative images of H&E sections of the colon from the various groups of mice with the resultant histological damage scores from control (con), oxazolone (3 mg ip, +72 hours) and oxazolone/Ad delete or Ad IL-5 (10^6 pfu, ip.)-treated mice (mean ± SEM, n = 9–18 from 3 to 5 experiments; *P < 0.05 compared to control and **P < 0.05 compared to oxazolone, respectively; m, muscle; l, lumen; arrowhead, friable epithelium; arrow, inflammatory infiltrate; asterisk, ulceration; original magnification, ×200).

Figure 8. Coculture with HL-60 differentiated eosinophils (1 × 10^5) and LPS (100 ng/ml) results in increased permeability of monolayers of the human colon-derived T84 epithelial cell line as measured by (A) transepithelial electrical resistance (TER: a measure of the paracellular barrier to the passive flux of ions) and (B) the apical to basolateral flux of horse-radish peroxidase (HRP, a 44-kDa protein) 48 hours later [mean ± SEM; n = 6 epithelial monolayers from two experiments; starting TER = 790–2460 Ω/cm²; *P < 0.05 compared to control (i.e., T84 cells only); **P < 0.05 compared to other groups].

Figure 9. Bar graph showing IL-10 production from spleen cells and lamina propria lymphocytes (LPLs) 48 hours after ConA (2 μg/ml) stimulation (mean ± SEM; n = 3 from a representative experiment; *P < 0.05 compared to control and oxazolone only; nd, not detected).
treated mice, aspects of an antiinflammatory effect of *H. diminuta* still occur.

The beneficial effect of infection with *H. diminuta* in DNBS-induced colitis raises the question as to why infection with this parasite is so detrimental in oxazolone-induced colitis. Increases in IL-4 that are superseded by increases in IL-13 production and a mild colonic eosinophilia are characteristic of oxazolone-induced colitis. We showed that the exaggerated disease in *H. diminuta* + oxazolone – treated mice was accompanied by increases in IL-5 and CCL11 mRNA and enhanced eosinophilia in the colon. Here we extend these observations by showing that infection with *H. diminuta* results in increased bone marrow production of eosinophils, a blood eosinophilia, and that systemic and local levels of IL-5 can be increased via activation of spleen cells and LPLs. These data raise the possibility that the severe colitis in *H. diminuta* + oxazolone–treated mice could be via production of IL-5 and subsequent eosinophil differentiation and activation.

In vivo neutralization of IL-5 (using an antibody strategy) inhibited the *H. diminuta*–induced exaggeration of oxazolone–colitis (*H. diminuta*) was not observed in the small intestine of the anti–IL-5 AB–treated mice, indicating that key signal transducer and activator of transcription 6–driven events required for the expulsion of the worm were intact. Moreover, the amelioration of colitis in these animals was accompanied by reduced numbers of circulating and colonic eosinophils, supporting the postulate that *H. diminuta* exaggeration of oxazolone–induced colitis is at least partially due to the recruitment of eosinophils to the gut. Lack of IL-5 attenuated the accumulation of eosinophils in the colon of dextran-sodium sulfate–treated mice; however, the degree of colitis was similar in wild-type and IL-5 KO mice, likely reflecting redundancy of proinflammatory events in DSS–colitis. This increase in eosinophils in DSS–induced colitis may explain, at least partially, the finding that infection with *H. diminuta* alleviated the ion transport abnormalities but not the histopathology in this model of colitis. However, colitis in oxazolone and oxazolone + anti–IL-5 AB–treated mice were similar. Thus, while colonic eosinophilia is characteristic of this model of colitis, these cells may play only a small role in the disease such that the normal variation between mice in their response to oxazolone is large enough to mask any role of eosinophils that could potentially be revealed by neutralization of IL-5.

Mice deficient in CCL11 develop less colitis than normal animals treated with DSS. Treatment of CCL11/CCL22 KO with oxazolone resulted in an obvious colitis that was not appreciably different from that which develops in normal mice. However, in contrast to normal BALB/c mice, infection with *H. diminuta* reduced the degree of colitis in KO mice. These findings are compatible with eosinophil participation in *H. diminuta* enhancement of oxazolone–induced colitis and suggest that any antiinflammatory effect (ie, the increase in IL-10 production) of infection with *H. diminuta* is superseded by the eosinophil effect. This underscores the complexity of the inflammatory process in the gut, where the outcome will be dependent on the interplay between numerous immune, stromal and neuronal cells.

Consistent with the inhibition of colitis in *H. diminuta* + oxazolone treated mice by anti–IL-5 AB, overexpression of IL-5 significantly increased the severity of oxazolone–induced colitis. The question remains as to precisely how IL-5/eosinophils enhance oxazolone–induced disease. In vitro coculture studies reveal that eosinophils can reduce the barrier properties of epithelial cell monolayers and that this is enhanced by the addition of the bacterial tri-peptide, fMet-Leu-Phe. Furthermore, major basic protein was found to increase murine colonic epithelial permeability and exaggerate colitis. Also, preliminary evidence suggests that eosinophil activation results in increases in epithelial permeability in biopsy specimens from patients with ulcerative colitis. Thus, the current study (Figure 8) adds to a small body amount of data suggesting that one mechanism by which eosinophils could enhance intestinal inflammation is by the induction of decreases in epithelial barrier function.

Having implicated IL-5, a cytokine considered indispensable for eosinophil differentiation, survival, and function, in the exaggeration of oxazolone–induced colitis, other mechanistic possibilities should not be overlooked. Thus, participation of i) IL-4 and IL-13 [cytokines mobilized in response to infection with helminth parasites and oxazolone (and possibly important in ulcerative colitis also)], ii) NK and NK T cells, and iii) TGFβ should be assessed in the *H. diminuta* + oxazolone model and any putative interaction with IL-5 or eosinophils defined.

Some patients with ulcerative colitis experienced benefit when given viable ova of the nematode, *Trichuris suis*, while allergic rhinitis was unaffected by infection with this helminth. If eosinophils are increased in ulcerative colitis, how can these clinical observations be reconciled with the data herein? Eosinophil levels were not documented in the clinical trial. Thus, the disease in this cohort of patients may not have had significant eosinophil involvement or *T. suis* may not have evoked a large eosinophilia in patients with a positive response to the helminth therapy. Also, it is important to emphasize that the essence of parasitism is host–specificity, and so the local and systemic responses to *H. diminuta* (a cestode) in mice need not be identical to those in humans in response to *T. suis* (a nematode); thus, host–parasite specific interactions could explain the disparity between the human and animal model investigations. In addition, oxazolone–induced colitis recapitulates neither the etiology of ulcerative colitis nor all of the nuances of the human condition. Nevertheless the analysis of the *H. diminuta* + oxazolone model indicates that an eosinophilic response could undermine the benefit of antiinflammatory events mobilized subsequent to infection with helminth parasites.

In summary, this study confirms that infection with the rat tapeworm, *Hymenolepis diminuta*, exaggerates disease in a TH2–biased murine model of acute colitis and that IL-5—and by inference eosinophils—participates in the enhanced inflammation. However, an anticolitic effect of infection with *H. diminuta* was observed in CCL11/CCL22 KO mice, highlighting that antiinflammatory or
immunoregulatory events that accompany infection with helminth parasites could be exploited therapeutically. Finally, the impact of infection with a helminth parasite is contextual, depending on the health status of the host and the nature of any concomitant disease, and a consideration of these parameters could be paramount in the assessment of the potential use of a therapeutic helminth.

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References