

# Mechanical stretch regulates microRNA expression profile via NF- $\kappa$ B activation in C2C12 myoblasts

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Received September 13, 2015; Accepted September 23, 2016

DOI: 10.3892/mmr.2016.5907

**Abstract.** MicroRNAs (miRNAs/miRs) and nuclear factor (NF)- $\kappa$ B activation are involved in mechanical stretch-induced skeletal muscle regeneration. However, there are a small number of miRNAs that have been reported to be associated with NF- $\kappa$ B activation during mechanical stretch-induced myogenesis. In the present study, C2C12 myoblasts underwent cyclic mechanical stretch *in vitro*, to explore the relationship between miRNA expression and NF- $\kappa$ B activation during stretch-mediated myoblast proliferation. The results revealed that 10% deformation, 0.125 Hz cyclic mechanical stretch could promote myoblast proliferation. The miRNA expression profile was subsequently altered; miR-500, -1934, -31, -378, -331 and -5097 were downregulated, whereas miR-1941 was upregulated. These miRNAs were all involved in stretch-mediated myoblast proliferation. Notably, the expression of these miRNAs was reversed following treatment of 0.125 Hz mechanically stretched C2C12 cells with NF- $\kappa$ B inhibitors, which was accompanied by C2C12 cell growth suppression. Therefore, the present study is the first, to the best of our knowledge, to demonstrate that the NF- $\kappa$ B-dependent miRNA profile is associated with mechanical stretch-induced myoblast proliferation.

## Introduction

In recent years, the incidence of skeletal muscle injury has increased. It is well known that skeletal muscles only possess limited self-renewal capacity. Skeletal muscles respond to mechanical load, and this mechanical stimulation has been

reported to promote myogenesis by provoking muscle precursor cell activation, proliferation and differentiation (1). In the field of regenerative medicine, appropriate mechanical stimulation has been applied to muscle-derived stem cells in order to promote skeletal muscle repair (2). Furthermore, it has been suggested that external mechanical stimuli are transformed into intracellular signals through certain mechanisms, in order to regulate myoblast activity (3,4). However, the mechanisms underlying the transformation of external mechanical stimuli into intracellular signals remain unclear.

Nuclear factor (NF)- $\kappa$ B, which belongs to the Rel family, refers to several transcription factors. The mammalian NF- $\kappa$ B family comprises seven members, including p65 (RelA), c-Rel, RelB, p50/p105 (NF- $\kappa$ B1) and p52/p100 (NF- $\kappa$ B2). NF- $\kappa$ B usually exists in the form of hetero- or homodimers, and the typical NF- $\kappa$ B complex is composed of a p50 and p65 heterodimer. In non-stimulated cells, the NF- $\kappa$ B complex is located in the cytoplasm and is inhibited by the inhibitor of NF- $\kappa$ B (I- $\kappa$ B $\alpha$ ). Once stimulated, the degradation of I- $\kappa$ B $\alpha$  and phosphorylation of NF- $\kappa$ B induce the translocation of NF- $\kappa$ B into the nucleus where it binds to target DNA (5). The translocation of NF- $\kappa$ B has been reported to regulate several cellular processes, including immunity and inflammation, cellular proliferation and differentiation (6-8). The role of NF- $\kappa$ B is indispensable and complex due to its presence in both proliferation and differentiation stages during the myogenic process (9). As for stretch-induced skeletal muscle alterations, the NF- $\kappa$ B signaling pathway has been shown to be activated in murine diaphragm muscles subjected to longitudinal mechanical stretch, resulting in upregulated expression of the stretch-response gene, Ankyrin repeat domain 2 (10). NF- $\kappa$ B activation by phosphorylation of p65 NF- $\kappa$ B has also been detected during cyclic stretch-induced myogenic differentiation (11). A previous study revealed that NF- $\kappa$ B transcriptional activity is beneficial to myoblast proliferation under appropriate stretch induction (12).

MicroRNAs (miRNAs/miRs) are small non-coding RNAs, some of which are highly evolutionarily conserved. miRNAs regulate gene expression at the post-transcriptional level by inhibiting translation of target mRNA or promoting target mRNA degradation (13). miRNAs have been identified as novel components in the gene regulatory network of myogenesis (14). A previous study reported that miRNAs are essential for the generation and maintenance of skeletal

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**Key words:** mechanical stretch, microRNAs, nuclear factor- $\kappa$ B, myoblasts, myogenesis

muscles (15). In particular, miRNAs specifically expressed in muscles, including miR-1 and miR-206, are able to promote skeletal myoblast differentiation (16,17). Conversely, another muscle-specific miRNA, miR-133, serves an opposite role in myogenesis by maintaining myoblasts in a proliferative state (16). Furthermore, local injection of miR-1, miR-133 and miR-206 can upregulate myogenin and myogenic differentiation 1 (MyoD) expression in injured skeletal muscles, and thus accelerates muscle regeneration (18). miRNAs that are ubiquitously expressed can also influence skeletal muscle development. miR-181 inhibits skeletal myogenesis, whereas miR-146a and miR-26a positively regulate myogenesis (19-21). However, interactions between the NF- $\kappa$ B signaling pathway and miRNAs during mechanical stretch-induced myoblast proliferation and differentiation remain unclear.

The present study subjected C2C12 myoblasts to cyclic stretch to investigate the interactions between NF- $\kappa$ B activation and miRNA expression during mechanical stretch-induced myoblast proliferation. The study focused on the myoblast proliferation process, since it represents the early stages of muscle regeneration. The computer-controlled Flexcell system was used to apply cyclic mechanical strain to C2C12 myoblasts. The appropriate mechanical stretch conditions that could promote C2C12 myoblast proliferation were initially confirmed, and the miRNA expression alterations during mechanical stretch-induced myoblast proliferation were determined using high-throughput sequencing and reverse transcription-quantitative polymerase chain reaction (RT-qPCR) methods. In addition, the interactions between NF- $\kappa$ B activation and miRNA expression were explored by treating stretch-induced myoblasts with or without an NF- $\kappa$ B inhibitor.

## Materials and methods

**Cell culture.** C2C12 mouse myoblast cells were purchased from American Type Culture Collection (Manassas, VA, USA). The frozen stock was thawed and maintained as monolayer cultures in Dulbecco's modified Eagle's medium supplemented with 10% fetal bovine serum (both Gibco; Thermo Fisher Scientific, Inc., Waltham, MA, USA), 100 U/ml penicillin and 1,000 U/ml streptomycin at 37°C in a humidified atmosphere containing 5% CO<sub>2</sub>. The medium was changed twice per week.

**Mechanical stretch stimulation of C2C12 cells.** The model of *in vitro* stretching of C2C12 myoblasts was established using the computer-controlled vacuum stretch system (FX-5000 Tension system; FlexCell International Corporation, Burlington, NC, USA). Briefly, C2C12 cells were seeded into type-I collagen-coated flexible-bottom six-well plates (BioFlex collagen I plates; FlexCell International Corporation) at a density of 1x10<sup>5</sup> cells/well in culture medium. The cells in the flexible-bottom six-well plates were then incubated at 37°C in a humidified 5% CO<sub>2</sub> atmosphere overnight. Subsequently, C2C12 cells underwent cyclic strain (10% deformation) at 0.125, 0.25 and 0.5 Hz for 2 h per day for 4 consecutive days. Cells cultured under the same conditions without cyclic strain were considered the control group.

**Cell viability assay and cell cycle analysis.** C2C12 cell viability was assessed 1, 2, 3 and 4 days after initiation of cyclic strain

using the Cell Counting kit-8 (CCK-8) assay. Briefly, C2C12 cells in the control group, and 0.125, 0.25 and 0.5 Hz groups were digested with 0.25% pancreatic enzyme (trypsin; Gibco; Thermo Fisher Scientific, Inc.) containing 0.02% EDTA. The cells were then reseeded in 96-well plates at a density of 5x10<sup>3</sup> cells/well in culture medium (8 duplicate wells/group). The plates were incubated for 24 h at 37°C, after which the cell medium was replaced with 100  $\mu$ l culture medium containing 10  $\mu$ l CCK-8 reagent (Dojindo Molecular Technologies, Inc., Kumamoto, Japan). Following a further incubation for 3 h at 37°C, the optical density (OD) values were measured at 450 nm using a microplate reader (Thermo Fisher Scientific, Inc.). Cell cycle analysis was performed by flow cytometry (FCM). Briefly, stretched C2C12 cells (10% deformation; 0.125 Hz) at day 4 and control cells were collected. Subsequently, the cells were permeabilized with 70% (v/v) ethanol and incubated overnight at 4°C. Finally, the cells were stained with 500  $\mu$ l propidium iodide (100  $\mu$ g/ml) at 37°C for 30 min and analyzed by FCM. The relative DNA proliferation index (DPI) was calculated using the following formula: [DPI = (S% + G<sub>2</sub>/M%) / (S% + G<sub>2</sub>/M% + G<sub>0</sub>/G<sub>1</sub>%)], according to our previous report (7).

**High-throughput sequencing.** High-throughput sequencing was applied to detect differentially expressed miRNAs in C2C12 cells between the control group and the 0.125 Hz stretch group at day 4. Total RNA was extracted from the C2C12 cells using TRIzol® (Invitrogen; Thermo Fisher Scientific, Inc.) according to the manufacturer's protocol. Small RNAs (18-75 nt) were separated from total RNAs by polyacrylamide gel electrophoresis and were purified. The small RNAs were ligated with 5' and 3' adapters using T4 RNA Ligase (Thermo Fisher Scientific, Inc.), and were reverse transcribed to cDNA using SuperScript II (Life Technologies; Thermo Fisher Scientific, Inc.). Generation of double-stranded cDNA was performed prior to PCR amplification. High-throughput sequencing of the purified DNA samples was performed using the Illumina Genome Analyzer IIX (Illumina, San Diego, CA, USA). To select differentially expressed miRNAs between the two groups, the frequency of miRNAs was normalized to calculate the ratio of 0.125 Hz stretched cells to control cells. A differentially expressed miRNA was indicated by a stretched/control ratio >2 and a statistically significant result, as determined by independent t-test with Bonferroni correction.

**NF- $\kappa$ B inhibition.** Two selective and irreversible NF- $\kappa$ B inhibitors: BAY11-7082 (BAY; Beyotime Institute of Biotechnology, Jiangsu, China) and pyrrolidine dithiocarbamate (PDTC; Beyotime Institute of Biotechnology) were used in the present study. In order to determine the effects of NF- $\kappa$ B inhibition on the stretched myoblasts, C2C12 cells that underwent 0.125 Hz cyclic strain (10% deformation) for 2 h per day for 4 consecutive days were simultaneously treated with 2.5  $\mu$ M BAY, 5  $\mu$ M BAY or 10  $\mu$ M PDTC. The effects of NF- $\kappa$ B inhibition on stretched C2C12 myoblasts were subsequently detected according to the aforementioned CCK-8 method and FCM analysis at corresponding time points.

**Western blot analysis.** Cells were lysed with radioimmuno-precipitation assay buffer (EMD Millipore, Billerica, MA, USA) and protein concentration was determined using the

bicinchoninic acid assay kit (Thermo, Fisher Scientific, Inc.). Protein samples (50  $\mu\text{g}$ ) were separated by SDS-PAGE (5% stacking gel, 10% separating gel), and were then transferred to polyvinylidene fluoride (PVDF) membranes (Bio-Rad Laboratories, Inc., Hercules, CA, USA). The membranes were then rinsed with TBS-0.05% Tween-20 (TBST) and blocked with 5% fat-free dried milk solution at room temperature for 1.5 h. Primary antibodies: Rabbit anti-NF- $\kappa\text{B}$  (ab16502; 1:2,000), rabbit anti-phosphorylated (P)-NF- $\kappa\text{B}$  (ab86299; 1:1,000), mouse anti-I- $\kappa\text{B}\alpha$  (ab211340; 1:1,000) and mouse anti-GAPDH (ab8245; 1:2,000) (all Abcam, Cambridge, MA, USA) were added, and the membranes were incubated overnight at 4°C. Subsequently, the PVDF membranes were rinsed 3 times with TBST (5 min/wash) and were incubated with horseradish peroxidase-conjugated goat anti-mouse IgG (ab97023; 1:3,000 and 1:5,000 for anti-I- $\kappa\text{B}\alpha$  and anti-GAPDH primary antibodies, respectively) and goat anti-rabbit IgG (ab6721; 1:5,000 and 1:3,000 for anti-NF- $\kappa\text{B}$  and anti-P-NF- $\kappa\text{B}$  primary antibodies, respectively) secondary antibodies from Abcam, prior to visualization by enhanced chemiluminescence. Experiments were repeated three times. Relative expression levels were determined using ImageJ software, version 2.1.4.7 (National Institutes of Health, Bethesda, MD, USA) and were normalized to loading controls.

**RT-qPCR.** Total RNA was extracted from C2C12 cells in the control group, 0.125 Hz stretch group and NF- $\kappa\text{B}$  inhibitor + 0.125 Hz stretch group at day 4 using TRIzol<sup>®</sup> according to the manufacturer's protocol. RT was performed using an ABI 9700 PCR amplification machine (Applied Biosystems; Thermo Fisher Scientific, Inc.) and a miR RT kit (Takara Bio, Inc., Otsu, Japan). Briefly, aliquots (2  $\mu\text{g}$ ) of total RNA were transcribed into cDNA in a total volume of 20  $\mu\text{l}$  with 5X PrimeScript buffer, 1 mmol/l nucleotide mixture, 50 pmol/l primer, 100 U PrimeScript Reverse Transcriptase, and 20 U RNase inhibitor. The RT reaction was conducted at 37°C for 15 min, 85°C for 5 sec and 4°C prior to qPCR analysis. qPCR was performed in triplicate using a PCR detection system (Stratagene Mx3000P qPCR system; Agilent Technologies, Santa Clara, CA, USA). miRNA specific primer sequences are listed in Table I. All primers were synthesized by Guangzhou Vipotion Biotechnology Co., Ltd. (Guangzhou, China). qPCR was conducted using the Taq PCR kit (Takara Bio, Inc.) in a 20  $\mu\text{l}$  reaction volume containing 2  $\mu\text{l}$  miRNA RT product, 0.2  $\mu\text{l}$  1 U/ $\mu\text{l}$  Taq DNA polymerase, 10  $\mu\text{l}$  5X PCR Buffer, 0.6  $\mu\text{l}$  5  $\mu\text{M}$  primer and deionized water. Reaction conditions were as follows: Initial denaturation at 95°C for 3 min, 40 cycles of 95°C for 12 sec and 62°C for 60 sec, followed by an extension step at 72°C for 1 min. The relative miRNA expression levels were calculated using the comparative threshold cycle  $2^{-\Delta\Delta\text{Ct}}$  method (22).

**Statistical analysis.** Data were analyzed by one-way analysis of variance and the post-hoc Bonferroni test was used for pair-wise comparisons. Statistical analyses were performed using SPSS 19.0 (IBM SPSS, Armonk, NY, USA) and the data were presented as the mean  $\pm$  standard deviation, error bars were representative of three independent experiments.  $P < 0.05$  was considered to indicate a statistically significant difference.

Table I. miR specific primer sequences.

Primer name	Sequence (5'-3')
mmu-mir-500	AAUGCACCUGGGCAAGGGUUCA
mmu-mir-1934	AGGAUGACGGUGGGGCGUGGUGA
mmu-mir-31	AGGCAAGAUGCUGGCAUAGCUG
mmu-mir-378a	ACUGGACUUGGAGUCAGAAGG
mmu-mir-331	GCCCCUGGGCCUAUCCUAGAA
mmu-mir-5097	GUUCAUGUCCUGUUCAGGCGCCA
mmu-mir-1941	AGGGAGAUGCUGGUACAGAGGCUU
miR, microRNA.	

## Results

**Alterations to the miRNA expression profile during mechanical stretch-induced myoblast proliferation.** To investigate the effects of various stretch frequencies on myoblast viability, the CCK-8 assay was performed 1, 2, 3 and 4 days after the cells underwent cyclic stretching. At day 3, the mean OD value of the 0.125 Hz stretch group (10% deformation) was significantly higher compared with in the control group; 1.65 vs. 0.71 respectively ( $P < 0.05$ ; Fig. 1A). At day 4, the mean OD value of the 0.125 Hz stretched cells remained higher compared with the control cells; 1.63 vs. 0.81 ( $P < 0.05$ ). Furthermore, cell cycle progression differed between the control cells and the 0.125 Hz stretched myoblasts (10% deformation) at day 4. As shown in Fig. 1B, the number of cells in the proliferative (S or G<sub>2</sub>) phase was markedly higher in the 0.125 Hz stretch group compared with in the control group. At day 4, the mean DPI in the 0.125 Hz stretch group was significantly increased compared with in the control cells (Fig. 1B). These results suggest that 10% deformation cyclic strain at 0.125 Hz can enhance myoblast viability and stimulate myoblast entry into the proliferative phase, thus promoting C2C12 cell proliferation. Conversely, 10% deformation cyclic strain at 0.5 Hz markedly inhibited C2C12 cell viability at day 4 compared with the control (Fig. 1A). Therefore, 10% deformation cyclic mechanical stretch at 0.125 Hz were considered the appropriate stretch conditions.

Alterations to the miRNA profile between the control group and the 0.125 Hz mechanical stretch group were determined by high-throughput sequencing. The hierarchical clustering analysis of differentially expressed miRNAs is presented in Fig. 2. Within the miRNA expression profiles, 10 miRNAs exhibited  $>2$ -fold variation in the 0.125 Hz mechanical stretch group compared with in the control group. Seven (miR-500, -1934, -31, -378a, -3473b, -331 and -5097) of which were downregulated, whereas three (miR-340, -449c and -1941) were upregulated in the mechanical stretch group (Fig. 3A). Among them, miR-500, miR-1934, miR-31, miR-378a, miR-331, miR-5097 and miR-1941, were validated through the RT-qPCR method (Fig. 3B).

**Differentially expressed miRNAs induced by cyclic mechanical stretch are reversed by NF- $\kappa\text{B}$  inhibition in C2C12 myoblasts.** The present study aimed to explore the association

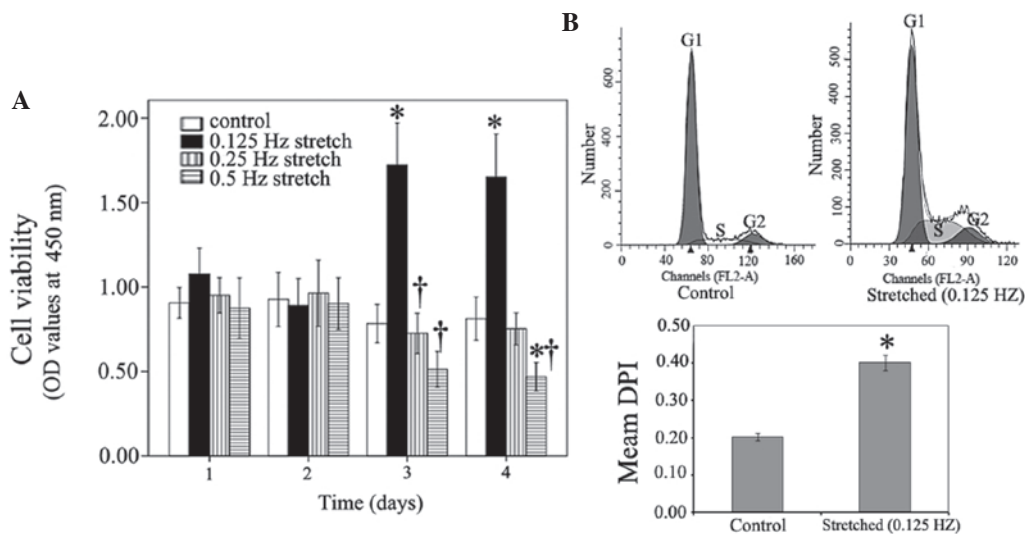


Figure 1. Appropriate mechanical stretch promotes myoblast proliferation. (A) Effects of mechanical stretch frequency on the viability of C2C12 cells, as detected by Cell Counting kit-8 assay. (B) Flow cytometric analysis of cell cycle progression in 0.125 Hz stretched myoblasts and control cells at day 4. \* $P < 0.05$  vs. the control group. † $P < 0.05$  vs. the 0.125 Hz group.  $n = 3$ . OD, optical density; DPI, DNA proliferation index.

between NF- $\kappa$ B activation and miRNA expression during stretch-induced myoblast proliferation. As shown in Fig. 4A, 0.125 Hz mechanical stretch increased the expression levels of NF- $\kappa$ B p65 and P-NF- $\kappa$ B p65, whereas the expression levels of the inhibitory factor I- $\kappa$ B $\alpha$  were decreased compared with the control group ( $P < 0.05$ ). However, treatment with 2.5 or 5  $\mu$ M NF- $\kappa$ B inhibitor BAY inhibited NF- $\kappa$ B p65 and P-NF- $\kappa$ B p65 expression, and increased the expression levels of I- $\kappa$ B $\alpha$  compared with the stretch group. Treatment with 5  $\mu$ M BAY resulted in a more significant inhibition of NF- $\kappa$ B activation ( $P < 0.05$ ). These results suggest that NF- $\kappa$ B is activated in the process of stretch-mediated myoblast proliferation, and the NF- $\kappa$ B inhibitor BAY can effectively inhibit activation of NF- $\kappa$ B p65 in stretched C2C12 cells.

Compared with 2.5  $\mu$ M BAY, 5  $\mu$ M BAY displayed an increased inhibitory effect on NF- $\kappa$ B activation. Therefore, 5  $\mu$ M BAY was selected as the dose for NF- $\kappa$ B inhibition in subsequent experiments regarding miRNA expression. As shown in Fig. 4B, 0.125 Hz stretch-induced downregulation of miR-500, miR-1934, miR-31 and miR-378a in C2C12 cells was reversed by 5  $\mu$ M BAY. Furthermore, 0.125 Hz stretch-induced reduced expression of miR-331 and miR-5097 was significantly reversed by 5  $\mu$ M BAY ( $P < 0.05$ ). Conversely, the 0.125 Hz mechanical stretch-induced expression of miR-1941 was significantly reduced following treatment with 5  $\mu$ M BAY ( $P < 0.05$ ). These data indicate that 5  $\mu$ M BAY can reverse mechanical stretch-induced miRNA expression alterations.

The effects of the NF- $\kappa$ B inhibitor BAY were also determined on mechanical stretch-induced myoblast proliferation, as assessed by CCK-8 assay and FCM. The 0.125 Hz mechanical stretch-induced increased OD value was significantly decreased in C2C12 cells following treatment with 2.5  $\mu$ M BAY on days 2, 3 and 4. Treatment with 5  $\mu$ M BAY was able to inhibit stretched myoblast viability on the first day, and sustain the inhibitory effect between days 2 and 4 ( $P < 0.05$ ; Fig. 5A). FCM revealed that stretched C2C12 cells treated with 5  $\mu$ M BAY were arrested in G<sub>1</sub> phase at day 4,

and the mean DPI in stretched C2C12 cells treated with 5  $\mu$ M BAY was significantly decreased compared with the control cells ( $P < 0.05$ ; Fig. 5B). These results suggest that the NF- $\kappa$ B inhibitor BAY may inhibit stretch-induced myoblast proliferation.

The present study used another NF- $\kappa$ B inhibitor, PDTC, to confirm the effects of NF- $\kappa$ B inhibition on stretched myoblasts. The results, as shown in Fig. 6, revealed that 10  $\mu$ M PDTC elicited similar effects to BAY. Treatment of C2C12 cells with 10  $\mu$ M PDTC inhibited activation of NF- $\kappa$ B, reversed alterations in stretch-induced miRNA expression and inhibited stretch-induced myoblast proliferation.

## Discussion

Variations in the duration, magnitude and frequency of mechanical stretch have discrepant influences on myoblasts. Appropriate external mechanical stimulation can promote the proliferation of myoblasts cultured *in vitro* (23,24). In addition, it has previously been reported that mechanical stretch (15% strain) may induce conspicuous injury to muscle cells (25), whereas cyclic tensile stretch (10% strain) at 1 Hz for 1 h followed by 23 h of relaxation could enhance myoblast proliferation (24). The results of the present study revealed that only 0.125 Hz mechanical stretch could promote myoblast proliferation when C2C12 myoblasts were exposed to 10% tensile cyclic strain at various frequencies (0.125, 0.25 and 0.5 Hz). Similar findings have been reported in human bladder smooth muscle cells cultured *in vitro*; 5% equibiaxial stretch at 0.1 Hz promoted smooth muscle cell proliferation and decreased apoptotic rate (25).

In the present study, seven mechanical stretch-associated miRNAs were detected in C2C12 cells: MiR-500, miR-1934, miR-31, miR-378a, miR-331, miR-5097 and miR-1941, which were identified using high-throughput sequencing and RT-qPCR analysis. Among them, miR-500 has previously been detected in the dorsal root ganglia of rats and is thought to be involved in nerve reconstruction (26). Furthermore, miR-31

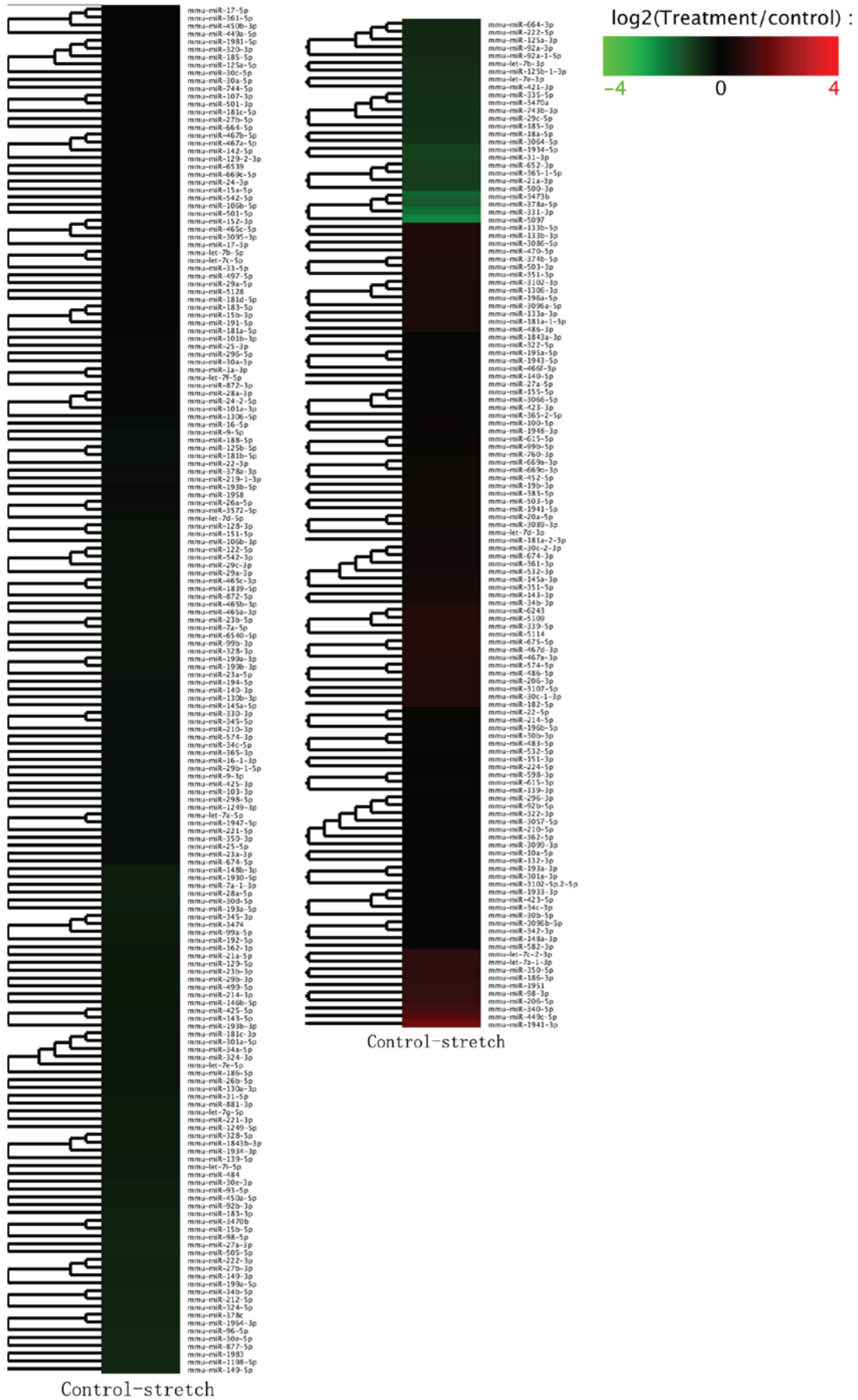


Figure 2. Hierarchical clustering analysis of the log<sub>2</sub> values of differentially expressed miRNAs between the 0.125 Hz mechanical stretch group and the control group. Green color indicates downregulated expression in the mechanical stretch group compared with in the control group; red color indicates upregulated expression in the mechanical stretch group compared with in the control group; black color indicates no significant difference in the expression between the two groups. miR, microRNA.

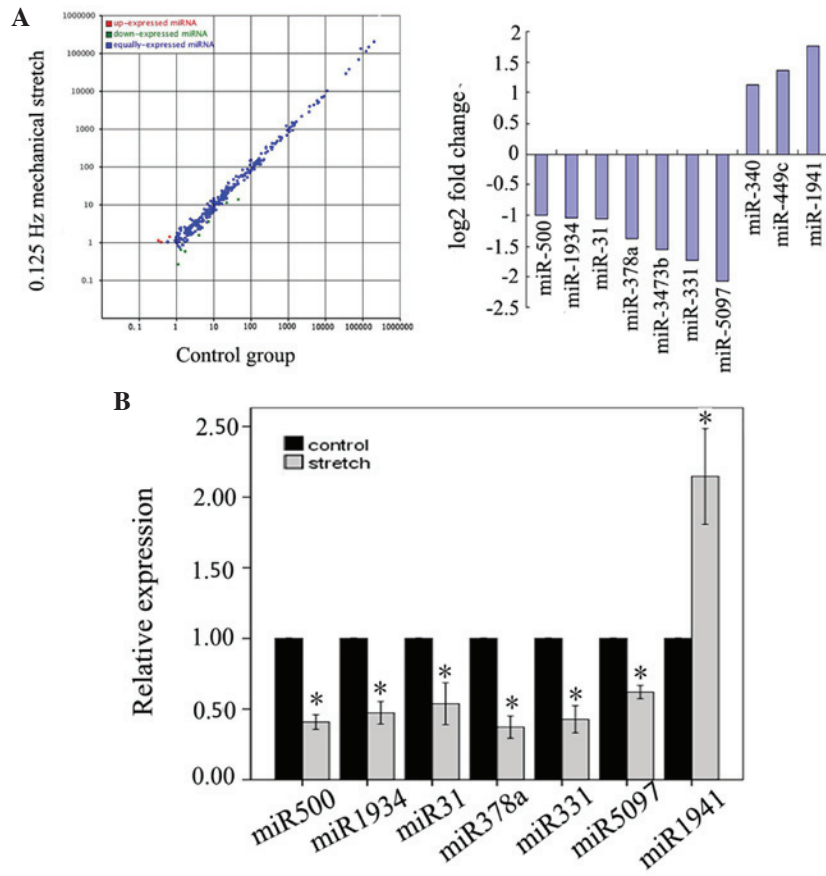


Figure 3. (A) Differentially expressed miRNAs in the 0.125 Hz mechanical stretch group compared with the control group. (B) Relative expression levels of miR-500, miR-1934, miR-31, miR-378, miR-331, miR-5097 and miR-1941 in the control and 0.125 Hz mechanical stretch groups. The expression levels in the 0.125 Hz mechanical stretch group were normalized to the control group. \*P<0.05 vs. the control group. n=3. miR, microRNA.

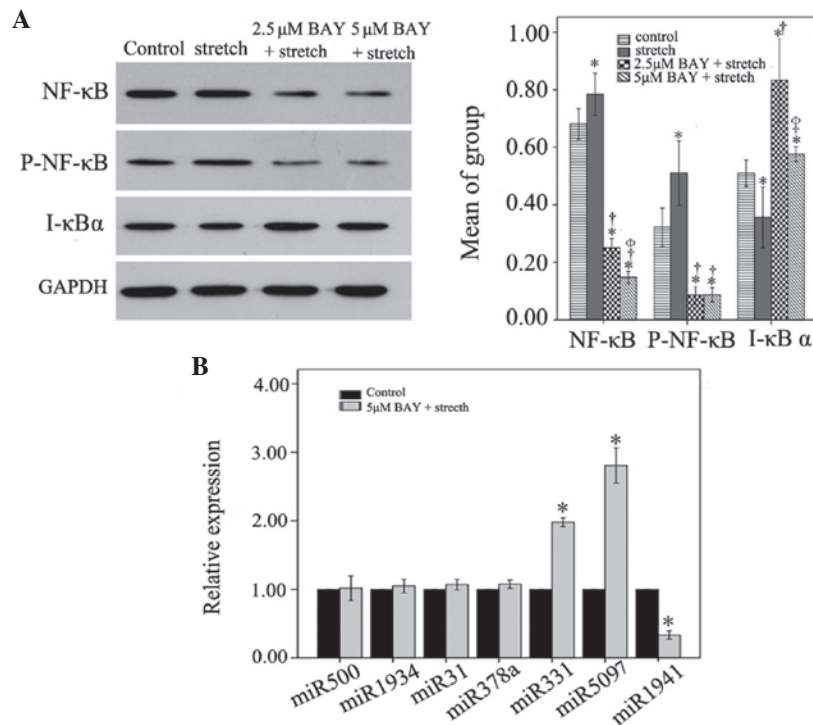


Figure 4. (A) Effects of NF- $\kappa$ B inhibition on the expression levels of NF- $\kappa$ B p65, P-NF- $\kappa$ B p65, and I- $\kappa$ B $\alpha$  proteins in 0.125 Hz stretched C2C12 cells. \*P<0.05 vs. the control group. #P<0.05 vs. the stretch group. P#P<0.05 vs. the 2.5  $\mu$ M BAY group. n=3. (B) Relative expression levels of miR-500, miR-1934, miR-31, miR-378, miR-331, miR-5097 and miR-1941 in control and 5  $\mu$ M BAY + mechanical stretch groups. Expression levels in the 5  $\mu$ M Bay + 0.125 Hz mechanical stretch group were normalized to the control group. \*P<0.05 vs. the control group. n=3. miR, microRNA; NF- $\kappa$ B, nuclear factor- $\kappa$ B; P-, phosphorylated; I- $\kappa$ B $\alpha$ , inhibitor of NF- $\kappa$ B; BAY, BAY11-7082.

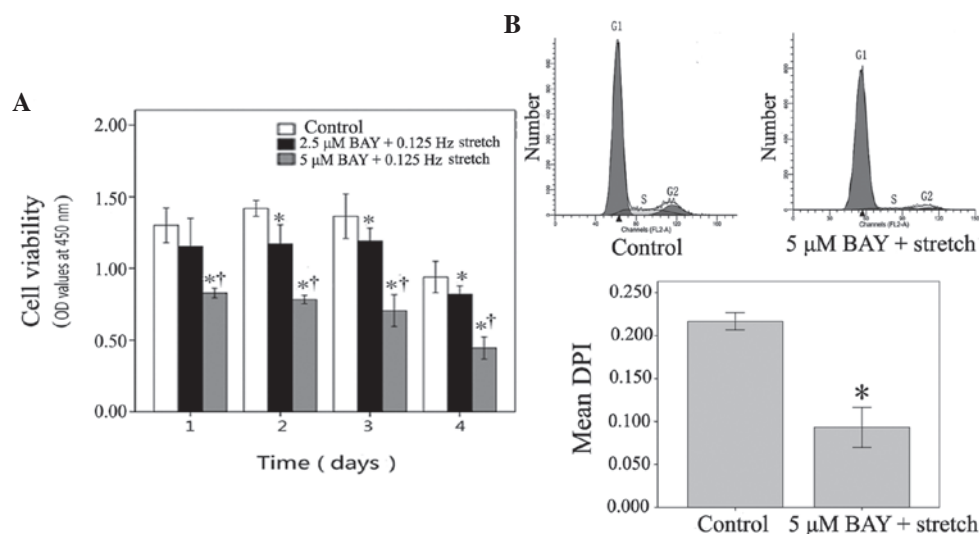


Figure 5. NF- $\kappa$ B inhibitor BAY inhibits stretch-induced C2C12 cell proliferation. (A) Effects of NF- $\kappa$ B inhibition on the viability of 0.125 Hz-stretched C2C12 cells, as detected by Cell Counting kit-8 assay. \* $P$ <0.05 vs. the control group. † $P$ <0.05 vs. the 2.5  $\mu$ M BAY group. (B) Flow cytometric analysis of cell cycle progression in stretched myoblasts treated with 5  $\mu$ M BAY and control cells at day 4. \* $P$ <0.05 vs. the control group. n=3. NF- $\kappa$ B, nuclear factor- $\kappa$ B; BAY, BAY11-7082; DPI, DNA proliferation index.

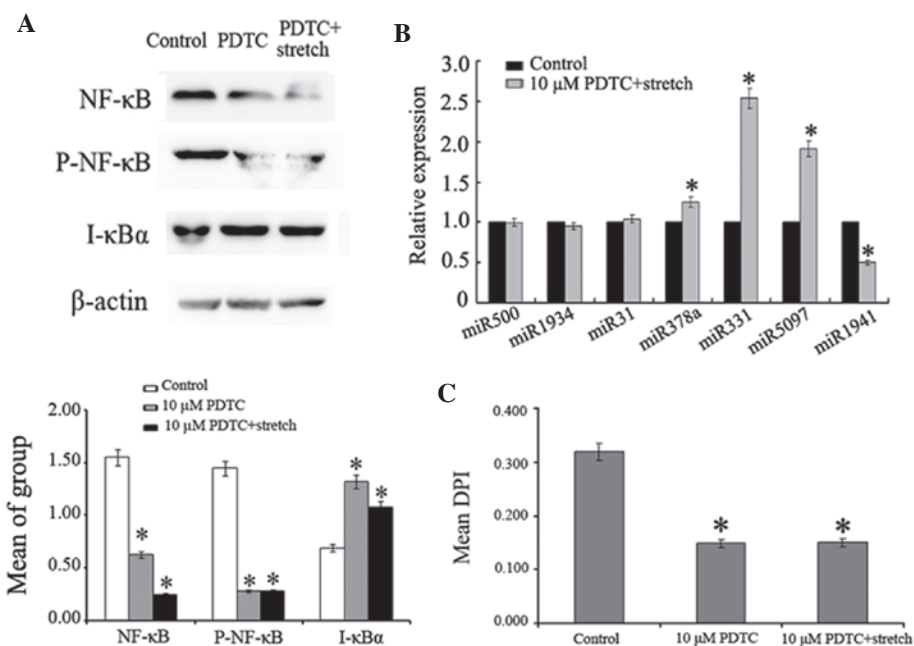


Figure 6. (A) Effects of 10  $\mu$ M PDTC on the expression levels of NF- $\kappa$ B p65, P-NF- $\kappa$ B p65 and I- $\kappa$ B $\alpha$  proteins in 0.125 Hz stretched C2C12 cells, as detected by western blotting. (B) Relative expression levels of miR-500, miR-1934, miR-31, miR-378, miR-331, miR-5097 and miR-1941 in control and 10  $\mu$ M PDTC + mechanical stretch groups. (C) Effects of 10  $\mu$ M PDTC on 0.125 Hz stretch-induced myoblast proliferation. \* $P$ <0.05 vs. the control group. n=3. miR, microRNA; NF- $\kappa$ B, nuclear factor- $\kappa$ B; P-, phosphorylated; I- $\kappa$ B $\alpha$ , inhibitor of NF- $\kappa$ B; PDTC, pyrrolidine dithiocarbamate; DPI, DNA proliferation index.

is able to promote the expression of myogenic factor 5 and myoblast proliferation. Targeted inhibition of miR-31 has been reported to be an effective therapeutic strategy for dystrophin rescue in human Duchenne muscular dystrophy (27,28). In addition, miR-378 may suppress proliferation of myoblasts by modulating the mitogen-activated protein kinases (MAPK) pathway and bone morphogenetic proteins (29,30). The over-expression of miR-378 promotes myoblast differentiation by strengthening MyoD activity (30). Increased expression of miR-331 has also been detected during myoblast differentiation (31). These findings are similar to the results of the

present study; the expression levels of miR-31, miR-378 and miR-331 were reduced during the process of 10% strain, 0.125 Hz mechanical stretch-induced myoblast proliferation. In addition, to the best of our knowledge, miR-1941, miR-1934, miR-5097 and miR-500 may be considered novel candidates for stretch-induced myogenesis, which have not been reported previously.

The present study further explored the effects of NF- $\kappa$ B, which participates in the enhanced proliferative process elicited by mechanical stretch. There are several signaling pathways associated with mechanical stretch in skeletal muscle cells,

including protein kinase C (32), MAPK (33), phosphatidylinositol-3 kinase/Akt (34), NF- $\kappa$ B (12), calcineurin (35) and nitric oxide synthase (36). However, findings published over the past few years have suggested that these signaling pathways have a close connection and serve crosstalk roles in inflammatory and immune responses (37-39). Notably, MAPK appears to be an upstream regulator of the NF- $\kappa$ B signaling pathway, which can successfully induce NF- $\kappa$ B-dependent myogenesis (8-11). In the present study, the NF- $\kappa$ B signaling pathway was activated by mechanical stretch, as determined by the markedly increased protein expression levels of NF- $\kappa$ B p65 and P-NF- $\kappa$ B p65. Following treatment with NF- $\kappa$ B inhibitors, translocation of NF- $\kappa$ B in stretch-loaded myoblasts was inhibited, which was accompanied by the reversal of miRNA expression and the suppression of proliferation in C2C12 myoblasts. Therefore, these results indicated that the NF- $\kappa$ B signaling pathway is activated and exerts direct effects in stretch-induced myoblasts. These results are concordant with the findings reported by Kumar *et al*, which indicated that NF- $\kappa$ B is indispensable for stretch-induced proliferation of C2C12 cells (12).

The results of the present study identified a link between NF- $\kappa$ B and miRNA expression. The expression levels of miR-500, miR-1934, miR-31, miR-378a, miR-331 and miR-5097 were downregulated in myoblasts following mechanical stretch, whereas NF- $\kappa$ B inhibition increased the expression of these miRNAs. Conversely, the upregulation of miR-1941 was markedly reduced by NF- $\kappa$ B inhibition in mechanically stretched C2C12 cells. Similarly, bioinformatics analysis in a previous study suggested the presence of an NF- $\kappa$ B binding element in the promoter region of miR-26a; inhibition of NF- $\kappa$ B in cardiac fibroblasts increased miR-26a expression, whereas overexpression of miR-26a inhibited NF- $\kappa$ B activity, thus indicating a feedback loop in cardiac fibroblasts (40). The results of the present study are the first, to the best of our knowledge, to reveal that miRNA expression is NF- $\kappa$ B-dependent during stretch-induced myoblast proliferation, thus suggesting a possible mechanism regarding the transformation of external mechanical stimuli into intracellular signals.

The present study used two structurally different NF- $\kappa$ B inhibitors in order to confirm the effects of the NF- $\kappa$ B signaling pathway on stretched myoblasts. BAY is a specific inhibitor of NF- $\kappa$ B, due to its inhibitory effect on the activation of NF- $\kappa$ B and the phosphorylation of I- $\kappa$ B $\alpha$  (41). PDTC is able to inhibit activation of NF- $\kappa$ B specifically by suppressing the release of the inhibitory subunit I- $\kappa$ B from the latent cytoplasmic form of NF- $\kappa$ B (42-44). The results of the present study revealed that the two NF- $\kappa$ B inhibitors were able to inhibit activation of NF- $\kappa$ B, reverse stretch-induced miRNA profile alterations in C2C12 myoblasts, and suppress stretch-induced myoblast proliferation. Accordingly, these data confirmed the interaction between miRNAs and the NF- $\kappa$ B signaling pathway during stretch-induced myogenesis.

In conclusion, cyclic stretch (10% deformation; 0.125 Hz) was able to activate the NF- $\kappa$ B signaling pathway, reduce the expression levels of miR-500, -1934, -31, -378, -331 and -5097, increase the expression of miR-1941, and promote proliferation of C2C12 cells. However, treatment with NF- $\kappa$ B inhibitors inhibited NF- $\kappa$ B activation in mechanically stretched C2C12

cells, which was accompanied by reversed expression of miRNAs and suppressed cell proliferation. These results demonstrated that cyclic stretch-mediated cell proliferation and miRNA profile alterations in C2C12 myoblasts occur via activation of the NF- $\kappa$ B signaling pathway, thus providing a possible mechanism during stretch-induced myogenesis.

### Acknowledgements

The present study was supported by the National Natural Science Foundation of China (grant no. 31100700) and the Guangdong Medical Research Foundation (grant no. A2015412).

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