Osteosarcoma, one of the most frequent primary malignant tumours, is regarded as a life-threatening solid malignancy of bone for children and adolescents predominantly. Osteosarcomas originate from primitive mesenchymal bone cells and usually occur in long bones instead of soft tissue.\(^1,2\) The morbidity of osteosarcomas is almost 11 million in the pediatric population. The combination of surgery, radiotherapy and chemotherapy improves the prognosis of 60%-75% patients with OS.\(^3,4\) However, a large portion of OS patients with pulmonary metastasis or relapse have a dismal prognosis on account of untimely diagnosis and drug resistance.\(^5\) Therefore, it is in urgent need to explore the molecular mechanism of tumour genesis, metastasis and drug resistance, which might provide novel strategy for clinical treatment and improve the outcome of advanced OS patients.

Circular RNAs (circRNAs), one kind of endogenous non-coding RNA, are characterized by closed continuous loop in shape, and
without 5'–3' polarity and polyA tail, which give them strong resistance to degradation from enzyme. Through the biological function of circRNAs is ignored in the past, emerging evidences have indicated that circRNAs play an important role in the various processes including physiological development and disease occurrence. The function of circRNAs in the tumorigenesis are multidinous. Most studied circRNA ciRS-7 was reported to exhibit high expression profile in different human cancers and could regulate the tumorigenesis by sponging for miR-7. CircFoxo3 has been demonstrated to function as proteins scaffolds through binding to ubiquitin enzyme MDM2 and its substrates p53 to inhibit angiogenesis and cell cycle progression. As to osteosarcoma, a series of differentially expressed circRNAs have been found by microassay expression analysis of circRNAs in OS patients. However, the molecular mechanism of circRNA in OS has not been uncovered.

In our study, we implicated a novel circRNA hsa_circ_0002137, whose expression was abnormally up-regulated in osteosarcoma tumours compared with normal bone tissue. Furthermore, the role of circ_0002137 was investigated via down- or up-regulated experiments and found that circ_0002137 inhibition suppressed cell invasion and cell proliferation of OS cell lines, meanwhile we found knockdown of circ_0002137 could induce more apoptotic OS cells. Furthermore, the correlation between circ_0002137 and miR-433-3p was predicted and confirmed through RNA pull-down assay and dual-luciferase reporter assay. Interestingly, we found that the inhibitory effect of circ_0002137 on OS was dependent of IGF1R. In conclusion, it was demonstrated that circ_0002137 could restrain the progression of OS through regulating miR-433-3p/IGF1R axis, providing a comprehensive landscape of circ_0002137 in the tumorigenesis and development of OS.

2 MATERIALS AND METHODS

2.1 Clinical samples collection

All clinical samples were collected from 15 untreated OS patients in the Affiliated Huaian NO.1 people's Hospital of Nanjing Medical University after surgical resection. All patients presented at the hospital. They all were provided written consent authorizing the use of clinical samples for research. Fresh samples were stored in liquid nitrogen. All experiments in this study were approved by ethic committee of the Affiliated Huaian NO.1 people's Hospital of Nanjing Medical University.

2.2 Antibodies

All antibodies used in this study were as following: anti-IGF1R antibody (Ab39675, Abcam), anti-GAPDH antibody (MA1-16757, Thermofisher), horseradish peroxidase-conjugated secondary antibodies goat anti-rabbit IgG (H + L) (G-21234, Thermofisher), goat anti-mouse IgG (G-21040, Thermofisher).

2.3 Cell culture and transfection

All OS cell lines including MG-63, U2OS, 143B and G292 and normal bone cells line hFOB 1.19 were purchased from Type Culture Collection of the Chinese Academy of Sciences. Cells were cultured at 37 or 34°C with 5% CO2. Cell medium was Dulbecco’s modified Eagle’s medium (DMEM, Hyclone) or F12 medium (Hyclone) supplemented with foetal bovine serum (FBS; Gibco) and 10% penicillin and streptomycin (100 U/mL P/S; Gibco). Cell transfection was performed following users guide of Lipofectamine 2000 reagent (Invitrogen).

2.4 Reverse transcription-quantitative polymerase chain reaction (RT-qPCR) assay

Trizol reagent (Thermo Fisher, Inc) was applied in total RNA extraction from tissues and cell lines. The quality of RNA was determined by using Nanodrop 2000 (Thermo Scientific). If necessary, RNA was incubated with RNase R (Epigenome Biotechnologies) for 15 minutes at 37°C. 3·5 μg total RNA was reverse-transcribed to cDNA using PrimeScript First Strand cDNA Synthesis Kit (Takara), The Primers for cDNA synthesis was the mix of oligodT and Random hexamers. Then qPCR was conducted used TB Green® Premix Ex Taq™ GC (Takara) on ABI 7500 Real-time PCR detection system (Applied Biosystems). The PCR setup was step 1 (95°C 5 minutes); Step 2 (95°C 10 seconds, 60°C 30-34 seconds, 40 cycles); Melt curve: 95°C 1 minute, 55°C 1 minute, 55-98°C (10 s/cycle 0.5°C/cycle). The level of RNA was quantified using 2^ΔΔq method. The Internal control was GAPDH or U6. The relative expression level of RNA was normalized using U6 and GAPDH. The primes were designed using circinteractome and NCBI.

GAPDH forward, 5'-CCGTCTAGAAAAACCTGC C-3'; and reverse, 5'-GCCAAATTGTTGTGCATACC-3'; U6, forward 5'-CTCGCTTCCGGCAGCACA-3'; and reverse 5'-AAC GCTTCAAGAATTTTGCT-3'; F: IGF1R, forward 5'-GTCGAAG AATCGCATCATCA-3'; and reverse 5'-GCACTCTGGCCCATCA-3'; CTC-3'; and reverse 5'-GGAAAGAATACGGTGAGCCGTG-3'; and reverse 5'-GACACCCAGGGGCACCCAAT-3'. Circ_0002137 primer 1: forward 5'- CAGAAGGAGGCCTGGTGTAA-3'; and reverse 5'- CTGGCCATACCATACCA-3'; primer 2: forward 5'- TACTGTCAGAAGGAGGCCTG-3'; and reverse 5'- GCGCACACAGGAAACATAG-3'.

2.5 Cell proliferation assay

Cell Counting Kit-8 (CCK8; Dojindo) assay was performed to detect the growth of OS cells. After transfection for 24 hours, 2000 cells were cultivated into 96-well plate. 10 μL CCK8 reagent was added
into each well at different time point then incubated for 2 hours. OD value was recorded at 450 nM on Tecan Infinite Pro M1000 (Tecan).

2.6 | Transwell assay

After transfection for 48 hours, cells were starved in serum-free medium for 24 hours. Then 1E5 cells were resuspended with cell medium containing 10% FBS and added into the transwell chamber coated with diluted Matrigel (Yeason, China). Meanwhile, DMEM cell medium containing 20% FBS was added into basolateral chamber. After cultured for 24 hours, the cells were immobilized using 4% polyformaldehyde then stained with 0.5%-1% crystal violet solution. The images were captured by microscope (Olympus).

2.7 | Migration assay

Cell migration was detected using wound-healing assay. Transfected cells were added into 6-well plate at the density of 1.5E5 cells per well. After being cultured for 12 hours, a straight scratch was scraped using pipette tips (200 μL) in each well and scratch position was marked using dark line. The images were captured at 24 and 48 hours, respectively. Image J software (NIH) were applied to calculate the ratio of wound-healing area.

2.8 | RNA pull-down assay

The RNA pull-down assay using biotinylated RNA was conducted as previous study described by wang et al.20 E7 OS cells were washed with PBS followed by being harvested and lysed in a lysis buffer (20 mM Tris, pH 7.5, 200 mM NaCl, 2.5 mM MgCl2, 0.05% Igepal, 60 U/mL Superase-In (Ambion), 1mM DTT, protease inhibitors (Roche)). The cell lysates were centrifuged for reserve. Biotinylated circRNA and control probe was incubated with M-280 streptavidin magnetic beads (Sigma). Before incubation, the beads were coated with RNase-free BSA and yeast tRNA (Sigma) to prevent unspecific binding with RNA. Then the biotinylated-RNA coated beads were incubated with cell lysate at 4°C for 3 hours on shaking table and washed twice. The RNA complexes bound to the beads were extracted with RNeasy mini Kit (Qiagen), then qRT-PCR assay was performed to qualify the level of RNA.

2.9 | Flow cytometry

Transfected cells were collected and washed by PBS. For cell cycle distribution, suspended cells were dealt with 70% ethanol (500 μL) and fixed at 4°C overnight. Then, RNase A (Sigma) was added into cell suspension for 20 minutes. Cell cycle was detected with FACS flow cytometry (BD). For cell apoptosis detection, cell suspension was treated with annexin V and propidium iodide dye (PI) (Sigma), then apoptotic cells were calculated by flow cytometry.

2.10 | Dual-luciferase reporter assay

The relationship between circ_0002137 and miR-433-3p/IGF1R was determined by dual-luciferase reporter assay. The luciferase reporter vectors with wildtype or mutant 3’-UTR of circ_0002137 or IGF1R were constructed. Cells were co-transfected with miR-433-3p mimics or miR-NC using Lipofectamine 3000. After 48 hours, the luciferase activity was determined using a kit called dual-luciferase reporter assay (Promega).

2.11 | Animal model

4 weeks-old nude mice were purchased from Shanghai Model Organisms Center (Shanghai, China). 1E7 stable transfected OS cell suspension MG63 or U2OS were subcutaneously injected into the lower dorsal flank on mice. There were six mice per group. After injection, the tumour volume was measured and calculated each 5 days until 5 weeks. Then the animals were sacrificed and nodules were collected and weighed. All animals’ related experiments were approved by ethical committee in the Affiliated Huaian NO. 1 people’s Hospital of Nanjing Medical University.

2.12 | Statistical analysis

GraphPad Prism 6.0 (GraphPad software) was used to make statistics. For the comparison of two group, F test was used to compare variances and Kolmogorov-Smirnoff test was used to test Gaussian distribution. If both of P values were more than 0.10, unpaired T-test was used. If the data didn’t meet the condition, non-parametric was used. When more than two groups are compared, Gaussian distribution was test using Kolmogorov-Smirnoff test, only the P value >.10, we thought the data was normal distribution. Meanwhile, the homogeneity of variance was tested using F test. One-way ANOVA or non-parametric test was chosen based the result of Kolmogorov-Smirnoff test and F test. If one-way ANOVA was applied, multiple comparisons were tested using Turkey test. All data were represented as mean ± SD, P value was indicated using asterisks. Only $P \leq .05$ was considered statistically significant.

3 | RESULTS

3.1 | Up-regulated expression pattern of Circ_0002137 is identified in OS patients

To order to search for potential biomarkers in osteosarcoma, the RNA sequencing was performed by Zhao and their finding indicated circular RNA circ_0002137 was differentially expressed in OS tissue and paired adjacent tissues from OS patients.17 Subsequently, our study demonstrated that circ_0002137 was remarkably up-regulated in OS patients compared with surrounding
non-tumour tissue using real-time RT-PCR (Figure 1A). Meanwhile, abnormal expression pattern of circ_0002137 was also identified in the transcriptional profiling of OS cell lines (U2OS, 143B, SJSA1 and MG-63) and normal bone cell hFOB 1.19 (Figure 1B). To explore the underlying role of circ_0002137 in OS, we constructed the plasmids to interfere the expression of circ_0002137 in OS cell lines and rechecked the efficiency of RNA interference. The result showed that si-circ_0002137 reduced the level of circ_0002137 significantly compared with si-NC group (Figure 1C).

### 3.2 | Down-regulation of Circ_0002137 retrained the proliferation of OS cell lines

With the purpose of demonstrating whether circ_0002137 could play a role in the tumorigenesis of OS, we conducted the CCK8 assay and found that knockdown of circ_0002137 decelerated the growth of OS cell lines (Figure 2A). To further investigate the negative effect on proliferation caused by circ_0002137, we performed the flow cytometry analysis. The result showed that circ_0002137 inhibition arrested OS cell line in G0/G1 phase (Figure 2B,C). Meanwhile, apoptotic cells were also detected through staining annexin V and PI. The data indicated that knockdown of circ_0002137 promoted the cell apoptosis of OS cell lines (Figure 2D,E). In conclusion, those findings demonstrated that circ_0002137 inhibition suppressed cell proliferation and enhanced cell apoptosis of OS cell lines.

### 3.3 | Circ_0002137 inhibition suppressed the metastasis and invasion of OS line

It was known that pulmonary metastasis or relapse usually took place on advanced OS patients, which led to unfavourable prognosis and high mortality. Therefore, it was important to detect whether circ_0002137 had therapeutic potential on inhibiting the metastasis of OS cell line. For this purpose, transwell assay showed that down-regulation of circ_0002137 significantly decreased the invasion ability of OS cell lines (Figure 3A,B). Moreover, we conducted wound-healing assay to determine cell migration of OS cells in si-circ_0002137 group, result showed the migration ability of cells was reduced in si-circ_0002137 compared with si-NC group (Figure 3C). To sum up, these results revealed circ_0002137 might be the inhibitor of the migration of OS tumour cell lines.

### 3.4 | Knockdown Circ_0002137 Suppresses Tumorigenesis of OS in Xenograft Model

Based on those data in vitro, we postulated that circ_0002137 might have influence on the tumour activity in vivo. Thus, OS cell lines were injected subcutaneously into nude mice after being transfected with si-circ_0002137 and si-NC to establish Xenograft model. Results revealed that circ_0002137 suppression significantly retarded tumour growth in vivo in accordance with tumour growth curve and tumour weight (Figure 4A,B). Additionally, knockdown of circ_0002137 inhibited the level of Ki67 and promoted the expression of caspase 3.

---

**FIGURE 1**  Circ_0002137 is highly expressed in OS patients. A, qRT-PCR assay determined relative circ_0002137 expression in OS tissues and adjacent non-tumour tissues, ***P < .001. B, Relative circ_0002137 expression in OS cell lines (U2OS, 143B, G292 and MG-63) and normal bone cell hFOB 1.19, ***P < .001. C, qRT-PCR was analysed for circ_0002137 expression in U2OS and MG-63 cells transfected with si-circ0002137, ***P < .001

**FIGURE 2**  Circ_0002137 inhibition retrained the proliferation of OS cell lines. A, Cell proliferation was detected using CCK8 assay on OS cell lines U2OS and MG-63 after transfected with si-circ0002137, ***P < .001. B-C, Cell cycle arrested was observed using flow cytometry. Si-circ0002137 group exhibited G0/G1 arrested compared with si-NC group, ***P < .001, *P < .05. D, Cell apoptosis was determined through staining apoptotic cells with Annexin V-FITC and PI, ***P < .001.
**FIGURE 3** Circ_0002137 inhibition suppressed the metastasis and invasion of OS line. A-B, Circ_0002137 inhibition reduced the invasion ability of OS cells through transwell assay. ***P < .001; **P < .01. C, Down-regulation of circ_0002137 decelerated the speed of wound healing in OS cells. ***P < .001; **P < .01; *P < .05

**FIGURE 4** Circ_0002137 Knockdown Suppresses Tumorigenesis of OS in Xenograft Model. A, Tumour growth curves of circ_0002137 inhibitor and si-NC transfected U2OS cells and MG-63 in nude mice. ***P < .001. B, The graph of tumour weight indicated circ_0002137 could suppress the tumour growth in OS model. ***P < .001; **P < .01. C, D, qRT-PCR was used to evaluate the mRNA level of Ki-67 and caspase 3 in nodules. ***P < .001; **P < .01
in si-circ_0002137 expressing nodules (Figure 4C,D). Therefore, it was demonstrated that circ_0002137 inhibition could restrain the OS's development in vivo.

3.5 | Circ_0002137 bond to miR-433-3p in OS cell line

Emerging evidences have proved that circular RNAs might function as miRNAs' sponge to regulate the transcription of linear RNAs and the expression of related protein. To further study the molecular mechanism of regulatory effect triggered by circ_0002137, we predicted the downstream molecule of circ_0002137 using circinteractome and circBase and found the potential target might be miR-433-3p. Putative-binding sites between circ_0002137 and miR-433-3p were shown in Figure 5A. To evaluate the correlation between circ_0002137 and miR-433-3p, dual-luciferase reporter assay was conducted and proved that overexpression of miR-433-3p mimics significantly inhibit the activity of circ_0002137 wt-luc rather than circ_0002137 mut-luc (Figure 5B). Furthermore, we confirmed the direct interaction between circ_0002137 and miR-433-3p through RNA pull-down experiment (Figure 5C). Subsequently, we detected the mRNA level of miR-433-3p in OS tissue. RT-QPCR data indicated that miR-433-3p had lower level than normal tissue rounding tumour (Figure 5D). We also observed circ_0002137 exhibited negative relationship with miR-433-3p in OS cell lines (Figure 5E). We detected the level of miR-433-3p when up-regulating circ_0002137 in OS cell lines. Lower miR-433-3p was observed in OS cells transfected with si-circ_0002137, suggesting the negative regulation between circ_0002137 and miR-433-3p. Those findings demonstrated that circ_0002137 could bind to miR-433-3p directly and regulate the level of miR-433-3p in OS cell lines.

3.6 | MiR-433-3p reversed the inhibitory influence caused by circ_0002137 in OS cell lines

Next, we explored whether circ_0002137 took part in the development of OS through controlling miR-433-3p. To verify the status of cell proliferation, CCK8 assay and cell cycle assay were performed. The final results proved that knockdown of miR-433-3p could alleviate the effect caused by circ_0002137 inhibition (Figure 6A,B). Meanwhile, the result of cell apoptosis showed that co-transfection of si-circ_0002137 and si-miR-433-3p in OS cells induced less apoptotic cells than OS cells transfected with si-circ_0002137 alone (Figure 6C,D). In conclusion, we made the conclusion that

![Figure 5](image-url)

**Figure 5** Circ_0002137 bond to miR-433-3p in OS cell line. A, Predicted binding sites between circ_0002137 and miR-433-3p using circInteractome (NIH, USA). B, Dual-luciferase reporter assay was conducted to detect the interaction between circ_0002137 and miR-433-3p. C, RNA pull-down assay was performed in order to verify the direct binding of circ_0002137 and miR-433-3p. D, The result of qRT-PCR demonstrated higher level of miR-433-3p in OS tissue compared with normal tissue. E, The level of miR-433-3p was negatively related with circ_0002137.
FIGURE 6 MiR-433-3p reversed the inhibitory influence caused by circ_0002137 in OS cell lines. A, Cell proliferation using CCK-8 indicated that miR-433-3p inhibition could restrain the inhibitory effect caused by circ_0002137. B, Cell cycle distribution determined by flow cytometry showed that G0/G1 phase arrest could be rescued by miR-433-3p inhibition. C-D,Cell apoptosis ratio demonstrated that higher ratio of apoptotic cells was reduced in si-circ_0002137+sh-miR-433-3p group.
circ_0002137 modulated the occurrence and progression of OS cell through regulating miR-433-3p.

3.7 IGF1R as the downstream functional target gene of circ_0002137/ miR-433-3p axis

To explore the underlying mechanism for circ_0002137 in OS, we tried to search for potential partner of circ_0002137/miR-433-3p. Firstly, bioinformatic tools (Targetscan and Starbase) were applied to predict IGF1R (insulin-like growth factor-1 receptor) might have putative-binding sites with miR-433-3p (Figure 7A). Subsequently, luciferase assay indicated that miR-433-3p mimic could affect the expression of IGF1R through regulating the activity of IGF1R promoter (Figure 7B). In addition, mRNA level of IGF1R was reduced with transfecting miR-433-3p mimic and the inhibitory effect induced by miR-433-3p mimic could be alleviated by co-expressing circ_0002137 (Figure 7C). As we expected, the protein level of IGF1R was determined using western blot. Results proved that IGF1R was also controlled by circ_0002137/ miR-433-3p (Figure 7D,E). Moreover, the relationship between IGF1R and miR-433-3p exhibited negatively correlated (Figure 7F). In conclusion, we made the conclusion that circ_0002137 could act as sponger of miR-433-3p and modulate the occurrence and development of OS through IGF1R.

4 DISCUSSION

IGF1R, one kind of transmembrane tyrosine kinase receptor, has been accepted its role in modulating cell differentiation, proliferation and apoptosis in several cancers. Meanwhile, previous studies have demonstrated insulin-like growth factors IGF1 and IGF2 were the ligands of IGF1R and their interaction could activate two major oncogenic signalling cascades: PI3K pathway and MAPK pathway. Intriguingly, IGF1R overexpression was observed in multiple kinds of tumour tissue, especially in OS patients. In addition, poor prognosis and survival rate was observed among the OS patients with elevated IGF1R. Thus, IGF1R was considered as an important target for OS treatment. In present, several clinical trials targeting IGF1R have been launched. Most of ongoing clinical trials were aimed to blocking IGF1R’s function via taking advantage of inhibitors of IGF1R, such as monoclonal antibody.
circ_0002137, which was reported to be differentially expressed in OS patients; however, its molecular mechanism was still unclear. By analysing those clinical data from medicine trials, we found that blocking bioactivity of IGF1R or IGF ligands using antibody was not appropriate because antibodies could not eliminate the overexpression of IGF on the surface of tumour cells. Thereby, other formats of IGF1R inhibitor need to be developed.

In our study, we reported a novel modulator of IGF1R: circ_0002137, which was reported to be differentially expressed in OS patients; however, its molecular mechanism was still unclear. Then we found the direct target of circ_0002137 might be miR-433-3p, which was identified to be a tumour suppressor in multiple tumours. For instance, the level of miR-433 was associated with unfavourable prognosis for the overall survival for patients with gastric carcinoma. In addition, miR-433 also functioned in liver cancer cell through inhibiting cell migration. Notably, the role of miR-433-3p in osteosarcoma also was reported by Mao et al. In our study, we found miR-433-3p might be a crucial partner of circ_0002137. Considered the relationship between miR-433-3p and IGF1R, we proposed the hypothesis that circ_0002137 regulated the target effects of RNA drugs needed to be solved in the future study.

CONFLICT OF INTEREST
The authors declare that they have no conflict of interest or personal relationships that could have appeared to influence the work reported in this paper.

AUTHOR CONTRIBUTION
Meng Zhang: Data curation (lead); Formal analysis (equal); Investigation (equal); Methodology (equal); Resources (equal).
Guang-Yang Yu: Data curation (supporting); Investigation (equal); Methodology (equal); Validation (equal).
Wei-Dong Liu: Project administration (equal); Supervision (equal); Writing-original draft (equal); Writing-review & editing (equal).
Gang Liu: Project administration (equal); Supervision (equal); Writing-original draft (equal); Writing-review & editing (equal).

DATA AVAILABILITY STATEMENT
The data used to support the findings of this study are available from the corresponding author upon request.

ORCID
Gang Liu https://orcid.org/0000-0002-5455-4362

REFERENCES
1. Jo VY, Doyle LA. Refinements in sarcoma classification in the current 2013 World Health Organization classification of tumours of soft tissue and bone. Surg Oncol Clin N Am. 2016;25:621-643.
2. Saha D, Saha K, Banerjee A, Jash D. Osteosarcoma relapse as pleural metastasis. South Asian J Cancer. 2013;2:56.
3. Heare T, Hensley MA, Dell’Orfano S. Bone tumors: osteosarcoma and Ewing’s sarcoma. Curr Opin Pediatr. 2009;21:365-372.
4. Kunz P, Fellenberg J, Moskovsky L, et al. Improved survival in osteosarcoma patients with atypical low vascularization. Ann Surg Oncol. 2015;22:489-496.
5. Wu PK, Chen WM, Chen CF, Lee OK, Haung CK, Chen TH. Primary osteogenic sarcoma with pulmonary metastasis: clinical results and prognostic factors in 91 patients. Jpn J Clin Oncol. 2009;39:514-522.
6. Hsu MT, Coca-Prados M. Electron microscopic evidence for the circular form of RNA in the cytoplasm of eukaryotic cells. Nature. 1979;280:339-340.
7. Cocquerelle C, Mascrez B, Hétuin D, Bailleul B. Mis-splicing yields circular RNA molecules. FASEB J. 1993;7:155-160.
8. Li Y, Zheng Q, Bao C, et al. Circular RNA is enriched and stable in exosomes: a promising biomarker for cancer diagnosis. Cell Res. 2015;25:981-984.
9. Memczak S, Jens M, Elefsinioti A, et al. Circular RNAs are a large class of animal RNAs with regulatory potency. Nature. 2013;495:333-338.
10. Guo JU, Agarwal V, Guo H, Bartel DP. Expanded identification and characterization of mammalian circular RNAs. Genome Biol. 2014;15:409.
11. Li RC, Ke S, Meng FK, et al. CIRI-S7 promotes growth and metastasis of esophageal squamous cell carcinoma via regulation of miR-7/HOXB13. Cell Death Dis. 2018;9:838.
12. Zhang J, Hu H, Zhao Y, Zhao Y. CDR1as is overexpressed in laryngeal squamous cell carcinoma to promote the tumour’s progression via miR-7 signals. Cell Prolif. 2018;51:e12521.
13. Zhang S, Liao K, Miao Z, et al. CircFOXO3 promotes glioblastoma progression by acting as a competing endogenous RNA for NFAT5. Neuro Oncol. 2019;21:1284-1296.
14. Yang W, Du WW, Li X, Yee AJ, Yang BB. Foxo3 activity promoted by non-coding effects of circular RNA and Foxo3 pseudogene in the inhibition of tumour growth and angiogenesis. Oncogene. 2016;35:3919-3931.
15. Wang C, Ren M, Zhao X, Wang A, Wang J. Emerging roles of circular RNAs in osteosarcoma. Med Sci Monit. 2018;24:7043-7050.
16. Liu W, Zhang J, Zou C, et al. Microarray expression profile and functional analysis of circular RNAs in osteosarcoma. Cell Physiol Biochem. 2017;43:969-985.
17. Xi Y, Fowdur M, Liu Y, Wu H, He M, Zhao J. Differential expression and bioinformatics analysis of circRNA in osteosarcoma. Biosci Rep. 2019;39:BSR20181514.
18. Panda AC, Dudekula DB, Abdelmohsen K, Gorospe M. Analysis of circular RNAs using the web tool circinteractome. Methods Mol Biol. 2018;1724:43-56.
19. Dudekula DB, Panda AC, Grammatikakis I, De S, Abdelmohsen K, Gorospe M. CircInteractome: a web tool for exploring...
circular RNAs and their interacting proteins and microRNAs. RNA Biol. 2016;13:34-42.
20. Wang K, Long B, Liu F, et al. A circular RNA protects the heart from pathological hypertrophy and heart failure by targeting miR-223. Eur Heart J. 2016;37:2602-2611.
21. Liu JP, Baker J, Perkins AS, Robertson EJ, Efstratiadis A. Mice carrying null mutations of the genes encoding insulin-like growth factor I (Igf-1) and type 1 IGF receptor (Igf1r). Cell. 1993;75:59-72.
22. Neuzillet Y, Chapeaublanc E, Krucker C, et al. IGF1R activation and the in vitro antiproliferative efficacy of IGF1R inhibitor are inversely correlated with IGFBP5 expression in bladder cancer. BMC Cancer. 2017;17:636.
23. Shali H, Ahmadi M, Kafil HS, Dorosti A, Yousefi M. IGF1R and c-met as therapeutic targets for colorectal cancer. Biomed Pharmacother. 2016;82:528-536.
24. Andersson MK, Aman P, Stenman G. IGF2/IGF1R signaling as a therapeutic target in MYB-positive adenoid cystic carcinomas and other fusion gene-driven tumors. Cells. 2019;8:913.
25. Li Z, Pan W, Shen Y, et al. IGFl/IGF1R and microRNA let-7e down-regulate each other and modulate proliferation and migration of colorectal cancer cells. Cell Cycle. 2018;17:1212-1219.
26. Scheffold A, Jebaraj BMC, Tausch E, et al. IGFlR as druggable target mediating PI3K-delta inhibitor resistance in a murine model of chronic lymphocytic leukemia. Blood. 2019;134:534-547.
27. Rodrigues Alves APN, Fernandes JC, Fenerich BA, et al. IGFlR/IRS1 targeting has cytotoxic activity and inhibits PI3K/AKT/mTOR and MAPK signaling in acute lymphoblastic leukemia cells. Cancer Lett. 2019;456:59-68.
28. Kuijjer ML, Peterse EF, van den Akker BE, et al. IR/IGF1R signaling as potential target for treatment of high-grade osteosarcoma. BMC Cancer. 2013;13:245.
29. Hassan SE, Bekarev M, Kim MY, et al. Cell surface receptor expression patterns in osteosarcoma. Cancer. 2012;118:740-749.
30. Wang Y, Lipari P, Wang X, et al. A fully human insulin-like growth factor-1 receptor antibody SCH 717454 (Robatumumab) has antitumor activity as a single agent and in combination with cytotoxics in pediatric tumor xenografts. Mol Cancer Ther. 2010;9:410-418.
31. Malempati S, Weigle B, Ingle AM, et al. Phase I/II trial and pharmacokinetic study of cixutumumab in pediatric patients with refractory solid tumors and Ewing sarcoma: a report from the Children’s Oncology Group. J Clin Oncol. 2012;30:256-262.
32. Schwartz GK, Tap WD, Qin LX, et al. Cixutumumab and temsirolimus for patients with bone and soft-tissue sarcoma: a multicentre, open-label, phase 2 trial. Lancet Oncol. 2013;14:371-382.
33. Gao J, Chesebrough JW, Cartlidge SA, et al. Dual IGF-1/II-neutralizing antibody MEDI-573 potently inhibits IGF signaling and tumor growth. Cancer Res. 2011;71:1029-1040.
34. Friedbichler K, Hofmann MH, Kroez M, et al. Pharmacodynamic and antineoplastic activity of BI 836845, a fully human IGF ligand-neutralizing antibody, and mechanistic rationale for combination with rapamycin. Mol Cancer Ther. 2014;13:399-409.
35. Luo H, Zhang H, Zhang Z, et al. Down-regulated miR-9 and miR-433 in human gastric carcinoma. J Exp Clin Cancer Res. 2009;28:82.
36. Ueda T, Volinia S, Okumura H, et al. Relation between microRNA expression and progression and prognosis of gastric cancer: a microRNA expression analysis. Lancet Oncol. 2010;11:136-146.
37. Yang Z, Tsuchiya H, Zhang Y, Hartnett ME, Wang L. MicroRNA-433 inhibits liver cancer cell migration by repressing the protein expression and function of cAMP response element-binding protein. J Biol Chem. 2013;288:28893-28899.
38. Hou XK, Mao JS. Long noncoding RNA SNSHG14 promotes osteosarcoma progression via miR-433-3p/FBXO22 axis. Biochem Biophys Res Commun. 2020;523:766-772.

How to cite this article: Zhang M, Yu G-Y, Liu G, Liu W-D. Circular RNA circ_0002137 regulated the progression of osteosarcoma through regulating miR-433-3p/IGF1R axis. J Cell Mol Med. 2021;00:1-11. https://doi.org/10.1111/jcmm.16166