The role of IncRNA MALAT1 in the regulation of hepatocyte proliferation during liver regeneration

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Abstract. Exploring the biological functions of long non-coding RNAs (lncRNAs) has come to the foreground in recent years. Studies have indicated that the lncRNA metastasis-associated lung adenocarcinoma transcript 1 (MALAT1) not only regulates tumorigenesis in hepatocellular carcinoma, but also controls cell cycle progression in hematopoietic cells. The present study was designed to investigate the biological role of IncRNA MALAT1 in liver regeneration. We carried out a series of assays during liver regeneration following 2/3 partial hepatectomy in mice. We explored the functions of IncRNA MALAT1 with a series of functional analyses in vitro. We found that MALAT1 was upregulated during liver regeneration. Moreover, MALAT1 accelerated hepatocyte proliferation by stimulating cell cycle progression from the G1 to the S phase and inhibiting apoptosis in vitro. In addition, our findings also demonstrated that MALAT1 was regulated by p53 during liver regeneration, and that p53 may be a key upstream regulator of MALAT1 activity. Mechanistically, we found that MALAT1 activated the Wnt/β-catenin pathway by inhibiting the expression of Axin1 and adenomatous polyposis coli (APC), and subsequently promoting the expression of cyclin D1. On the whole, the findings of this study suggest that MALAT1 is a critical molecule for liver regeneration. Pharmacological interventions targeting MALAT1 may thus prove to be therapeutically beneficial in liver failure or liver transplantation by promoting liver regeneration.

Introduction

Liver regeneration refers to the liver repair process that occurs following the loss of part of the liver due to surgery, trauma, poisoning, infection and necrosis, or liver transplantation. When liver regeneration occurs, hepatocytes in the G0 phase are stimulated by various feedback signals to enter a period of rapid growth. Liver regeneration has valuable implications for hepatobiliary surgeries, such as the resection of a liver mass, liver transplantation and split liver transplantation, as the above-mentioned surgical procedures depend largely on the ability of the liver to regenerate. Poor liver regeneration may lead to the death of the patient, for example, via small-for-size syndrome (SFSS) (1-4). For that reason, understanding the mechanisms of liver regeneration has great implications for clinical practice. A series of interacting factors, including inflammatory cytokines (5), hepatocyte growth factor (HGF) (6), cell cycle regulators, cyclin-dependent kinases and metabolic regulators control the hepatocyte cell cycle to ensure the rapid and orderly progression of liver regeneration (7). Studies have indicated that microRNAs (miRNAs) or miRs play a key role in liver regeneration (8-10). Despite the identification of many cytokines, miRNAs and growth factors that regulate the expression of genes controlling proliferation during liver regeneration (11), the exact molecular mechanisms which are responsible for liver regeneration remain unclear, thus prompting our interest to investigate the molecular mechanisms underlying liver regeneration.

In recent years, the field of RNA biology has been dominated by the study of miRNAs; however, several major biological processes are also regulated by long non-coding RNAs (lncRNAs), including carcinogenesis, development and differentiation (12,13). lncRNA molecules are >200 nucleotides in length and do not encode proteins. Studies have reported that some lncRNAs, including IncRNA H19 and IncRNA LALR1, play key roles during liver regeneration (14,15). IncRNA H19 promotes hepatocyte proliferation during liver regeneration in both the rat and mouse (15). Improving the effectiveness of current regeneration techniques may help to identify a potential therapeutic target to stimulate liver regeneration.

Metastasis-associated lung adenocarcinoma transcript 1 (MALAT1) is one of the most widely studied lncRNAs and is frequently overexpressed in malignant tumors; therefore,
MALAT1 is frequently regarded as a biomarker for a variety of tumors. A previous study demonstrated that MALAT1 localizes in dynamic structures known as nuclear speckles, which are key elements in the collection and recruitment of splicing factors (16). Researchers have also found that MALAT1 plays a variety of biological functions in hematopoeitic cells, including the regulation of the cell cycle (17). However, the biological role of the IncRNA MALAT1 in liver regeneration and its mechanisms of action regarding the regulation of hepatocyte proliferation remain unclear.

In this study, we focused on the mechanisms underlying the MALAT1-mediated modulation of hepatocyte proliferation. We investigated the role of MALAT1 in the regulation of liver regeneration with a murine model of 2/3 partial hepatectomy (PH) and an in vitro hepatocyte proliferation assay. The comprehensive dynamic changes in MALAT1 expression were examined at multiple time points (including 0, 2, 6, 12, 15, 18, 24, 36, 48, 72, 120, 144 and 168 h) following 2/3 PH; we then aimed to elucidate the mechanisms through which MALAT1 regulates liver regeneration in the mouse. We found that MALAT1 promoted hepatocyte proliferation and increased the expression of cyclin D1 by activating the Wnt/β-catenin pathway. We also found that the relative expression level of p53, which is a negative regulator of MALAT1, was the opposite of MALAT1 expression following 2/3 PH. Our study may have identified a potential therapeutic target for use during liver regeneration.

Materials and methods

Animals and model of 2/3 PH. A total of 60 male BALB/c mice (8-12 weeks old, weighing 23±2 g) were purchased from the Central Laboratory of Animal Science, Wuhan University (Wuhan, China) and were maintained in a specific pathogen-free facility with a 12-h light/dark cycle and were fed standard laboratory chow with free access to water ad arbitrium. All the animal protocols used in the study were reviewed and approved by the Animal Experiment Center of Wuhan University/Animal Biosafety Level-III Laboratory. 2/3 PH was performed on 8-12-week-old mice using a previously described method (18). Following a laparotomy, the left and median liver lobes were resected in two steps. Five mice were used for each 2/3 PH time point (0, 2, 6, 12, 15, 18, 24, 36, 48, 72, 120 and 168 h) following 2/3 PH; we then aimed to elucidate the mechanisms through which MALAT1 regulates liver regeneration in the mouse. At different time points following 2/3 PH, the animals were sacrificed and the liver tissues were harvested. Liver tissues were obtained from the remnant livers and then fixed immediately in 10% buffered formalin phosphate solution, embedded in paraffin and were then cut into 5-mm-thick sections. The sections were stained with hematoxylin and eosin (H&E). The H&E staining kit was obtained from Beijing Solarbio Science & Technology Co., Ltd. (Beijing, China). Motic Images 2000 (Motic China Group Co. Ltd, Guangzhou, China) was used for the characterization of histopathological changes.

Histological analysis. At different time points following 2/3 PH, the animals were sacrificed and the liver tissues were harvested. Liver tissues were obtained from the remnant livers and then fixed immediately in 10% buffered formalin phosphate solution, embedded in paraffin and then cut into 5-mm-thick sections. The sections were stained with hematoxylin and eosin (H&E). The H&E staining kit was obtained from Service Bio Co., Ltd. (Wuhan, China Cat. no. G0105). All tissue sections were examined with an Olympus BH-2 microscope (Olympus Optical Co. Ltd., Beijing, China). Motic Images 2000 (Motic China Group Co. Ltd, Guangzhou, China) was used for the characterization of histopathological changes.

ELISA. The mice were sacrificed and blood samples were collected via the post-orbital venous plexus at the mentioned observation time points. Serum samples were separated and stored at -80°C until further use. The serum levels of alanine aminotransferase (ALT), aspartate aminotransferase (AST) and albumin (ALB) were measured using respective ELISA kits (Nanjing Jiancheng Biological Technology, Inc., Nanjing, China) as previously described (19).

Table I. Primer and target sequences used in this study.

| Genes                  | Sequences or target sequences (5’→3’) |
|------------------------|--------------------------------------|
| MALAT1-F               | TGCAGTTGTCACATGTGTTGTCG              |
| MALAT1-R               | GGCCAGCTGCAACATTCAA                  |
| GAPDH-R                | GTGAAAAGGTCTTGTTGTAACGGC             |
| GAPDH-R                | CTGCGCTTGGAAAGATGTTGAGCC             |
| HGF-F                  | TCATTGGTAAAGAGAGCGCA                 |
| HGF-R                  | GTCACAGACTTTCTGACT                   |
| β-catenin-F            | ATGGCTTTGGAATGAGACT                 |
| β-catenin-R            | TGAGGTCTGGGCGTGTC                    |
| Cyclin D1-F            | GAGGAGCAGAAGTGCGCGAGA                |
| Cyclin D1-R            | GCCGGGTAAGTGTCAGTG                   |
| Axin1-F                | GCTGTATTGGAGACTGCT                   |
| Axin1-R                | GTACCCGGCCATTGACT                    |
| GSK-3β-F               | TTTATTGCCGCTAGTTC                    |
| GSK-3β-R               | AAGCCGGCTGTTCCTGCCGT                |
| APC-F                  | CCGTGGCAGAGGAAACC                    |
| APC-R                  | CCTGCTGACATCTACTGTGT                 |
| p53-F                  | CAGGCCCTCCTCAGTACTG                 |
| p53-R                  | ACCCTATGAGGGCCCAAGAT                 |
| MALAT1-siRNA-1         | GCCUGAGGGUCUUGCAUAUATT               |
| MALAT1-siRNA-2         | UAUUGUGACAAAGUCUCAGGTCT             |
| MALAT1-siRNA-3         | GCAGAGAGAUUCUGCUCAUUTT              |
| MALAT1-siRNA-scramble  | AAGAUGAAGACACUCUUCUGGCTT             |
|                         | UUCUCGAGACGUGUCAGGUTT               |
|                         | ACUGCAGACGUCUGGAGATT                |

HGF, hepatocyte growth factor; APC, adenomatous polyposis coli.
Table II. Information regarding the antibodies used in this study.

| Antibody | WB | IHC | IF | Specificity | Company |
|----------|----|-----|----|-------------|---------|
| GAPDH (10494-1-AP) | 1:5,000 | | | Rabbit polyclonal | Proteintech |
| β-catenin (no. 8480) | 1:1,000 | 1:100 | | Rabbit monoclonal | Cell Signaling Technology |
| Cyclin D1 (no. 2978) | 1:1,000 | | | Rabbit monoclonal | Cell Signaling Technology |
| Axin1 (no. 2087) | 1:1,000 | | | Rabbit monoclonal | Cell Signaling Technology |
| c-Myc (sc-788) | 1:200 | | | Rabbit polyclonal | Santa Cruz Biotechnology |
| Cyclin B1 (55004-1-AP) | 1:2,000 | | | Rabbit polyclonal | Proteintech |
| Cyclin E1 (11554-1-AP) | 1:1,000 | | | Rabbit polyclonal | Proteintech |
| GSK-3β (no. 12456) | 1:1,000 | | | Rabbit polyclonal | Cell Signaling Technology |
| APC (sc-896) | 1:200 | | | Rabbit polyclonal | Santa Cruz Biotechnology |
| p53 (sc-6243) | 1:200 | 1:200 | | Rabbit polyclonal | Santa Cruz Biotechnology |
| PCNA (10205-2-AP) | 1:2,000 | 1:200 | | Rabbit polyclonal | Proteintech |
| Histone H3 (no. 4499) | 1:2,000 | | | Rabbit polyclonal | Cell Signaling Technology |

WB, western blot analysis; IHC, immunohistochemistry; IF, immunofluorescence; GSK-3β, glycogen synthase kinase-3β; APC, adenomatous polyposis coli; PCNA, proliferating cell nuclear antigen.

Reverse transcription-quantitative PCR (RT-qPCR). Total RNA was isolated from the frozen tissues or cells using TRIzol reagent (Invitrogen Life Technologies, Carlsbad, CA, USA) according to the manufacturer's instructions. For RT-qPCR, a reverse transcription kit (Takara Biotechnology Co., Ltd., Dalian, China) was used to synthesize cDNA. A SYBR-Green PCR kit (Toyobo, Osaka, Japan) was used to complete the RT-qPCR analyses. PCR reactions were performed on a CFX96 system (Bio-Rad CFX96; Bio-Rad Laboratories, Inc., Hercules, CA, USA). The results of RT-qPCR were normalized to glyceraldehyde 3-phosphate dehydrogenase (GAPDH), and the 2^ΔΔCT values were normalized to the GAPDH levels. All the RT-qPCR experiments were repeated at least 3 times. The primer sequences used in this study are listed in Table I.

Cell culture and transfection. NCTC 1469 and BNL CL.2 cells were obtained from the China Center for Type Culture Collection (Wuhan, China). The NCTC 1469 cells were cultured with Dulbecco's modified Eagle's medium (DMEM H-21, 4.5 g/l glucose) supplemented with 15% horse serum. BNL CL.2 cells were cultured with DMEM supplemented with 10% fetal bovine serum (FBS). The growth media used in the present study contained 5 U/ml penicillin and 5 µg/ml streptomycin sulfate. Full-length MALAT1 (GenBank: NR_002847.2) was cloned into the pcDNA3.1 vector (Invitrogen Life Technologies, Carlsbad, CA, USA). An empty vector (pcDNA3.1) was used as a control. siRNAs against MALAT1 were designed and synthesized by Beijing View Solid Biotechnology (Beijing, China). The siRNA target sequences used in this study are listed in Table I. The MALAT1 expression levels were detected by RT-qPCR. Scramble siRNA was used as a control. Transfection was conducted using a Lipofect™ in vitro DNA and siRNA transfection kit (both from SigmaGen Laboratories, Gaithersburg, MD, USA) according to the manufacturer's instructions. The hepatocytes were fixed with 4% paraformaldehyde 48 h after transfection for further analysis.

NCTC 1469 cells were treated with HGF (Cat. no. #315-23; PeproTech, Rocky Hill, NJ, USA) using a concentration range of 5-20 ng/ml. It is recommended to reconstitute the lophilized HGF in sterile 18MQ·cm H2O not less than 100 µg/ml, which can then be further diluted to other aqueous solutions.

Flow cytometric analysis. The cells transfected with siRNA or the pcDNA3.1 vector were collected at 48 h post-transfection. Following double-labeling with Annexin V-FITC and propidium iodide (PI), apoptosis was analyzed using a flow cytometer (FACScan®; BD Biosciences, Franklin Lakes, NJ, USA) equipped with CellQuest 3.3 software. For the cell cycle analysis, 1x10⁵ cells were plated into a 60-mm dish and incubated for 24 h. The cells were subsequently stained with 1 ml of staining solution [PI (50 µg/ml) and RNase A (20 µg/ml)] for 15 min using the Cycletest™ PLUS DNA reagent kit (BD Biosciences) according to the manufacturer's instructions.

Western blot analysis. The liver tissue samples and hepatocytes were homogenized in lysis buffer (Beyotime Institute of Biotechnology, Haimen, China) and incubated for 30 min on ice. For the detection of β-catenin, cytoplasmic and nuclear protein extracts were prepared from the cells using the NE-PER Nuclear and Cytoplasmic Extraction Reagent kit (Cat. no. 78835; Thermo Fisher Scientific, Inc., Waltham, MA, USA) according to the manufacturer's instructions. Total protein was electrophoresed by sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) and then transferred onto polyvinylidene fluoride (PVDF) membranes (Millipore, Billerica, MA, USA). The membranes were incubated with the corresponding primary antibodies overnight at 4°C. The following day, the membranes were incubated with secondary antibodies [goat anti-rabbit IgG (H+L)-HRP (LK2001; Sungene Biotech Co., Ltd., Tianjin, China)] for 1-2 h. Proteins were detected using the ECL immunoblotting kit (Beyotime Institute of Biotechnology). The fluorescent bands were observed in a dark room with infrared light. The above procedure was performed according to the manufacturer's instructions. The information for the antibodies is listed in Table II.
Cell proliferation assay. A Cell Counting Kit-8 kit (CCK-8; Dojindo, Kumamoto, Japan) was used for the cell proliferation assays. Approximately 3-5x10^4 cells were plated in triplicate in 96-well plates and cultured in growth medium. The plate was incubated at 37°C and 5% CO₂, for an appropriate length of time (4-6 h) until cell adherence. Ten microliters of CCK-8 solution were then added to each well of the plate. The absorbance was measured at 450 nm using a microplate reader (168-1000XC; Bio-Rad Laboratories, Inc.) at the indicated time points.

5-Ethynyl-2′-deoxyuridine (EdU) incorporation assay. The EdU incorporation assay was performed using an EdU Apollo DNA in vitro kit (C10310-1; Guangzhou RiboBio Co., Ltd., Guangzhou, China) according to the manufacturer’s instructions. The hepatocytes were stained with EdU 48 h following transfection. First, the cells were incubated with 100 µl of 50 mM EdU/well for 2 h. The cells were then fixed with 4% paraformaldehyde for 15 min at room temperature. Following fixation and washing with phosphate-buffered saline (PBS), the cells were incubated with 50 µl of 2 mg/ml glycine for 5 min. One hundred microliters of permeabilization buffer (PBS containing 0.5% Triton X-100) was added to each well, and the cells were incubated with 100 µl 1X Apollo solution for 30 min at room temperature in the dark. Then, 100 µl 1X Hoechst 33342 solution was added dropwise into each cell climbing slice for 15 min. The cell climbing slices were washed, dried, mounted and then observed under a fluorescence microscope (Olympus BX53; Olympus, Tokyo, Japan).

Immunohistochemistry (IHC) and immunofluorescence (IF). Paraffin-embedded tissue was cut into 4-µm-thick sections and processed for IHC in accordance with a previously described protocol (20). For IF, the cells were fixed with 4% paraformaldehyde, permeabilized using 0.2% Triton X-100 and incubated with primary and secondary antibodies according to the manufacturer’s instructions. The cells were counterstained with 4,6-diamidino-2-phenylindole (DAPI) (Calbiochem, San Diego, CA, USA) and imaged using a confocal laser scanning microscope (Olympus FV1000; Olympus).

Statistical analysis. Quantitative data are presented as the means ± standard deviation (SD) from at least 3 independent experiments. Statistical significance was assessed using the Student’s t-test, χ² test or Fisher’s exact test, which were performed using SPSS 22.0 statistical software (IBM SPSS, Inc., Chicago, IL, USA). Differences were considered statistically significant when P-values were <0.05.

Results

MALAT1 is upregulated during liver regeneration. To determine the impact of MALAT1 on liver regeneration, we first generated mouse model of 2/3 PH. To assess rehabilitation from acute liver injury, the serum levels of ALT, AST and ALB were determined 0, 24, 36, 48, 72 and 168 h following 2/3 PH. As shown in Fig. 1A-C, all the indicators returned to normal at 168 h following 2/3 PH. H&E staining of the liver was carried out at different time points. Following hepatectomy, we observed obvious cell edema at 15 and 36 h. Mitotic figures were clearly observed at 72 h, and the hepatic lobe structure was close to normal at 120 h (Fig. 1D). Mouse hepatocyte proliferation was determined in vivo by staining for proliferating cell nuclear antigen (PCNA) (Fig. 1D) using IHC. We found that the number of PCNA-positive nuclei reached a peak at 36 h following 2/3 PH. The cell proliferative ability was reduced at 72 and 120 h, and the proliferation at 24 and 168 h was not significant (Fig. 1D). The remnant liver/body weight ratio is shown in Fig. 1E.

2/3 PH in male BALB/c mice caused a ~5-fold increase in MALAT1 expression. MALAT1 expression increased from 2 h, peaked at 15 h following PH, and then returned to almost normal levels 120 h following 2/3 PH (Fig. 1F). These results suggested that MALAT1 was specifically overexpressed in hepatic tissue following 2/3 PH. Taken together, this sharp increase indicates that MALAT1 may play a role in liver regeneration.

IncRNA MALAT1 accelerates cell cycle progression in hepatocytes and promotes proliferation in vitro. As it has been previously demonstrated that hepatocytes from adult male mice enter the S phase at 36 h following 2/3 PH (18) and that the expression of MALAT1 reaches a maximum after 12-18 h of liver regeneration, we hypothesized that MALAT1 may play an important role in regulating the cell cycle, as the expression of MALAT1 peaked before 36 h.

First, we analyzed MALAT1 expression in BNL CL.2 and NCTC 1469 cells. MALAT1 expression was significantly higher in the BNL CL.2 cells than in the NCTC 1469 cells (Fig. 2A). As shown in Fig. 2B, siRNA-2 was the most effective at silencing MALAT1. MALAT1 expression was upregulated almost 18-fold following transfection of the NCTC 1469 cells with the pcDNA3.1 vector carrying MALAT1. CCK-8 (Fig. 2C) and EdU immunofluorescence assays (Fig. 2D) were used to detect changes in the proliferative ability of the MALAT1-silenced BNL CL.2 cells and MALAT1-overexpressing NCTC 1469 cells. The experimental results revealed that cell proliferation was reduced when MALAT1 was silenced in the BNL CL.2 cells and was increased when MALAT1 was overexpressed in the NCTC 1469 cells. As shown in Fig. 2D, the MALAT1-overexpressing NCTC 1469 cells had more EdU-positive cells than the control group, while the MALAT1-silenced BNL CL.2 cells had fewer EdU-positive cells than the control cells. Moreover, FACS analysis indicated that the MALAT1-silenced BNL CL.2 cells were arrested in the G0/G1 phase of the cell cycle. Furthermore, the number of cells in the G0/G1 phase decreased significantly and the number of cells in the S phase increased significantly when MALAT1 was overexpressed in the NCTC 1469 cells (Fig. 2E). At the same time, the rate of apoptosis increased when the BNL CL.2 cells were transfected with MALAT1-siRNA-2, but decreased when the NCTC 1469 cells were transfected with pcDNA3.1-MALAT1 (Fig. 2F). To further examine the role of MALAT1 in cell cycle progression, the expression of cyclin E1 and B1 was examined in the MALAT1-silenced BNL CL.2 cells. The results revealed that the G0/G1 phase was extended when MALAT1 was silenced, while the G2/M phase was shortened when MALAT1 was overexpressed (Fig. 2F). These results indicated that MALAT1 facilitates liver cell proliferation by accelerating cell cycle progression and blocking liver cell apoptosis.
HGF increases MALAT1 expression. As MALAT1 expression increased during the early stages after 2/3 PH and HGF rapidly increases following PH and plays a substantial role during liver regeneration (6), we hypothesized that the increase in MALAT1 expression may be mediated by HGF. To confirm our hypothesis, the NCTC 1469 cells were treated with 5, 10 or 20 ng/ml HGF to induce hepatocyte proliferation. As shown in Fig. 3A, MALAT1 expression gradually increased following treatment with increasing concentrations of HGF. The increased expression of HGF was detected in the BALB/c mice subjected to 2/3 PH (Fig. 3B). Moreover, as shown in Fig. 3C, we identified a correlation between the HGF and MALAT1 levels in the liver tissue at different time points following 2/3 PH.

MALAT1 activates Wnt/β-catenin signaling by inhibiting Axin1 and adenomatous polyposis coli (APC) in vitro. It has been demonstrated that the Wnt/β-catenin pathway is associated with proliferation and differentiation (21). Thus, to determine whether MALAT1 accelerates hepatocyte proliferation by activating the Wnt/β-catenin pathway, we evaluated the levels of cyclin D1, β-catenin, the β-catenin-degradation component, Axin1, glycogen synthase kinase (GSK)-3β and APC in MALAT1-silenced BNL CL.2 cells and MALAT1-overexpressing NCTC 1469 cells. The results revealed an increase in the protein and mRNA levels of β-catenin and cyclin D1 in the MALAT1-overexpressing NCTC 1469 cells (Fig. 4A, C and E). The protein and mRNA levels of Axin1 and APC decreased when MALAT1 was overexpressed in the NCTC 1469 cells (Fig. 4B and C); however, there was no significant change in the level of GSK-3β (Fig. 4B and C). The level of c-Myc increased when MALAT1 was overexpressed, but declined following the knockdown of MALAT1 (Fig. 4C).

As shown in Fig. 4D and E, inactive β-catenin was located in the cytoplasm, and total β-catenin staining decreased when MALAT1 was silenced in the BNL CL.2 cells. In addition, western blot analysis of β-catenin demonstrated that the level of β-catenin in the nucleus also decreased significantly when MALAT1 was silenced. These effects were reversed when MALAT1 was overexpressed. There was no significant difference in the level of cytoplasmic β-catenin when MALAT1 was silenced or overexpressed (Fig. 4E). β-catenin phosphorylation occurs in a multiprotein complex that includes the GSK-3β, β-catenin, Axin1 and APC proteins; this phosphorylation leads to the degradation of cytoplasmic β-catenin (22,23).

On the whole, our data demonstrate that MALAT1 promotes cell cycle progression and accelerates hepatocyte proliferation during liver regeneration by activating Wnt/β-catenin signaling.
Figure 2. Metastasis-associated lung adenocarcinoma transcript 1 (MALAT1) accelerates the hepatocyte cell cycle and inhibits apoptosis in vitro. (A) The expression of MALAT1 in BNL CL.2 cells and NCTC 1469 cells was determined by RT-qPCR. The level of MALAT1 in NCTC 1469 cells was set to a value of 1. (B) RT-qPCR analysis of MALAT1 expression after MALAT1 was knocked down by siRNA in BNL CL.2 cells. MALAT1 expression was determined by RT-qPCR after MALAT1 was overexpressed in NCTC 1469 cells. (C) Cell proliferation was assessed by CCK-8 assay. (D) Cell proliferation was assessed using 5-ethyl-2-deoxyuridine (EdU) immunofluorescence staining. (E) FACS analysis showing that the number of cells in the G1/G0 phase increased significantly in MALAT1-silenced BNL CL.2 cells and that the number of cells in the S phase increased in MALAT1-overexpressing NCTC 1469 cells. (F) FACS analysis showing a significant increase in the number of apoptotic cells when MALAT1 was silenced in BNL CL.2 cells and a decrease in the number of apoptotic NCTC 1469 cells following transfection with pcDNA3.1-MALAT1. The percentage of apoptotic cells is also shown. (G) Western blot analysis of the expression of cyclin B1 and E1 in MALAT1-silenced BNL CL.2 cells at different time points during the cell cycle. The data are presented as the means ± SD of 3 independent experiments. *p<0.05, **p<0.01 vs. the control or the 0 h time point.

Figure 3. Hepatocyte growth factor (HGF) stimulates the expression of long non-coding RNA (lncRNA)-metastasis-associated lung adenocarcinoma transcript 1 (MALAT1). (A) The expression of MALAT1 increased in NCTC 1469 cells following treatment with increasing HGF concentrations. (B) Relative mRNA expression of HGF at different time points following 2/3 partial hepatectomy (PH). (C) The expression of MALAT1 and HGF in mouse liver samples was detected by RT-qPCR at various time points following 2/3 PH. The data are presented as the means ± SD of 3 independent experiments. *p<0.05, **p<0.01 vs. the control (no treatment) or the 0 h time point.
By suppressing Axin1 and APC, MALAT1 indirectly reduces the stability of the β-catennin degradation complex, decreasing the level of inactive β-catennin, while increasing the level of active β-catennin. Active β-catennin is released and begins functioning. Subsequently, active β-catennin translocates to the nucleus to control the transcription of target genes, such as c-Myc and cyclin D1. Finally, MALAT1 enhances hepatocyte proliferation by promoting cell cycle progression in vitro.

Discussion

The key finding of the present study was that MALAT1 is upregulated during mouse liver regeneration. MALAT1 may act as a regulatory factor in the cell cycle. Our results also demonstrated that both HGF and p53 are closely associated with MALAT1 during liver regeneration. MALAT1 can activate Wnt/β-catennin signaling in vitro. Axin1 and APC were suppressed by MALAT1, and the stability of the β-catennin degradation complex became weak. Active β-catennin translocated from the cytoplasm to the nucleus, where it then enhanced the transcription of c-Myc and cyclin D1. Ultimately, MALAT1 facilitated hepatocyte proliferation by promoting cell cycle progression from the G1 to the S phase (Fig. 6).

By binding to the promoter of MALAT1, p53 can negatively regulate the expression of MALAT1 (17,27). MALAT1 is

\[\text{CGGCCATGGCCGCCAAGGTCGCCGT}\]

Figure 4. Metastasis-associated lung adenocarcinoma transcript 1 (MALAT1) activates the Wnt/β-catennin pathway by suppressing Axin1 and APC. (A) RT-qPCR analysis revealing the levels of β-catennin and cyclin D1 in MALAT1-silenced BNL CL.2 cells and MALAT1-overexpressing NCTC 1469 cells. (B) Changes in the expression of the β-catennin destruction complex (Axin1, APC and GSK-3β) after the silencing or the overexpression of MALAT1 were detected by RT-qPCR. (C) Changes in the expression of the β-catennin destruction complex (Axin1, APC and GSK-3β) and downstream target genes (cyclin D1 and c-Myc) after the silencing or overexpression of MALAT1 were detected by western blot analysis. The value below each band indicates the fold change in the protein level. (D) Changes in the expression and distribution of β-catennin in the cytoplasm and nucleus were observed by immunofluorescence (IF) staining after the silencing or overexpression of MALAT1. Yellow arrows indicate the translocation of β-catennin (scale bars, 20 µm). (E) The levels of nuclear, cytoplasmic and total β-catennin protein after the silencing or overexpression of MALAT1 are shown. The data are presented as the mean ± SD of 3 independent experiments. *p<0.05, **p<0.01 vs. the respective control.
an essential factor in maintaining the proliferative potential of early-stage hepatocytes. Combined with the results of our study, these results indicate that MALAT1 may be transcriptionally regulated by p53 during liver regeneration following 2/3 PH.

Although a number of cytokines, growth factors (such as HGF) and miRNAs regulate proliferation-related genes during liver regeneration (28,29), the role of lncRNAs in liver regeneration remains largely unknown. A comprehensive analysis of lncRNA expression during mouse liver regeneration has revealed changes in the expression of multiple lncRNAs (14). In this study, we found that HGF increased MALAT1 expression in vitro; however, the peak in HGF expression was delayed compared to the peak in MALAT1 expression. The above-mentioned phenomenon indicates that HGF may function as a regulatory factor for MALAT1. Between 15 and 24 h following PH, the anti-MALAT1 effects override the HGF-mediated increase in MALAT1, indicating that additional signaling molecules and transcription factors may regulate the expression
of MALAT1; we aim to continue to investigate the mechanisms underlying the association between HGF and MALAT1 in future studies. The HGF and Wnt/β-catenin pathways both play key roles in accelerating hepatocyte proliferation and can cooperate to stimulate hepatocyte proliferation, which may be crucial in liver regeneration following PH (30).

The complete liver regeneration process takes approximately 7 days in mice; however, the mechanisms through which the liver initiates and terminates regenerative growth are not yet fully understood (31). Exploring the biological functions of IncRNAs has recently come to the foreground (12,32). Previous studies investigating the non-coding RNAs in the liver have mainly focused on the association between the presence of these RNAs and the development of hepatocellular carcinoma (HCC) or the differential expression of these RNAs in carcinomas and the adjacent tissue (33). Some genes associated with hepatocyte proliferation and liver regeneration are also expressed abnormally in HCC (34,35). Liver regeneration is a specific cellular process that is both similar to and different from HCC (36). To date, it remains unclear as to whether the IncRNAs regulating cell proliferation in HCC participate in liver regeneration. The overall mechanisms warrant further investigation. Although partial liver transplantation can minimize organ scarcity, the risk of death of a healthy donor, the development of SFSS in both the donor and recipient and technical complexity limit its applicability. Compounds that interfere with the signaling pathways that regulate proliferation are receiving increased attention and may be used to prevent or reverse SFSS (37). In conclusion, to the best of our knowledge, our research group is the first to report a biological role of MALAT1 in hepatocyte proliferation. MALAT1 is expected to become an indicator of prognosis for a variety of liver diseases and liver resection.

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