

Original Article

Scinderin is a novel transcriptional target of BRMS1 involved in regulation of hepatocellular carcinoma cell apoptosis

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Abstract: Tumor metastasis suppressor factor BRMS1 can regulate the metastasis of breast cancer and other tumors. Here we report scinderin (SCIN) as a novel transcriptional target of BRMS1. SCIN protein belongs to the cytoskeletal gelsolin protein superfamily and its involvement in tumorigenesis remains largely illusive. An inverse correlation between the expression levels of BRMS1 and SCIN was observed in hepatocellular carcinoma (HCC) cells and tissues. On the molecular level, BRMS1 binds to *SCIN* promoter and exerts a suppressive role in regulating *SCIN* transcription. FACS analysis and caspase 9 immunoblot reveal that knockdown of *SCIN* expression can sensitize HCC cells to chemotherapeutic drugs, leading to suppression of tumor growth *in vivo*. Consistently, overexpression of SCIN protects cells from apoptotic death, contributing to increased xenografted HCC cell growth. In summary, our study reveals SCIN as a functional apoptosis regulator as well as a novel target of BRMS1 during HCC tumorigenesis. Inhibition of SCIN might bring a potential cancer therapy approach.

Keywords: BRMS1, scinderin, cell apoptosis, hepatocellular carcinoma, transcriptional regulation

Introduction

Human breast cancer metastasis suppressor 1 (BRMS1) was first discovered and cloned in 2000 by Seraj et al. [1]. Later on, it has been demonstrated that BRMS1 plays a suppressive role in regulating tumor metastasis in breast cancer, melanoma, non-small cell lung cancer, hepatocellular carcinoma, among others [2-5]. In consistent with these findings, the expression level of BRMS1 has been found to be significantly down-regulated in multiple metastatic tumor tissues [6-9]. It has been shown that BRMS1 mainly acts as a component of Sin3a•HDAC complex, which regulates chromatin status and further affects gene transcription efficiency [10, 11]. We have previously reported two well-defined tumor-metastasis related genes, osteopontin (*OPN*) and death-associated protein kinase 1 (*DAPK1*), which were under transcriptional control of BRMS1 in hepatocellular carcinoma (HCC) cell lines [9, 12].

Herein, we further identified Scinderin (*SCIN*, also known as adseverin) as a novel transcriptional target of BRMS1. *SCIN* is located in 7p21.3 and encodes a Ca²⁺-dependent actin filament severing and capping protein. SCIN protein was discovered in chromaffin cells in 1990 but its biological functions remain largely illusive [13]. As a member of the gelsolin superfamily, SCIN has the core structure of gelsolin-like domains and is involved in vesicle transport, exocytosis via regulating the polymerization and disassembly of F-actin network [14-16]. In addition, SCIN also exhibits activities in regulating other cellular processes such as cell differentiation, osteoclastogenesis, among others [17-19]. The involvement of SCIN in tumor development was first investigated in the megakaryoblastic leukemia cells [20]. Miura et al. later found that SCIN expression is significantly increased in the cisplatin-resistant urothelial cancer patients by comparison with cisplatin-sensitive control group [21]. Knockdown of *SCIN* led to an increase of mitochondria-medi-

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ated cell apoptosis *in vitro*. Recently, the expression of *SCIN* has been shown to be up-regulated in human prostate cancer, lung carcinoma and gastric cancer, suggesting an oncogenic role of *SCIN* in regulating tumor growth [22-24]. However, the molecular mechanism accounting for abnormal *SCIN* expression in tumor remains unknown.

In our study, several lines of evidence demonstrated that *SCIN* expression is transcriptionally suppressed by *BRMS1* in HCC cells. The association relationship between *BRMS1* and *SCIN* expression in HCC tissues and cells was studied through western blot analysis and the transcriptional mechanism was illuminated through dual luciferase assay and ChIP experiment. To further investigate the function of *SCIN*, lentivirus carrying shRNA against *SCIN* or recombinant plasmid overexpressing *SCIN* were utilized. It is found that knockdown of endogenous *SCIN* sensitized HCC cells to apoptosis, whereas expression of *SCIN* protected cells from apoptotic cell death. Moreover, *SCIN* promoted HCC cell growth in xenografted tumor mice. Taken together, our findings characterized *SCIN* as another functional downstream effector of *BRMS1*, providing another molecular pathway accounting for *BRMS1*'s tumor suppressive role in HCC cells.

Materials and methods

Tumor specimens

Fresh surgical specimens of HCC, including tumor tissues and the neighboring pathologically nontumorous liver tissues, were obtained from liver cancer patients at Zhongshan Hospital, Shanghai, China. Written informed consent was obtained from all these patients. Tissue samples were immediately frozen in liquid nitrogen after surgery and later stored at -80°C.

Quantitative real-time PCR (qRT-PCR)

RNA extracted from tissues or cultured cells using TRIzol reagent (Life Technologies, USA) was used for reverse transcription using PrimeScript™ RT reagent and gDNA Eraser kit (Takara, Japan). Quantitative PCR analysis was performed with the CFX Connection detection system (Bio-Rad, USA) using SYBR Green Supermix kit (Takara). Cycle parameters were

95°C for 5 min hot start and 40 cycles of 95°C for 5 sec, 58°C for 10 sec and 72°C for 20 sec. Samples with no cDNA templates were used as negative control to rule out contamination in each run. Melting curves were analyzed to confirm the specificity of the PCR product. Primers for *BRMS1*: forward, 5'-ACTGAGTCAGCTGCG-GTTGCGG-3'; reverse, 5'-AAGACCTGGAGCTGCTC TGGCGTGC-3'. Primers for *SCIN*: forward, 5'-TCTGCGTTCCTGACTGTTCC-3'; reverse, 5'-GACCTCCTTCTTTGATGTTCC-3'. 18S rRNA was used as the internal control. Primers for 18S rRNA: forward, 5'-GTAACCCGTTGAACCCATT-3'; reverse, 5'-CCATCCAATCGGTAGTAGCG-3'.

Cell culture and transfection

HCC cell lines SK-Hep1, Huh7, QGY-7701, Hep-3B, SMMC-7721 and human embryonic kidney cell line 293T (HEK293T) were cultured with Dulbecco's modified Eagle's medium (DMEM), supplemented with 10% fetal bovine serum (Gibco, USA) in 5% CO₂ humidified atmosphere at 37°C. Cells of 70% confluency were transfected with indicated plasmids or siRNA using Lipofectin 2000 (Invitrogen, USA) according to the manufacturer's instructions. Small interfering RNA against *BRMS1* as previously described was used [9] and the target sequence of si-*SCIN* was: 5'-CGAGATGAGCTGACA-ACAT-3'. In experiments evaluating cells' sensitivity to apoptosis stimulation, etoposide (Sigma, USA) and cisplatin (Abcam, USA) were directly added to complete medium with indicated dosages.

Western blot analysis

Protein samples were separated by different concentrations of SDS-PAGE and then transferred to PVDF membranes. Membranes were incubated in 5% fat-free milk to avoid non-specific binding. After blocking, membranes were incubated with specific primary antibodies against different proteins at 4°C overnight. Membranes were then washed in TBST for four times and incubated with HRP-conjugated secondary antibody for 1 h at room temperature. Later on, membranes were washed in TBST for four times before being visualized by enhanced chemiluminescence (Pierce, USA) on a molecular imager ChemiDoc XRS+ system (Bio-Rad). Antibodies used include anti-*BRMS1* (Abcam), anti-*SCIN* (Abcam), anti-caspase 9 (cell signaling technology, USA), anti-β-actin (Sigma), anti-

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Myc (Sigma), anti-Flag (Sigma), peroxidase-conjugated goat anti-mouse IgG and goat anti-rabbit IgG (Jackson, USA).

Chromatin immunoprecipitation (ChIP)

The ChIP assay was performed according to our previous work [12]. Briefly, HEK293T cells were seeded at 1×10^7 /dish in 100 mm dishes and then transfected with Myc-tagged BRMS1 plasmid or empty vector. At 36 h after transfection, cells were treated with 4% formaldehyde (Amresco, USA) to make them crosslinked and fixed. Cell nuclei were first released by lysis buffer in pH 8.1 and then sonicated. Genomic DNA fragment was immuno-precipitated by anti-Myc antibody at 4°C overnight after treating with protein A-agarose beads. Samples containing mouse IgG were used as negative control to exclude genomic contamination. Treated genomic DNA fragments were collected, washed, de-crosslinked and then digested with Proteinase K. PCR was applied to examine the quantity of these DNA fragments recovered by phenol/chloroform extraction using the following primers: forward, 5'-TCGTCTCCTCTGTCAA-CT-3'; reverse, 5'-TTATTCGCCGCGCCACTTTA-3'.

Dual-luciferase assay

Cells were seeded into 24-well plates at a density of 1×10^5 per well and transfected with indicated SCIN promoter segments or pGL3-Basic empty vector. Internal control pRL-TK vector (20 ng/well) was used for normalization. Primers for SCIN promoter were: forward, CGGCTAGCTTGCCCAATGAAATGACAGA; reverse, CCAAGCTTAGTTGCGGTCCCCTTTACT. SCIN-P1 primers were: forward, CGGCTAGCTTGCCCAATGAAATGACAGA; reverse, CCAAGCTTTCTTTGTGCATACCCATCA. pGL3-SCIN promoter 2 primers were: forward, CGGCTAGCCCTGCTGCTCTCGGTTTAGT; reverse, CCAAGCTTAGTTGCGGTCCCCTTTACT; pGL3-SCIN promoter 3 primers were: forward, CGGCTAGCACAAAGGAGTGCCAGCAGT; reverse, CCAAGCTTGAGCAGCAGGAGAACCTTA.

Flow cytometry analysis

For cell cycle phase analysis, cells were harvested and resuspended in 0°C 70% ethanol to fix overnight. DNA was stained with propidium iodide (50 µg/mL) and treated with RNase (100 µg/mL) and then analyzed by FACSCalibur (BD

Biosciences, CA, USA). The apoptotic cell population corresponds to cells in sub-G1 phase. Each result was representative of three independent experiments with triplicate samples for each condition.

Cell proliferation assay

Cells were plated in 96-well plates at a density of 1500 cells/well. In the serum deprivation assay, medium was replaced by DMEM after 24 hours' culture with complete medium. Cell proliferation was detected using CCK8 assay. Generally, culture medium was replaced by 0.5 mg/mL CCK8 (Dojindo, Japan) diluted in DMEM. Cells were then kept in 37°C for 4 hours and absorbance at a wavelength of 490 nm was read by a microtiter reader (Bio-RAD, USA). The final growth curve was constructed using the absorbance data.

Tumor formation in nude mice

Three-week-old BALB/c female nude mice were obtained from Shanghai Laboratory Animal Co. Ltd. (SLAC, China) and maintained on standard laboratory chow. Cells were harvested and resuspended in DMEM to a final concentration of 2×10^7 cells/ml. Cells were then subcutaneously injected into each mouse. Tumors were visible around five days after injection. Mice were sacrificed till the average size of the tumors was around 800 mm³. Photos were taken, and weight of tumors was measured (n = 5).

Histological studies

Xenografted tumor tissues were embedded in paraffin and cut into sections (4 µm thickness) for immunohistochemistry studies. The avidin-biotin complex method was used in Ki-67 immunostaining. Briefly, endogenous peroxidase was blocked with H₂O₂. After BSA blocking, sections were incubated with anti-Ki-67 antibody (Abcam, USA) at 4°C overnight, followed by biotinylated secondary antibody. The sections were further treated with diaminobenzidine and counterstained with hematoxylin. TUNEL (terminal deoxynucleotide transferase mediated dUTP nick end labeling) assay was performed using the *in situ* Cell Apoptosis Detection kit (Boster, China) according to the manufacturer's instructions.

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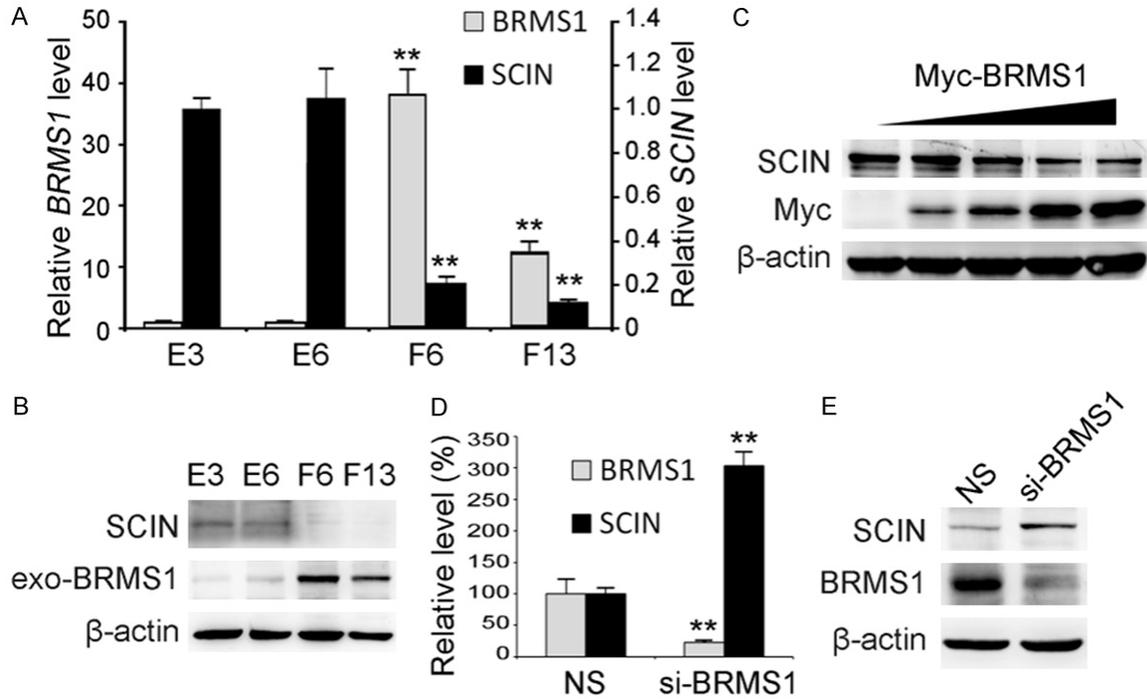


Figure 1. The expression level of SCIN was suppressed by BRMS1. (A) Relative *BRMS1* and *SCIN* mRNA levels in SK-Hep1 cell clones stably expressing exogenous *BRMS1* (F6 and F13) and control cell clones (E3 and E6) were analyzed through qRT-PCR. Values were normalized to internal control 18S rRNA and expressed as mean \pm SD, n = 3. (B) The protein levels of exogenous *BRMS1* (exo-*BRMS1*) and endogenous SCIN in SK-Hep1 cell clones were investigated through western blot. β -actin was used as a loading control. (C) Myc-tagged *BRMS1* (Myc-*BRMS1*) and endogenous SCIN protein levels in QGY-7701 cells transiently expressing different dosages of Myc-*BRMS1* were investigated through western blot. Relative longer exposure of endogenous SCIN is shown. Endogenous *BRMS1* and SCIN expression levels in Huh-7 cells infected with lentivirus expressing siRNA targeting *BRMS1* (siBRMS1) or non-silencing control siRNA (NS) were analyzed through qRT-PCR (D) and western blot (E).

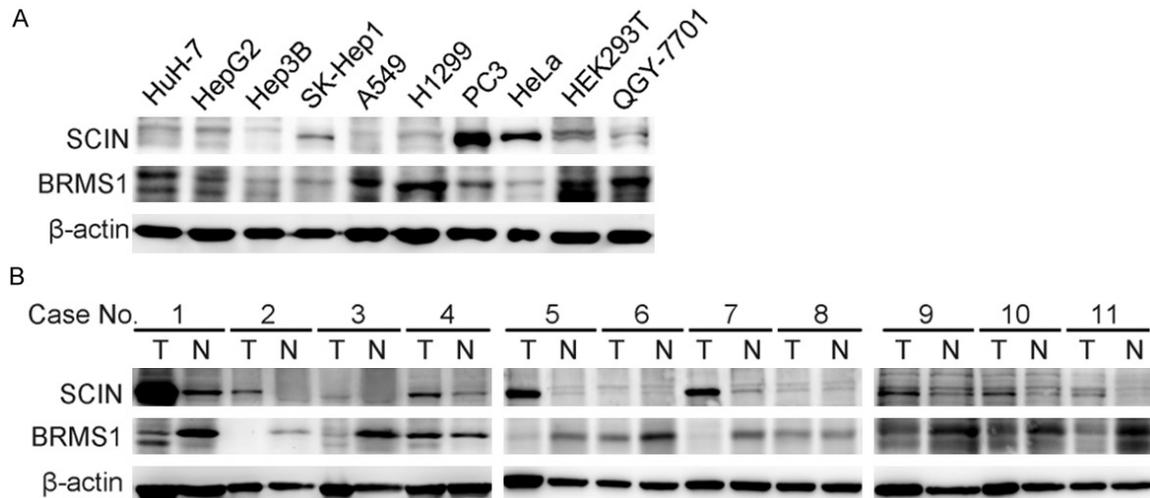


Figure 2. The protein level of SCIN was inversely correlated with BRMS1 level in HCC cells and tissues. A. Western blot analysis of BRMS1 and SCIN levels in HEK293T, five HCC cell lines (Huh-7, HepG2, Hep3B, SK-Hep1, QGY-7701), two lung carcinoma cell lines (A549, H1299), one prostate cancer cell line (PC3) and one cervical carcinoma cell line (HeLa). B. Western blot analysis of BRMS1 and SCIN levels in 11 paired HCC specimens. T, tumor tissues; N, corresponding non-tumorous tissues.

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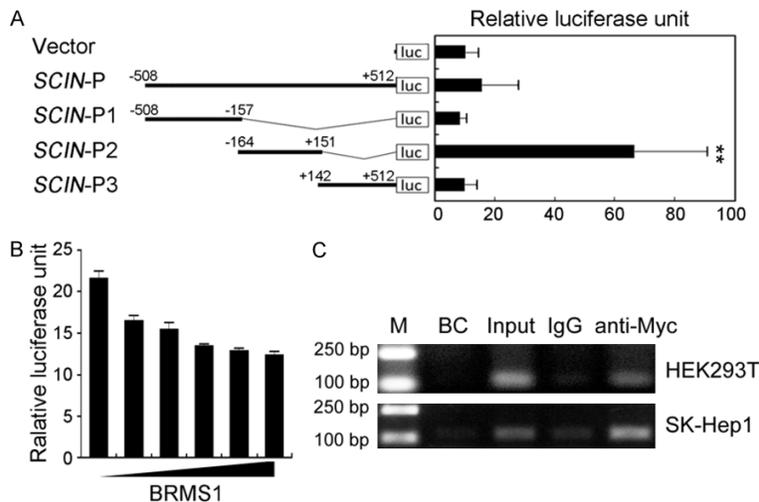


Figure 3. BRMS1 could transcriptionally suppress the activity of the promoter of SCIN. A. The full length SCIN promoter (SCIN-P) and three deletion mutants (SCIN-P1, P2 and P3) were constructed into luciferase reporter plasmid. Schematic representations of SCIN promoter constructs are presented in the left panel. Relative reporter activities in HEK293T cells were investigated through dual luciferase assay (right panel). Values are presented as mean \pm SD, $n = 3$. B. Relative reporter activities of SCIN-P2 were examined after co-transfection with different dosages of Myc-tagged BRMS1. Values are presented as mean \pm SD, $n = 3$. C. Gel electrophoresis photos of the ChIP assay to detect the binding ability between BRMS1 protein and SCIN promoter in HEK293T and SK-Hep1 cells. Soluble chromatin was prepared from cells transfected with Myc-tagged BRMS1 and immunoprecipitated by an anti-Myc antibody or IgG control. Specific primers were utilized in the PCR analysis. Blank controls (BC) with no DNA template were utilized to rule out contamination and soluble chromatin before immunoprecipitation was utilized as input control.

Statistical analysis

Categorical data were analyzed by Fisher's exact test. Comparisons of quantitative data were analyzed by Student's *t*-test. We considered two groups with a *p* value less than 0.05 to be different (*), and with a *p* value less than 0.01 to be significantly different (**).

Results

SCIN and BRMS1 levels were inversely correlated in HCC cells and tissues

Previously, we have applied microarray analysis of whole genome expression to screen for novel transcriptional targets of BRMS1 [12]. One rarely-studied gene, SCIN, was found to be suppressed in two stable SK-Hep1 cell clones overexpressing exogenous BRMS1 by comparison with control cell clones (Figure 1A). Western blot analysis result in consistent with this was shown in Figure 1B. To ensure that

SCIN expression was specifically impaired by BRMS1, different dosages of recombinant BRMS1 plasmid were transiently introduced into QGY-7701 cells. As shown in Figure 1C, endogenous SCIN expression was gradually decreased upon Myc-BRMS1 overexpression in a dosage-dependent manner. Furthermore, Huh-7 cells with relative higher endogenous BRMS1 levels were subjected to RNA interference. As shown in Figure 1D and 1E, both mRNA and protein levels of endogenous SCIN were increased after endogenous BRMS1 was successfully knocked down in Huh-7.

In order to further evaluate the relationship between BRMS1 and SCIN, different cancer cell lines and HCC tissues were investigated using western blot. As shown in Figure 2A, the BRMS1 and SCIN levels in most cancer cell lines were inversely correlated. In addition, we found

that SCIN protein was up-regulated while BRMS1 was mainly suppressed in HCC tissues (T) compared with the corresponding non-tumorous tissues (N). Of note, the expression pattern of SCIN was inversely correlated with BRMS1 in 7 paired HCC tissues such as H1, H2, H5, among others. This finding provides another piece of evidence that BRMS1 probably exerted a transcriptional suppression effect on SCIN expression during tumorigenesis.

BRMS1 was able to transcriptionally suppress SCIN expression

To investigate the regulatory mechanism between BRMS1 and SCIN, the promoter region (-508 to +512 bp) of human SCIN gene was cloned from human genomic DNA and a series of deletion mutants, including SCIN-P1 (-508 to -157 bp), SCIN-P2 (-164 to +151 bp), SCIN-P3 (+142 to +512 bp), were constructed subsequently. The above four DNA fragments were then inserted into pGL3-Basic vector to detect

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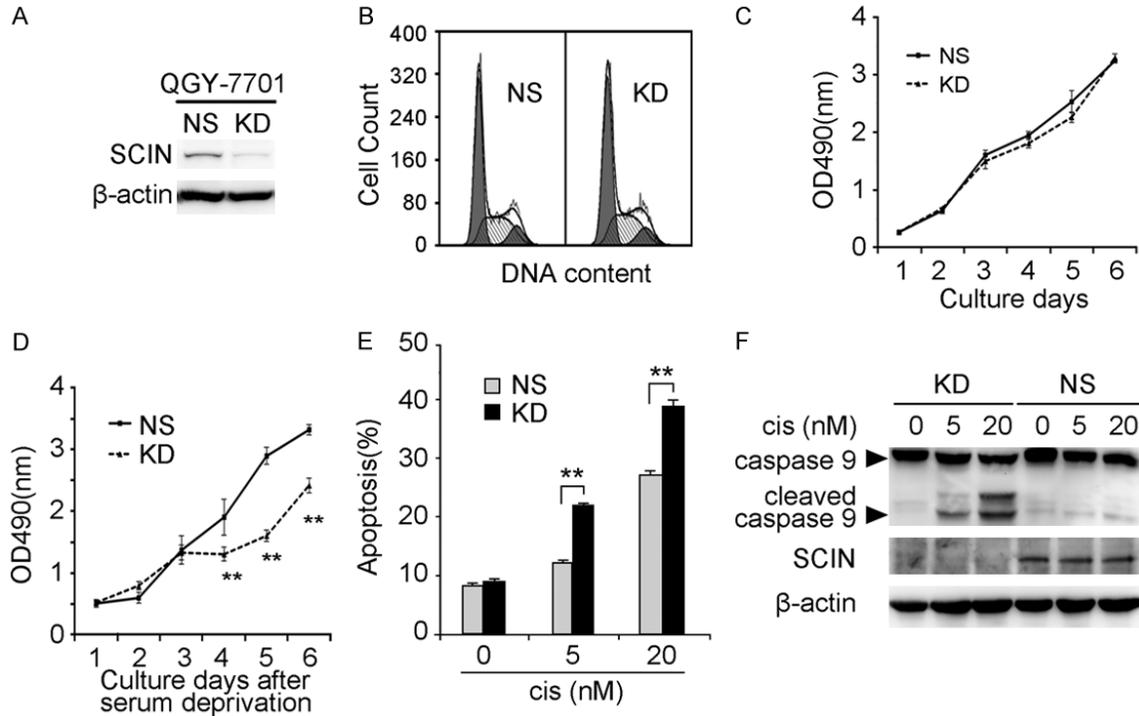


Figure 4. Effects of SCIN knockdown on QGY-7701 cells. (A) Endogenous SCIN level in QGY-7701 cells infected with lentivirus expressing shRNA targeting SCIN (KD) or non-silencing shRNA (NS) was investigated by western blot. β -actin was used as a loading control. (B) Representative flow cytometric images of lentivirus-infected QGY-7701 cells from cell cycle analysis. Cell growth curves of lentivirus-infected QGY-7701 cells under normal culture condition (C) and serum deprivation (D). Values are indicated as mean \pm SD, $n = 5$. Lentivirus-infected QGY-7701 cells treated with indicated dosages of cisplatin (cis) were subjected to cell apoptotic analysis through sub-G1 analysis (E) and caspase 9 immunoblot (F). Values are indicated as mean \pm SD, $n = 3$.

their transcriptional activity using luciferase assay. As shown in **Figure 3A**, *SCIN*-P2 exhibited the highest report gene's activity among all the promoter constructs. Next, when cells were co-transfected with different dosages of *BRMS1* plasmid, the luciferase activity of *SCIN*-P2 gradually decreased upon *BRMS1* expression in a dosage-dependent manner (**Figure 3B**). To further examine whether *BRMS1* could bind to this *SCIN* promoter region, chromatin immunoprecipitation was carried out with cells overexpressing myc-tagged *BRMS1*. As shown in **Figure 3C**, the +38 to +134 bp region of *SCIN*-P2 was specifically immunoprecipitated by anti-Myc antibody, but not IgG control, in both HEK293T and SK-Hep1 cells. Taken together, these data suggested a transcriptional suppression relationship between *BRMS1* and *SCIN*.

SCIN was involved in regulating HCC cells' sensitivity to apoptotic stimulus

To elucidate the biological functions of *SCIN* during HCC development, we utilized lentivirus-

mediated RNA interference in QGY-7701 cells. As shown in **Figure 4A**, the specific *SCIN*-targeting siRNA successfully knocked down endogenous *SCIN* expression (KD) by comparison with non-silencing siRNA control (NS). Next, we investigated several cellular behaviors in QGY-7701 cells under different conditions. No significant difference was found in the cell cycle distribution or cell growth under normal culture conditions (**Figure 4B, 4C**). However, when cells were deprived of serum, cell growth capacities were significantly impaired in *SCIN* knockdown cells compared with control cells (**Figure 4D**), indicating that *SCIN* knockdown might sensitize HCC cells to apoptotic stimulus. To confirm this hypothesis, the chemotherapeutic drug cisplatin was utilized in the apoptosis assay as reported previously [21]. Not surprisingly, flow cytometry analysis revealed a higher level of apoptosis in *SCIN* knockdown cells compared with control cells upon cisplatin treatment (**Figure 4E**). Moreover, cleavage of caspase 9 was specifically activated in *SCIN* knockdown cells, but not in the control cells,

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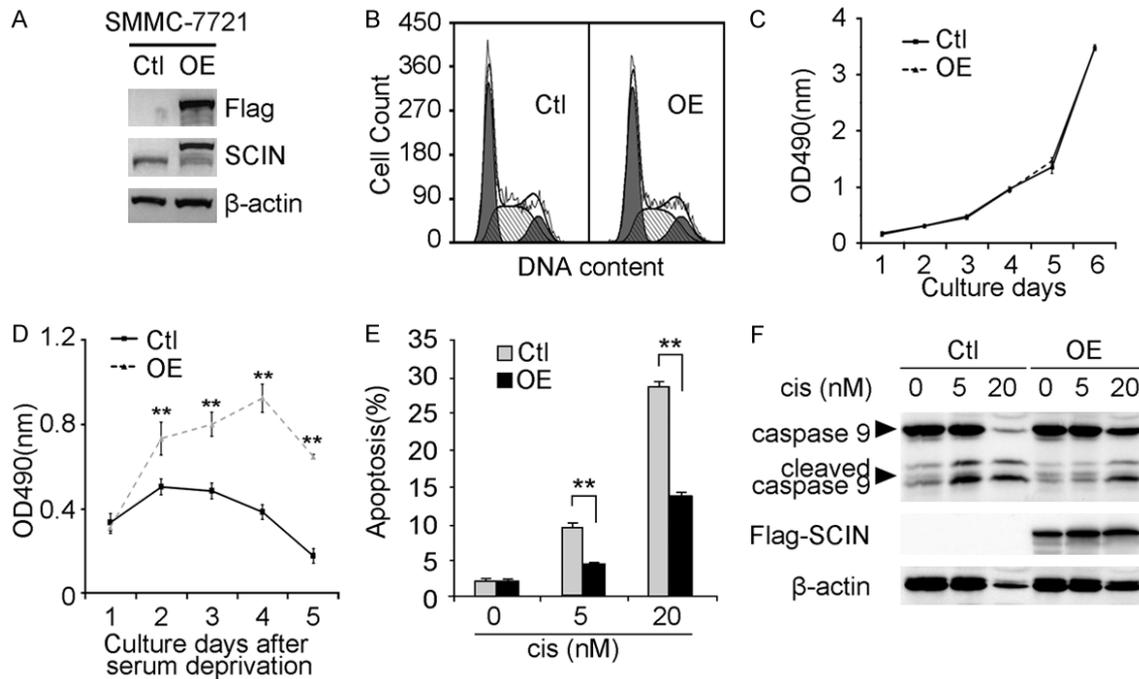


Figure 5. Effects of SCIN overexpression on SMMC-7721 cells. (A) Exogenous SCIN level in SMMC-7721 cells infected with lentivirus overexpressing flag-tagged SCIN (OE) or empty vector (Ctl) was investigated by western blot. β -actin was used as a loading control. (B) Representative flow cytometric images of lentivirus-infected SMMC-7721 cells determined from cell cycle analysis. Cell growth curves of lentivirus-infected SMMC-7721 cells under normal culture conditions (C) and serum deprivation (D). Values are indicated as mean \pm SD, $n = 5$. Lentivirus-infected SMMC-7721 cells treated with indicated dosages of cisplatin (cis) were subjected to cell apoptotic analysis through flow cytometry (E) and caspase 9 immunoblot (F). Values are indicated as mean \pm SD, $n = 3$.

after cisplatin treatment in a dosage-dependent manner (Figure 4F).

To further confirm the function of SCIN in regulating cell apoptosis, we established SMMC-7721 stable clone overexpressing flag-tagged SCIN (OE) and the control clone (Ctl) in parallel (Figure 5A). In consistent with the above findings, while no difference was observed in cell cycle distribution or cell growth under normal conditions between two clones (Figure 5B, 5C), cells overexpressing SCIN are less sensitive to serum deprivation compared with control cells (Figure 5D). Furthermore, SCIN overexpression protected SMMC-7721 clone from cisplatin-induced cell apoptosis and caspase 9 cleavage (Figure 5E, 5F). Collectively, these data strongly implied that SCIN level could modify HCC cells' sensitivity to apoptotic stimulus.

SCIN was able to regulate HCC cell growth *in vivo*

To explore the role of SCIN on HCC cell growth *in vivo*, tumorigenicity assay in BALB/c nude

mice was performed. We first subcutaneously injected equal amounts of QGY-7701 KD cells and NS cells into nude mice separately. Tumors were visible in NS group from eight days after injection and they increase in size stably. However, no visible tumor was observed in all mice injected with KD cells throughout the whole experiment (22 days in total) (Figure 6A). Next, we investigated the effect of SCIN overexpression on xenografted tumor growth. As shown in Figure 6B, SMMC-7721 OE cells display dramatically increased tumor growth in nude mice compared with Ctl cells. The average tumor weight of the OE cells was 3.47-fold heavier than that of Ctl cells.

Furthermore, xenografted tumor tissues were embedded in paraffin and subjected to immunohistochemistry studies. As shown in Figure 6C, there is no obvious difference in the fraction of Ki-67-positive cells, suggesting SCIN has less effect on regulating cell proliferation *in vivo*. By contrast, the TUNEL assay revealed a lower apoptotic cell population in OE group than in Ctl group (Figure 6D), indicating SCIN

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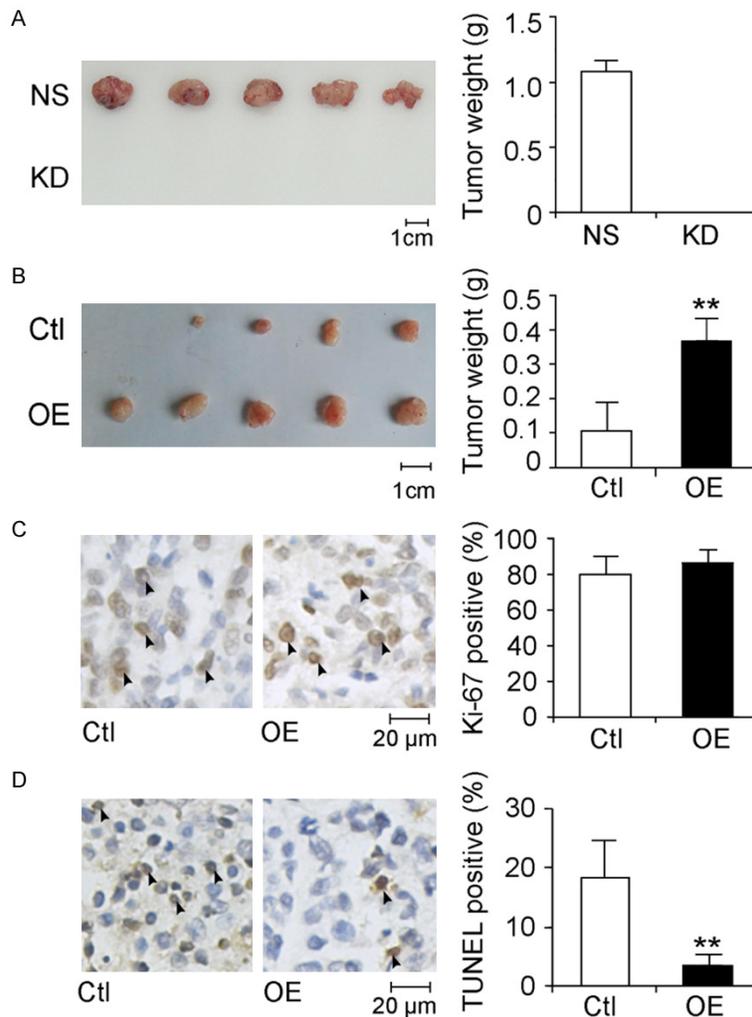


Figure 6. Evaluation of the function of SCIN on HCC cell growth in vivo. Photographs (left panels) and tumor weight measurements (right panels) of tumors dissected from nude mice engrafted with lentivirus-infected QGY-7701 cells (A) and SMMC-7721 cells (B). Scale bars, 1 cm. Values are indicated as mean \pm SD, $n = 5$. (C) Immunohistochemistry staining against Ki-67 was performed in tumor tissues derived from lentivirus-infected SMMC-7721. Representative images of tumor sections are given in the left panel and typical Ki-67-positive cells are indicated with arrowheads. Scale bar, 20 μ m. Ki-67 positive cell populations were statistically analyzed in the right panel. Values are indicated as mean \pm SD, $n = 8$. (D) TUNEL assays were performed on tumor sections. Representative images of tumor sections are given in the left panel, and typical apoptotic cells are indicated with arrowheads. Scale bar, 20 μ m. Apoptotic cell populations are statistically analyzed in the right panel. Values are indicated as mean \pm SD, $n = 8$.

overexpression protected cells from apoptosis *in vivo*.

Discussion

Our study identified SCIN as a novel transcriptional target of BRMS1 in HCC cells and SCIN was able to promote tumor cell growth through modulating the apoptotic levels in cells. Ac-

cumulating evidence demonstrated that BRMS1 has an essential role in regulating gene transcription mainly through interacting with Sin3•HDAC chromatin remodeling complexes or NF- κ B transcription factor [10, 25, 26]. Although BRMS1 suppresses several metastasis-related genes' expression through deacetylation of RelA/p65 subunit of NF- κ B, such as *OPN*, *uPA* and *CXCR4* [3, 26, 27], we ruled out the possibility that NF- κ B was involved in the transcriptional regulation of *SCIN* mainly because there is no NF- κ B binding site within the promoter of *SCIN*. On the other hand, Sin3•HDAC is a complex associated with a number of DNA-binding proteins and mediates transcriptional regulation of a variety of protein-coding genes and miRNAs implicated in diverse biological functions [28]. After the ChIP assay, which revealed a binding relationship between BRMS1 protein and the +38 to +134 bp region of *SCIN* promoter, we performed *in silico* prediction and found a putative GC binding factor (GCF) binding site (GCGCCGCT) within the binding region. GCF is able to bind to GC-rich sequence and suppress the expression of several tumor-related genes [29, 30]. It is interesting to note that the expression of the growth factor receptor (EGFR) gene is under the control of both GCF and Sin3•HDAC

[11, 30]. Together with our data, we raised the hypothesis that Sin3•HDAC might act in concert with GCF on *SCIN* expression. However, this still remains to be experimentally investigated in our future work.

Although the association of SCIN with cancer has been revealed for a long time, multiple reports addressing the role of SCIN in tumor

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development were controversial among different investigators. Expression of SCIN in megakaryoblastic leukemia cells induced cell apoptosis, inhibited cell proliferation and tumorigenesis [20]. By contrast, silencing of SCIN in gastric cancer cells suppressed cell invasion and tumor metastasis [24]. In addition, different changes of cell cycle distribution and different cyclins affected by SCIN were reported in 3 different studies [22, 23, 31]. Although heterogeneity of tumor cell lines might partially explain these disparities, more comprehensive studies are required to uncover the relationship between SCIN and cancer. In our present work, an anti-apoptotic role of SCIN has been revealed in HCC cells by different approaches both *in vitro* and *in vivo*, while no obvious changes were shown in cell cycle distribution or cell proliferation. Our finding was in consistent with the report in human bladder cancer cell, where SCIN was identified as the cisplatin-resistant marker via interacting with VDAC and further inhibiting cell apoptosis [21]. In addition, SCIN has 63% homology with gelsolin, and gelsolin superfamily has been demonstrated to play an important role in cell apoptosis by modulating dynamic actin assembly [32, 33]. Overexpression of gelsolin in Jurkat cells strongly inhibited caspase-3 activation and cell apoptosis level triggered by different reagents [34]. Consistently, gelsolin or villin knockout mice exhibited higher apoptotic cell population, shorter survival and enhanced caspase activation in liver after exposure to stimulatory Fas antibody [35-37]. Taken together, the results presented here highlight that SCIN might also be another important gelsolin superfamily member mediating the crosstalk between apoptosis pathway and cytoskeleton reorganization.

Furthermore, abnormal apoptosis has been demonstrated to affect HCC growth and metastasis [38, 39]. As we previously reported, BRMS1 also exhibits a strong activity in regulating HCC cell apoptosis, which further contributes to its suppressive role in tumorigenesis and tumor metastasis [5, 9]. A few proteins, including OPN, Bim and Bcl-2, have been demonstrated to partially account for the proapoptotic effect of BRMS1 in breast cancer cells [40, 41]. Given that overexpression of SCIN has a similar effect as knockdown of BRMS1 in regulating HCC cells' sensitivity to

apoptotic stimulus, we raised the hypothesis that SCIN might be another downstream molecule employed by BRMS1 in modulating cancer cell apoptosis pathway. Investigations of how these different transcriptional targets of BRMS1 collaborate to regulate cell apoptosis are therefore of great importance and interest in the future.

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Disclosure of conflict of interest

None.

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