MicroRNA-20b promotes the accumulation of CD11b+Ly6G+Ly6Clow myeloid-derived suppressor cells in asthmatic mice

HUA MA1*, SHUJUN GUO1*, YULAN LUO1, YIMENG WANG1, HE LONG WANG1, JING HE1, JIE TANG1, LIN SHEN2, CHUANWANG SONG1

1Department of Immunology, Anhui Provincial Key Laboratory of Infection and Immunity, Bengbu Medical College, Bengbu, Anhui, China
2Scientific Research Center, Bengbu Medical College, Bengbu, Anhui, China

*These authors contribute equally to this work.

Abstract

miR-20b is a member of the miR-106a-363 gene cluster, which has been shown to play an important role in a variety of diseases, including cancer, inflammation, and autoimmune diseases. Our previous study indicated that miR-20b has an inhibitory effect on airway inflammation in asthmatic mice, but the exact mechanism is unclear. In this study, we report that the ratio of CD11b+Ly6G+Ly6Clow cells, but not the amount of CD11b+Ly6C+Ly6G– cells, was increased in the lung tissue of asthmatic mice after intranasal instillation with miR-20b mimics, while Th2-type cytokines (interleukin (IL)-4 and IL-13) were significantly decreased in the bronchoalveolar lavage fluid. In addition, the transcription factor CREB regulated the expression of miR-20b. Our findings suggest that miR-20b can induce the accumulation of myeloid-derived suppressor cells in the lungs of asthmatic mice, which may be a mechanism by which miR-20b inhibits airway inflammation in asthmatic mice. Thus, miR-20b may be used as a target for the effective treatment of asthma in the future.

Key words: asthma, MiR-20b, myeloid-derived suppressor cells (MDSCs), allergic airway inflammation.

Introduction

Bronchial asthma (here referred to as asthma) is chronic airway inflammation caused by many kinds of cells and inflammatory factors. Airway hyper-responsiveness, chronic inflammation, and airway remodeling are characteristic features of asthma [1, 2]. Epidemiological studies have found that the incidence and mortality of asthma are increasing year by year, but its pathogenesis is still unclear at present [3]. MicroRNAs (miRNAs) are non-coding RNAs which are about 22 nucleotides in length. They regulate the expression of target genes by inhibiting their translation or causing mRNA degradation [4]. An increasing number of studies has found that many miRNAs may participate in asthma pathogenesis [5]. In an asthmatic mouse model, the upregulation of miR-221 and miR-485-3p may play an important role in the pathogenesis of asthma by downregulating the expression of Spre-2 [6]. Collison et al. demonstrated that the expression of miR-126 was upregulated in the lung tissues of asthmatic mice, blocking miR-126 significantly reduced airway eosinophil cell infiltration [7]. Panganiban et al. showed that the expression of miR-1248 was upregulated in the serum of asthmatic patients, and it positively regulates the expression of interleukin (IL)-5, which is a key factor for chemotaxis, survival, and maturation of eosinophils [8]. In severe asthma, blood CD8+ T cells are activated at the same time, accompanied by reduced miR-146a/b and miR-28-5p expression [9]. In our previous study, we found that the expression of miR-20b in the lung tissues of asthmatic mice was significantly lower than that of normal mice [10]. In addition, miR-20b inhibited airway inflammation in asthmatic mice, but the mechanism remains unclear [11]. Since myeloid-derived suppressor cells (MDSCs) effectively inhibit airway inflammation in asthmatic mice [12], in this study we analyzed the effect of miR-20b on MDSCs content in the lung tissue of asthmatic mice.

Material and methods

Mice

Female BALB/c mice (2- to 3-week-old) were purchased from the Center of Experimental Animals of Bengbu Medical College, Donghai Avenue 2600, Bengbu, Anhui, 233030 Bengbu, China, e-mail: chuanwangsong@163.com

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Medical College. This study was approved by the Animal Care and Ethics Committee of Bengbu Medical College.

Establishment of experimental asthma model and treatment

Female BALB/c mice were randomly divided into four groups, a normal control group (Control), an asthmatic group (Asthma), a miR-20b-mimic treatment group (MiR-20b mimic) and a scrambled miR-20b treatment group (MiR-20b scrambled), with seven mice in each group. For all groups except the Control group, mice were sensitized by intraperitoneal injection with 200 μl liquid (including 50 μg of OVA and 2 mg of aluminum hydroxide) on days 0, 7, and 14. On day 21 from the start of the experiment, the mice were placed in an atomization inhalation box and were challenged with 5% OVA solution inhalation with nebulization for 30 min, continuously for 4 weeks. In addition, the MiR-20b mimic group was treated with 20 μg miR-20b mimics in 40 μl doses by nasal drip starting from the 20th day, once every three days for a total of 10 doses. The MiR-20b scrambled group received scrambled miR-20b instillation to replace the treatment with miR-20b mimics (Fig.1).

Preparation of bronchoalveolar lavage fluid (BALF)

On day 49, the mice in each group were sacrificed and tracheal intubations were performed. Mice were lavaged 6 times with 0.8 ml PBS. The BALF obtained (in total about 4.8 ml) was centrifuged at 500 × g for 5 min and the supernatant was stored at −20°C for later use in an enzyme-linked immunosorbent assay (ELISA).

Oligonucleotides, siRNAs, and primers

MiR-20b mimics, scrambled miR-20b, CREB siRNA, and CREB control siRNA were obtained from GenePharma (Shanghai, China). The miRNA sequences used were as follows: miRNA-20b mimics, sense: 5'-CAA AGU GCU CAU AGU GCCU GCA GGU AG-3', antisense: 5'-ACC UGC ACU AUG AGC ACU UUG UU-3'; scrambled miR-20b, sense: 5'-UUC UCC GAA CGU GUC AGC UTT-3', antisense: 5'-ACG UGA CAC GUU CGG AGA ATT-3'; CREB siRNA, sense: 5'-GUC UCC ACA AGU CCA AAC ATT-3', antisense: 5'-UGU UUG GAC UUG UGG AGA CTT-3'; and CREB control siRNA, sense: 5'-UUC UCC GAA CGU GUC AGC UTT-3', antisense: 5'-ACG UGA CAC GUU CGG AGA ATT-3'. PCR forward and reverse primers were synthesized by Sangon Biotech (Shanghai, China). The primer sequences used were: GAPDH primer, sense: 5'-GGC AAA TTC AAC GGC ACA-3', antisense: 5'-TCC AGC ACA TAC TCA GCA CC-3'; CREB primer, sense: 5'-TGC CAC ATT AGC CCA GGT TCG TCA TAG TG-3', antisense: 5'-GTG CAG GGT CCG AGG T-3'; miR-20b RT primer: 5'-GGG ACG CCA TAA CAA CT-3'; CREB RT primer: 5'-AGG GGG AGA CGC TTC TAC AGC CA-3', antisense: 5'-ATG CCA AAG TGC TCA TCG TAG TG-3'; 5'-ATG CCA AAG TGC TCA TCG TAG TG-3'; CREB control RT primer: 5'-ACG UGA CAC GUU CGG AGA ATT-3', antisense: 5'-AAG GCT TCA CGA ATT TGC GT-3'; and MiR-20b RT primer: 5'-AAG GCT TCA CGA ATT TGC GT-3'.

Transient transfection

CREB siRNA and control siRNA (160 nM) were transfected using Lipofectamine 2000 (Invitrogen) according to the instructions of the manufacturer. A murine blood macrophage cell line (RAW 264.7 cells; 7.2 × 10^5 cells/well) was seeded into a six-well plate, and the following day, the transfected cells reached 70-90% confluence. The siRNAs and Lipofectamine 2000 were diluted with Opti-MEM with thorough mixing at room temperature for 20 min. The RAW 264.7 cells were incubated with this mixture for 48 h in standard medium. Finally, the cells were harvested for specific experiments.

Quantitative real-time PCR (qRT-PCR)

Total RNA was extracted using TRIZOL (Invitrogen). cDNA was synthesized with a TransScript First-strand cDNA Synthesis SuperMix kit from TransGen. The reaction mix included total RNA (500 ng), 1 μl anchored oligo (dT), 10 μl 2 × TS Reaction Mix, and 1 μl TransScript RT/RI Enzyme Mix (the anchored oligo (dT) was replaced with the RT primer corresponding to the target gene when
the miRNA cDNA first-strand was synthesized). The reaction conditions were 30 min at 42°C then 85°C for 5 min. Quantitative PCR (qPCR) was performed with the StepOnePlus System (ABI) using the TransStartTM EcoGreen qPCR SuperMix (TransGen). The amplification conditions were 94°C for 30 s, followed by 40 cycles of 94°C for 5 s, and 60°C for 30 s. U6 RNA or GAPDH were used as endogenous controls for miRNA or other mRNA levels. The relative gene expression was analyzed using the 2^{-ΔΔCT} method.

FACS analysis

On day 49, the lung tissues from each group were obtained. Single-cell suspensions were prepared from these lung tissues and incubated with fluorescein-conjugated antibodies for 30 min at 4°C in the dark. The stained cells were analyzed using a flow cytometer (BD FACS Caliber).

ELISA

The amount of cytokines in the serum or the BALF was determined with commercial sandwich ELISA kits (CUSABIO) according to the manufacturer’s instructions.

Western blot analysis

Briefly, RAW 264.7 cells were lysed with NP-40 lysis buffer with 1 mM PMSF. Cell lysates were separated by 10% sodium dodecyl sulfate-polyacrylamide gel electrophoresis and transferred to nitrocellulose membranes. The membranes were blocked with 5% BSA in TBST and probed with the respective antibodies. The blots were detected using enhanced chemiluminescence according to the manufacturer’s protocol (Beyotime biotechnology).

Statistical analysis

Data are expressed as mean ± standard deviation. SPSS16.0 was used to perform single-factor analysis of variance for multiple groups. A P-value < 0.05 was considered statistically significant.

Results

MiR-20b suppresses Th2 cytokine production in the BALF of asthmatic mice

To understand the mechanism by which miR-20b inhibits allergic airway inflammation, we studied the effect of miR-20b on the production of Th1/Th2-type cytokines in mouse BALF. The results are shown in Fig. 2. Compared with the normal control group, the levels of the Th2-type cytokines IL-4 and IL-13 were significantly increased in the BALF of the asthmatic mice. However, the Th1-type cytokine IFN-γ was not significantly decreased in the asthmatic mice. The secretion of IL-13 and IL-4 was significantly decreased after treatment with miR-20b mimics in asthmatic mice (p < 0.01).

In contrast, the production of IL-10 in each group was not significantly different in this experimental system.

MiR-20b promotes the accumulation of CD11b+Ly6G+Ly6Clow MDSCs in asthmatic mice

Since MDSCs have significant immunosuppressive activity, their relationship with asthma has recently attracted attention. To discern whether the inhibitory effect of miR-20b on airway inflammation involves MDSCs, the MDSC profiles in the lung tissues of mice in each group were assayed (Fig. 3). The results showed that there was not a significant increase of MDSCs in the asthmatic group compared with the normal group. After treatment with miR-20b mimics, the CD11b+Ly6G+Ly6Clow MDSC ratio was significantly elevated in asthmatic mice (p < 0.05), while the increase in CD11b+Ly6C+Ly6G–MDSCs was not statistically significant. Treatment with miR-20b-scramble did not have a significant effect on the content of MDSCs in the lung tissues of asthmatic mice.

MiR-20b increases CCL2 and TGF-β in asthmatic mice

In this study, the CCL2 content in each group was detected, and the results are shown in Fig. 4A. There was not a significant difference in CCL2 concentration between the asthmatic group and the normal group, while CCL2 levels in BALF were significantly higher after asthmatic mice treatment with miR-20b mimics. Therefore, miR-20b induced the accumulation of MDSCs accompanied by an increase in CCL2 concentration in the lungs of asthmatic mice. Because TGF-β is an important effector molecule of MDSCs, the concentration of TGF-β in the BALF and serum of each group of mice was assayed (Fig. 4B, C). As expected, relative to the other three groups, the concentration of TGF-β, whether in the serum or in the BALF, was increased in the miR-20b-mimic treatment group.

CREB regulates miR-20b expression

At present, there have been few studies on the regulation of miR-20b expression. We used two bioinformatics software packages, P-MATCH 1.0 (http://www.gene-regulation.com/cgi-bin/pub/programs/pmatch/bin/p-match.cgi) and AliBaba2 (http://www.gene-regulation.com/pub/programs/alibaba2/index.html), to predict the transcription factors that might be involved in the regulation of miR-20b expression, in which the CREB score was higher. Thus, we attempted to verify whether CREB regulates the expression of miR-20b. The expression of miR-20b in RAW 264.7 cells transfected with CREB siRNA was analyzed by qRT-PCR analysis (Fig. 5A, B). CREB siRNA decreased the miR-20b level to 0.375-fold of the control (p < 0.01). To confirm the activity of CREB siRNA, we examined the mRNA and protein level of CREB in RAW 264.7 cells after transfection. The level of CREB mRNA decreased by 66% after CREB
siRNA transfection for 48 h (Fig. 5C). The expression of the CREB protein was also decreased by 75% (Fig. 5D). These results indicate that the expression of miR-20b is regulated by CREB upstream in RAW 264.7 cells.

**Discussion**

Myeloid-derived suppressor cells are a group of myeloid precursor cells which fail to differentiate into mature immune cells. Having prominent immunosuppressive activity, MDSCs can significantly inhibit the function of a variety of T cells [13, 14]. Myeloid-derived suppressor cells can be divided into different subsets based on their surface molecules. In humans, MDSCs can be subdivided into the CD11b+CD33+HLA-DR–CD14+ and CD11b+CD15+HLA-DR–CD14– subgroups. In mice, MDSCs can be further subdivided into CD11b+Ly6G+Ly6Clow and CD11b+Ly6C+Ly6G– subsets according to Ly-6C/G ex-
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In this study, the proportion of CD11b+Ly6G+Ly6C<sup>low</sup> MDSCs increased in lung tissue from asthmatic mice after administration of nasal drops containing miR-20b mimics, but the ratio of CD11b+Ly6C+Ly6G<sup>-</sup> MDSCs was not changed. The latest research shows that many miRNAs are related to the biological functions of MDSCs, such as chemotaxis, aggregation, and functional activities [17]. Liu et al. demonstrated that miR-494 may activate the Akt pathway by targeting PTEN to regulate the accumulation and functional activity of MDSCs. In addition, miR-494 mediated the chemotaxis of MDSCs by CXCR4 chemokines in tumor tissues [18]. Li et al. showed that miR-155 and miR-21 increased GM-CSF and IL-6-induced bone marrow cell-derived MDSCs, which includes both G-MDSC and M-MDSC types. Furthermore, miR-155 and miR-21 activated STAT3 by targeting SHP-T and PTEN to synergistically induce the generation of MDSCs [19]. Mei et al. demonstrated that miR-200c contributed to the expansion and immunosuppressive activity of MDSCs in a mouse lung cancer and melanoma model [20]. Our previous study indicated that CCL2 signaling mediates the migration of MDSCs into the lung. In this study, we found that miR-20b can induce an increase in CCL2 concentration in the lungs of asthmatic mice [12]. Therefore, miR-20b may induce the accumulation of MDSCs by upregulating CCL2 chemokines in the lungs of asthmatic mice.

Once activated by inflammation- or tumor-driven factors, MDSCs can release a variety of biologically active mediators, including inducible nitric oxide synthase (iNOS), arginase 1, reactive oxygen species (ROS), IL-10, and transforming growth factor β (TGF-β) to inhibit the function of immune cells [21, 22]. Our preliminary results indicate that tumor-derived MDSCs inhibit the allergic Th2 response in a TGF-β-dependent manner. In this study, we also found that the elevation of MDSCs induced by miR-20b was accompanied by an increase in TGF-β in BALF and blood. Thus, miR-20b may inhibit airway inflammation in a TGF-β-dependent manner by inducing MDSCs in asthmatic mice.

Th1/Th2 cytokines play a key role in regulating airway inflammation in asthmatic mice [23]. It is noteworthy that the administration of miR-20b suppressed Th2 cytokines in the BALF of asthmatic mice in the current study. Recent studies have shown that miRNAs play an important role in regulating the balance of the Th1/Th2 immune response [24]. Lu et al. showed that lung eosinophil infiltration was decreased and the Th1-type cytokine IFN-γ was significantly increased when miR-21 was blocked in a murine asthma model. Meanwhile, DC with miR-21 deficiency increased the production of IL-12 after stimulation with LPS. miR-21-deficient mice had significantly increased Th1 delayed-type hypersensitiv-

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**Fig. 3.** miR-20b promotes the accumulation of CD11b+Ly6G+Ly6C<sup>low</sup> MDSCs in asthmatic mice. Lung tissues of mice were surgically removed on day 49 after baseline. Single-cell suspensions were prepared and stained with FITC-Ly6G, PE-Ly6C, and PE-cy5-CD11b. Stained cells were analyzed using flow cytometry (A). The proportion of CD11b+Ly6G+Ly6C<sup>low</sup> cells (B) and CD11b+Ly6C+Ly6G<sup>-</sup> cells (C) in lung tissues were calculated. Data are expressed as mean ± standard deviation; *p < 0.05 compared with the Asthma group.
ity [25]. These studies emphasize that miR-21 mainly regulates Th1 response rather than the Th2 reaction. Meanwhile, Mohnle et al. showed that miR-146 phosphorylated STAT4 by targeting PKCε, thus resulting in Th1-cell differentiation [26]. However, our study showed that miR-20b inhibited the Th2-type response in asthmatic mice, but had no effect on Th1-type response. A study by Mattes et al. also showed that the function of Th2 cells was inhibited, and the symptoms of asthma were reduced, in mice with miR-126 ablation, and blocking miR-126 may lead to the increased expression of POU domain class 2 associating factor 1, which can negatively regulate the expression of the Th2-specific regulatory factor GATA3 [27].

This study also demonstrated that the transcription factor CREB can regulate the expression of miR-20b at the upstream level. Since the pathogenesis of asthma is closely related to miRNAs, research on the regulation of miRNAs may shed light on new asthma treatments [28]. At present, research on the regulation of miRNA expression is relatively rare. Recent studies show that the CCCTC-binding factor (CTCF) can not only control the expression of miR-125b1 and miR-375, but can also adjust the level of the miR-290 cluster in tumor cells [29]. Chavali et al. demonstrated that binding of the HMGI/Y protein to matrix attachment regions (MARs) induced histone acetylation, resulting in the expression of the miR-17-92 cluster.
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and individual miRNAs (miR-221, 93, 17, and let-7b) in neuroblastoma cells but not in fibroblasts [30]. Zhang et al. showed that two members of the E-2b family (PEA3 and ELK-1) regulated the expression of miR-200b. Furthermore, ELK-1 inhibited the expression of miR-200b, and PEA3 promoted the expression of miR-200b [31]. Since CREB can regulate the expression of miR-20b, regulating CREB upstream of miR-20b may provide a therapeutic target for asthma treatment.

In summary, we show an important role for miR-20b in the pathogenesis of asthma, inducing the accumulation of CD11b+Ly6G+Ly6C<sub>low</sub> cells in lung tissues. This may be a mechanism by which miR-20b suppresses airway inflammation in asthmatic mice. This study provides an experimental basis for the future clinical therapy of asthma using miR-20b as a target point of treatment.

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The authors declare no conflict of interest.
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