PGRN protects against colitis progression in mice in an IL-10 and TNFR2 dependent manner

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This study was aimed to determine the role and regulation of progranulin (PGRN) in the pathogenesis of inflammatory bowel diseases (IBD). Dextran sulfate sodium (DSS)-, picrylsulfonic acid (TNBS)-induced, bone marrow chimera and CD4+CD45Rbhi T cell transfer colitis model were established and analyzed in wild-type and several genetically-modified mice, including PGRN, IL-10 and TNFR2 deficient mice. Elevated levels of PGRN were found in colitis samples from human IBD patients and mouse colitis models in comparison to the corresponding controls. PGRN-deficient mice became highly susceptible to DSS- and TNBS-induced colitis, whereas recombinant PGRN ameliorated the pathology and reduced the histological score in both DSS and TNBS colitis models. In addition, hematopoietic-derived PGRN was critical for protection against DSS-induced colitis, and lack of PGRN signaling in CD4+ T cells also exacerbated experimental colitis. PGRN-mediated protective effect in colitis was compromised in the absence of IL-10 signaling. In addition, PGRN’s effect was also largely lost in the TNFR2-deficient colitis model. Collectively, these findings not only provide the new insight into PGRN’s anti-inflammatory action in vivo, but may also present PGRN and its derivatives as novel biological agent for treating IBD.

Inflammatory bowel diseases (IBD) results from a continuum of complex interactions between a quart of host-derived and external elements. IBD including both Crohn’s disease (CD) and Ulcerative colitis (UC), is characterized by an imbalance in the mucosal intestinal immune system and a shift towards the proinflammatory side. Nearly 4 million people worldwide are affected by either Ulcerative colitis or Crohn’s disease. It has been well established that TNFα plays a crucial role in the pathogenesis of IBD, and TNF-blockers represent a well-accepted therapeutic option in the treatment of IBD. Since the TNF blockade only show clinical remission in 50–70% of patients of IBD, alternative biological therapies can be an attractive candidate for further research.

Progranulin (PGRN), also known as granulin epithelin precursor (GEP), PC-cell-derived growth factor (PCDFG), proepithelin, and acrogranin, is a 593-amino-acid secreted growth factor. PGRN plays a critical role in a variety of physiologic and disease processes, including early embryogenesis, wound healing, inflammation, and host defense. PGRN also function as a neurotrophic factor. PGRN exerts its therapeutic effect in inflammatory arthritis through binding to TNFR and DR3, and in turn disturbing TNFα/TNFR and TL1A/DR3 interactions. PGRN−/− bone marrow-derived macrophages (BMDMs) reduced IL-10 production and increased amounts of pro-inflammatory cytokines. PGRN also suppresses TNFα and IL-6 secretion from activated microglia. Recombinant PGRN was able to block TNFα-induced respiratory robust from neutrophils.

It was reported that PGRN antibodies occurred frequently in patients with CD and UC. Cytotoxicity assays showed a proinflammatory effect of PGRN antibodies on human colon HT29 cells. Moreover, PGRN-antibodies, opposite to recombinant PGRN, enhanced TNF-α-induced downregulation of FOXP3 in CD4+CD25hi Tregs. In this study, we investigated the role of PGRN in intestinal inflammation and the potential molecular mechanism involved.

Methods

Human biopsies. Collection of biopsies from human colons was carried out in accordance with the guidelines and approved by IRB, and informed consent was obtained from all subjects during colonoscopy procedures at the Mount Sinai Hospital, New York in collaboration with Dr. Lloyd Mayer.
**Results**

PGRN is elevated in the colon from IBD patients and mice colitis model. To determine the expression of PGRN in intestinal tissue during inflammatory conditions we performed immunohistochemistry in the human intestinal tissue samples. As shown in Fig. 1A, the expression of PGRN is significantly elevated in the intestinal tissue of IBD patients compared with non-inflammatory tissue controls. PGRN was detectable in normal colon tissues (day 0), and its expression was significantly increased at day 1 after TNBS injection (Fig. 1B). Stronger PGRN staining was also observed in epithelial cells and inflammatory cell infiltrating areas at day 4 (Fig. 1B). We performed real-time PCR to examine PGRN gene expression pattern in immune cells sorted from normal wild-type mice spleens. We found that PGRN gene was highly expressed in CD11b+ cells and Ly6G+ cells compared to other immune cells including T and B cells (Fig. 1C). To determine whether PGRN is functional on the intestinal cells, we stimulated the human intestinal epithelial Caco2 cells with 1 µg/ml recombinant PGRN, we found that PGRN activated Erk phosphorylation (Fig. 1D). These data suggests that PGRN is induced in the IBD inflammatory conditions and is functional on the intestinal cells.

PGRN deficient mice develop more severe colitis in comparison to the wild-type mice in chemical-induced colitis model. To study the contribution of endogenous PGRN to colitis development, we assessed the age- and sex-matched wild-type and PGRN−/− mice after oral administration of 3% DSS in drinking water. In a survival study, no wild-type mice died until day 10 after DSS induction, but a mortality higher than 60% was observed in the PGRN−/− group (Fig. 2A). The PGRN−/− mice suffered from significant body weight loss from day 4 (day 4: 97.8 ± 1.2% vs. 91.6 ± 5.2%, p = 0.02; day 5: 96.1 ± 0.9% vs. 87.3 ± 4.7%, p = 0.002; day 6: 90.3 ± 1.5% vs. 79.5 ± 4.6%, p = 0.0008; day 7: 86.3 ± 1.7% vs. 75.6 ± 2.2%, p = 0.0005) (Fig. 2B). In addition, the stool consistency scores of PGRN−/− mice became significantly worse compared to those of DSS-fed wild-type mice (day 5, p = 0.005; day 6, p = 0.04; day 7, p = 0.005) (Fig. 2C). And PGRN−/− mice displaying significantly elevated bleeding scores (day 4, p = 0.03; day 5, p = 0.005; day 6, p = 0.013; day 7, p = 0.009) (Fig. 2D) relative to DSS-induced wild-type mice. The evaluation of colon length is one of the parameter with the lowest variability in DSS-induced colitis model. We found that the colons of PGRN−/− mice were on average 20% shorter than those of wild-type mice treated with DSS (7.5 ± 0.5 vs. 6.5 ± 0.5, p = 0.03) (Fig. 2E). Colitis tissue from DSS-administered mice were examined to determine whether clinical signs of colitis were correlated with histological severity. Marked histopathological changes were seen in

**Immunoblotting.** The human intestinal epithelial Caco2 cells were lysed in RIPA lysis buffer containing protease inhibitors. Proteins were resolved on a 10% SDS-polyacrylamide gel and electroblotted onto a nitrocellulose membrane. After blocking with 5% non-fat dry milk in Tris buffer-saline-Tween 20 (10 mM Tris-HCl, pH 8.0; 150 mM NaCl; and 0.5% Tween 20), blots were incubated with anti-Erk, -Phospho-Erk (Cell Signaling Technology) antibody for 1 hour. After washing, the secondary antibody (horseradish peroxidase-conjugated) anti-rabbit immunoglobulin (1: 2000 dilution) was added, and bound antibody was visualized using an enhanced chemiluminescence system (Amer sham Life Science, Arlington Heights, IL, USA).

**Flow cytometry.** Cells were stained with anti-CD16/32, -CD3, -CD4, -CD8, -CD11c, -CD19, -CD25, -MHCI, -mPDCA-1 (bTraffic). Intracellular staining for Foxp3 (bioscience staining buffer set) and IL-17 and IFN-γ were conducted according to manufacturers’ instructions (Fixation and Fixation and Permeabilization Solution Kit Briege and eBioscience, respectively). Data were acquired on a LSRII (bBiosciences) and analyzed with FlowJo software (Tree star, Ashland OR).

**Statistical analysis.** Results represent mean ± SEM. Differences in groups were analyzed with unpaired, 2-tailed t tests. P values less than 0.05 were considered significant.
Figure 1 | PGRN expression is increased in intestinal samples from human IBD patients and mouse colitis models. (A) Immunohistochemistry for PGRN was conducted on sections of noninflammatory bowel disease and IBD patient’s intestinal tissue. (B) PGRN expression was detected in colonic sections from wild-type (C57BL/6) spleens. (D) Human intestinal Caco2 cells were treated with PGRN at different time-points, and the Erk signaling was examined.

colonic sections of DSS-fed PGRN−/− mice characterized by severe transmural inflammation with focal areas of extensive ulceration and necrotic lesions, along with infiltration of inflammatory cells (Fig. 2F). Semiquantitative scoring of these histological parameters confirmed that colitis severity in PGRN−/− mice was obviously higher than that in wild-type mice (p < 0.01) (Fig. 2G).

To determine whether these observations were specific to DSS-induced colitis model, another acute colitis model which resembles Crohn’s disease was established through intrarectal administration of 150 mg/kg picrylsulfonic acid (TNBS) in ethanol. Compared with WT mice, PGRN−/− mice significantly lost more body weight at day 3 and 4 after TNBS induction (day 3: 88.7 ± 4.6% vs. 82.3 ± 4.6%, p = 0.039; day 4: 94.7 ± 3.8% vs. 86.1 ± 2.3%, p = 0.005) (Fig. 3A). In addition, the histological features of PGRN−/− mice were significantly different from wild-type mice (Fig. 3B). Taken together, these results suggest that PGRN-dependent signaling is critical for protection against DSS- and TNBS-induced acute colitis.

Hematopoietic-derived PGRN is critical for protection against DSS-induced colitis. PGRN is highly expressed in epithelial cells as well as in immune cells17. To investigate whether immune cell-derived PGRN plays a role in DSS-induced colitis model, we generated chimera mice made by reconstituting irradiated wild-type mice with bone marrow cells from KO/WT donors, then subjected them to 3% DSS solution for 5 days. The wild-type mice reconstituted with PGRN+/− bone marrow cells lost more body weight compared to wild-type mice transplanted with wild-type bone marrow (day 4: 101.4 ± 2.1% vs. 98.8 ± 2.2%, p = 0.03; day 5: 100.5 ± 0.8% vs. 98.4 ± 2.9%, p = 0.05; day 6: 99.5 ± 0.7% vs. 96.5 ± 3.1%, p = 0.019; day 7: 97.3 ± 1.1% vs. 94.6 ± 1.8%, p = 0.011) (Fig. 4A). HE stained colon sections of wild-type mice that received PGRN+/− bone marrow displayed severe epithelial ulceration and more inflammatory lymphocytes infiltration (Fig. 4B) and significantly higher scores (p < 0.01, Fig. 4C). We found that wild-type mice transplanted with wild-type bone marrow were less sensitive to DSS-induced colitis. Collectively, these results demonstrate that hematopoietic cell-derived PGRN protects against DSS-induced colitis.

Lack of PGRN signaling in CD4+ T cells also exacerbates experimental colitis. To determine whether PGRN played a role in chronic colitis model, we established CD4+CD45RB+ T cells transfer colitis model. The results demonstrated that transfer of PGRN−/−CD4+CD45RB+ T cells led to an accelerated onset of body weight loss at d28 and d34 (p < 0.01) (Fig. 5A). RAG1−/− mice that received PGRN−/− T cells had a greater rate of mortality (Fig. 5B) and showed marked alterations of the colon (Fig. 5C). In line with the findings of accelerated body weight loss, lack of PGRN signaling in CD4+ T cells leads to more severe signs of intestinal inflammation.

The severity of colonic inflammation is not resulted from the impairment of intestinal barrier function or the alternations of colonic microbiota in DSS-fed PGRN−/− mice. Impairment of the intestinal barrier further drives the pathogenesis of intestinal inflammation in DSS-induced colitis and human IBD30. The results revealed that no significant changes were observed between wild-type and PGRN−/− DSS mice (water: 19.3 ± 3.0 vs. 18.2 ± 5.7, p = 0.39; DSS model: 131.5 ± 25 vs. 131.8 ± 20, p = 0.495) (Fig. 6A). These results suggest that PGRN deficiency does not affect the function of the intestinal barrier at either physiologic or disease conditions.

Intestinal microbiota plays a crucial role in the pathogenesis of inflammatory bowel disease31. To determine whether lack of PGRN signaling affects intestinal bacterial overgrowth and leads to bacteremia, we performed bacteria assay. Unexpectedly, we did not observe any significant changes in the number of bacteria between naive and DSS-treated wild-type and PGRN−/− mice (Fig. 6 B–E). These results demonstrate that the severity of colitis in PGRN−/− mice is not caused by bacterial overgrowth or due to the alteration of the composition of intestinal microbiota.
Recombinant PGRN ameliorates colitis syndrome. In order to determine whether recombinant PGRN is able to ameliorate the colitis syndrome in wild-type mice, a therapeutic group of mice (n = 8) were treated with 100 μg PGRN every two days beginning at day 1 after DSS induction, whereas control group (n = 8) were treated with PBS. The PBS group exhibited the great amounts of body weight loss and were unable to recover as rapidly in comparison to the PGRN-treated group (day 6: 93 ± 3.7% vs. 98 ± 3.5%, p = 0.02; day 7: 90 ± 5.0% vs. 95 ± 3.2%, p = 0.02; day 8: 88.6 ± 6.7% vs. 96.8 ± 4.7%, p = 0.009) (Fig. 7A). And the colon was shorter in control group than in mice treated with recombinant PGRN (p, 0.05) (Fig. 7B). PGRN treatment significantly increased the IL-10 release in colonic explants from DSS colitis mice (p < 0.01) (Fig. 7C). Histological results also showed that recombinant PGRN protected against experimentally induced colonic hyperplasia and leukocyte infiltration in colonic tissues (Fig. 7D).

DSS-induced colitis was also observed in immunodeficient mice 32. In a separate group, we treated colitic Rag1−/− mice with PBS and PGRN as described above. Colitic Rag1−/− mice treated with recombinant PGRN lost significantly less body weight when compared with those receiving PBS (day 6: 82 ± 8.7% vs. 91 ± 5.9%, p = 0.04; day 7: 81.1 ± 9.7% vs. 91.4 ± 4.5%, p = 0.02) (Fig. 8A). Colons of PBS-treated DSS-fed mice were significantly shorter than those of recombinant PGRN-treated colitic mice (Fig. 8B). Furthermore, HE staining of colonic sections confirmed the amelioration in colitis severity in PGRN-treated colitic mice characterized by focal areas of reduced ulceration and less leukocyte infiltration (Fig. 8C).

IL-10 signaling is required for PGRN-mediated protection from intestinal inflammation. IL-10 plays a critical role in regulating inflammatory response in intestine since the most prominent phenotype observed in IL-10-deficient mice is inflammation in the intestine 33. PGRN−/− mice macrophages produce less IL-10 than wild-type mice, and the serum IL-10 level is significantly elevated in PGRN−/− mice 18. To determine whether IL-10 signaling is required for PGRN-mediated protection from colitis, we blocked the IL-10 signaling with a specific anti-IL-10...
receptor antibody (clone 1B1.3A). Control group mice (n = 5) were injected with recombinant PGRN and isotype antibody, whereas blocking group mice (n = 5) received PGRN and 200 μg of anti-IL10R mAb per mice every 2 days. As shown in Fig. 9A, the anti-IL10R mAb blocking group mice suffered from significant body weight loss from day 4 (day 4: 97.8 ± 2.2% vs. 91.6 ± 2.0%, p = 0.001; day 5: 99.1 ± 1.1% vs. 93.8 ± 1.6%, p = 0.0002; day 6: 98.8 ± 3.5% vs. 90.4 ± 0.6%, p = 0.007; day 7: 97.4 ± 4.7% vs. 86.9 ± 5.2%, p = 0.013; day 8: 97.4 ± 6.1% vs. 85 ± 6.6%, p = 0.007; day 9: 98.6 ± 6.8% vs. 90.4 ± 0.6%, p = 0.007; day 8: 97.4 ± 6.1% vs. 85 ± 6.6%, p = 0.007). These results demonstrated that anti-10R mAb offsets the protection of PGRN against body weight loss. Marked inflammatory cell infiltration, colonic ulceration and changes of crypt architecture were also observed in intestinal sections of anti-10R treatment mice (Fig. 9B). In addition, PGRN’s therapeutic effect was lost in IL-10−/− mice, at least in part (day 3: 87.1 ± 2.5% vs. 84.4 ± 5.1%, p = 0.16; day 4: 86.7 ± 2.2% vs. 88.3 ± 4.8%, p = 0.026; day 5: 88.2 ± 3.3% vs. 90.5 ± 4.2%, p = 0.20) (Fig. 9C). Intestinal sections of the PGRN group and PBS group have similar histological changes (Fig. 9D). No significant changes were observed in Foxp3, IFN-γ and IL-17 level in T cells isolated from MLN between PGRN and PBS group (CD4+IL-17+ cells: 2.52 ± 0.5% vs. 2.07 ± 0.8%, p = 0.21; CD4+IFN-γ+ cells: 1.51 ± 0.48% vs. 1.62 ± 0.69%, p = 0.21; CD4+Foxp3+ cells: 10.5 ± 1.7% vs. 9.88 ± 2.1%, p = 0.49) (Fig. 9E). These results suggest that IL-10 signaling is required for PGRN-mediated protection from intestinal inflammation.

Discussion

The objective of this study was to examine the potential role of PGRN in the pathogenesis of IBD as well as the molecular mechanism involved. It was reported that PGRN expression was increased during colonic inflammation and PGRN was a critical protein involved in colonic injury repair. We further revealed that PGRN expression was elevated in the patients with IBD, and its level was induced in the course of mouse experimental colitis (Fig. 1), suggesting that PGRN might be an important growth factor involved in mucosal inflammatory response and damage. Although PGRN−/− mice have no evidence of immune system activation and spontaneous colitis

Figure 3 | PGRN−/− mice are highly susceptible to TNBS-induced colitis. (A) Body weight of WT (n = 6) and PGRN−/− mice (n = 5). (B) HE staining of representative colonic sections. Scale bars, 400 μm.

Figure 4 | PGRN signaling in hematopoietic cells is important for protection against DSS-induced colitis. WT/WT (n = 6) and WT/PGRN−/− (n = 6) mice were fed a 3% DSS solution in drinking water for 5 days, followed by normal drinking water for 2 days. (A) Body weight, (B) Histopathological changes in colon tissue. (C) Semiquantitative scoring of histopathology. Scale bars, 200 μm.
phenotype (Fig. 11), our findings demonstrate that PGRN deficiency exacerbates the severity of acute mucosal inflammation induced by chemical agents. PGRN−/− mice were significantly more susceptible to colitis-associated body weight loss, diarrhea, mortality, colon length and histological changes (Figs. 2, 3), indicating a critical role for the PGRN in protection against DSS- and TNBS-induced colitis. In addition, recombinant PGRN significantly reduced the severity of DSS-induced colonic injury assessed by clinical, biological and histological parameters (Fig. 7).

Bone marrow chimera mice results demonstrate that hematopoietic cell-derived PGRN was sufficient for protection against colitis (Fig. 4). PGRN also exerts its beneficial effect even in the absence of both T and B cells (Fig. 8). However, this finding does not belittle the role of T cells- and B cells-derived PGRN in chronic DSS colitis and other IBD models. Several recent articles have indicated that PGRN contributes to the pathogenesis of chronic autoimmune diseases19,39,40. In CD4+CD45Rbhigh T cells transfer colitis model, transfer of PGRN−/− CD4+CD45Rbhigh T cells leads to more severe intestinal inflammation (Fig. 5). Intriguingly, PGRN is highly expressed in CD11b+ cells and Ly6G+ cells (Fig. 1C), and adoptive transfer of CD11b+Ly6G+ myeloid-derived suppressor cells (MDSCs) has been reported to decrease intestinal inflammation41. Studies are warranted to elucidate the role of PGRN expressed in CD11b+ and Ly6G+ cells in the pathogenesis of colitis.

Figure 5 | Lack of PGRN signaling in CD4+ T cells executes experimental colitis. (A) Body weight following adoptive transfer of CD4+CD25−CD45Rbhi T cells from WT and PGRN−/− donors into Rag1−/− recipients. (B) Survival rate of Rag1−/− mice reconstituted with WT and PGRN−/− CD4+CD25−CD45Rbhi T cells. (C) Representative colonic sections from Rag1−/− mice reconstituted with WT and PGRN−/− CD4+CD25−CD45Rbhi T cells. All data represent means ± SE of a representative experiment. *p < 0.05; **p < 0.01. Scale bars, 200 μm.

Figure 6 | Intestinal barrier function and the microbiota are not altered in PGRN−/− mice. (A) Intestinal permeability was examined by detection of a FITC-dextran in the serum from DSS-induced mice. (B–E) Q-PCR analysis of intestinal microbiota in naive or DSS-treated wild-type and PGRN−/− mice. Values are shown as a relative ratio to total bacterial 16 s rDNA measured by 2−ΔΔCT method. All data represent means ± SE of three independent experiments. *p < 0.05; **p < 0.01.
Figure 7 | Recombinant PGRN attenuates the inflammation in DSS-induced colitic mice. (A) Body weight of PBS treated versus PGRN treated DSS induced colitis mice. Per group 8 mice were used. (B) Representative macroscopic pictures of colonic sections from mice treated with PBS or PGRN. (C) Levels of IL-10 were examined after colonic tissue explants were harvested at d8 and cultured ex vivo at 37°C for 24 h. (D) Representative HE-stained colon sections from mice treated with PBS and PGRN. All data represent means ± SE of three independent experiments. *p < 0.05; **p < 0.01. Scale bars, 200 μm.

Figure 8 | Recombinant PGRN attenuates the severity of DSS-induced colitis established in Rag1−/− mice. (A) Body weight. Rag1−/− mice were subjected to DSS-induced colitis and their body weight was measured. 5 mice per group were used. (B) Statistics analysis of colon length of each group. (C) Representative HE-stained colon sections of mice treated with PBS and PGRN. All data represent means ± SE of three independent experiments. *p < 0.05; **p < 0.01. Scale bars, 200 μm.
PGRN binds to TNFR2 with higher binding affinity and TNFR2 has been shown to be important for PGRN activities. TNFR2 has also been found to be required for PGRN-mediated protection of LPS-induced lung injury. In addition, PGRN-stimulated bone formation and fracture healing also depended on TNFR2 signaling. Our finding that PGRN lost its therapeutic effect, at least in part, in TNFR2-deficient mice (Fig. 10) corresponds with previous reports and supports the concept that TNFR2 plays an important role in various PGRN-mediated pathophysiological conditions. Interestingly, a lack of TNFR2 expression within CD4+ T cells was also reported to exacerbate the development of T cells transfer colitis in mice. It is noted that Chen and colleagues agreed that PGRN stimulation of regulatory T cells (Treg) was mediated by TNFR2. They reported that PGRN stimulates mouse Tregs through enhancing TNFα-induced Treg. The effect of TNFα on Tregs from mice and humans remains to be controversial. The data from Chen lab suggest that TNFα promotes murine Treg in vitro, whereas in humans, TNFα inhibits the suppressive function of Tregs through negative regulation of Foxp3 expression. Thus, it appears that PGRN stimulates the sup-
pressive function of both human and mouse Tregs, although the exact molecular mechanism remains to be further delineated. Although the effect of TNFα on Treg function remains controversial, the beneficial and therapeutic effects of Tregs in autoimmune diseases have been well-accepted\textsuperscript{48–50}. In addition, TNF inhibitors, including Enbrel, Remicade, Humira, have been accepted as the most effective anti-inflammatory therapeutics and the most successful biotech drugs\textsuperscript{51,52}. Importantly, recent studies reported that PGRN antibodies occurred frequently in patients with both CD and UC, and had significant neutralizing effects on PGRN plasma levels\textsuperscript{22}. These studies suggest that PGRN and its derivatives may be more effective alternatives to market TNF inhibitors for treating the IBD patients who are diagnosed to be PGRN antibodies positive.

The spontaneous colitis in IL-10R\textsuperscript{2}/\textsuperscript{2} mice and Blimp\textsuperscript{2}/\textsuperscript{2} mice, which exhibit a defect in IL-10 signaling\textsuperscript{33,34}, suggests the importance of IL-10 in preventing intestinal inflammation. IL-10 has been identified as a potential therapeutic molecule in Crohn’s disease and ulcerative colitis\textsuperscript{1,55,56}. In line with the report that PGRN-treated mice showed a significant increase in IL-10 production in collagen-induced arthritis model\textsuperscript{18}, we found that PGRN treatment also increased the levels of IL-10 in colitis models, and PGRN largely lost its protective effect in colitis when IL-10 signaling was blunted by IL-10R blocking antibody or in IL-10-deficient mice. This further confirms that upregulation of IL-10 production is one of the mechanism involved in the anti-inflammatory effects of PGRN in various inflammatory conditions. To better understand the roles of TNFR2 signaling and IL-10 in PGRN-mediated anti-inflammatory effect, we proposed a model illustrated in Fig 12. In brief, PGRN binds to TNFR2 in target cells, exemplified with Tregs, leading to the secretion of IL-10, an anti-inflammatory cytokine known to be involved in

Figure 11 | PGRN\textsuperscript{2}/\textsuperscript{2} mice have normal immune cells numbers and do not develop spontaneous colitis. (A) DC cells from the spleen of wild-type and PGRN\textsuperscript{2}/\textsuperscript{2} mice were gated with CD11c and MHCI\textsubscript{I}, and then further examined with CD11b and CD8 markers using flow cytometry. (B) Spleen and MLN cells from WT and KO mice were assessed for CD4\textsuperscript{+}Foxp3\textsuperscript{+} T cells. (C) Splenic plasmacytoid DC (mPDCA-1\textsuperscript{+}CD3\textsuperscript{−}CD19\textsuperscript{−}) were assessed by flow cytometry. (D) Histopathological changes in colon tissue from WT and KO mice were examined by H&E staining. (n = 3 mice/genotype, mean ± SD, *p < 0.05). Scale bars, 200 μm.
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Author contributions

F.H.W. designed and performed experiments, and wrote the paper. Y.Y.Z. performed experiments, acquired data, and drafted the manuscript. J.J.L. and J.J.M. performed immunohistochemistry assays. Q.Y.T. collected and analyzed data. J.Q.L. assisted in construction of bone marrow chimera model. J.J.L., W.T., W.M.Z. and X.P.Y. assisted in analyzing the data and editing the manuscript. C.J.L. supervised this study, analyzed data, and edited the manuscript.

Additional information

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