Research Article

Wogonin Strengthens the Therapeutic Effects of Mesenchymal Stem Cells in DSS-Induced Colitis via Promoting IL-10 Production

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Inflammatory bowel diseases (IBD) are prevalent and debilitating diseases; their clinical remedy is desperately unmet. Mesenchymal stem cells (MSCs) are pluripotent stem cells, possessing self-renew ability and multidifferentiation potential function [3]. MSCs showed therapeutic features including angiogenesis, tissue repair, and immunomodulation, among which IL-10 was an important effector molecule [4]. Several researches supported that IL-10 is highly relevant to IBD, and IL-10−/− mice would spontaneously develop colitis [5].

1. Introduction

Inflammatory bowel disease (IBD) is a chronic, relapsing-remitting, inflammatory gastrointestinal disease with a rising incidence worldwide in the past two decades [1]. However, current therapeutic options are far from satisfactory. Recently, cell therapies have been explored in IBD, including mesenchymal stem cells (MSCs) [2]. MSCs are pluripotent stem cells, possessing self-renew ability and multidifferentiation potential function [3]. MSCs showed therapeutic features including angiogenesis, tissue repair, and immunomodulation, among which IL-10 was an important effector molecule [4]. Several researches supported that IL-10 is highly relevant to IBD, and IL-10−/− mice would spontaneously develop colitis [5].

In humans, polymorphisms in IL-10 [6] have been found to be correlated with very early onset of colitis. Our previous studies on rodent models of IBD also showed that MSC administration could significantly improve intestinal inflammation via IL-10 [7, 8]. Consistently, some strategies have been proposed to improve the therapeutic efficacy of MSC, and engineering IL-10 overexpressing MSCs have achieved better therapeutic efficacy in immune relative diseases [9–11].

Accumulating researches indicated the extracts of herbal plants have diverse biological activities [12]. Scutellaria baicalensis is widely used as Chinese herbal medicine, and Wogonin is one of its major bioactive compounds [13]. Wogonin is a flavonoid-like compound which possesses anticancer and immunomodulatory effects [14]. Although
2. Material and Methods

2.1. Animals. Male C57BL/6 mice aged 8-12 weeks were purchased from the Nanjing Model Experimental Animal Center and housed in pathogen-free conditions at Sun Yat-sen University. The age- and weight-matched of mice were approved by the Zhongshan School of Experimental Animals Ethics Committee, Sun Yat-sen University, Guangzhou, China, and in strict compliance with the corresponding guidelines.

2.2. Reagents and Antibodies. Reactive Oxygen Species Detection Assay Kit was purchased from BioVision (San Francisco, CA, USA). Wogonin was purchased from MedChemExpress (Israel Shekel), and LPS was purchased from Sigma-Aldrich (St. Louis, MO, USA). The following antibodies were used for cell surface staining: CD34-PE, CD44-FITC, CD29-PE-Cy7, Sca-1-AF700, CD45-AF700, and CD49e-APC are all from BioLegend and Mouse IL-10 ELISA Set and Zombie Green™ Fixable Viability Kit were purchased from BD Biosciences. Mouse IL-10 MAb was purchased from R&D Systems.

2.3. Induction and Assessment of Colitis. Dextran sulfate sodium salt (DSS) was purchased from MP Biomedicals (United States) which was used to induce acute colitis in mice. 3% (w/v) DSS was dissolved in sterile water and supplied to the mice in drinking water for 7 days, followed by regular drinking water until day 8. Mice were randomly divided into five groups, and there were 6 mice in each group: the mice were fed regular water as the control group, were exposed to 3% DSS water as the IBD group, were administered 1 × 10⁶ MSCs via peritoneal injection as the MSC group, and were injected 10 mg/kg Wogonin via i.p. as the Wogonin group, and the combination of MSC and Wogonin group received both 1 × 10⁶ MSCs and 10 mg/kg Wogonin. Meanwhile, body weights of animals were scaled and recorded daily. Eight days later, the animals were sacrificed, and the length of the colon from the cecocolic junction to the anal verge was measured. Severity of the disease was assessed by weight loss, colon length, and histopathology scores of the colon. Colon tissues were processed for histological analysis by hematoxylin and eosin (H&E) stain. Histology was used to evaluate the inflammation (I) (0, none; 1, slight; 2, moderate; 3, severe), extent (E) (0, none; 1, mucosa; 2, mucosa and submucosa; 3, transmural), regeneration (R) (4, no tissue repair; 3, surface epithelium not intact; 2, regeneration with crypt’s depletion; 1, almost complete regeneration; 0, complete regeneration or normal tissue), crypt damage (C) (0, none; 1, basal 1/3 damaged; 2, basal 2/3 damaged; 3, only surface epithelium intact; 4, entire crypt and epithelium lost), and percentage involvement (P) (1, 1%–25%; 2, 26%–50%; 3, 51%–75%; 4, 76%–100%) according to previous reports [21]. The division animal groups and the histological scoring system are listed in Table 1.

There is evidence that both MSCs and Wogonin could alleviate IBD [15–20], it is still unclear whether combining MSCs with Wogonin be more superior to either therapy alone.

In this research, we demonstrated that Wogonin extended the therapeutic efficiency of MSCs on DSS-induced murine colitis in vivo. It was due to the increased IL-10 expression in the intestinal tissue and peritoneal cavity. Mechanistic studies suggested that Wogonin enhanced IL-10 production via AKT/GSK3 signal pathway. Furthermore, in vivo, the data exhibited that neutralizing anti-IL-10 antibody can abrogate the therapeutic effects of a combination of MSCs and Wogonin on colitis.

2.4. HE Staining. Colon tissue was removed and then immersed in 4% paraformaldehyde. After dewaxing with dehydrated alcohol gradient, the tissue was embedded in paraffin and stained with hematoxylin and eosin (H&E).

2.5. MSC Differentiation Assays. MSCs were isolated from the bone marrow of 6-8 weeks C57BL/6 mice, according to the improved low-density culture method [22]. The StemPro™ Osteogenesis Differentiation Kit (Thermo Fisher Scientific) and StemPro™ Adipogenesis Differentiation Kit (Thermo Fisher Scientific) were used for osteogenic and adipogenic differentiation in vitro, according to the manufacturer’s recommended procedures. All cells at 6-8 passage were used in this study.

<table>
<thead>
<tr>
<th>Animal groups</th>
<th>Tissue inflammation</th>
<th>Extent</th>
<th>Regeneration</th>
<th>Crypt damage</th>
<th>Percentage involvement</th>
<th>Histological score</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control/</td>
<td>None</td>
<td>None</td>
<td>Complete regeneration or normal tissue</td>
<td>None</td>
<td>—</td>
<td>0</td>
</tr>
<tr>
<td>IBD/</td>
<td>Slight</td>
<td>Mucosa</td>
<td>Almost complete regeneration</td>
<td>Basal 1/3 damaged</td>
<td>1%–25%</td>
<td>1</td>
</tr>
<tr>
<td>MSCs/</td>
<td>Moderate</td>
<td>Mucosa and submucosa</td>
<td>Regeneration with crypt’s depletion</td>
<td>Basal 2/3 damaged</td>
<td>26%–50%</td>
<td>2</td>
</tr>
<tr>
<td>Wogonin/</td>
<td>Severe</td>
<td>Transmural</td>
<td>Surface epithelium not intact</td>
<td>Only surface epithelium intact</td>
<td>51%–75%</td>
<td>3</td>
</tr>
<tr>
<td>Wogonin+MSCs</td>
<td>—</td>
<td>—</td>
<td>No tissue repair</td>
<td>Entire crypt and epithelium lost</td>
<td>76%–100%</td>
<td>4</td>
</tr>
</tbody>
</table>

Table 1: The division animal groups and the histological scoring system.
2.6. **ELISA.** Supernatants of cultured cells and mouse peritoneal lavage fluid samples were collected for detection of mouse IL-10 by enzyme-linked immunosorbent assay (ELISA Kit IL-10 from BD Biosciences, San Jose, CA, USA). The sensitivities of ELISA kits were 31.2 pg/ml for IL-10. ELISA assays were performed according to the manufacturer’s instructions.

2.7. **RNA-seq.** After 24 h Wogonin treatment, RNA was extracted from both untreated MSCs and treated MSCs using Trizol (Invitrogen, Carlsbad, CA). RNA-seq data were normalized through the DESeq2 pipeline, and gene heat map was fulfilled with packages pheatmap under R 4.0.2. The gene ontology analysis was performed based on clusterProfiler written by Yu et al. [23].
2.8. Western Blot. Cells or homogenized tissue were lysed in ice-cold RIPA buffer (50 mM Tris-HCl pH 8.0, 150 mM sodium chloride, 1% NP-40, 0.5% sodium deoxycholate, 0.1% sodium dodecyl sulfate, 2 mM EDTA containing 1x protease inhibitor cocktail, and phosphatase inhibitor (Roche)). Equivalent total protein extracts were separated by SDS-polyacrylamide gel electrophoresis (PAGE) and transferred to nitrocellulose membrane (GE Healthcare Life Sciences, Germany). After that, membranes were blocked with 5% bovine serum albumin (BSA) for 1 h at room temperature. Primary antibodies were incubated, and a horseradish peroxidase-conjugated secondary antibody was followed.

The primary antibodies used were as follows: antibodies for TNF-α antibodies (#11948T), HIF-1α antibodies (#4914S), p-GSK3β (#9323T), GSK3β (#9315S), p-AKT (#4060S), AKT (#4691S), p-Stat3 (#9145S), Stat3 (#9404S), p-JNK (#4671), JNK (#9258), and GAPDH (#2118) were obtained from Cell Signaling Technology; all those antibodies were diluted with antibody solution and used at a dilution of 1 : 2000. IL-10 (sc-365858) was obtained from Santa Cruz and employed in a 1 : 500 dilution. The secondary antibodies were used at dilution 1 : 5000. All proteins were visualized by Chemiluminescent HRP Substrate (Millipore, WBKLS0500), and chemical luminescence of membranes was detected by.

![Western Blot Image](image-url)

**Figure 2:** MSCs and Wogonin increased IL-10 levels in both colon tissue and peritoneal lavage fluid. Mice were administered 3% DSS in drinking water for 7 days, followed by regular drinking water. On the 2nd day, mice received MSCs, Wogonin, or a combined treatment, respectively. The colon tissue and peritoneal lavage fluid were collected on 8 d. (a) The expression of TNF-α and IL-10 in the colon was detected by western blot, and the results represented three independent experiments. (b) TNF-α and (c) IL-10 expressions were calculated separately. (d) The IL-10 levels of peritoneal lavage fluid were measured by ELISA. Data represent mean values ± SEM. The statistical significance of difference between two means was calculated with an unpaired. n = 6/group. A one-way ANOVA was performed, *P < 0.05; **P < 0.01; ***P < 0.001; ****P < 0.0001; ns: no significance.
Figure 3: Continued.
and dead/live streaming antibodies for 30 min at 4°C and 0.05% sodium azide and labeled with surface markers. Cells were washed with PBS buffer containing 0.1% BSA.

2.9. Flow Cytometry. For cell surface marker staining, the cells were washed with PBS buffer containing 0.1% BSA and 0.05% sodium azide. All the above stained cells were assayed by FACS Aria II (Becton Dickinson, San Jose, USA), and the data were analyzed by FlowJo software (TreeStar, San Carlos, USA).

For ROS detection, CS&T beads (BD, 661414) were used for machine calibration and performance tracking. BD rainbow beads were for PMT voltage optimization (BD, 559123). MSCs were treated with Wogonin (0, 12.5, 25, and 50 μM) in the presence of LPS. After 24 h, cells were incubated with 20 μM DCF-DA at 37°C for 20 min, then detected by flow cytometry. Fluorescence minus one (FMO) was applied for positive cell gating. Anti-rat Igk negative control compensation particle set (BD, 552845) was used for fluorochrome compensation.

2.10. IL-10 Blockade Experiment In Vivo. Eight to twelve weeks C57BL/6 male mice were exposed to 3% DSS water for 7 days, then fed the regular water. On the 2nd day, all mice were peritoneal injected 1 × 10⁶ MSCs and 10 mg/kg Wogonin. Three of them additionally received neutralizing anti-mouse IL-10 antibody (R&D Systems) at a dose of 100 μg/mouse via peritoneal injection. On day 8, the colonic length and pathological lesions of the colon were detected.

2.11. Statistical Analysis. All statistical tests were performed with GraphPad Prism 7 (GraphPad Software Inc., San Diego, USA). Significant differences between data sets were performed with either the unpaired Student’s t test when comparing two groups, one-way ANOVA for more than two groups, or two-way ANOVA for two variables (GraphPad Software Inc., San Diego, CA, USA). Data were represented as mean or mean ± SEM. **P < 0.001; ***P < 0.001; **P < 0.01; *P < 0.05; and P > 0.05, not significant, as stated in figure legends.

3. Results

3.1. Wogonin Enhanced the Therapeutic Effects of MSCs on DSS-Induced Colitis via Increasing IL-10. To explore the influence of Wogonin on the therapeutic effects of MSCs, we compared the effect of MSCs or Wogonin alone, and the effect when combined. From day 5, weight loss was observed in all mice except the control mice. As shown in Figure 1(a), the IBD mice lost most weight, but the combination of MSC and Wogonin group had the least weight loss. In this study, colon length was also recorded as an index of mouse colitis. On the 8th day, all colon tissues were collected and measured. The results indicated that the length of colon tissues from all mice exposed to DSS was shorter than those from the control groups. Interestingly, the colons from the combination group were significantly longer than those from the Wogonin or MSC groups (Figures 1(b) and 1(c)). This implied that a combination of Wogonin and MSCs possessed better therapeutic effects on murine colitis than using them separately. Meanwhile, colon histopathological examination consistently showed less intestinal inflammation in the combination group comparing to the other treatment groups (Figure 1(d)). The pathological score of the combination of MSCs and Wogonin-treated mice was the lowest in all treated mice (Figure 1(e)).

To investigate the possible therapeutic mechanisms, the expression of TNF-α and IL-10 in colon was determined by western blot. The data suggested that DSS administration significantly enhanced the expression of proinflammatory cytokine TNF-α and reduced the expression of anti-inflammatory cytokine IL-10 in the colon tissue. No matter
whether MSCs and Wogonin are used together or alone, they could significantly reduce local TNF-α and increased local IL-10 (Figures 2(a)–2(c)). To further inspect the change of cytokines, the levels of TNF-α and IL-10 in peritoneal lavage fluids were detected by ELISA. However, the level of TNF-α was too low to be detected. In line with western blot data, IL-10 significantly increased in peritoneal lavage fluids from the combination group comparing to the MSC or Wogonin groups. This result suggests that IL-10 may be the key factor for Wogonin strengthening the therapeutic effect of MSC (Figure 2(d)).

3.2. Wogonin Promoted IL-10 Production of MSCs. To understand how Wogonin strengthened the therapeutic effects of
MSCs, we compared the difference of gene expression profiles between untreated MSCs and Wogonin-treated MSCs in the presence of LPS. After 24 h stimulation, untreated MSCs and Wogonin-treated MSCs were collected, and the transcriptome analysis was performed. The transcriptome analysis identified the upregulation of 1000 genes and downregulation of 2000 genes in Wogonin-treated MSCs (Figure 3(a)). GO analysis indicated that the upregulated genes were highly enriched in the cell cycle (Figure 3(b)). Furthermore, RNA-seq analysis also revealed that Wogonin enhances the expression of various chemokines and IL-10 in MSCs (Figure 3(c)).

To further verify the influence of Wogonin on the IL-10 production of MSCs, we activated MSCs with different concentrations of Wogonin in the presence of LPS in vitro. After 24 h stimulation, the supernatant was collected, and the IL-10 level was detected by ELISA. As shown in Figure 4(a), the IL-10 secretion from MSCs gradually increased in a dose-dependent manner, reached the peak at 25 μM, then decreased when continuously increasing Wogonin dosage. It was illustrated that 25 μM was the optimal dose of Wogonin to increase the IL-10 production in MSCs.

To clarify whether Wogonin affects the characteristics of MSCs, we firstly analyzed surface markers of untreated MSCs and 25 μM Wogonin-treated MSCs. Comparing to untreated MSCs, MSCs expressed the same panel of surface markers after Wogonin treatment, including CD29, CD44, CD49e, and Sca-1, and did not express the hematopoietic stem cell markers CD34 or CD45 (Figure 4(b)). Additionally, osteogenic differentiation and adipogenic differentiation were tested. As confirmed by Alizarin red S staining and Oil O staining, respectively, Wogonin-treated MSCs showed no change in mesodermal differentiation capacity compared with untreated MSCs (Figures 4(c) and 4(d)).

3.3. Wogonin Increased ROS Level in MSCs. To investigate the effects of Wogonin on ROS level in MSCs, different doses of Wogonin were used to stimulate MSCs in vitro, and the ROS levels of MSCs were detected by DCFDA staining and analyzed by flow cytometry. The results showed that Wogonin treatment obviously increased the ROS level of MSCs in a dose-dependent manner, being greatest at 25 μM (Figures 5(a) and 5(b)). When Wogonin dose was increased to 50 μM, ROS level decreased instead, which was consistent with the IL-10 expression of MSCs (Figures 3(a) and 3(b)). These data implied that Wogonin treatment regulated the ROS level of MSCs.

3.4. Effect of Wogonin on the GSK3β/AKT Signaling Pathway of MSCs. To elucidate the potential signaling pathways of Wogonin increasing IL-10 production in MSCs after Wogonin treatment, we tested protein expression level IL-10 of MSCs that were cultured with LPS in the absence or presence of Wogonin at different dose firstly. A role for transcript factor HIF-1α involved in IL-10 expression was well established; therefore, the expression of HIF-1α was also detected. Consistent with IL-10 in MSC supernatant, we found that HIF-1α and IL-10 gently upregulated in the presence of Wogonin treatment, significantly on 25 μM dose, but decreased on 50 μM dose (Figures 6(a) and 6(b)). Next, signaling pathways involved in IL-10 production were assessed, and western blot analysis showed a remarkable increase of phosphorylation of glycogen synthase kinase 3β (GSK3β) and phosphorylation of AKT in the presence of Wogonin with 25 μM, while total GSK3β and AKT changed slightly (Figures 6(a), 6(c), and 6(d)). Phosphorylation of Stat3 signaling kept intact on Wogonin treatment (Figures 6(a) and 6(e)). These results suggested that Wogonin upregulates the IL-10 production involved in GSK3β/AKT signaling pathways.
The image shows a continuation of Figure 6, which compares the relative protein expression of various proteins under different conditions. The figure highlights the effects of LPS (lipopolysaccharide) and a compound on the expression levels of HIF-1α, IL-10, p-GSK3β, GSK3β, p-AKT, AKT, p-Stat3, and Stat3. The graphs illustrate changes in protein expression levels normalized to GAPDH, with different concentrations of the compound and LPS treatment groups. The data suggests modulation of these proteins under various experimental conditions.
Figure 6: Wogonin improves the IL-10 production in MSCs via GSK3β/AKT pathway. MSCs were stimulated by Wogonin at a different dose (12.5, 25, and 50 μM) in the presence of LPS. After 24 h stimulation, western blot was performed. The cell extract probed with antibodies as indicated, and GAPDH as loading control. Relative protein expression was calculated, respectively. (a) The representatives of the protein expression of HIF-1α, IL-10 as well as GSK3β, AKT, and Stat3 and their phosphorylated forms (p-GSK3β, p-AKT, and p-Stat3). The effects of Wogonin on the expression of HIF-1α and IL-10. (b) The effects of Wogonin on GSK3β. (c) AKT (d) and Stat3 (e) signaling pathways. All the western blotting experiments were repeated three times. Values are means ± SEM. The statistical significance of difference between two means was calculated with two-way ANOVA followed by multiple comparisons with t test and Bonferroni correction, *P < 0.05.
3.5. IL-10 Blockade Abrogated the Therapeutic Effects of a Combination of MSCs and Wogonin on Colitis.

To confirm the crucial role of IL-10 on ameliorating mouse colitis, neutralizing IL-10 antibody was used for the combination of MSCs and Wogonin mice. As shown in Figure 7, IL-10 blockade abolished the improvement of the combination on murine intestinal inflammation. After IL-10 neutralization, the shorter colon length was observed, compared with the no neutralization mice (Figure 7(a)). HE staining data also suggested that more serious intestinal epithelial structure damage and more inflammatory cell infiltration were observed in the colon tissues from the IL-10 neutralization mice than those from no IL-10 neutralization mice (Figure 7(b)). These results further demonstrated that the increased production of IL-10 in MSCs is a key factor for Wogonin to improve the therapeutic efficacy of MSC on DSS-induced colitis.

4. Discussion

Ulcerative colitis (UC), along with Crohn’s disease (CD), is chronic and debilitating inflammatory bowel disease (IBD). IBD has been reported to have high prevalence with increasing trends in the rate of incidence [24]. The hygiene hypothesis is the most popular explanation for this growing incidence; meanwhile, the particular etiological factors were still enigma [25]. The impairment of colon mucus layer barrier and epithelium and change of luminal microbial diversity (dysbiosis) were commonly observed in the pathogenesis of UC. TNF-α and multitude of Th2-derived cytokines were
reported to be involved in this process [26]. The lack of a cure or effective long-term treatment options has resulted in substantial morbidity of IBD [27], which highlighted the urgent demand of investigation of a new therapeutic regime.

Mesenchymal stem cells (MSCs) are multipotent stromal cells which can differentiate into different cell types according to their residential niches. The cells sprouted into the clinic stage center due to their features of versatile development potential, self-renewal, allogenic compatibility, and immunomodulatory effects. MSCs are currently being widely tested as a cellular therapy in chronic intestinal diseases, including IBD, and showed promising clinical outcomes with unmet optimization and improvement [28–30], such as in vivo low level of MSC recruitment, and persistence decreased their therapeutic efficacy [31].

To overcome these hurdles, IFN-γ, TGF-β, IGF-1C, IL-1β, and PGE2 were applied to precondition MSCs before they were administered onto different experimental models, and pretreated MSCs showed an enhanced therapeutic effect, with different limitations [32–36]. All of these promising explorations encouraged us to search further alternatives.

Wogonin is a natural flavonoid component extracted from the dried root of Scutellaria baicalensis. Various studies showed its direct anti-inflammatory effects and immunoregulation on immune cells [37]. Interestingly, Wogonin was proved as a particular therapeutic adjuvant for anti-inflammation [14], which inspired our explorations of the potential role of Wogonin in MSC medication on IBD.

On our solid and highly repeatable DSS-induced mouse colitis model, MSCs and Wogonin showed an additive or synergistic effect on the mitigation of colitis. As expected, TNF-α amount in inflammatory colon tissue was decreased by MSCs, Wogonin alone, and their combinations. This was consistent with the alleviated intestinal inflammation by pathology score, which echoed by previous reports.

As another key player of inflammation, IL-10 is an anti-inflammatory factor, playing a critical role in the suppression of autoimmune reactivity and in termination of the inflammatory response. IL-10 from T cell have been established an anti-inflammatory effect in IBD [38], and IL-10 secreted from MSCs have been proved to attenuate inflammation in mice [9–11, 39, 40]. Our recent report showed Wogonin regulated IL-10 production [41]. Hence, it is interesting to study the effect of Wogonin on IL-10 production from MSC in our mouse model.

Strikingly, IL-10 amount in peritoneal lavage fluid was increased by Wogonin and MSCs additively, which did not show in intestinal tissues. Comparably, TNF-α amount in peritoneum was under detectable range. This indicated that Wogonin might directly alter peritoneal MSCs on-site.

To verify this hypothesis and examine the full profile of the effect of Wogonin on MSCs from a different level, LPS-activated MSCs cocultured with/without Wogonin in vitro.

Firstly, Wogonin enhanced abundant proliferation-associated gene expression in MSCs, but not those development-associated genes. This was consistent with no surface, and differentiation markers were changed by Wogonin culture. Meanwhile, both TNF-α and IL-10 mRNA amounts were augmented along with the activation of numerous chemokine genes. This echoed with the increased protein amount of peritoneal IL-10.

Secondly, there is a clear dose-effect of Wogonin enhancing IL-10 production from MSCs. 25 μM Wogonin showed optimal augment effect, and higher concentration suppressed further IL-10 secretion. Reactive oxygen species (ROS) generation by NADPH oxidase has been shown to interact with IL-10 production in macrophages [42, 43]. Hence, it is not surprising to repeat this on MSCs. All these results indicated the adjuvant feature of Wogonin on MSC activation.

To further investigate the molecular machinery of Wogonin on MSCs, TLR4 signaling pathways were explored. Strikingly, the TLR4/AKT/GSK3β pathway was triggered by Wogonin. It is still difficult to pinpoint the target receptor on MSCs due to the herb derivative, and their discovery of this machinery on MSCs suggested the new strategy in IBD therapy [44]. Our in vivo IL-10 neutralization data reinforced this concept.

To our knowledge, our study firstly showed Wogonin enhanced IL-10 production from MSCs directly, which restrained DSS-induced colitis. This implied new therapeutic strategy for IBD.

5. Conclusions

Although MSCs show promising therapeutic potential in treating inflammatory bowel disease (IBD), their efficacy is influenced by multiple factors. Wogonin, a bioactive compound from *Scutellaria baicalensis*, has diverse immunoregulatory effects. Our present study demonstrated that Wogonin extended the therapeutic efficiency of MSCs on DSS-induced murine colitis in vivo, which was due to the increased IL-10 expression in the intestinal tissue and peritoneal cavity. RNA sequencing analysis suggested Wogonin enhanced IL-10 expression, which was confirmed via in vitro experiments. Furthermore, western blot data implied that Wogonin upregulated transcript factor HIF-1α in MSCs via AKT/GSK3β signal pathway. Summarily, our study found that combining MSCs with Wogonin was superior to MSC therapy alone, which provides a novel strategy for MSC clinical application on IBD.

Data Availability

The data used to support the findings of this study are included within the article.

Conflicts of Interest

The authors declared no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

Authors’ Contributions

Qiongli Wu, Shujuan Xie, and Yanwen Peng designed experiments, performed research, and interpreted data. Yinhong Zhu established a mouse IBD model. Yinhong Zhu, Jiatong Tian, and Jingrou Chen performed cell culture
and differentiation assays. Qiongli Wu contributed to flow cytometry analysis. Shujuan Xie performed western blot. Changyou Wu and Shiqiu Xiong substantially revised the manuscript. Yanwen Peng and Yujin Ye supervised research and interpreted data. Yanwen Peng, Qiongli Wu, Shujuan Xie, and Yujin Ye wrote the manuscript. Qiongli Wu and Shujuan Xie contributed equally to this work. All authors have read and approved the final manuscript.

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