

H3K9me2 regulates early transcription factors to promote mesenchymal stem-cell differentiation into cardiomyocytes

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Abstract. Studies have shown that histone H3 at lysine 9 (H3K9me2) is an important epigenetic modifier of embryonic development, cell reprogramming and cell differentiation, but its specific role in cardiomyocyte formation remains to be elucidated. The present study established a model of 5-Azacytidine-induced differentiation of rat bone mesenchymal stem cells (MSCs) into cardiomyocytes and, on this basis, investigated the dimethylation of H3K9me2 and its effect on cardiomyocyte formation by knockdown of H3K9me2 methylase, euchromatic histone-lysine N-methyltransferase 2 (G9a) and H3K9me2 lysine demethylase 3A (KDM3A). The results demonstrated that, in comparison with the normal induction process, the knockdown of G9a could significantly reduce the H3K9me2 level of the MSCs in the induced model. Reverse transcription-quantitative (RT-q) PCR demonstrated that the expression of cardiac troponin T (cTnT) was significantly increased. In addition, flow cytometry demonstrated that the proportion of cTnT-positive cells was significantly increased on day 21. With the knockdown of KDM3A, the opposite occurred. In order to explore the specific way of H3K9me2 regulating cardiomyocyte formation, western blotting and RT-qPCR were used to detect the expression of key transcription factors including GATA binding protein 4 (GATA-4), NK2 Homeobox 5 (Nkx2.5) and myocyte enhancer

factor 2c (MEF2c) during cardiomyocyte formation. The decrease of H3K9me2 increased the expression of transcription factors GATA-4, Nkx2.5 and MEF2c in the early stage of myocardial development while the increase of H3K9me2 inhibited the expression of those transcription factors. Accordingly, it was concluded that H3K9me2 is a negative regulator of cardiomyocyte formation and can participate in cardiomyocyte formation by activating or inhibiting key transcription factors of cardiomyocytes, which will lay the foundation for the optimized induction efficiency of cardiomyocytes in *in vitro* and clinical applications.

Introduction

Bradyarrhythmias are various life-threatening conditions such as sinus syndrome and atrioventricular blockage and are caused by cardiac pacing or conduction dysfunction (1). Implanted electronic pacemakers are the preferred treatment for such diseases (2). However, electronic pacemakers still produce adverse reactions such as infection, metal allergy, electrode dislocation, electronic interference and lack of response to neurohormones. Therefore, biological pacemakers provide new options for the treatment of arrhythmia. Bone marrow mesenchymal stem cells (MSCs) are among the main sources of cardiac repair and regenerative cell therapy because these cells are readily available and have little inherent immunogenicity to any adverse immune response (3). *In vitro*, 5-Azacytidine (5-AZA) was found to induce the differentiation of a variety of the MSCs into cardiomyocytes (4-7). *In vivo*, the transplantation of MSCs has been shown to improve cardiac function in damaged hearts by myocardial cell regeneration and paracrine effects that promote angiogenesis, reduce cardiac fibrosis and prevent host myocardial cell apoptosis (8). However, just a small percentage of the MSCs express cardiac-specific proteins during differentiation, which significantly limits their potential clinical application (3,9). Induction efficiency can be increased by transfection of key genes and transcription factors or simulating specific myocyte growth microenvironment (10-12). However, these methods have a number of

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disadvantages including complicated operating procedures, high costs and a low induction efficiency.

Dimethylation of histone H3 at lysine 9 (H3K9me2) has, in recent years, been found to be closely related to the process of cell differentiation, proliferation and reprogramming (13). Following H3K9me2 methylase, euchromatic histone-lysine N-methyltransferase 2 (G9a) was inhibited and H3K9me2 was decreased, the levels of cardiac transcription factors GATA binding protein 4 (GATA-4), NK2 Homeobox 5 (Nkx2.5) and myocardial protein produced by Wnt11 treatment were 2.6-5.6 times greater compared with those of the untreated group (14). Zhang *et al.* (15) found that H3K9me2 lysine demethylase 3A (KDM3A) could control pathological cardiac hypertrophy and fibrosis. In their study, JIB-04 inhibits KDM3A and suppresses the transcription of fibrotic genes that overlap with genes down-regulated in Kdm3A-KO mice vs. wild-type controls. This suggests that KDM3A may be involved in cardiac development.

As part of the present study, in a system for the 5-AZA induction of MSCs into cardiomyocytes, the expression of H3K9me2 was affected by the knockdown of G9a and KDM3A, resulting in a change in differentiation efficiency during induction. The role of H3K9me2 in the differentiation of the MSCs into cardiomyocytes was identified, thus optimizing the induction efficiency of cardiomyocytes *in vitro* and laying a foundation for clinical applications.

Materials and methods

Culture, expansion and characterization of rat MSCs. A total of 20 healthy male and 20 female Wistar rats (weight, 150-200 g; age, 6-8 weeks) were obtained from Yangzhou University and three rats were used in each of three individual experiments. Rats received standard care under a 12 h dark/light cycle (25°C with humidity of 60%) and given a free access to food and water. Before the experiment, adaptive feeding was carried out for one week. The animal's health and behavior were monitored daily until mortality. All procedures involving the care and use of animals conformed to the guidelines of Animal Research: Reporting of *in vivo* Experiments (ARRIVE guidelines 2.0; <https://arriveguidelines.org/arrive-guidelines>) and were approved by the animal care guidelines of the Laboratory Animal Management and Experimental Animal Ethics Committee of Yangzhou University (approval no. 201903478). Bone mesenchymal stem cells (MSCs) were obtained as described by Huang *et al.* (16) and Nippert *et al.* (17). All rats were anesthetized with 20% sodium urethane (0.2 g/ml; 1.0 g/kg intraperitoneal injection) and subsequently sacrificed by cervical dislocation. Animal mortality was verified by ascertaining cardiac and respiratory arrest. MSCs were extracted from the tibias and femurs. The cell suspension from three rats was mixed together and the cells were dispersed by pipetting, following which they were placed in a centrifuge at 1,100 x g for 4 min at 37°C. The supernatant and adipose tissue were removed. The cell suspension was transferred to a 15 ml centrifuge tube containing 5 ml Percoll (1.073 g/ml; Sigma-Aldrich; Merck KGaA). Cells were dispersed by pipetting again and centrifuged at 1,500 x g for 30 min at 4°C. The mononuclear cells in the middle layer were obtained, washed three times with phosphate buffered saline (PBS) and then cultured in Dulbecco's Modified Eagle's

Medium (DMEM) (Thermo Fisher Scientific, Inc.) containing 10% fetal bovine serum (FBS; Biowest) at 37°C in a 5% CO₂ incubator. During the expansion and proliferation of MSCs, the culture medium was replaced every three days and cells were passaged once they reached 70% confluency. For some experiments, MSCs were cultured in the presence or absence of 1 μM BIX01294 (MedChemExpress) for 12 h.

For cell phenotypic characterization, MSCs of the third passage were incubated overnight at 4°C with mouse anti-CD29 (1:100; cat. no. sc-9970; Santa Cruz Biotechnology, Inc.), anti-CD44 (1:100; cat. no. sc-7297; Santa Cruz Biotechnology, Inc.) and anti-CD45 (1:100; cat. no. sc-1178; Santa Cruz Biotechnology, Inc.) and then with an appropriate secondary antibody conjugated to PE (1:100; cat. no. sc-516141; Santa Cruz Biotechnology, Inc.) for 1 h at room temperature. The cells were subsequently detected using a BD FACSCanto™ II flow cytometer (BD Biosciences). The results were analyzed and processed by FlowJo version 10.0 (FlowJo LLC).

Construction of G9a and KDM3A lentivirus interference vectors. According to the CDS-region sequences of G9a and KDM3A provided by National Center of Biotechnology Information (<https://www.ncbi.nlm.nih.gov>), three interference targets were designed. Meanwhile, an unrelated sequence was designed as a negative control (Table I). After the company (Genomeditech) synthesized the upstream and downstream sequences, an oligo double chain was formed by the annealing process and then three recombinant interference plasmids were obtained by means of connection with pGMLV-SC5 RNA interference vectors through *Bam*HI and *Eco*RI double-enzyme digestion, respectively. These were called short hairpin (sh)1-G9a, sh2-G9a, sh3-G9a, sh1-KDM3A, sh2-KDM3A, sh3-KDM3A and sh-negative control (NC). The recombinant interfering plasmids were then subjected to sequencing for further verification. Following successful sequencing, the vector with the strongest interference activity was delivered to Shanghai Jiman Co., Ltd. for lentivirus parcel (titer: 2.5x10⁸ TU/ml) after verification of activity.

Cell transfection. MSCs were cultured in DMEM supplemented with 10% FBS and the cells were kept at 37°C and 5% CO₂ containing atmosphere. The cells were plated in 24-well plates, grown to a confluency of 70-80% and were transfected with 1 μg/plate shRNA-G9a or shRNA-KDM3A lentivirus, which were diluted in polybrene (Sigma-Aldrich; Merck KGaA). sh-NC were transfected with plasmids that were non-targeting and indistinguishable for KDM3A and G9A. The cells that were not treated were used as controls. Following treatment, the cells were observed daily through the use of a fluorescent microscope (magnifications, x100, 200 and 400; TE2000; Nikon Corporation). After 72 h, the cells were collected for further experiments.

Cardiac differentiation induction. The MSCs were divided into four separate groups: i) MSCs group: MSCs were untreated; ii) control group: MSCs were transfected with control lentivirus; iii) shKDM3A group: MSCs were transfected with lentivirus encoding shRNA-KDM3A; and iv) shG9a group: MSCs were transfected with lentivirus encoding shRNA-G9a. All groups treated with 10 μM of 5-AZA (Sigma-Aldrich;

Table I. Primers used for plasmid construction.

Gene	Primer sequence (5'-3')
sh1-G9a	F: GATCCGCCTGTACTATGACGCGTACTCTCGAGAGTACGCGTCATAGTACAGGCTTTTTT R: AATTA AAAAAGCCTGTACTATGACGCGTACTCTCGAGAGTACGCGTCATAGTACAGGCG
sh2-G9a	F: GATCCGGTTTTGCACTGCAGCTCAATCCTCGAGGATTGAGCTGCAGTGCAAACCTTTTTT R: AATTA AAAAAGGTTTTGCACTGCAGCTCAATCCTCGAGGATTGAGCTGCAGTGCAAACCG
sh3-G9a	F: GATCCGCGGCTGCTCCAGGAGTTTAACTCGAGTTAACTCCTGGAGCAGCCGCTTTTTT R: AATTA AAAAAGCGGCTGCTCCAGGAGTTTAACTCGAGTTAACTCCTGGAGCAGCCGCG
sh1-KDM3A	F: GATCCGCAAGTCTTCTGAGAATAATGCTCGAGCATTATTCTCAGAAGACTTGCTTTTTT R: AATTA AAAAAGCAAGTCTTCTGAGAATAATGCTCGAGCATTATTCTCAGAAGACTTGCG
sh2-KDM3A	F: GATCCGCAGCCAATTCTCCACCTAACCTCGAGGTTAGGTGGAGAATTGGCTGCTTTTTT R: AATTA AAAAAGCAGCCAATTCTCCACCTAACCTCGAGGTTAGGTGGAGAATTGGCTGCG
sh3-KDM3A	F: GATCCGCAGGTGTCACTAGGCTTAATCTCGAGATTAAGCCTAGTGACACCTGCTTTTTT R: AATTA AAAAAGCAGGTGTCACTAGGCTTAATCTCGAGATTAAGCCTAGTGACACCTGCG
sh-NC	F: UUCUCCGAACGUGUCACGUTT R: ACGUGACACGUUCGGAGAATT

sh, short hairpin; NC, negative control (an unrelated sequence designed as a negative control). F, forward; R, reverse; G9a, euchromatic histone-lysine N-methyltransferase 2 KDM3A, H3K9me2 lysine demethylase 3A.

Merck KGaA). Following overnight incubation, the medium was aspirated and the cells were washed with PBS three times for three to five min. New culture medium was added and was then replaced every three days. Following 5-AZA treatment, cell morphology was observed daily using a phase-contrast microscope (DMIL-PH1; Leica Microsystems GmbH).

Western blotting. Cells from different treatment groups were lysed with RIPA lysis buffer (Applygen Technologies, Inc.) and samples containing 25–50 µg of total protein lysates (after Bradford estimation) were separated by 10% SDS-PAGE and transferred onto a PVDF membrane (EMD Millipore), which were then semi-dry blocked by TBS-0.05% Tween 20 containing 5% fetal calf serum (Gibco; Thermo Fisher Scientific, Inc.) for 1 h at room temperature. The membranes were incubated overnight at 4°C with primary antibodies targeted against: H3K9me2 (1:1,000; cat. no. 39239; Active Motif, Inc.), H3 (1:3,000; cat. no. H0164; Sigma-Aldrich; Merck KGaA), cTnT (1:3,000; cat. no. MABT368; Sigma-Aldrich; Merck KGaA), GATA-4 (1:3,000; cat. no. SAB4501128; Sigma-Aldrich; Merck KGaA), Nkx2.5 (1:5,000; cat. no. 13921-1-AP; ProteinTech Group, Inc.), MEF2c (1:3,000; cat. no. SAB2103534; Sigma-Aldrich; Merck KGaA) and β-actin (1:1,000; cat. no. SAB3500350; Sigma-Aldrich; Merck KGaA). The blots were then washed and incubated for 2 h with an appropriate secondary antibody (1:3,000, cat. no. A9169; 1:5,000, cat. no. AP162P; 1:5,000, cat. no. AP160P; all Sigma-Aldrich; Merck KGaA). The membranes were then washed and enhanced chemiluminescence (ECL) detection was performed using Pierce ECL kit (Thermo Fisher Scientific, Inc.). Densitometric analysis was performed using Image Lab software (version 3.0, Bio-Rad Laboratories, Inc.).

Total RNA isolation and reverse transcription-quantitative (RT-q) PCR. A RNeasy Mini kit (Qiagen GmbH) was used

Table II. Primers used for RT-qPCR.

Gene	Primer sequence (5'-3')
cTnT	F: TTCGACCTGCAGGAAAAGTT R: GTGCCTGGCAAGACCTAGAG
Nkx 2.5	F: ACCGCCCTACATTTTATCC R: GACAGGTACCGCTGTTGCTT
GATA4	F: TCTCACTATGGGCACAGCAG R: CCGAGCAGGAATTTGAAGAG
MEF2c	F: CTCCCCTGTGGACAGTCTGA R: CAGAGGGGCTTTCTCTGTCC
G9a	F: GCTACCATGACTGCGTTCTG R: TCCCGGCAGATGATCTTCTC
KDM3A	F: GGCAGTTCAAGCTCTTCTCG R: TGGACAGATGGGCTTCACAT
GAPDH	F: GGAAAGCTGTGGCGTGATGG R: GTAGGCCATGAGGTCCACCA

cTnT, cardiac troponin-T; Nkx2.5, NK2 Homeobox 5; GATA-4, GATA binding protein 4; MEF2c, myocyte enhancer factor 2c; G9a, euchromatic histone-lysine N-methyltransferase 2; KDM3A, H3K9me2 lysine demethylase 3A; F, forward; R, reverse.

to extract the total RNA from the MSCs of different treatment groups post-seeding, according to the manufacturer's instructions. cDNA was amplified by PCR through the use of a Qiagen PCR kit according to the manufacturer's instructions at 50°C for 30 min, 95°C for 3 min, 95°C for 15 sec and 60°C for 30 sec. GATA-4, Nkx2.5 and MEF2c expressions were detected by RT-qPCR. The 20 µl PCR amplification reaction included 2 µl cDNA, 10 µl SYBR Taq, 0.8 µl forward primer,

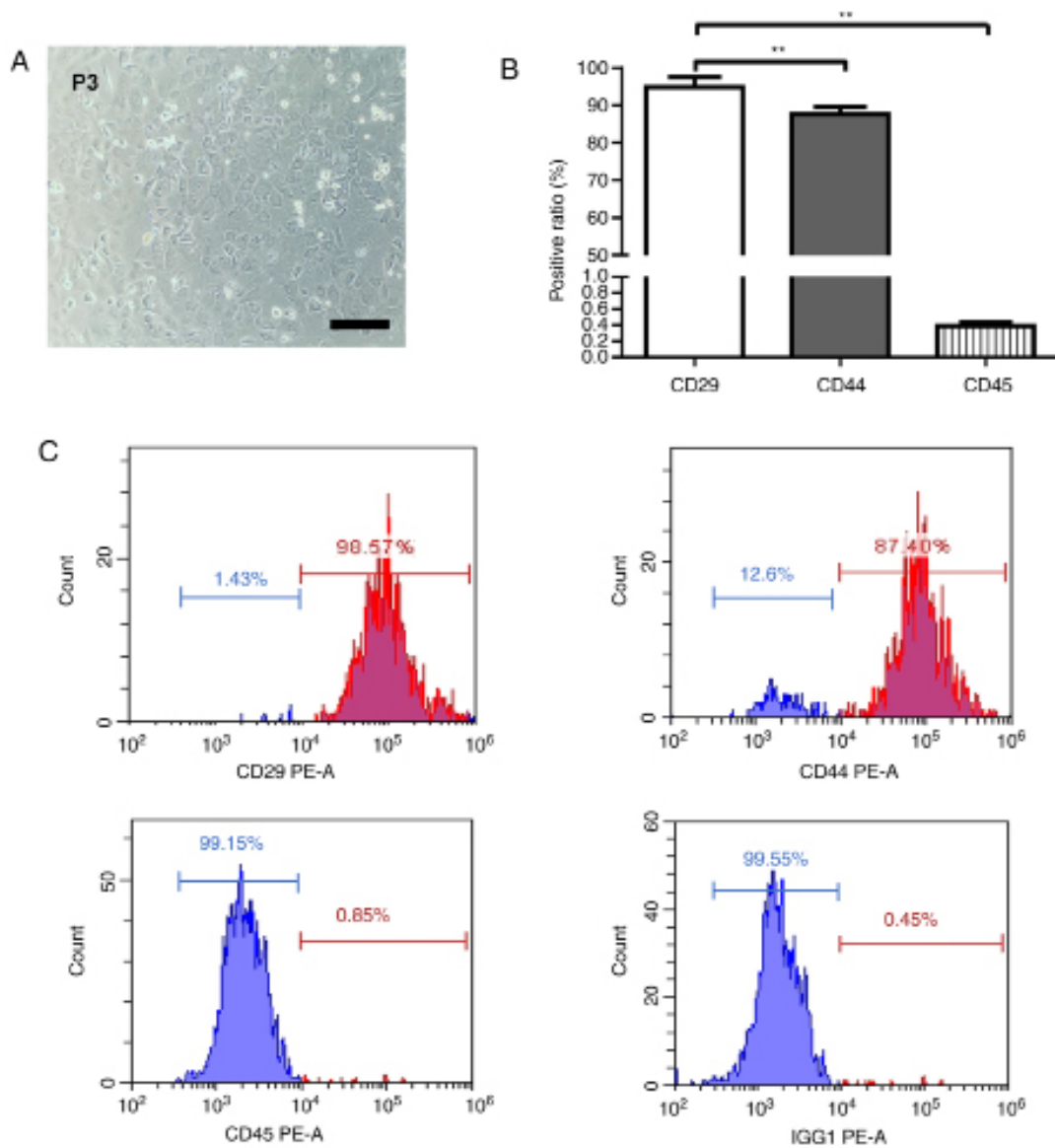


Figure 1. Morphology and characterization of isolated MSCs. (A) The morphology of the MSCs at passage 3 shows an adherent fibroblast-like shape and the initial formation of cell colonies after 10 days of culturation. Scale bar=50 μ m. (B) Histogram showing the average positive percentage of different antibodies from three independent experiments. Columns represent mean values and error bars represent standard deviation. ** $P < 0.01$ vs. CD45 expression. (C) Fluorescence-activated cell sorting analysis of CD29, CD44 and CD45 expression in MSCs. IgG1 was blank control. MSCs, rat mesenchymal stem cells; P3, passage 3.

0.8 μ l reverse primer, 0.4 μ l RoxII and 6 μ l double-distilled water. Subsequently, PCR reaction was achieved on the basis of the two-step procedure (95°C for 15 min; 95°C for 10 sec and 60°C for 32 sec) and procedure was repeated 40 times. Primers applied were shown in Table II. The PCR instrument for RT-qPCR was ABIPRISM 7500 (Applied Biosystems; Thermo Fisher Scientific, Inc.). Each experimental condition was repeated in triplicate. The relative mRNA quantities were determined using the $2^{-\Delta\Delta C_q}$ method (18).

Flow cytometry. Subsequent to treatment with 5-AZA, the expression of cTnT was detected in four groups on different days. Cells were incubated overnight at 4°C with anti-rat cTnT (1:200; cat. no. 15513-1-AP; ProteinTech Group, Inc.). Then, goat anti-rabbit fluorescein isothiocyanate (PE)-labeled mouse anti-rabbit IgG (1:500; cat. no. sc-3753; Santa Cruz Biotechnology, Inc.) was used as the secondary antibody and

incubated with the cells for 1 h at room temperature. The percentage of fluorescent protein-positive cells was detected by flow cytometry using a BD FACSCanto™ II flow cytometer (BD Biosciences). The results were analyzed and processed by FlowJo version 10.0 (FlowJo LLC).

Statistical analysis. The results were expressed as mean \pm standard error. Data were analyzed by one way ANOVA and Tukey's post hoc test using SPSS v23 (IBM Corp.). Unpaired t-test was used to evaluate significance between two groups. $P < 0.05$ was considered to indicate a statistically significant difference.

Results

Culture and characterizations of isolated rat MSCs. The obtained cells were cultured for 48 h. Under the microscope,

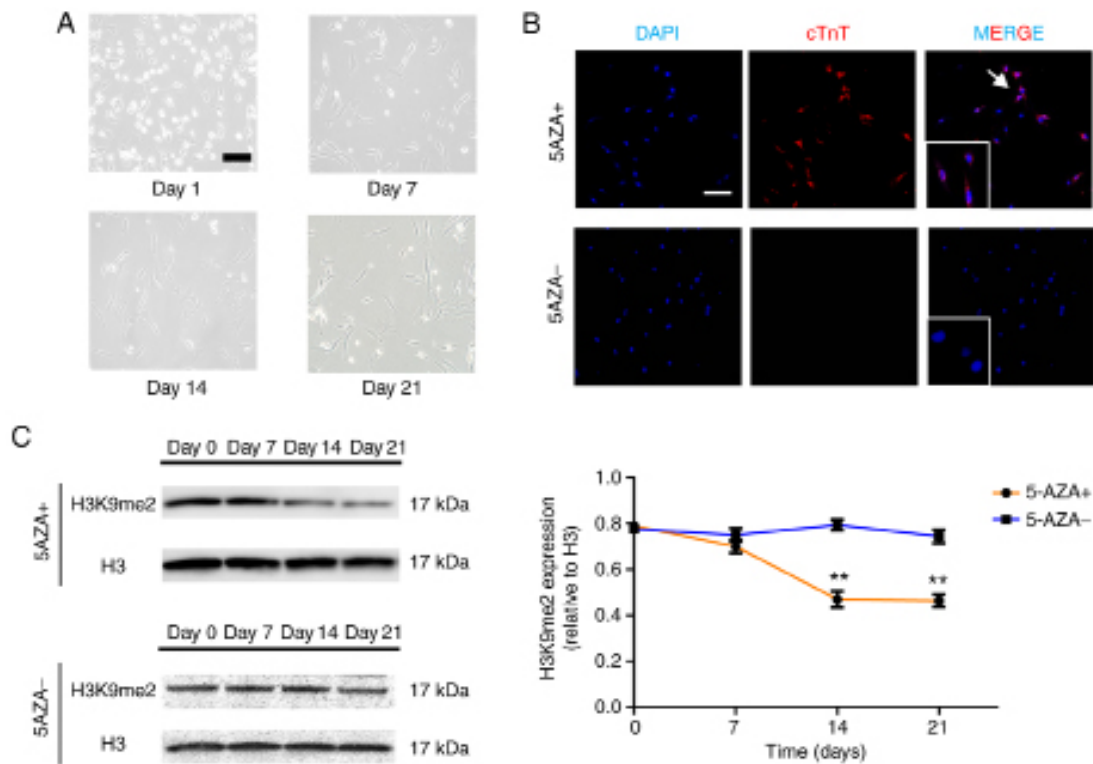


Figure 2. Morphological changes in the mesenchymal stem cells treated with 5-AZA. (A) Morphology of P3 MSCs treated with 5-AZA for 21 days. (B) Cardiomyocyte differentiation of the MSCs with 5-AZA treatment was detected by immunocytochemistry, which was demonstrated cTnT protein expression was upregulated on day 21. The white arrow indicates positive cells. The white box area indicates a partial enlargement of the image. Scale bars=50 μ m. (C) Following the treatment of the MSCs with 5-AZA, western blotting demonstrated the downregulation of H3K9me2 protein expression. H3 was used as an internal control. ** $P < 0.01$ vs. the untreated group. 5-AZA, 5-Azacytidine; P3, passage 3; MSCs, rat mesenchymal stem cells; cTnT, cardiac troponin-T; H3K9me2, histone H3 at lysine 9.

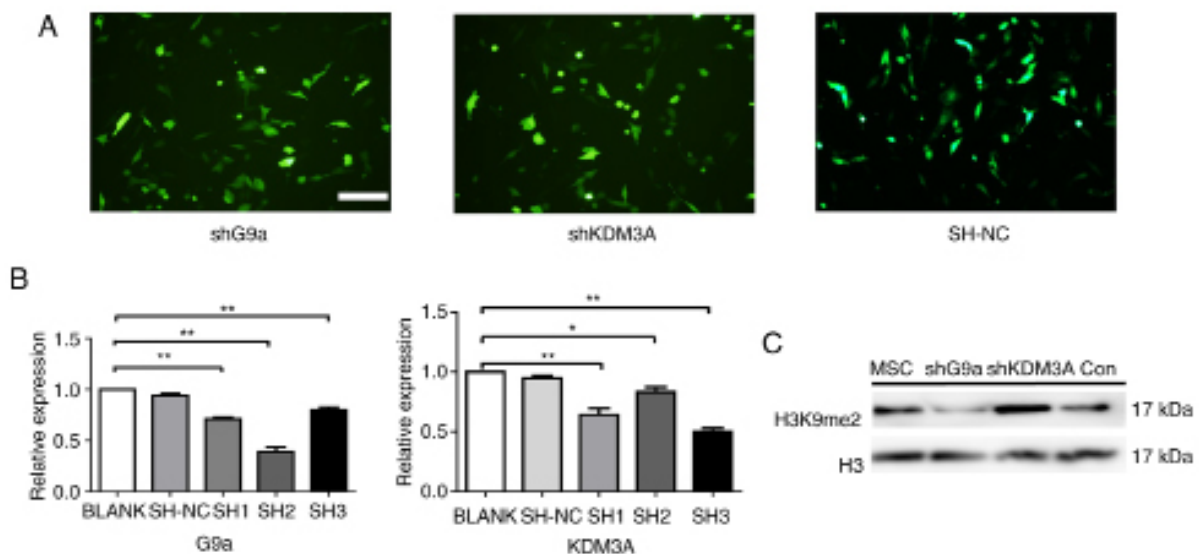


Figure 3. The screening and activity verification of interfering vectors. (A) Cell observation showing the fluorescence expression, confirming the vectors are transfected into the MSCs. Scale bar=50 μ m; (B) The relative mRNA expression of G9a and KDM3A with shRNA-mediated knockdown of the indicated factors in MSCs, confirming the most suitable interference vector. The mean \pm standard deviation is shown; n=3 independent experiments. * $P < 0.05$ and ** $P < 0.01$. (C) Western blotting showing protein levels of H3K9me2 in the interference of G9a and KDM3A at 72 h. H3 was used as an internal control. MSCs, rat mesenchymal stem cells; G9a, euchromatic histone-lysine N-methyltransferase 2; KDM3A, H3K9me2 lysine demethylase 3A; sh, short hairpin.

some adherent cells with fusiform growth could be seen and a few cells were polygonal. After 7 to 10 days, cell growth reached 80-90% confluence and was arranged in a spiral pattern (Fig. 1A). To characterize the MSCs, they were

analyzed with antibodies against CD44, CD29 and CD45 by means of flow cytometry. As shown in Fig. 1B and C, the majority of cells expressed CD29 and CD44 at moderate to high levels while these cells were negative for CD45 ($P < 0.01$).

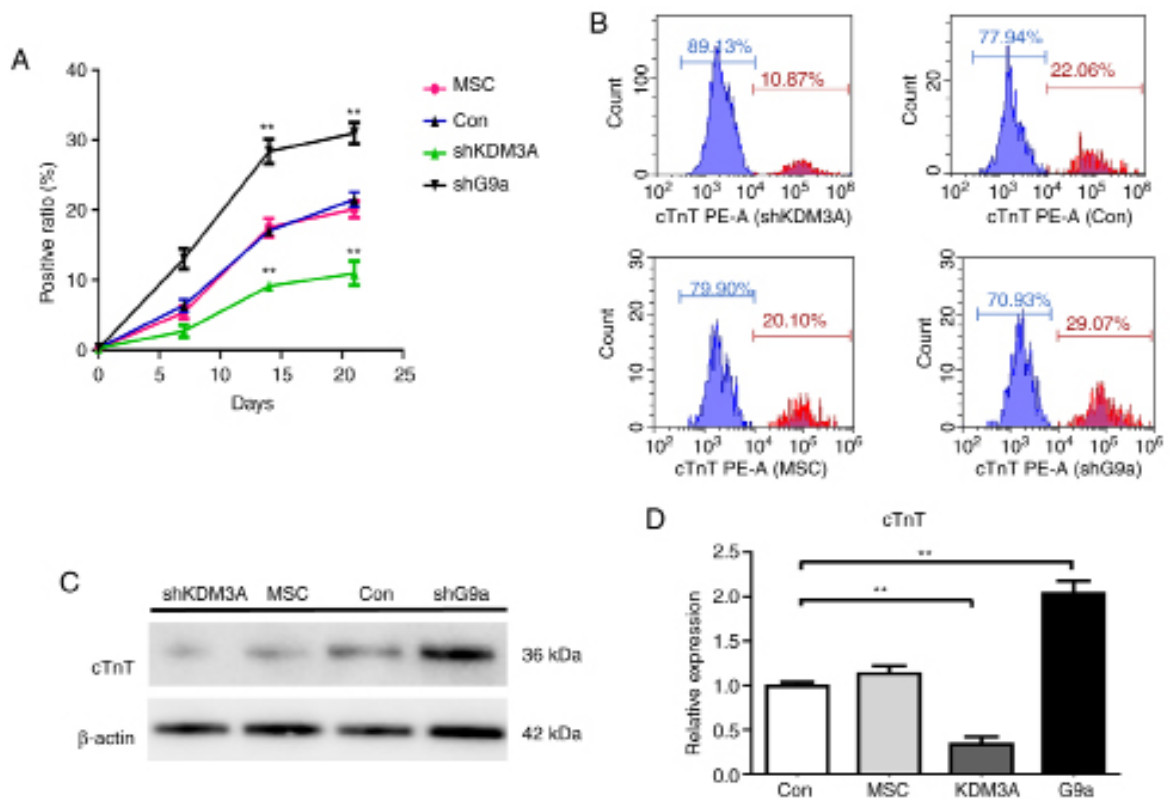


Figure 4. The levels of cTnT following 5-AZA treatment. (A) Flow-cytometry analysis demonstrated that cTnT-positive cells rate gradually increased in all groups following 5-AZA induction. The results on different days demonstrated that cTnT-positive cell rate in the G9a group was the highest and the KDM3A group was the lowest. The mean \pm standard deviation is shown; $n=3$ independent experiments. P by t-test. (B) Fluorescence intensity was detected by means of flow cytometry, which demonstrated that the proportion of cTnT-positive cells rate was $\sim 30\%$ in G9a knockdown group on day 21 and $\sim 11\%$ in the KDM3A knockdown group. (C) Cardiomyocyte differentiation of the MSCs with 5-AZA treatment was detected by western blot analysis, which demonstrated that cTnT protein levels were upregulated following G9A knockdown on day 21. β -actin was used as an internal control. (D) Cardiomyocyte differentiation of the MSCs with 5-AZA treatment was detected by reverse transcription-quantitative PCR, which demonstrated that G9a knockdown increased the expression of cTnT gene mRNA by 1.5 times compared with the control group. The opposite result was found when KDM3A was knocked down ($n=3$, $^{**}P<0.01$ vs. the control group). cTnT, cardiac troponin-T; 5-AZA, 5-Azacytidine; G9a, euchromatic histone-lysine N-methyltransferase 2; KDM3A, H3K9me2 lysine demethylase 3A; MSCs, rat mesenchymal stem cells; sh, short hairpin.

This confirmed that the MSCs isolated in this study expressed cell markers consistently with those reported previously (9).

Cardiac differentiation of the MSCs with 5-AZA induction.

In order to construct a system for the differentiation of MSCs into cardiomyocytes, the cells were treated with 5-AZA; $\leq 50\%$ of the MSCs died and detached from the plate. On the seventh day following 5-AZA treatment, the cell volume increased significantly and gradually became short columnar with the same direction of arrangement. With the extension of culture time, the cell morphology was varied, the cytoplasm was rough and the myofilament structure appeared after two weeks. After three weeks it was observed that the cells were long, and spindle shaped or short and column shaped and were closely connected with adjacent cells (Fig. 2A). After three weeks the cell proliferation slowed down, and a portion of the cells died. No cell pulsation was observed during this process. Immunohistochemical staining results demonstrated that cTnT positive cells appeared following 5-AZA treatment (Fig. 2B). These data indicated that the system for the differentiation of MSCs into cardiomyocytes was successfully constructed.

To observe the changes of H3K9me2 in the process of induced differentiation, western blotting was used to detect the

expression of H3K9me2. It was found that H3K9me2 decreased gradually with the increase in days, but there was no significant change in untreated group (Fig. 2C). This suggests that H3K9me2 may be involved in the process of differentiation.

The interference vectors of G9a and KDM3A genes can affect the expression of H3K9me2. In order to investigate the function of H3K9me2 in the differentiation of the MSCs into cardiomyocytes, three interference carriers of G9a and KDM3A, respectively were constructed (Fig. 3A). The expressions of G9a and KDM3A were detected 48 h following transfection of the MSCs with the interference vector thus constructed. Among them, RT-qPCR demonstrated the strongest interference activity with respect to sh3-KDM3A. Similarly, sh2-G9a was screened for the best interference ($P<0.05$; Fig. 3B). Thus, sh3-KDM3A and sh2-G9a were selected for further study.

To observe H3K9me2 expression following H3K9me2 methylation modification enzyme knockdown, H3K9me2 expression was detected at 72 h following lentivirus transfection of rat MSCs. Western blotting results demonstrated that, compared with the control group, the H3K9me2 was significantly reduced when G9a was knocked down but the level of H3K9me2 was significantly increased following KDM3A knockdown (Fig. 3C). This suggests that the knockdown of

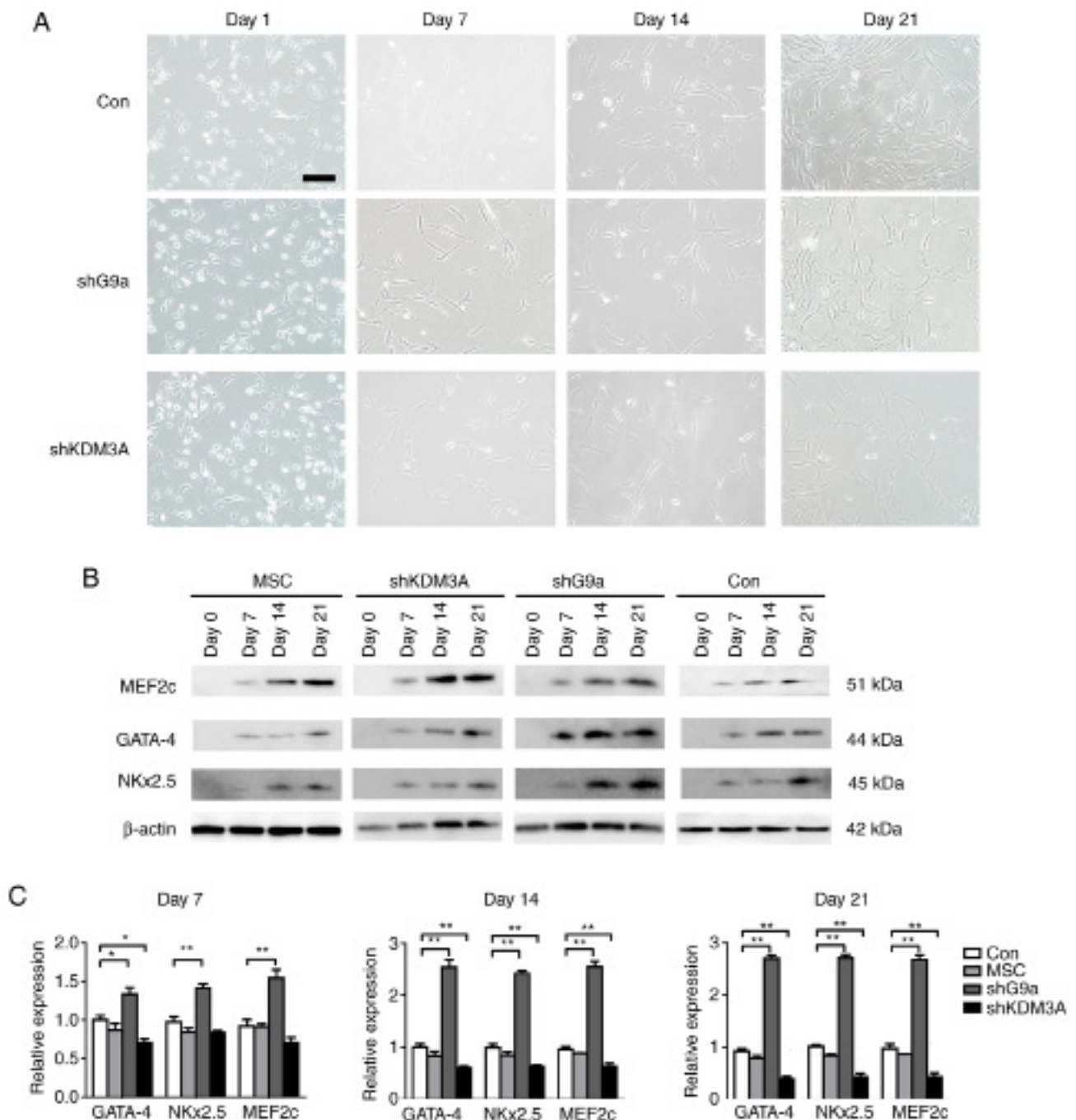


Figure 5. Morphological and early transcription factor expression changes in the mesenchymal stem cells treated with 5-AZA. (A) The cells in all groups partly succumbed following 5-AZA treatment and the surviving cells began to proliferate and differentiate. After a week the cells were enlarged, partially fusiform and regularly arranged and after three weeks they formed a myotubular-like structure by connecting with adjacent cells. Scale bar=50 μ m. (B) Cardiomyocyte differentiation of the MSCs with 5-AZA treatment was detected by western blotting, which demonstrated the upregulation of MEF2c, GATA-4 and Nkx2.5 mRNA expression on days 7, 14 and 21 in all groups. β -actin was used as a reference gene. (C) Cardiomyocyte differentiation of the MSCs with 5-AZA treatment was detected by reverse transcription-quantitative PCR. When G9a was knocked down, GATA-4, Nkx2.5 and MEF2c mRNA expression was upregulated on days 7, 14 and 21 compared with those of the control group. By contrast, GATA-4, Nkx2.5 and MEF2c mRNA expression were decreased following KDM3A knockdown. GAPDH was used as a reference gene. * P <0.05 and ** P <0.01 vs. control group. 5-AZA, 5-Azacytidine; MSCs, rat mesenchymal stem cells; MEF2c, myocyte enhancer factor 2c; GATA-4, GATA binding protein 4; Nkx2.5, NK2 Homeobox 5; sh, short hairpin; MSCs, untreated MSCs; control, MSCs transfected with control lentivirus; shG9a group, MSCs transfected with lentivirus encoding shRNA-G9a; KDM3A group, MSCs transfected with lentivirus encoding shRNA-KDM3A; G9a, euchromatic histone-lysine N-methyltransferase 2; KDM3A, H3K9me2 lysine demethylase 3A.

H3K9me2 methylating enzyme could affect the expression of H3K9me2.

The knockdown of G9a enhances cardiac differentiation of the MSCs treated with 5-AZA. In order to compare the effect

of H3K9me2 on the final induction efficiency, the expression of cardiac protein cTnT was detected by means of flow cytometry, western blotting and RT-qPCR. Following 5-AZA treatment, cTnT-positive cells in all groups increased gradually as more days passed. However, there were significant differences in the

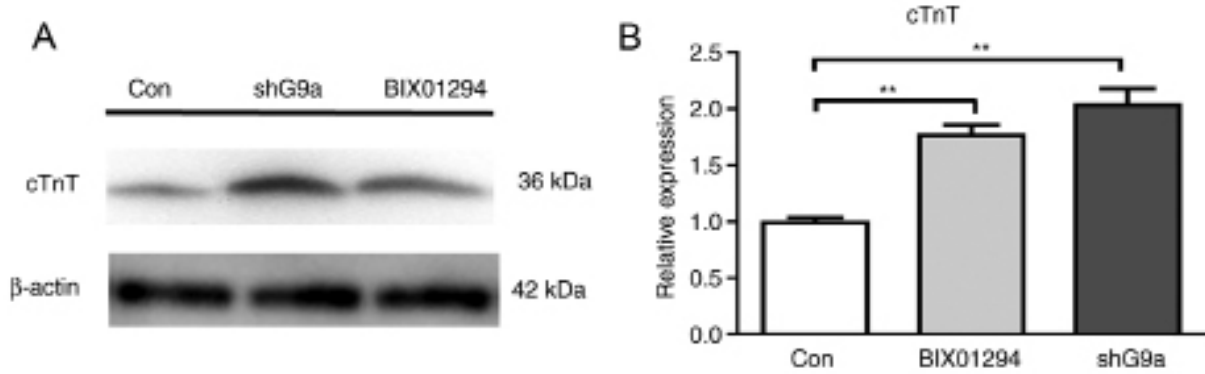


Figure 6. The expression level of cTnT following pretreatment with BIX01294. (A) Cardiomyocyte differentiation of MSCs with 5-AZA treatment was detected by western blotting analysis, which demonstrated that cTnT protein levels were upregulated in shG9a and BIX01294 groups on day 21. β -actin was used as an internal control. (B) Reverse transcription-quantitative PCR demonstrated that knockdown G9a increased the expression of cTnT gene mRNA vs. the control group. (n=3, **P<0.01 vs. the control group). cTnT, cardiac troponin-T; MSCs, rat mesenchymal stem cells; 5-AZA, 5-Azacytidine; sh, short hairpin; G9a, euchromatic histone-lysine N-methyltransferase 2.

percentages of cTnT-positive cells between the groups with different treatments as shown in Fig. 4A (P<0.05). In the control group, less than 20% of the differentiating cells that were cTnT positive after three weeks of culturation. When KDM3A were knocked down, slightly more than 10% of cells were positive for cTnT after three weeks of differentiation. However, almost 30% of the cells stained positive for cTnT following G9a knockdown (Fig. 4B). Western blotting demonstrated the same results (Fig. 4C). Similarly, the mRNA expression of the shG9a group was significantly upregulated for the specific cardiomyocyte markers on day 21 (P<0.05; Fig. 4D). The results demonstrated that the reduction of H3K9me2 by interfering with G9a could increase the efficiency of the MSCs induced differentiation into cardiomyocytes.

The knockdown of G9a and KDM3A can affect the expression of myocardial specific transcription factors. To investigate the effect of H3K9me2 on MSCs-induced differentiated cardiomyocyte, changes in cell morphology and the expression of early transcription factors GTA-4, Nkx2.5 and MEF2c following the knockdown of G9a and KDM3A were observed. There was no significant difference in cell morphology between the groups (Fig. 5A). The results of western blotting demonstrated that the myocardial specific transcription factors GTA-4, Nkx2.5 and MEF2c were increased in all groups (Fig. 5B). Compared with the MSCs and control groups, RT-qPCR demonstrated that, following the knockdown of G9a, the GATA-4, Nkx2.5 and MEF2c mRNA expression was higher on days 7, 14 and 21 (P<0.05). Following the knockdown of KDM3A, the expression levels of GTA-4, Nkx2.5 and MEF2c were significantly downregulated (P<0.05; Fig. 5C). These results suggest that inhibition of H3K9me2 by knockdown G9a can increase the expression of cardiac early transcription factors.

BIX01294 stimulates differentiation of MSCs. The present study investigated the effect of BIX01294 on the process of differentiation to demonstrate that inhibiting G9a can improve the efficiency of the differentiation of MSCs into cardiomyocytes. MSCs were divided into three groups: Treated with 1 μ M BIX01294 for 12 h, transfected with shRNA-G9a and control. All groups were induced by 5-AZA. The expression

of cTnT on the 21st day was detected by western blotting and RT-qPCR (Fig. 6A and B). The results demonstrated that both BIX01294 group and shG9a group can increase the expression of cTnT, but there was no significant difference between two groups. The results further demonstrated that H3K9me2 was involved in the process of cardiomyocyte' differentiation.

Discussion

Compared with implantable electronic pacemakers, biological pacemakers are more suitable for treating chronic arrhythmia and pathological sinus syndrome (2). Currently, the efficiency of stem cell induction into cardiomyocytes is low, which limits its clinical application. Our previous study treated very small embryonic-like stem cells with 5-AZA and the positive expression rates of cTnT and α -actin were 18.41 and 19.43%, respectively on day 14 (19). In the present study, epigenetics was used to improve the differentiation efficiency of stem cells.

5-AZA, widely used in the cardiac differentiation of stem cells, can regulate the histone demethylation and DNA methylation (20). A number of stem cells have been used to differentiate into cardiomyocytes by 5-AZA (19-23). The mechanism of 5-AZA may be related to CpG base-pair demethylation and regulation of early myocardial transcription factors (24-27). However, the efficiency of 5-AZA in inducing the differentiation of stem cells into cardiomyocytes varies between different laboratories. Antonitsis *et al* (24) induced MSCs into cardiomyocytes with an efficiency of 15%. After embryonic stem cells were treated with 5-AZA, Choi *et al* (28) found that the proportion of sarcomeric α -actin positive cells accounted for 6.48% on day 15 measured by means of flow cytometry. In the present study, MSCs isolated by density centrifugation expressed CD29 and CD44, but not CD45, which is consistent with previously reported expression profiles of the MSCs (12). By using 5-AZA, cTnT-expressing cells were successfully induced, and it is suggested that the present study successfully constructed a system of 5-AZA inducing MSCs into cardiomyocytes. In the present study, cTnT-positive cells were 19% following 5-AZA induction, which was not different from previous studies. The low efficiency with respect to the induced cardiomyocyte differentiation of the MSCs could

possibly be due to the reduced response to other factors produced in the microenvironment that could be involved in the process of cardiac differentiation following the treatment of 5-AZA (12). Therefore, an effective way to improve the induction efficiency is required.

It is becoming clear that histone modifications regulate transcription and control cell fate during cell development. During cardiac development, Histone acetylation/methylation and DNA methylation were both involved in regulating GATA binding protein 4 (GATA-4) expression (29), while H3K27me1, H3K27me3, H3K36me3 and H3K4me3 are associated with the common histone methylation modifications of H3K9me3/me2. H3K27me3 and H3K9me3/me2 are associated with gene silencing and H3K27me1, H3K4me3 and H3K36me3 are associated with gene activation (30). During the maturation of cardiomyocytes, the expression of H3K9me2 can be regulated to affect the expression of genes related to cardiac development and protect the heart from pathological hypertrophy caused by overexpression of this gene (20). In the present study, H3K9me2 decreased gradually with the increase of induction days, suggesting that H3K9me2 may be involved in the differentiation of the MSCs into cardiomyocytes. In addition, the changes of H3K9me2 expression were not significant on the 7th day, but were from the 14th day, which was consistent with the changes in the expression of cTnT. This may be because the differentiation of MSCs into cardiomyocytes is a dynamic process. The results of the present study are consistent with previous studies (9,20). Unlike G9a, the mechanism by which 5-AZA regulates histone methylation remains unclear. A previous study also demonstrated that 5-AZA inhibits H3K9me2 rather than G9a/Glp expression (31).

It was hypothesized that the differentiation efficiency could be affected by regulating the expression of H3K9me2. G9a and KDM3A are important H3K9me2 enzymes for methylation modification. G9a can modify histones at H3K9 and H3K27 sites in promoter regions of some genes through its histone lysine methyltransferases activity, thus regulating gene transcriptional silencing (32). Previous studies have found that knockdown of G9a or use of G9a inhibitor can significantly reduce the level of H3K9me2 *in vivo* (33,34). In the present study, knockdown of G9a also reduced the expression of H3K9me2, which was consistent with their results. Qin *et al* (35) found that the knockout of KDM3A significantly increased H3K9me2/me1 levels. The present study found that knockdown of KDM3A increased the expression of H3K9me2, which was consistent with previous research results (15,35). However, the knockdown of G9a and KDM3A did not cause changes in cell morphology during the induction of 5-AZA. It was considered that the changes in H3K9me2 expression caused by knockdown of G9a and KDM3A only caused changes in the transcription level of genes regulated by chromatin, which did not involve changes in DNA sequencing.

Additionally, the present study compared the final cTnT-positive cell rate and the differentiation efficiency increase following G9a knockdown, which was greater compared with the previous efficiency following treatment with 5-AZA alone (19,24,28). The same effect can be achieved with G9a inhibitors BIX012094. 5-AZA can induce the differentiation of bone marrow mesenchymal stem cells into cardiomyocytes by participating in the gene expression

regulation of the transcription factors GATA-4 and/or Nkx2.5 in early myocardial development (36,37). In the present study, following 5-AZA treatment, the early transcription factors GATA-4, NKx2.5 and MEF2c increased in all groups of the MSCs, indicating that the treated cells had reached the myocardial phenotype, consistent with the results of differentiation induced by different inducers (9,12,16). The results of the present study found that the expressions of GATA-4, NKx2.5 and MEF2c were highest following G9a knockdown but that the expression of transcription factors in early cardiac development was decreased following KMD3A knockdown, indicating that H3K9me2 may be involved in the differentiation of MSCs into cardiomyocytes. The final results of the present study demonstrated that the reduction H3K9me2 by G9a knockdown can improve the efficiency of 5-AZA inducing the differentiation of MSCs into cardiomyocytes. This may be because H3K9me2 recruits inhibitory factors, such as members of the HP1 family, leading to tighter binding of DNA and histones, thereby inhibiting transcription of key genes in cardiac cell development (38). Unfortunately, the present study did not compare the efficiency of 5-AZA and G9a against H3K9me2 inhibition directly. Therefore, in the next stage of the study, further research will be conducted on this aspect.

In conclusion, the present study suggested that H3K9me2 may be involved in the process of differentiation MSCs into cardiomyocytes. Interference with G9a is an effective way to increase 5-AZA induction of the MSCs into cardiomyocytes *in vitro*.

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Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Authors' contributions

XS, HL, PX, ML and XG designed the study, collected the data, performed the statistical analyses and contributed to the writing of the manuscript. QZ and BL helped supervise the experiments. PX and YZ helped with the data collection, study design and supervised the study. QZ and BL participated in the study design and helped to critically revise the manuscript. XS and XG confirm the authenticity of all the raw data. All authors read and approved the final manuscript.

Ethics approval and consent to participate

All procedures involving the care and use of animals conformed to the Animal Research: Reporting of *in vivo*

Experiments (ARRIVE) guidelines and were approved by the Laboratory Animal Management and Experimental Animal Ethics Committee of Yangzhou University (approval no. 201903478).

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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