**ABSTRACT**
Circular (circ)RNA is a special type of endogenous RNA consisting of a covalently closed loop structure without 5’ to 3’ polarity and a polyadenylated tail. Accumulating evidence suggests that circRNAs play important roles in the development and progression of human cancers. However, the role of circRNAs in the progression of hepatocellular carcinoma (HCC) is largely unknown. This was addressed in the present study using high-throughput sequencing to identify aberrantly expressed circRNAs in HCC patient tissue and cell lines. We found that circ-baculoviral IAP repeat-containing (BIRC)6 was upregulated in HCC tissue samples and cells; this was associated with the overall survival of HCC patients. circ-BIRC6 knockdown reduced HCC cell proliferation, migration, and invasion and enhanced their apoptosis. Additionally, circ-BIRC6 overexpression negatively regulated the expression of microRNA miR-3918, which was identified as an inhibitor of B cell lymphoma (Bcl)2. The tumor-suppressive effect of circ-BIRC6 deletion was abrogated by inhibiting miR-3918. These results indicate that circ-BIRC6 functions as a competing endogenous RNA that regulates Bcl2 expression by sponging miR-3918, and may serve as a prognostic biomarker and therapeutic target for the treatment of HCC.

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**Introduction**
Hepatocellular carcinoma (HCC) is one of the most common malignancies [1] and the second-leading cause of cancer-related death worldwide [2]. Despite recent progress in the development of therapeutic strategies, the prognosis of most HCC patients remains poor because they are typically diagnosed at advanced stages of the disease [3]. Clarifying the mechanisms underlying the malignant progression of HCC and identifying suitable biomarkers for early detection and for monitoring therapeutic efficacy can improve patient outcome.

Circular (circ)RNAs are a large class of endogenous non-coding (nc)RNAs that covalently link 5’ and 3’ ends of RNA to generate a circular structure [4], and play important roles in many biological processes [5]. Recent developments in high-throughput sequencing technology and bioinformatics approaches have permitted the identification and characterization of many circRNAs in human cells [6], with most being up- or downregulated in specific tissues [7]. circRNAs play an important role in the progression of various malignancies [8] including pancreatic [9], bladder [10], and cervical [11] cancers. circRNAs contribute to cancer development by acting as micro (mi)RNA sponges to regulate gene and protein expression [12], and may serve as biomarkers for diagnosing and predicting the prognosis of cancer patients [13]. As such, elucidating the mechanisms underlying circRNA function can provide a basis for the development of novel cancer treatments.

To this end, in this study we compared the expression profiles of circRNAs in HCC and normal tissues by high-throughput sequencing, and show that the circRNA circ-baculoviral IAP repeat-containing (BIRC)6 is upregulated in HCC tissues, which is associated with HCC patient prognosis. According to circBase [14], circ-BIRC6 (ID hsa_circ_0003288) is located on chromosome 2 (32703702–32718734) and is generated by back-splicing of the BIRC6 transcript (NM_016252). We found that circ-BIRC6 knockdown inhibited HCC cell proliferation, migration, and invasion and promoted their apoptosis; moreover, circ-BIRC6 acted as a sponge of the...
miRNA miR-3918, which targets the 3′ untranslated region (UTR) of B cell lymphoma (Bcl)2 transcript. These results indicate that circ-BIRC6 enhances Bcl2 expression by sponging miR-3918 and is a potential biomarker and therapeutic target for HCC treatment.

Materials and methods

Patient samples

HCC and matched adjacent non-tumor tissues (n = 55) were obtained from Qianfoshan Hospital of Shandong Province with patient consent and Ethics Committee approval. HCC was diagnosed by pathological examination. None of patients received chemotherapy, radiotherapy, or any other medical intervention before surgery. The tissue samples were flash frozen and stored in liquid nitrogen until use.

Cell culture and transfection

HepG2, Bel-7402, and Huh-7 human HCC cell lines were obtained from the American Type Culture Collection (Rockville, MD, USA). SMMC-7721 human hepatocarcinoma cells and the L02 normal human hepatic cell line were purchased from the Typical Culture Reserve Center of China (Shanghai, China). 293T cells were obtained from Chinese Academy of Science Typical Culture Collection (Shanghai, China). The cells were cultured in Dulbecco’s Modified Eagle’s medium (HyClone, Logan, UT, USA) supplemented with 10% fetal bovine serum (FBS; Gibco, Grand Island, NY, USA), 100 U/ml penicillin, and 100 μg/ml streptomycin in a humidified incubator at 37°C and 5% CO₂.

Small interfering (si)RNA against circ-BIRC6 (5′-CUGAAAGGUUCUUCACGT-3′) and a negative control siRNA along with circ-BIRC6 overexpression sequence (supplementary Table1), miR-3918 inhibitor (5′-AGUCUCCACUCGCGCCCUG U-3′) and mimic (5′-ACAGGGCCGACAUGG AGACU-3′) were purchased from GenePharma (Shanghai China). The siRNAs and miRNAs were transfected into HCC cells using Lipofectamine 2000 reagent (Invitrogen) according to the manufacturer’s protocol.

Cell proliferation assay

Cell proliferation was evaluated with Cell Counting Kit (CCK)-8 (Dojindo Laboratories, Kumamoto, Japan). Transfected cells were seeded in a 96-well plate at a density of 5 × 10³ cells/well and cultured for 24, 48, 72, and 96 h. A 10-μl volume of CCK-8 reagent was added to each well. The absorbance at 450 nm was measured on a Spectra Max 190 instrument (Molecular Devices, Sunnyvale, CA, USA). Experiments were performed with triplicate samples.

Transwell migration and invasion assays

A transwell cell migration system (8-μm pore size; BD Biosciences, Franklin Lakes, NJ, USA) and 24-well transwell plates with Matrigel (BD Biosciences) were used to evaluate cell migration and invasion, respectively. Transfected cells were resuspended in serum-free medium to a density of 2.0 × 10⁵cells/ml. A 200-μl aliquot of cells was added to the upper chamber of the transwell insert, and 600 μl complete medium containing 10% FBS were added to the lower well, followed by incubation at 37°C and 5% CO₂ for 24 h. The cells were fixed with methanol and stained with 0.1% crystal violet. The cells that passed through the filter were photographed and counted under a microscope.

Flow cytometry analysis

After transfection for 48 h, HepG2 and Bel7402 cells (2 × 10⁵ cells per well) were collected and stained with Annexin V-fluorescein isothiocyanate (FITC) and propidium iodide (PI) (Dojindo Laboratories) according to the manufacturer’s instructions. Apoptosis was assessed by flow cytometry (FACSAria II; BD Biosciences).

Fluorescence in situ hybridization (FISH)

FISH was performed to detect the presence of circ-BIRC6 using a Cy3-labeled probe (5′-CY3-ACCTTCAACTGAAAAGGTCTTGCACGCT-3′) and miR-3918 using a FAM-labeled probe (5′-FAM-AGTCTCCATCTGCGGCCCTGT-FAM-3′) respectively. FISH probes were designed by Servicebio.
Cells were cultured on coverslips until they reached 60%–70% confluence, then washed three times with phosphate-buffered saline (PBS) for 5 min and fixed with 4% paraformaldehyde for 15 min at room temperature. After washing with PBS, the cells were incubated overnight at 37°C with the FISH probes in hybridization buffer in a humid chamber according to the manufacturer’s protocol. The following day, the cells were washed and stained with 4’,6-diamidino-2-phenylindole for 10 min in the dark. Images were obtained using a fluorescence microscope (Hitachi, Tokyo, Japan).

Reverse transcription and quantitative real-time (qrt)-pcr

RNA was extracted from tissues or cells using TRIzol reagent (Invitrogen) according to the manufacturer’s protocol. Total RNA was reverse transcribed using the FastQuant RT kit (Tiangen Biotech, Beijing, China) according to the manufacturer’s protocol, and qRT-PCR analysis of circ-BIRC6, Bcl2, and miR-3918 expression was performed using SYBR Select Master Mix (Tiangen Biotech) on an ABI ViiA7 system (Applied Biosystems, Foster City, CA, USA), with glyceraldehyde 3-phosphate dehydrogenase (GAPDH) and U6 used as internal controls. The following forward and reverse primers were used for qRT-PCR:
- circ-BIRC6, 5’-TGAAAGGTTCCTTGCACGCAT-3’ and 5’-GCTGGGGTTCGTCCACACATC-3’; BI RC6 mRNA, 5’-TCAACCTCCGCCTACACAGTCG-3’ and 5’-ACCGCCTCCATCCACCTCTTC-3’;
- Bcl2 mRNA, 5’-GCATTTCCGCCAGATGTCAG-3’ and 5’-CGTTTCAGTACTCAGTCACTCC-3’;
- miR-3918, 5’-ACAGGGCCGAGATGG-3’; and U6, 5’-ACGCAAATTCGTGAAGCGTTC-3’.

Western blot analysis

HCC cells were lysed in cold radioimmunoprecipitation lysis buffer (Beyotime Institute of Biotechnology, Shanghai, China) containing 1 mM phenylmethylsulfonyl fluoride. Protein concentration was quantified with a BCA Protein Assay kit (Beyotime Institute of Biotechnology). Equal amounts of protein were separated by 10% sodium dodecyl sulfate polyacrylamide gel electrophoresis and transferred to a polyvinylidene difluoride membrane (Millipore, Billerica, MA, USA) that was incubated overnight at 4°C in a solution of non-fat milk and antibody (1:1000 dilution; Abcam, Cambridge, MA, USA), followed by incubation with horseradish peroxidase-conjugated anti-rabbit IgG secondary antibody (Beyotime Institute of Biotechnology) for 2 h at room temperature. Protein bands were visualized with enhanced chemiluminescence detection reagent (Millipore), and signal intensity was quantified using Image Lab software (Bio-Rad, Hercules, CA, USA).

Luciferase reporter assay

The sequences of circ-BIRC6 and Bcl2 3’ UTR containing wild-type (WT) or mutant (Mut) miR-3918 binding sites were cloned into the luciferase vector (Promega). The generated vectors were transfected into 293T cells along with miR-NC and miR-3918 mimic using Lipofectamine 2000. The Dual-luciferase Reporter Assay (E1910; Promega, Madison, WI, USA) was used to measure luciferase activity in cell lysates 48 h after transfection.

Rna-binding protein immunoprecipitation (RIP) assay

The RIP assay was performed using the EZMagna RIP kit (Millipore) according to the manufacturer’s protocol. Cells were lysed in RIP lysis buffer, followed by incubation with RIP buffer containing magnetic beads conjugated with anti-human Argonaute (Argo)-2 antibody (Millipore) or normal mouse IgG (Millipore) as a negative control. Immunoprecipitated proteins were digested with proteinase K and the associated RNA was purified and analyzed by qRT-PCR using appropriate primers.

Tumor xenograft model

Animal experiments were approved by the Animal Care and Use Committee of Shandong University. Short hairpin RNAs (shRNAs)
targeting circBIRC6 (5′-TTTCAACTGAAAG GTTCTTGCAC-3′), negative control (NC) (5′- TTCTCCGAACGTGTCACGTTTC-3′), over-expressed circ-BIRC6 (oe-circ-BIRC6) (supplementary Table1) and corresponding negative control (Vector) were cloned into the lentiviral vectors and lentiviral supernatant was collected to infected Bel7402 cells. Stably transfected Bel7402 cells were subcutaneously injected into the flank of 4-week-old BALB/c nude mice. Tumor size was measured every 3 days using calipers and tumor volume was calculated using the formula length × width²/2. After 4 weeks, mice were sacrificed and tumor weight was measured.

Statistical analysis

Data were analyzed with SPSS v.17.0 software (IBM, Armonk, NY, USA) and are shown as mean ± SD. The Student’s t test was used to compare the means of two groups, and one-way analysis of variance was used for multiple-group comparisons. The χ² test or Fisher’s exact test was used to analyze the relationship between circ-BIRC6 and miR-3918 expression levels and the clinicopathological features of HCC patients. Correlations were evaluated using Pearson’s correlation coefficient. Survival was evaluated by Kaplan–Meier analysis and compared with the log-rank test. P < 0.05 was considered statistically significant.

Results

Circ-BIRC6 upregulation in HCC is associated with HCC patient prognosis

To investigate the relationship between circRNAs and HCC progression, we performed high-throughput sequencing and found that many circRNAs were up- or downregulated in HCC as compared to adjacent normal tissue (Figure 1(a)). One of these, circ-BIRC6, was overexpressed in HCC tissue; this was confirmed by qRT-PCR analysis of HCC and normal tissues (Figure 1(b)). To further examine the correlation between circ-BIRC6 expression and HCC severity, we assessed circ-BIRC6 expression in HCC at different stages of the disease. circ-BIRC6 level was associated with tumor-node-metastasis (TNM) stage in HCC (Figure 1(c)), and was higher in samples with as compared to those without vascular invasion (Figure 1(d)). A qRT-PCR analysis confirmed that circ-BIRC6 expression was higher in HCC cell lines – i.e., HepG2, Bel-7402, SMMC-7721, and Huh7 cells – than in normal L02 cells (Figure 1(e)). Given its upregulation in HCC tissue, we investigated whether circ-BIRC6 expression is associated with HCC patient prognosis. The Kaplan-Meier analysis using median circ-BIRC6 level as the cut-off point showed that HCC patients with high circ-BIRC6 expression had shorter overall survival (OS) than those with low circ-BIRC6 expression (Figure 1(f)). These data indicate that circ-BIRC6 is aberrantly expressed in HCC tissue and may serve as a biomarker for poor prognosis.

Relationship between circ-BIRC6 expression and clinicopathological features of HCC patients

Given that high circ-BIRC6 expression was correlated with poor prognosis in HCC patients, we analyzed the relationship between circ-BIRC6 level and clinicopathological features of HCC patients including age, gender, serum alpha-fetoprotein level, tumor size, vascular invasion, and TNM stage. circ-BIRC6 expression was associated with tumor stage (P = 0.0143) and vascular invasion (P = 0.0151), but was unrelated to other clinical characteristics (Table 1).

Uni- and multivariate Cox regression analyses of clinical variables showed that circ-BIRC6 expression (P = 0.005), vascular invasion (0.006), tumor size (0.045), Edmondson-Steiner histological grade (P = 0.024), and TNM stage (P = 0.001) were associated with the OS of HCC patients (Table 2). We performed a multivariate analysis to assess whether circ-BIRC6 expression is an independent prognostic factor and found that circ-BIRC6 level along with vascular invasion and TNM stage was associated with OS. These data imply that circ-BIRC6 contributes to HCC progression.

Circ-BIRC6 knockdown inhibits cell proliferation, migration, and invasion and promotes apoptosis in HCC cells

We investigated the function of circ-BIRC6 in HCC using HepG2 and Bel7402 cells transfected with siRNA against circ-BIRC6. A qRT-PCR
analysis showed that circ-BIRC6 expression was downregulated in transfected HepG2 and Bel7402 cells, while BIRC6 mRNA had no significant changes, which indicated si-circ-BIRC6 specifically targeted the circRNA transcript but not the linear RNA one (Figure 2(a)). We also evaluated cell proliferation with the CCK-8 assay and found that circ-BIRC6 knockdown suppressed cell proliferation (Figure 2(b)). Additionally, circ-BIRC6 silencing enhanced HepG2 and Bel7402 cell apoptosis, as determined by flow cytometry (Figure 2(c)) and inhibited migration and invasion in the transwell assays (Figure 2(d,e)). These results indicate that circ-BIRC6 plays an oncogenic role in HCC by positively regulating cancer cell proliferation, migration, and invasion and negatively regulating apoptosis.

Table 1. Relationship between circ-BIRC6 expression and clinicopathological factors.

| Characteristics     | Low expression (n = 27) | High expression (n = 28) | P value |
|---------------------|------------------------|--------------------------|---------|
| Sex                 |                         |                          |         |
| Female              | 11                      | 14                       | 0.5913  |
| Male                | 16                      | 14                       |         |
| Age (year) ≤ 60     | 14                      | 17                       | 0.5913  |
| > 60                | 13                      | 11                       |         |
| Serum AFP Negative  | 16                      | 11                       | 0.1810  |
| Positive            | 11                      | 17                       |         |
| Tumor size ≤ 5cm    | 18                      | 13                       | 0.1765  |
| > 5cm               | 9                       | 15                       |         |
| TNM stage I and II  | 20                      | 11                       | 0.0143  |
| III and IV          | 7                       | 17                       |         |
| vascular invasion   |                         |                          | 0.0151  |
| Negative            | 19                      | 10                       |         |
| Positive            | 8                       | 18                       |         |
**Circ-BIRC6 overexpression promotes proliferation, migration, and invasion while inhibiting apoptosis in HCC cells**

Based on the above results, we speculated that circ-BIRC6 overexpression in HepG2 and Bel7402 cells would enhance their malignant character. Upregulation of circ-BIRC6 in cells transfected with a circ-BIRC6 overexpression vector was confirmed by qRT-PCR (Figure 3(a)), circ-BIRC6 overexpression increased cell proliferation (Figure 3(b)) as well as migration and invasion (Figure 3(d,e)) while inhibiting apoptosis (Figure 3(c)), as determined with the CCK-8 assay, transwell assays, and by flow cytometry analysis, respectively. These results confirm that circ-BIRC6 has an oncogenic function in HCC.

**Circ-BIRC6 directly targets miR-3918**

circRNAs can act as an miRNA sponge in cancer [15]. To investigate the mechanism by which circ-BIRC6 promotes HCC progression, we screened potential targets of circ-BIRC6 by in silico analysis and identified miR-3918 as a candidate molecule. FISH experiments showed that circ-BIRC6 and miR-3918 were primarily localized in the cytoplasm of HepG2 cells (Figure 4(a)), suggesting that the two interact. A putative miR-3918 binding site was identified in the circ-BIRC6 sequence by bioinformatics analysis (Figure 4(b)); miR-3918 mimic suppressed luciferase activity in 293T cells transfected with the WT circ-BIRC6 sequence, while mutation of the predicted circ-BIRC6 binding site abrogated this effect (Figure 4(c)).

miRNAs suppress gene expression by forming the RNA-induced silencing complex (RISC) that includes Argonaute (Ago)2. We investigated whether circ-BIRC6 and miR-3918 are present in the same RISC complex with the RIP assay and found that both molecules were enriched in the complex immunoprecipitated using the anti-Ago2 antibody as compared to control IgG (Figure 4(d)). Furthermore, circ-BIRC6 knockdown enhanced miR-3918 expression in HepG2 and Bel7402 cells, as determined by qRT-PCR (Figure 4(e)). These results imply that circ-BIRC6 and miR-3918 directly interact.

We then analyzed miR-3918 expression in HCC and normal tissues and found that it was reduced in the former relative to the latter (Figure 4(f)). Moreover, miR-3918 level was inversely correlated with that of circ-BIRC6 in HCC tissue specimens (Figure 4(g)). We also analyzed the correlation between miR-3918 expression and clinicopathological features of HCC and found that it was associated with tumor stage (P = 0.0306), although it was unrelated to other clinical features (Supplementary Table 2).

**miR-3918 plays an anti-tumorigenic role in HCC**

To confirm the role of miR-3918 in HCC, HepG2 and Bel7402 cells were transfected with miR-3918 inhibitor or the negative control. The efficiency of miR-3918 silencing was confirmed by qRT-PCR (Supplementary Figure 1A). miR-3918 inhibition enhanced cell proliferation (Supplementary Figure 1B) as well as migration and invasion (Supplementary Figure 1D, E) while inhibiting apoptosis (Supplementary Figure 1C), which is similar to the effects of circ-BIRC6 overexpression.

To confirm the role of miR-3918 in HCC, HepG2 and Bel7402 cells were transfected with miR-3918 mimic (Supplementary Figure 2A). The results of the CCK-8 assay, flow cytometry analysis, and transwell assay showed that miR-3918 overexpression

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**Table 2. Univariate and multivariate cox regression analyses circ-BIRC6 for OS of patients in the cohort.**

| Variables                        | Univariate analysis | Multivariate analysis |
|----------------------------------|---------------------|-----------------------|
| HR (95%CI) HR (95%CI)            | P values            | P values              |
| Circ-BIRC6 (High vs. low)        | 2.820(1.225–5.026)  | 0.005                 |
| Age (>60 vs. ≤60 y)              | 1.001(0.974–1.029)  | 0.930                 |
| Gender (M vs. F)                 | 1.393(0.675–2.877)  | 0.370                 |
| HBsAg (positive vs. negative)    | 1.526(0.757–3.078)  | 0.237                 |
| AFP (>20ng/ml vs. <20ng/ml)      | 1.754(0.886–3.473)  | 0.107                 |
| Vascular invasion (positive vs. negative) | 2.727(1.324–5.615) | 0.006                 |
| Tumor size (>5cm vs. ≥5cm)       | 2.025(1.016–4.036)  | 0.045                 |
| Edmonson grade (IV+III vs. I+ II) | 2.207(1.110–4.389)  | 0.024                 |
| TNM stage (III+IV vs. I+ II)     | 3.098(1.543–6.217)  | 0.001                 |

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**Table 2. Univariate and multivariate cox regression analyses circ-BIRC6 for OS of patients in the cohort.**

| Variables                        | Univariate analysis | Multivariate analysis |
|----------------------------------|---------------------|-----------------------|
| HR (95%CI) HR (95%CI)            | P values            | P values              |
| Circ-BIRC6 (High vs. low)        | 2.343(1.088–5.044)  | 0.030                 |
| Age (>60 vs. ≤60 y)              | -                   | -                     |
| Gender (M vs. F)                 | -                   | -                     |
| HBsAg (positive vs. negative)    | -                   | -                     |
| AFP (>20ng/ml vs. <20ng/ml)      | -                   | -                     |
| Vascular invasion (positive vs. negative) | 2.530(1.100–5.814)  | 0.029                 |
| Tumor size (>5cm vs. ≥5cm)       | 1.188(0.532–2.653)  | 0.675                 |
| Edmonson grade (IV+III vs. I+ II) | 1.059(0.446–2.519)  | 0.896                 |
| TNM stage (III+IV vs. I+ II)     | 2.879(1.258–6.590)  | 0.012                 |
suppressed cell proliferation (Supplementary Figure 2B) and cell migration and invasion (Supplementary Figure 2D, E) while promoting apoptosis (Supplementary Figure 2C). These results indicate that miR-3918 and circ-BIRC6 have opposing functions in HCC.
**Figure 3.** circ-BIRC6 overexpression promotes HCC cell proliferation, migration, and invasion and inhibits apoptosis. (a) Relative expression of circ-BIRC6 in HepG2 and Bel7402 cells transfected with overexpression and empty vector. oeCirc, circBIRC6 overexpression vector; oeVec, control overexpression vector. (b) circ-BIRC6 overexpression promotes the proliferation in HepG2 and Bel7402 cells, as determined with the CCK-8 assay. (c) Cell apoptosis was inhibited in the circ-BIRC6 overexpression group relative to control cells in the flow cytometry analysis. (d, e) Migration and invasion of HepG2 and Bel7402 cells transfected with oeCirc or oeVec were evaluated with the transwell assay. Data are presented as mean± SD. *P < 0.05, **P < 0.01, ***P < 0.001.
circ-BIRC6 promotes Bcl2 expression by sponging miR-3918

miRNAs regulate gene expression by targeting the 3' UTR of mRNAs. We identified Bcl2 as a potential target of miR-3918 by bioinformatics analysis based on the presence of a putative miR-3918 binding site in the 3' UTR of the Bcl2 transcript (Figure 5(a)). miR-3918 overexpression reduced the luciferase activity of the WT Bcl2 3' UTR in 293T cells but had no effect when the predicted binding site was mutated (Figure 5(b)). Furthermore, miR-3918 overexpression and circ-BIRC6 silencing reduced Bcl2 mRNA (Figure 5(c, d)) and protein (Figure 5(e,f)) levels in HepG2 and Bel-7402 cells, as determined by qRT-PCR and western blotting, respectively. Conversely, miR-3918 inhibition combined with circ-BIRC6 overexpression had the opposite effects (Figure 5(g–j)). Furthermore, Bcl2 mRNA expression was negatively correlated with that of miR-3918 and positively correlated with that of circ-BIRC6 in HCC tissues (Figure 5(k,l)). These data suggest that circ-BIRC6 promotes Bcl2 expression by sponging miR-3918.

miR-3918 inhibition reverses the effect of circ-BIRC6 silencing

To verify whether circ-BIRC6 promotes HCC progression via regulation of miR-3918 and Bcl2, miR-3918 expression was suppressed in HCC cells transfected with an siRNA against circ-BIRC6. miR-3918 inhibition increased cell proliferation in circ-BIRC6-depleted HepG2 and Bel7402 cells (Figure 6(a)). miR-3918 inhibition decreased the expression of miR-3918 (Figure 6(b)); this was accompanied by an upregulation on mRNA and protein level of Bcl2 (Figure 6(c,g)). Moreover, miR-3918 inhibition abolished the effects of circ-BIRC6 knockdown on cell migration, invasion, and apoptosis (Figure 6(d–f)).
These data demonstrate that circ-BIRC6 promotes the malignant progression of HCC by modulating the miR-3918/Bcl2 axis.

**Discussion**

circRNAs have been implicated in a variety of biological processes [16] and human cancers [17]; they have been shown to play an important role in cancer initiation and progression and may function as...
Figure 6. miR-3918 inhibition reverses the effect of circ-BIRC6 depletion. (a) Cell proliferation was evaluated with the CCK-8 assay. (b, c) miR-3918 and Bcl2 expression in HepG2 and Bel7402 cells, as determined by qRT-PCR. (d) Detection of apoptosis by flow cytometry. (e, f) Migration and invasion of HepG2 and Bel7402 cells, as determined with the transwell assay. (g) The protein expression of Bcl2 in HepG2 and Bel7402 cells, as determined by western blotting. Data are presented as mean ± SD. *P < 0.05, **P < 0.01, ***P < 0.001.
molecular markers for cancer diagnosis and treatment [18]. For instance, the circRNA protein arginine methyltransferase 5 promotes metastasis of urothelial carcinoma of the bladder by sponging miR-30c to induce epithelial-to-mesenchymal transition [19], while RNA hsa_circ_0000977 silencing suppressed pancreatic ductal adenocarcinoma progression by stimulating miR-874-3p and inhibiting polo-like kinase 1 expression [20]. Downregulation of hsa_circ_0011946 suppressed the migration and invasion of MCF-7 breast cancer cells by targeting replication factor C subunit 3 [21]. In HCC, circMTO1 acts as a sponge of the miRNA miR-9 to suppress HCC progression [22]. Our study identified a novel circRNA – circ-BIRC6 – that plays an important role in HCC progression.

HCC is a common and aggressive cancer that is linked to a large number of cancer-related deaths worldwide [2]. Despite therapeutic advances, the prognosis of patients remains poor. In our study, we found that circ-BIRC6 is upregulated in HCC tissue and cell lines. Importantly, this was associated with the prognosis of HCC patients, with a multivariate Cox analysis revealing that circ-BIRC6 was an independent factor predicting poor outcome. circ-BIRC6 silencing inhibited the proliferation, migration, and invasion of HCC cells while promoting their apoptosis; circ-BIRC6 overexpression had the opposite effects. circRNAs function as miRNA sponges that modulate mRNA stability and regulate gene transcription or protein translation [12]. Previous studies showed that miRNAs plays a critical role in the development and progression of gastric [23], lung [24], colorectal [25], and cervical [26] cancers. We investigated whether circ-BIRC6 acts as a sponge of miR-3918, which was identified as a potential target of circ-BIRC6. We found that circ-BIRC6 and miR-3918 are enriched in the cytoplasm of HCC cells by FISH analysis, suggesting a functional interaction. The luciferase activity reporter and RIP assays confirmed the direct binding between circ-BIRC6 and miR-3918. circ-BIRC6 silencing also increased miR-3918 level, as determined by qRT-PCR. Although its precise function is unknown, miR-3918 was downregulated in HCC tissue and there was an inverse correlation between circ-BIRC6 and miR-3918 expression: when miR-3918 was overexpressed, HCC cell proliferation, migration, and invasion were suppressed and apoptosis was enhanced, whereas miR-3918 inhibition had the opposite effects, suggesting that miR-3918 has a tumor-suppressive role. We identified Bcl2 as a potential target of miR-3918, and
confirmed their interaction with the luciferase reporter assay. circ-BIRC6 knockdown and miR-3918 overexpression reduced Bcl2 mRNA and protein levels, and miR-3918 inhibition reversed the decreases in HCC cell proliferation, migration, and invasion and increase in apoptosis induced by circ-BIRC6 silencing. Thus, circ-BIRC6 promotes HCC progression via a regulatory axis that includes miR-3918 and Bcl2.

In conclusion, our study demonstrates for the first time that circ-BIRC6 is upregulated in HCC tissues and is associated with poor prognosis of HCC patients, and promotes HCC progression by targeting the miR-3918/Bcl2 axis. Based on these findings, circ-BIRC6 may be a useful prognostic biomarker in HCC and a promising therapeutic target for disease treatment.

Disclosure statement
No potential conflict of interest was reported by the authors.

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Ethical approval
This study was approved by the Ethics Committee on Research Involving Human Subjects of Faculty of Medicine, Qianfoshan Hospital, Shandong University.

Informed consent
Informed consent was obtained from all participants in the study.

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