MicroRNA-21 Promotes Allergic Airway Inflammation and AHR and Inhibits Mesenchymal Stem Cell Migration in Cockroach Allergen Induced Asthma Model

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Research

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Abstract

Background

Mesenchymal stem cells (MSCs) have been used to treat asthma in a mouse model. However, the efficacy and mechanism of MSCs are not elucidated. MicroRNAs (miRNAs) play a key role in asthma and related to the aim of this study was to illustrate the role of miR21 and its influence on MSC migration in asthma model.

Methods

A mouse model of asthma was established using cockroach extract (CRE), and miR-21 expression was examined. A miR-21 lentivirus construct was used to investigate the role of miR-21 in vivo and in vitro in mouse bone marrow-derived (BM) MSCs. A TOPFlash reporter gene assay was used to study the signaling downstream of miR-21. IL-4, IL-5, IL-13, IgE, and IgG1 levels in bronchoalveolar lavage fluids were determined by enzyme-linked immunosorbent assays.

Results

miR-21 was upregulated in CRE-induced asthmatic mice. MiR-21 promoted allergic airway inflammation and airway hyperreactivity by inhibiting BM-MSC migration. β-Catenin was found to act downstream of miR-21 as a negative regulator of miR-21. Rescue experiments verified that miR-21 inhibited BM-MSC migration by suppressing Wnt/β-catenin signaling.

Conclusion

MiR-21 promotes allergic airway inflammation and AHR and inhibits BM-MSC migration through Wnt/β-catenin signaling, which may serve as an effective therapeutic target for asthma.

1. Introduction

Asthma is a chronic inflammatory disorder characterized by airway obstruction, heterogeneous inflammation, mucus secretion, airway hyperresponsiveness (AHR), and high expression of immunoglobulin E (IgE) [1–4]. Various behavioral activities, environmental factors, stimulating allergies, and excessive T-helper type 2 (TH2) cell responses are considered major causes of asthma [5, 6]. Currently, the treatment of asthma mainly depends on allergen immunotherapy targeting allergic immunocytes including TH2 cells, B cells, T cells, B regulatory cells, T regulatory cells, mast cells, macrophages, and dendritic cells [7–10]. Due to diverse pathogens and evolutionary antigens, the body produces stronger resistance, and some steroids are being developed for the treatment of asthma [11]. Although current drugs and allergen immunotherapies are effective to a certain extent, they have numerous side effects, including decreased bone mineral density, skin thinning and bruising, immunologic tolerance, rejection, and anaphylaxis [12, 13]. And the morbidity of asthma is still increasing [14, 15]. Therefore, novel therapeutic strategies targeting pathophysiological events need to be further explored.

Mesenchymal stem cells (MSCs) have attracted considerable attention for the treatment of various diseases, both in basic medicine and pre-clinical research, owing to their self-renewing and differentiating properties [16–23]. Murine bone marrow-derived (BM-) MSCs inhibited allergic responses in a mouse model of asthma [24], and human BM-MSCs suppressed chronic airway inflammation in a murine asthma model [25]. Increasing studies have shown that MSCs ameliorate asthma in both mouse models, and the mechanisms underlying the therapeutic effects of MSCs have been explored [26–30]. However, little is known about the mechanism underlying MSC migration to target sites to exert curative effects. Migration is one of the vital characteristics of MSCs and is crucial for MSC therapy targeting inflammatory sites. It is fundamental to the application of MSC-based asthma treatments in future clinical research.

MicroRNAs (miRs) are small, non-coding RNAs that play roles in transcriptional and post-transcriptional gene expression regulation by targeting the 3′-untranslated regions of specific mRNAs [31–33]. Recent evidence suggests that miRs play a critical role in the diagnosis and treatment of asthma [34–36]. Accumulating evidence indicates that MSCs ameliorate asthma via miRs that mediate relevant signaling pathways [37–40]. Additionally, some studies have shown interactions between MSCs and miRs in asthma [37, 41, 42]. MiR-21, which is related to cell apoptosis, invasion, and migration, is one of the most widely researched miRs [43, 44]. It has been reported to suppress tumor cell migration into infected areas in several cancers [44–46]. Based on a literature review, we found that miR-21 is mostly overexpressed in asthma [1–3], which we validated in preliminary experiments. Therefore, we aimed to investigate the role and mechanism of miR-21 in asthma treatment with MSCs.

In this study, the miR-21 expression and its influence on allergic airway inflammation and AHR and MSC migration were assessed in cockroach extract (CRE)-induced mouse model of asthma. MiR-21 overexpression and silencing were used to evaluate the signaling pathway mediating MSC migration in vivo and in vitro.

2. Materials And Methods

2.1 Asthma mouse model establishment

Six- to eight-week-old C57BL/6 mice were purchased from SLAC Laboratories (Shanghai, China). The mice were housed under standard conditions with a 12-h/12-h light/dark cycle at the Animal Center of Hunan Cancer Hospital (Changsha, China) and had free access to food and water. All animal experiments and procedures were performed according to the guidelines of the Center for Medical Ethics, Central South University.

A German CRE (B46, Blattella germanica)-induced asthma mouse model was conducted. In brief, mice were sensitized by intratracheal inhalation of 20 μg CRE in 50 μL of PBS under anesthesia using isoflurane (Sigma, USA) on days (D) 0–4. The mice were challenged by intratracheal instillation of the same
amount of CRE on D10–D13. Control mice received the same volume of PBS during the sensitization and challenge phases. Mice were randomly assigned to the two treatment groups (n = 6 each) and were analyzed in a double-blind manner. On D14, the mice were sacrificed. The lung tissues were dissected for histological analyses, and bronchoalveolar lavage fluids (BALFs) were harvested to count total cells and inflammatory cells. A timeline of the mouse experiment is shown in Figure 1A.

2.2 Analysis of lung inflammation

Lung inflammation was assessed. Briefly, BALFs were centrifuged at 300 × g at 4°C for 10 min and washed with ice-cold PBS. Total cells and inflammatory cells including eosinophils (Eos), lymphocytes (Lym), macrophages (Mac), and neutrophils (Neu), were counted to evaluate the percentages of inflammatory cells in BALF by flow cytometry using a FACs Calibur cytometer (BD Biosystems).

IL-4, IL-5, and IL-13 in mouse BALFs were examined by ELISAs, using commercial kits (eBioscience, USA) according to the manufacturer’s recommendations. Serum CRE-specific IgE and IgG1 were examined by ELISAs.

2.3 Cell culture and lentivirus infection

MSCs were cultured in α-minimum Eagle’s medium (Hyclone, USA) supplemented with 10% fetal calf serum (Hyclone), 100 mg/mL streptomycin (Gibco, USA), 100 IU/mL penicillin (Gibco) in a humidified incubator with 5% CO₂ at 37°C (Thermo Fisher, USA). Cells were passaged in tissue-culture flasks (Coming, USA) in medium containing 0.25% trypsin-EDTA (Gibco) until approximately 80% confluence. Detailed cell culture procedures were reported in our previous study [47]. MSCs at passages below 8 were used in all experiments.

MiR-21, anti-miR-21, miR negative control (NC), and anti-miR-NC oligonucleotides (50 nM, RiboBio, Guangzhou, China) were transfected into MSCs using the riboFECT™ CP Transfection Kit (Ribo) according to the manufacturer’s protocol. The sequence of Pri-miR-21 was as follows: 5’-TCACAAGACATAAAGACGACACACATTTTGTACGCAAACACCAGATGTGGATGAATGTGAAAAACATCTTTGTATGG ATGTCACTAGATAAGAAAATGTCAGACAGCCACATCGACTGCTGTTGCCATGAGATTTCACAGTCAACATCGCTGTAAGTCCACCAAGGTGTGTCAGAGCAGATGTCATTGCTGTGATCCATGCTGTTGGGCTT-3’. The miR-21 sequence was 5’-GTGCATTGCTGTTGCATTGC-3’. BM-MSCs were seeded in a six-well plate (Corning) at 1 × 10⁵ cells/well and infected with lentiviral anti-miR-21 and miR-21 mimic, and miR-NC and anti-miR-NC as controls, in the presence of 10 mg/mL polybrene (Millipore, Germany).

Small interfering (si) RNA targeting the β-catenin gene (Ctnnb1) was synthesized at GenePharma (Shanghai, China). An unrelated sequence was used as a negative control, according to our previous report [48]. Briefly, siRNAs were transfected into BM-MSCs during the logarithmic growth phase using Lipofectamine 2000 (Invitrogen, USA) according to the manufacturer’s instructions. During the challenge period (D10–13), mice were intranasally administrated 2 × 10⁶ infectious units (IFUs) of miR-21 lentivirus, and an equal amount of empty lentiviral vector was used as a control.

2.4 BM-MSC migration assays in vivo and in vitro

To verify whether exogenous BM-MSCs can migrate to the lungs in asthmatic mice, 5 × 10⁵ GFP⁺ BM-MSCs in 0.2 mL of PBS were injected via the tail vein, and the same amount of PBS was injected as a control. To assay MSC migration in vitro, BM-MSCs (1 × 10⁵) were transfected with a plasmid expressing miR-21, anti-miR-21, and β-catenin using Lipofectamine 2000 (Invitrogen, USA) in serum-free minimum Eagle’s medium at 37°C in a 5% CO₂ atmosphere for 48 h. Then, the cells were added to the upper chambers of Transwell inserts with 8.0-µm pores in serum-free medium, while normal growth medium was placed in the lower chambers. After incubation for 24 h, the MSCs were fixed and stained with 20% methanol violet (Beyotime, China) and 0.1% crystal violet (Beyotime, China), and counted under a microscope.

2.5 Quantitative reverse-transcription (q-)RT PCR

To evaluate the expression of miR-21 and Ctnnb1 in BM-MSCs, total RNA was isolated from BM-MSCs using TRizol reagent (Invitrogen, USA). The RNA was treated with RNase-free DNase I (Roche, Basel, Switzerland). cDNA was synthesized using the All-in-One™ First Strand cDNA Synthesis Kit (AMRT-0050, Genecopoeia, USA) and miRNA First Strand cDNA Synthesis Kit (AMRT-0050, Genecopoeia). Real-time qPCR was run using the All-in-One™ qPCR mix (Genecopoeia) and All-in-One™ miRNA qRT-PCR Detection Kit (Genecopoeia) on an ABI 7300HT real-time PCR system (Applied Biosystems, USA). Primers for miR-21, Ctnnb1, Gapdh, and U6 were synthesized at Genecopoeia. Relative expression levels of miR-21 and Ctnnb1 were evaluated using the 2⁻∆∆CT method [49, 50]. U6 was used as an endogenous control for miR-21, and Gapdh was used as an internal control for Ctnnb1.

2.6 Western blotting

Lung tissues and BM-MSCs from the different treatment groups were collected and homogenized in RIPA lysis buffer (Beyotime, China) supplemented with 1 mM phenylmethylsulfonyl fluoride (Beyotime). Protein concentrations were determined using a BCA Protein Assay Kit (Beyotime), and 25 µg protein was loaded per lane. Western blotting was conducted as described previously [48]. The primary antibodies used were anti-β-catenin (D10A8, CST, 1: 1000) and anti-GAPDH (14C10, CST, 1: 1000), and goat anti-rabbit IgG (Invitrogen) was used as the secondary antibody. Immunoreactive bands were visualized with enhanced chemiluminescence reagent (Bio-Rad, USA), using a Tanon-4500 digital image system (Tanon Science & Technology, China).

2.7 Immunostaining and immunofluorescence

For the histological assessment of lung inflammation, mouse lungs were collected and soaked in 10 mL of ice-cold PBS. The samples were fixed in 4% formaldehyde (Sangon Biotech, Shanghai, China), and 5-µm sections were cut and stained with hematoxylin and eosin (HE) or with periodic acid Schiff
reagent (Sigma). The sections were examined and photographed under a microscope (Nikon, Japan). Detailed immunostaining procedures for lung histology were described in our previous report [48].

For immunofluorescence, lung sections were blocked with protein-blocking serum-free solution (Dako, Denmark), permeabilized with 0.1% Triton X-100 for 10 min, and blocked in 1% BSA at room temperature for 30 min. The sections were incubated with primary antibody against β-catenin (1:100 dilution) at 4 °C overnight. The sections were washed and incubated with specific secondary antibody in PBS (1:100 dilution) for 60 min. Nuclei were stained with DAPI (Beyotime) at room temperature for 10 min. Immunofluorescence was evaluated using a confocal microscope (Leica Microsystems, Germany). The intensity of co-staining was determined using image acquisition and analysis software (Image J), and the values are presented as mean fluorescence intensity per square micro.

2.8 TOPFlash reporter gene assay

To evaluate the signaling downstream of miR-21, we used a pair of TOPFlash/FOPFlash luciferase reporter constructs (Upstate Biotechnology, USA). BM-MSCs were seeded in a 24-well plate (Corning) at 1 × 10^5 cells/well and allowed to settle for 24 h. TOPFlash or FOPFlash plasmid (500 ng), pRL-TK Renilla luciferase plasmid (150 ng, Promega, USA), and miR-21 mimic (50 nM) or miR-NC (50 nM) were cotransfected into the BM-MSCs. After 24 h of incubation, the cells were stimulated with 50 ng/ml recombinant murine Wnt3a (PeproTech, USA) for another 24 h. Then, the cells were harvested and luciferase reporter activity was measured in the reporter lysis buffer of the Luciferase Assay System (Promega). TOPFlash and FOPFlash signals were normalized to Renilla luciferase activity, and data are represented as normalized TOPFlash/FOPFlash activity values.

2.9 Statistical analysis

Data are reported as the mean ± SD derived from at least three independent experiments. Means were compared using a two-tailed paired Student’s t-test or using the Chi-square test for categorical variables. *P < 0.05 was considered statistically significant.

3. Results

3.1 MiR-21 expression is increased in CRE-induced asthma model mice

MiR-21 expression was significantly higher in CRE-challenged mice than in PBS-treated mice (Figure 1B). Further, miR-21 was significantly upregulated in miR-21-transfected CRE model mice as compared to miR-NC-transfected mice (Figure 1C).

3.2 MiR-21 promotes lung inflammation in asthma

To understand the role of miR-21 in asthma, we conducted histological analysis. HE staining demonstrated that miR-21 accelerated inflammatory cell infiltration, goblet cell hyperplasia, and mucus overproduction (Figure 2A). Although the numbers of Lym, Mac, and Neu were not significantly different, there were significant increases in total cells and Eos in the miR-21+CRE group when compared with the miR-NC+CRE group (Figure 2B and 2C). The ability of AHR to resist methacholine at 10 mg/mL and 30 mg/mL was increased in CRE-challenged miR-21 mice when compared to CRE-challenged miR-NC mice (Figure 2D). Additionally, allergen-specific IgE (Figure 2E) and IgG1 (Figure 2F) levels were significantly increased in CRE-challenged miR-21 mice. Th2-secreted cytokines, including IL-4 (Figure 2G), IL-5 (Figure 2H), and IL-13 (Figure 2I), in BALFs were increased in CRE-challenged miR-21 mice. These results indicated that the aggravated Th2-dependent inflammatory reaction in asthma is related to miR-21 expression.

3.3 BM-MSCs improve allergic airway inflammation and AHR, whereas miR-21 aggravates inflammatory reactions by suppressing BM-MSC migration

A schematic diagram of MSC administration in the CRE-induced mouse model was shown in Figure 3A. In miR-21-transfected asthmatic mice, GFP⁺ BM-MSCs migrated into the lungs, but the number of BM-MSCs in the lungs was significantly lower than that in miR-NC-transfected asthmatic mice, indicating that miR-21 inhibits BM-MSC migration into the lungs (Figure 3B). Mean fluorescence intensities are shown in Figure S1. Next, we aimed to confirm that miR-21 promotes allergic airway inflammation and AHR by inhibiting BM-MSC migration. MiR-21-transfected asthmatic mice engrafted with GFP⁺ MSCs exhibited decreased airway inflammation cell infiltration, and goblet cell secretion when compared with miR-21-transfected asthmatic mice administered PBS, suggesting that BM-MSCs inhibit airway inflammation and infiltration (Figure 3C). These results were consistent with those of previous studies showing that MSCs suppress chronic/allergic airway inflammation in a murine asthma model [25, 27]. Interestingly, inflammatory cell infiltration in the lungs was significantly decreased in the miR-21 + CRE + BM-MSCs group when compared with the miR-21 + CRE + PBS group (Figure 3C). Although the numbers of Lym, Mac, and Neu showed no obvious differences, there were significant decreases in total cells and Eos cells after BM-MSC transplantation in asthmatic mice, whereas the numbers of total cells and Eos in the BALF were significantly increased in the miR-21-transfected group post BM-MSC injection (Figure 3D and 3E). Furthermore, IgE (Figure 3F) and IgG1 (Figure 3G) levels were significantly decreased after BM-MSC transplantation in asthmatic mice, whereas these levels were enhanced in miR-21-transfected mice. IL-4 (Figure 3H), IL-5 (Figure 3I), and IL-13 (Figure 3J) levels in the BALF were significantly decreased after BM-MSC transplantation in asthmatic mice, but were significantly increased in mice transplanted with miR-21. These results were consistent with the histological examination findings. Thus, these results corroborated that BM-MSC transplantation significantly improved the inflammatory reaction, whereas miR-21 aggravated the inflammatory reaction by inhibiting BM-MSC migration in asthma.

3.4 MiR-21 suppresses BM-MSC migration in vitro

MiR-21 expression in the different treatment groups was confirmed by RT-qPCR (Figure 4A). First, we conducted wound-healing assays to evaluate the effect of miR-21 on the migration ability of BM-MSCs in vitro. Compared with the miR-NC group, BM-MSC migration was suppressed in the miR-21 group at 24 h and 48 h (Figure 4B), and when anti-miR-21 was added, BM-MSC migration was obviously enhanced. These results indicated that miR-21 inhibits BM-MSC
migration in vitro. To assess the role of miR-21 in directed migration of BM-MSCs, we used a Boyden chamber to evaluate BM-MSC migration. As shown in Figure 4C, miR-21 suppressed BM-MSC migration, and the migration ability of BM-MSCs was significantly enhanced when anti-miR-21 was added. Detailed MSC numbers are shown in Figure 4D. Together, these data indicated that miR-21 suppresses BM-MSC migration in vitro.

3.5 MiR-21 inhibits Wnt/β-catenin signaling in BM-MSCs

To explore the molecular mechanism underlying miR-21-mediated blockade of BM-MSC migration, we investigated the involvement of the Wnt/β-catenin pathway. We found that Wnt3a was activated as indicated by increased TOPFlash signals (∼4-fold) in miR-NC-transfected BM-MSCs, suggesting that Wnt/β-catenin signaling was induced (Figure 5A). When BM-MSCs were treated with the miR-21 mimic, Wnt3a was unable to significantly increase TOPFlash activity, suggesting that miR-21 inhibits Wnt/β-catenin signaling in BM-MSCs (Figure 5A). Furthermore, overexpression of miR-21 significantly decreased Ctnnb1 expression, whereas knockdown of miR-21 significantly increased Ctnnb1 expression (Figure 5B). Similar results were obtained at the protein level (Figure 5C, 5D). These results indicated that miR-21 inhibits the Wnt/β-catenin pathway in BM-MSCs.

3.6 MiR-21 suppresses Wnt/β-catenin signaling to inhibit BM-MSC migration in vitro

To explore the correlation between BM-MSC migration and Wnt/β-catenin signaling further, we assessed the role of β-catenin in anti-miR-21-transfected BM-MSCs. We transfected BM-MSCs with β-catenin siRNA or siRNA-NC and anti-miR-21 and then measured β-catenin expression. β-Catenin protein expression was significantly increased after anti-miR-21 transfection (Figure 6A, 6B). Further, knockdown of β-catenin using siRNA significantly decreased BM-MSC migration, whereas anti-miR-21 promoted BM-MSC migration by upregulating β-catenin expression (Figure 6C). Additionally, cell staining revealed that knockdown of β-catenin reduced BM-MSC migration upon addition of anti-miR-21, whereas BM-MSC migration was enhanced by upregulating β-catenin expression (Figure 6D and 6E). These findings suggested that miR-21 downregulates Wnt/β-catenin and subsequently impairs BM-MSC migration in vitro.

3.7 MiR-21 inhibits BM-MSCs migration by suppressing Wnt/β-catenin signaling in vivo

After demonstrating the role of the Wnt/β-catenin pathway in the effect of miR-21 in suppressing BM-MSC migration in vitro, we further assessed the effects of miR-21 on BM-MSC migration in vivo. We found that Ctnnb1 mRNA expression was enhanced after BM-MSC transplantation in asthmatic mice when compared with the PBS control group, and it was obviously decreased in the miR-21 + CRE + BM-MSCs group as compared to the miR-NC + CRE + BM-MSCs group (Figure 3A and 7A), suggesting that miR-21 negatively regulates BM-MSCs in asthma. Similar findings were obtained at the protein level (Figure 7B, 7C, and 7D). Collectively, these results indicated that miR-21 inhibits BM-MSC migration by suppressing Wnt/β-catenin signaling in CRE-induced asthma in vivo.

4. Discussion

Accumulating evidence demonstrates that transplanted MSCs can repair lung injury and improve inflammatory airway disorders [26, 27, 51]. Recently, researchers have found that miRNAs are involved in airway allergic inflammation in a mouse model of asthma after BM-MSC transplantation [37]. These findings suggested that miRNAs may participate in MSC migration by targeting β-catenin signaling. In support of this hypothesis, we found that: (1) miR-21 promoted asthma by suppressing MSC migration in a CRE-induced asthma mouse model; (2) miR-21 and anti-miR-21 transfection indicated that miR-21 inhibited BM-MSC migration in vitro; (3) miR-21 inhibited BM-MSC migration by suppressing Wnt/β-catenin signaling both in vivo and in vitro. A summarizing diagram of the role of miR-21 in suppressing BM-MSC migration by decreasing Wnt/β-catenin signaling in asthma is shown in Figure 8. We found that miR-21 is involved in the regulation of β-catenin expression; however, the exact mechanism requires further study. The conventional mechanism of miRNA regulation of gene expression is the inhibition of translation. Sometimes, miRNAs regulate gene expression at both transcriptional and translational levels; in this way, both mRNA and protein expression will be inhibited. In addition, one miRNA can regulate multiple target genes, and target genes of other miRNAs form a complex regulatory network that is extensively involved in biological abnormalities [52]. Collectively, our findings provide a novel insight into the molecular mechanisms underlying the regulation of MSC migration and provide a new therapeutic target for treating asthma.

Several miRNAs are implicated in asthma pathogenesis [34, 53], and miRNA-21 is a key factor in allergic airway diseases [44, 45, 54]. Eosinophilic inflammation and IL-4 levels reportedly are reduced and accompanied by an increase in IFN-γ levels in ovalbumin-induced airway inflammation in miR-21-knockout mice [55]. Further, miR-21 downregulates the expression of phosphatase and tensin homolog, which antagonizes phosphoinositide 3-kinase (PI3K) activity in a severe steroid-insensitive asthma mouse model [43]. Kim et al. showed that treatment with a miR-21-specific antagonist or the PI3K inhibitor LY294002 reduced PI3K activity and restored HDAC2 levels, which led to inhibition of the AHR and restoration of steroid sensitivity in allergic airway disease, indicating that miR-21 is a novel therapeutic target for asthma [44]. More recently, Lee et al. found that inhibition of miR-21 ameliorated allergic inflammation in a mouse model of asthma [45]. In line herewith, we found that suppression of miR-21 can ameliorate allergic inflammation in asthma by reducing the infiltration of inflammatory cells, especially eosinophils, TH2 cytokines IL-4, IL-5, and IL-13 in the BALF, AHR, and specific IgE production in the BALF.

MSCs have shown a therapeutic effect in models of acute lung inflammation and fibrosis by targeting several inflammatory cells, including mast cells, natural killer cells, B cells, T cells, regulatory cells, and dendritic cells [56, 57]. MiRNAs can serve as downstream adapters to regulate MSC migration and mediate different diseases [26, 37]. Yue et al. found that miR-124 expression is markedly decreased in MSCs in response to hepatocyte growth factor stimulation via suppression of β-catenin signaling [58]. Lee et al. reported that miR-543 and miR-590-3p decrease AIMP3/P18 expression levels by modulating cellular aging in MSCs in vitro [42]. More recently, Qiu et al. demonstrated that miR-155 regulates oxidative stress and cyclooxygenase-2 in CRE-induced murine asthma [38]. Thus, several miRNAs can mediate MSCs to influence asthma development and may serve as therapeutic targets.

It is well known that MSC migration into the injured site is crucial for MSC therapy in various diseases. How exactly miR-21 regulates the inflammatory response to exacerbate asthma is still unknown. In the present study, we found that miR-21 expression was markedly increased in the asthma model, and increased miR-21 expression was associated with suppressed MSC migration. MiR-21 could suppress MSC immigration both in vivo and in vitro. Our results
strongly suggest that miR-21 promotes CRE-induced asthma by suppressing MSC migration. Therefore, miR-21 negatively regulates the chemotactic migration of BM-MSCs, which can serve as a potential target for treating asthma in future.

Wnt/β-catenin plays a key role in the development of airway inflammation. Activation of β-catenin signaling by both Wnt-dependent and Wnt-independent pathways has been demonstrated to contribute to airway remodeling [59]. Wnt/β-catenin signaling activation by Wnt3a or LiCl improves the migratory capacity of MSCs [60], and the Wnt inhibitor Dkk-1 promotes pathological Th2-mediated inflammation [61]. Activation of Wnt/β-catenin signaling in alveolar epithelial cells attenuates intercellular adhesion molecule 1/vascular cell adhesion molecule 1-mediated adhesion and inhibits lung inflammation [62]. The Wnt/β-catenin pathway is one of the major pathways associated with MSC migration [60]. The Wnt pathway is activated in inflammatory bowel disease and can be suppressed by the transplantation of MSCs, which differentiate into intestinal epithelium [63]. MiR-124 expression is markedly decreased in MSCs in response to hepatocyte growth factor stimulation [58]. It seems that Wnt/β-catenin signaling plays different roles in inflammatory diseases by targeting inflammatory cells and the immune microenvironment. In the present study, miR-21 was characterized as a novel regulator of the canonical Wnt/β-catenin signaling pathway, which is involved in the migration of MSCs. Based on our findings, we conclude that miR-21 inhibits MSC migration by targeting Wnt/β-catenin signaling. This conclusion was supported by the following evidence. First, miR-21-treated CRE-induced asthma model mice showed decreased β-catenin expression compared to miR-NC-treated mice. Second, miR-21-treated asthmatic mice showed less MSC migration than miR-NC-treated mice. Finally, miR-21 was found to be associated with β-catenin expression in influencing MSC migration both in vivo and in vitro. Taken together, our data indicate that miR-21 inhibits MSC migration by suppressing Wnt/β-catenin signaling.

5. Conclusions
Our study demonstrated that miR-21 significantly stimulates CRE-induced asthma by inhibiting MSC chemotactic migration both in vivo and in vitro. More importantly, we provided evidence that Wnt/β-catenin signaling is involved in the inhibitory effect of miR-21 on the migration of BM-MSCs. MiR-21 facilitates the migration of BM-MSCs, which is a potential target for MSC-based treatment of asthma. Other possible mechanisms involved in MSC migration with regard to the treatment of asthma remain to be explored.

Abbreviations
AHR: airway hyperresponsiveness; Ig: immunoglobulin; TH2: T helper type 2; IL: interleukin; MSC: mesenchymal stem cell; BM: bone marrow; WT: wild-type; CRE: cockroach extract; BALF: bronchoalveolar lavage fluid; NC: negative control; siRNA: small interfering RNA; IFU: infectious unit; Eos: eosinophil; Lym: lymphocyte; Mac: macrophage; Neu: neutrophile; ELISA: enzyme-linked immunosorbent assay; HE: hematoxylin and eosin; PI3K: phosphoinositol 3-kinase

Declarations
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Ethics approval and consent to participate
This study was submitted to and approved by the Center for Medical Ethics, Central South University. All protocols were approved by the Institutional Review Board of the Animal Center of Central South University.

Consent for publication
We confirm that the figures in the manuscript are original, and all authors agreed with publication.

Competing interests
The author declares no competing financial interests.

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Availability of data and materials
Detailed data produced and analyzed in the current study are available from the corresponding author upon reasonable request.

Authors' contributions
Tianli Cheng: Conceptualization, Project administration, Writing - Original Draft
Jianfu Heng: Project administration, Writing - Review & Editing, Formal analysis
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Figures

Figure 1

A

B

C

Figure 1

MiR-21 is increased in CRE-induced asthma model mice (A) Protocol for establishment of the CRE-induced asthma mouse model. In the challenge phase, mice were intranasally administered 2 × 106 infectious units (IFUs) of lentivirus carrying miR-21 or empty lentivirus. (B) RT-qPCR analysis of miR-21 expression in the lungs of the CRE-challenged asthmatic mice. (C) RT-qPCR analysis of miR-21 expression in the lungs of miR-21- or miR-NC-transfected CRE-challenged asthmatic mice. Data are representative of three independent experiments (n = 6). Data are presented as the mean ± SEM. *P < 0.05, **P < 0.01.
miR-21 promotes lung inflammation in asthma (A) Hematoxylin and eosin (HE) staining was used to evaluate airway inflammation and cell infiltration in PBS- and CRE-treated mice that were transfected with miR-21 or miR-NC. (B) Total cell numbers in BALFs. (C) Numbers of different inflammatory cells (Eos, Lym, Mac, and Neu) in BALFs. (D) Airway hyperresponsiveness to metacholine was analyzed in the different groups. ELISAs were used to evaluate CRE-specific IgE (E) and IgG1 (F) levels in BALFs. ELISAs were used to evaluate the Th2 cytokines IL-4 (G), IL-5 (H), and IL-13 (I) in BALFs. Data are representative of three independent experiments (n = 6). Data are presented as the mean ± SEM. *P < 0.05, **P < 0.01, ***P < 0.001.
Figure 3

BM-MSCs improve allergic airway inflammation and AHR, whereas miR-21 aggravates the inflammatory reaction by suppressing BM-MSC migration (A) Timeline of BM-MSC engraftment of CRE-induced asthmatic mice. The mice were injected intravenously with BM-MSCs or PBS on 9 day. (B) GFP immunofluorescence was used to demonstrate that BM-MSCs migrated to the lungs. (C) HE staining was used to evaluate airway inflammation infiltration in CRE-induced asthmatic mice treated with BM-MSCs or PBS. Total cell (D) and differentiated leukocyte (E) numbers in BALFs. ELISAs were used to evaluate CRE-specific IgE (F) and IgG1 (G) levels in BALFs. ELISAs were used to evaluate the Th2 cytokines IL-4 (H), IL-5 (I), and IL-13 (J) in BALFs. Data are representative of three independent experiments (n = 6). Data are presented as the mean ± SEM. *P < 0.05, **P < 0.01, ***P < 0.001.
miR-21 suppresses BM-MSC migration in vitro (A) RT-qPCR analysis of miR-21 expression after transfection of BM-MSCs with miR-21, miR-NC, anti-miR-21, or anti-miR-NC. (B) Wound healing assays of BM-MSCs transfected with miR-21, miR-NC, anti-miR-21, or anti-miR-NC. (C) BM-MSCs were transfected with miR-21, miR-NC, anti-miR-21, or anti-miR-NC and subjected to Transwell migration assay (magnification, 20×). (D) Quantification of migrated MSCs. Data are representative of three independent experiments. Data are presented as the mean ± SEM. *P < 0.05, **P < 0.01, ***P < 0.001.
Figure 5

miR-21 inhibits Wnt/β-catenin signaling in BM-MSCs (A) TOPFlash/FOPFlash luciferase activity in BM-MSCs treated with miR-21 mimic. (B) RT-qPCR analysis of Ctnnb1 expression in BM-MSCs transfected with miR-21, miR-NC, anti-miR-21, or anti-miR-NC. GAPDH was used as a reference gene. (C) Immunoblotting of β-catenin in BM-MSCs transfected with miR-21, miR-NC, anti-miR-21, or anti-miR-NC. (D) Protein quantification. All data are reported as the mean ± SD of three independent experiments. *P < 0.05.
Figure 6

miR-21 suppresses Wnt/β-catenin signaling to inhibit BM-MSC migration in vitro (A) β-Catenin protein in BM-MSCs transfected with anti-miR-21 and β-catenin-siRNA was detected by western blotting. (B) Protein quantification. (C) Wound healing assays of BM-MSCs transfected with anti-miR-21 and β-catenin-siRNA. (D) The Boyden chamber assay was used to evaluate the migration ability of BM-MSCs under different treatments. (E) Quantification of migrated cells in the different groups. All data are reported as the mean ± SD of three independent experiments. *P < 0.05, **P < 0.01, ***P < 0.001.

Figure 7

miR-21 inhibits BM-MSC migration by suppressing Wnt/β-catenin singling in vivo (A) RT-qPCR analysis of Ctnnb1 mRNA expression in the lungs after transfection of BM-MSCs into CRE-induced asthma model mice. (B) Immunohistochemistry of β-catenin expression in the lungs of CRE-induced asthma mice.
model mice. (C) Western blot analysis of β-catenin protein expression in the lungs of CRE-induced asthma model mice that received miR-21-transfected BM-MSCs. (D) Quantification of protein contents. Data are presented as the mean ± SEM. *P < 0.05, ***P < 0.001.