

# Study on Targeting Relationship Between miR-320b and FGD5-AS1 and Its Effect on Biological Function of Osteosarcoma Cells

This article was published in the following Dove Press journal:  
*Cancer Management and Research*

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**Objective:** To probe into the expression of FGD5-AS1 in osteosarcoma and its relationship with miR-320b.

**Methods:** The tissue and serum samples of 97 patients with osteosarcoma were collected, and the serum samples of 100 healthy subjects who concurrently underwent physical examination were selected as the control. FGD5-AS1 expression in tissues and serum was detected, and osteosarcoma cells were transfected to measure cell behaviors such as proliferation, invasion and apoptosis.

**Results:** FGD5-AS1 was highly expressed in osteosarcoma, and its elevated expression indicated poor survival of patients. Serum FGD5-AS1 was related to tumor size and clinical stage and could be used for the diagnosis of osteosarcoma. The study of osteosarcoma cell lines U2OS and SaOS-2 showed that after inhibiting FGD5-AS1, the viability and invasion capacity of osteosarcoma cells decreased statistically compared with the control group (CG), while the apoptosis ability could be improved by further regulating apoptotic proteins ( $P < 0.05$ ). Detection of EMT-related proteins identified that E-cadherin increased while N-cadherin decreased significantly after FGD5-AS1 inhibition ( $P < 0.05$ ). Correlation analysis revealed a negative correlation between miR-320b and FGD5-AS1 ( $r = -0.410$ ,  $P < 0.001$ ). Overexpression of miR-320b significantly inhibited cell viability, invasion and EMT ability, and increased the apoptosis rate, while inhibiting miR-320b expression produced the opposite results. The targeting relationship between miR-320b and FGD5-AS1 was confirmed through the biological prediction website, luciferase assay and RNA binding protein immunoprecipitation (RIP) assay. Inhibition of miR-320b could reverse the regulatory effect of FGD5-AS1 knockdown on osteosarcoma cells.

**Conclusion:** FGD5-AS1 is highly expressed in osteosarcoma and is involved in the biological progression of osteosarcoma by targeting miR-320b.

**Keywords:** miR-320b, FGD5-AS1, osteosarcoma, biological behavior

## Introduction

Osteosarcoma occurs predominantly in adolescents and is a primary bone malignancy.<sup>1</sup> Metastasis, especially in the lungs and bones, is present in approximately 15% to 20% of the patients when diagnosed, and develops rapidly.<sup>2</sup> Pain and swelling of the affected bones are the main clinical manifestations of the disease. Thanks to the continuous development of social medical technology and the implementation of comprehensive treatment methods including chemotherapy and limb salvage surgery, the survival rate of osteosarcoma patients has been increased

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in the past few decades.<sup>3</sup> It is reported that the 5-year survival rate of patients with osteosarcoma is about 70%.<sup>4</sup> Although our understanding of osteosarcoma has been improved with the deepening of research, there are still many patients with chemotherapy resistance and lung metastasis with a 5-year survival rate of less than 20% and an unfavorable prognosis.<sup>5,6</sup> And the molecular mechanisms underlying the development of carcinogenesis and metastasis are still unknown. Numerous studies have shown that microRNAs (miRNA) are involved in the progression of osteosarcoma,<sup>7</sup> and their abnormal expression profiles are related to the prognosis of patients.<sup>8</sup> It is also widely acknowledged that miRNAs are indispensable in regulating the self-renewal ability, invasion and migration of osteosarcoma cells.<sup>9</sup> Among them, miR-320 has been pointed out by a number of studies as a potential therapeutic target for cancer, and a prognostic indicator for multiple tumors.<sup>10,11</sup> Moreover, in osteosarcoma, miR-320 is reduced in cell lines, and inhibits the heterologous expression of cell proliferation and induces cell cycle arrest.<sup>11</sup> The study of Lv et al<sup>12</sup> suggested that high levels of XIST were negatively correlated with miR-320b, and could promote osteosarcoma progression by regulating miR-320b. All the preceding evidence suggests that miR-320b participates in the biological process of osteosarcoma.

The critical role of long non-coding RNAs (lncRNAs) in osteosarcoma has been already well established, but the role of the key lncRNAs in the pathogenesis of osteosarcoma has not been fully clarified. As a new type of lncRNA, FGD5-AS1 has been shown to be effective in promoting non-small cell lung cancer<sup>13</sup> and colorectal cancer.<sup>14</sup> Based on previous studies, we hypothesized that FGD5-AS1 was vital in tumor progression. In this study, we detected FGD5-AS1 and explored its effects on osteosarcoma cells by evaluating abnormal biological behaviors such as cell viability, apoptosis, and migration. In addition, as numerous studies indicated that lncRNA played a competitive role in endogenous RNA (ceRNA) and regulation of miRNA in various diseases,<sup>15,16</sup> the regulatory relationship between FGD5-AS1 and miR-320b was detected in this study, and later through luciferase reporter, we confirmed that FGD5-AS1 targeted the regulation of miR-320b. Overall, the purpose of our study is to demonstrate the role of FGD5-AS1 in osteosarcoma and its regulatory mechanism with miR-320b.

## Materials and Methods

### Sample Collection

A total of 97 patients with osteosarcoma treated in The First Hospital of Ningbo were collected. The ratio of male to female was 57:40, the age ranged from 12 to 28 years old, and the average age was 18.87±5.18 years old. Concurrently, 100 healthy subjects, aged 12–26 years, with an average age of 18.57±4.98 years old and the male-to-female ratio of 55:45, who underwent health examinations were selected as the control group (CG). There was no significant difference in general clinical data between the two groups ( $P>0.05$ ). The protocol of this study was approved by the Clinical Research Ethics Committee of The First Hospital of Ningbo, conducted in accordance with the Declaration of Helsinki, and all the participants had signed the informed consent. In this study, patients were followed up in the form of telephone interview, outpatient reexamination and medical record inquiry, and the overall survival (OS) was from the end of the first treatment to death or the end of follow-up.

### Cell Culture

Osteosarcoma cell lines U2OS, HOS, SaOS-2, SW1353 and human normal osteoblast cell line hFOB1.19 were all purchased from Aolu Biotechnology Co., Ltd., Shanghai, China. The cell lines were incubated in Dulbecco's Modified Eagle's Medium (DMEM: Chundu Biotechnology Co., Ltd., Wuhan, China) supplemented with 10% heat-inactivated fetal bovine serum in a 5% CO<sub>2</sub> constant temperature incubator at 37°C.

### Cell Transfection

Using Lipofectamine (Invitrogen, United States, Lipofectamine® 2000 Transfection Reagent), FGD5-AS1 inhibitor (si-FGD5-AS1), negative control (si-NC), miR-320b overexpression/inhibitor (miR-320b mimics, miR-320b inhibitor) and negative control (miR-NC) were transfected into U2OS and SaOS-2 cell lines, respectively, for observation and research.

### Quantitative PCR Detection

Total RNA was extracted from the collected cell, tissue and serum samples using TRIZOL reagent (Invitrogen, CA, United States). High capacity cDNA reverse transcription kit (Applied Biosystems, CA, United States) was used for reverse transcription of RNA samples, and the specific operation was carried out according to the

**Table 1** Primer Sequences

	Upstream	Downstream
FGD5-AS1	5'-CGTGGAGAAGAATTGGGC-3'	5'-CGTGGAGAAGAATTGGGC-3'
miR-320b	5'-TCCGAAACGGGAGAGTTGG-3'	5'-GTGCAGGGTCCGAGGT-3'
GAPDH	5'-ACCTGACCTGCCGTCTAGAA-3'	5'-GTCAAAGTGGAGGAGTGGG-3'
U6	5'-GCTTCGGCAGCACATATACTAAAT-3'	5'-CGCTTCACGAATTTGCGTGT-CAT-3'

instructions. Quantitative RT-PCR was performed by Fast Start Universal SYBR Green Master (Roche, CA, United States). Primers are shown in Table 1. The relative fold change of candidate genes was analyzed using the  $2^{-\Delta\Delta C_t}$  method (Table 1).

## Western Blot Detection

The collected cells were lysed using RIPA buffer (Sigma-Aldrich, MO, United States). Protein concentration was determined by BCA (Beyotime Biotechnology, Shanghai, China), and the protein samples were diluted with buffer solution. The same amount of extracts were separated by SDS-PAGE and then transferred to a polyvinylidene fluoride (PVDF) membrane (Millipore, Shanghai, China) after ionization. Then, the membrane was blocked with 5% skimmed milk for 2h, immersed in primary antibodies Bax, Caspase3, Bcl-2, E-cadherin, N-cadherin, and  $\beta$ -Actin primary antibody (Invitrogen, CA, United States), and then blocked at 4 °C overnight. After that, the primary antibody was removed by washing the membrane, and the horseradish peroxidase-labeled goat anti-rabbit secondary antibody (Invitrogen, CA, United States) was added for 1h of incubation at 37 °C. After rinsing the membrane three times with TBST, the band was analyzed using ECL Luminescence Reagent (Thermo Fisher, MA, United States).

## CCK-8 Determination for Cell Viability

Osteosarcoma cells were inoculated into 96-well plates ( $2 \times 10^3$ ) with complete medium, and then transfected with different cell lines. After transfection for 24h, 48h, 72h, and 96h, 10  $\mu$ L CCK-8 solution (Biosharp, Shanghai, China) was added to each well for 2h of culture at 37 °C. The optical density (wavelength 450nm) of each well was measured by enzyme-linked immunosorbent assay (ELISA; Multiskan™ FC microplate reader, Thermo Fisher Scientific, MA, USA), and the cell growth curve was plotted. The experiment was repeated in triplicate.

## Apoptosis Assay

AnnexinV-FITC apoptosis kit (Keygen, Nanjing, China) was used to detect the apoptosis. After transfection, the cells were digested with trypsin, washed with PBS, and then resuspended in 500  $\mu$ L binding buffer. AnnexinV-FITC and PI were then added to incubate in the dark for 15–20 minutes, and FC500MCL flow cytometry system was utilized for detection. The experiment was repeated 3 times.

## Invasion Experiment

The invasive ability of the cells was evaluated by transwell assay. The cells were inoculated into the upper chamber coated with matrix glue (BD Biosciences, CA, United States), and the medium with and without 10%FBS was added to the lower and upper wells, respectively, and cultured at 37 °C for 24 h. After wiping the stroma and cells of the upper chamber that did not pass through the membrane surface, the cells were immobilized and stained with crystal violet. The invasion of cells was observed under a microscope.

## Luciferase Reporter Gene Detection

Osteosarcoma cell line was seeded in 24-well plates. After incubation, the pmirGLO reporter vectors with wild-type (FGD5-AS1-WT) and mutant (FGD5-AS1-MUT) were co-transfected into cells with miR-320b-mimics and miR-NC. The luciferase intensity was detected using a dual-luciferase reporter gene system (Solarbio, CA, China), and the steps were carried out in strict accordance with the instructions.

## RNA Binding Protein

### Immunoprecipitation (RIP Experiment)

The RIP kit (Millipore, MA, United States) was used for detection, and the experimental procedures were carried out in strict accordance with the kit instructions. The whole cell protein extracts were incubated with RIP wash buffer containing magnetic beads bound to human anti-Ago2 antibody (Millipore, MA, USA) or mouse immunoglobulin G (IgG) controls. Protein in the sample

was digested with protease K, and immunoprecipitation RNA was extracted. The purified RNA was subjected to qRT-PCR analysis to prove the presence of binding targets.

## Statistical Analysis

All the experiments were carried out three times independently. GraphPad 7 was adopted to analyze and visualize the data. The counting data were expressed as percentage (%) and verified by chi-square test and denoted as  $\chi^2$ . Student's *t*-test was utilized for the analysis of the two groups, one-way analysis of variance (ANOVA) was used to confirm the differences among multiple groups, and LSD-*t*-test was adopted for pairwise post hoc comparisons. The diagnostic value was analyzed by receiver operator characteristic (ROC) curve. The overall survival of patients was plotted by K-M survival curve and analyzed by Log-rank test. Correlation analysis of gene expression was performed using Pearson correlation coefficients. When  $P < 0.05$ , there was a statistical difference.

## Results

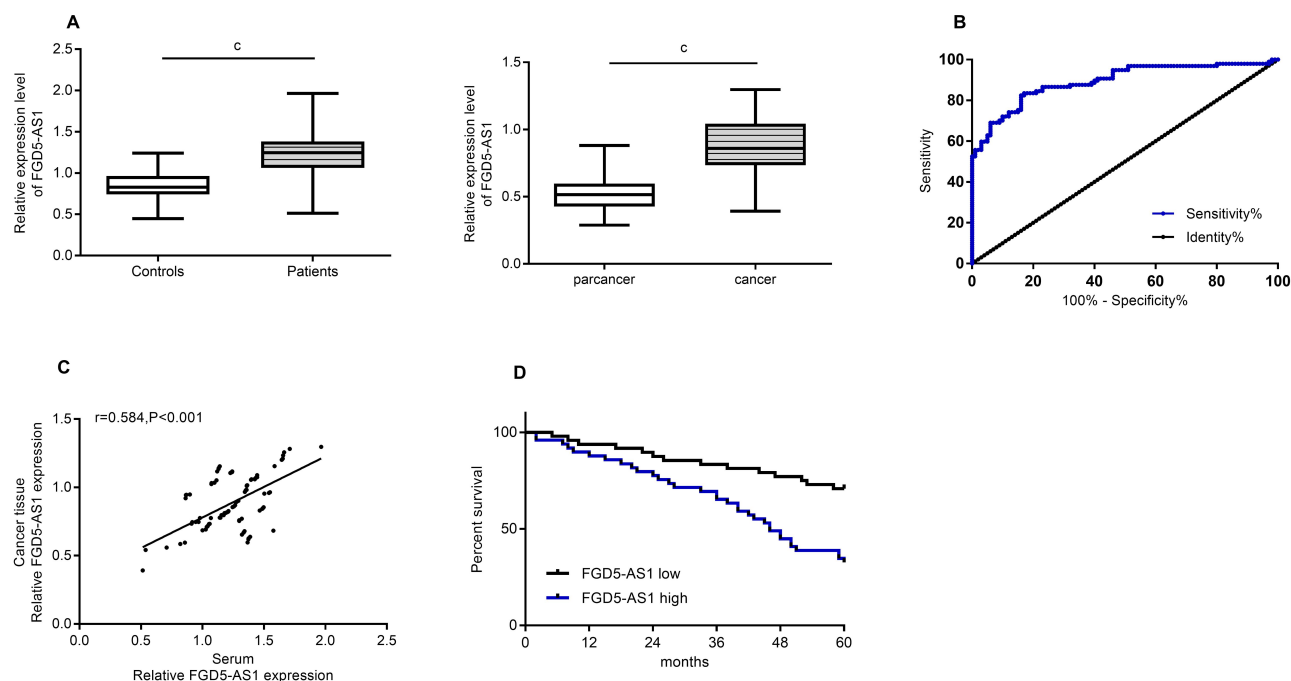
### FGD5-AS1 Expression in Osteosarcoma

Through the detection of patient samples, it was found that the serum FGD5-AS1 of osteosarcoma patients was higher than that of normal controls ( $P < 0.05$ ), and FGD5-AS1 was

higher in osteosarcoma tissues than in adjacent tissues (Figure 1). Correlation coefficient analysis showed that FGD5-AS1 in osteosarcoma tissue and serum was positively correlated, and serum FGD5-AS1 could be used in the diagnosis of osteosarcoma, with the AUC of 0.893. From Table 2, we can see that serum FGD5-AS1 was correlated with tumor size and clinical stage. Then, with the median value of FGD5-AS1 as the boundary, we classified patients into high and low expression groups, with 49 cases in each group. By analyzing the 5-year survival rate, it was found that the survival rate of patients with high FGD5-AS1 expression was significantly lower than that of patients with low expression ( $P < 0.05$ ). (Figure 1 and Table 2)

### Effects of Inhibiting FGD5-AS1 on Tumor Cells

In this section, we conducted cytological experiments to further explore the role of FGD5-AS1 in osteosarcoma cells (Figure 2). It was found that FGD5-AS1 was higher in osteosarcoma cell lines U2OS, HOS, SaOS-2 and SW1353 than in human normal osteoblast cell line hFOB1.19 ( $P < 0.05$ ), so we selected U2OS and SaOS-2 cell lines for follow-up study to construct anti-FGD5-AS1 cell lines. The results showed that compared with the CG, FGD5-AS1 inhibition remarkably decreased the viability



**Figure 1** FGD5-AS1 expression in osteosarcoma. (A) Expression of FGD5-AS1 in serum and tissue of osteosarcoma. (B) Serum FGD5-AS1 level was used for the diagnosis of patients with osteosarcoma. (C) Correlation of serum and tissue FGD5-AS1 levels in osteosarcoma. (D) Correlation between FGD5-AS1 and prognosis and survival of patients with osteosarcoma.

**Note:** <sup>c</sup>Represented the comparison between the two groups, <sup>c</sup> $P < 0.001$ .

**Table 2** Relationship Between FGD5-AS1 and Pathological Characteristics of Patients with Osteosarcoma n [%]

Pathological Data	Cases	FGD5-AS1		X <sup>2</sup>	P
		High Expression	Low Expression		
Gender				0.548	0.460
Male	57	27 (55.10)	30 (62.50)		
Female	40	22 (44.90)	18 (37.50)		
Age				0.825	0.364
>20 years old	42	19 (38.78)	23 (47.92)		
≤20 years old	55	30 (61.22)	25 (52.08)		
Tumor size (cm)				18.580	<0.001
>5	37	29 (59.18)	8 (16.67)		
≤5	60	20 (40.82)	40 (83.33)		
Clinical stage				14.390	<0.001
I-II	54	18 (36.73)	36 (75.00)		
III-IV	43	31 (63.27)	12 (25.00)		
Metastasis				0.657	0.418
With	30	17 (14.29)	13 (27.08)		
Without	67	32 (65.31)	35 (72.92)		
Site				1.500	0.221
Tibia/femur	65	30 (61.22)	35 (72.92)		
Others	32	19 (38.78)	13 (27.08)		

and invasion of osteosarcoma cells ( $P<0.05$ ), and enhanced the apoptosis ability of osteosarcoma cells by further regulating the apoptotic proteins ( $P<0.05$ ). Detection of EMT-related proteins revealed an increased E-cadherin and a decreased N-cadherin after FGD5-AS1 inhibition ( $P<0.05$ ). (Figure 2)

### Target Relationship Verification

We searched for the target of FGD5-AS1 through starBase, IncRNASNP2, and Diana prediction websites, and found four common miRNA targets, among which, miR-320b had a strong binding site with FGD5-AS1. Through the starBase website, we found the binding site of miR-320b and FGD5-AS1 (Figure 4). Luciferase assay identified that the fluorescence activity of FGD5-AS1-WT was inhibited after miR-320b overexpression. Through RIP assay, it was observed that the miR-320b and FGD5-AS1 expression bound to Ago2 was higher than that bound to immunoglobulin (Ig) G. After transfection of osteosarcoma cells, it was noticed that miR-320b increased after inhibiting FGD5-AS1 ( $P<0.05$ ). miR-320b presented low expression in osteosarcoma tissues, and correlation analysis revealed

showed that miR-320b was negatively associated with FGD5-AS1 ( $r = -0.410$ ,  $P<0.001$ ). (Figure 3)

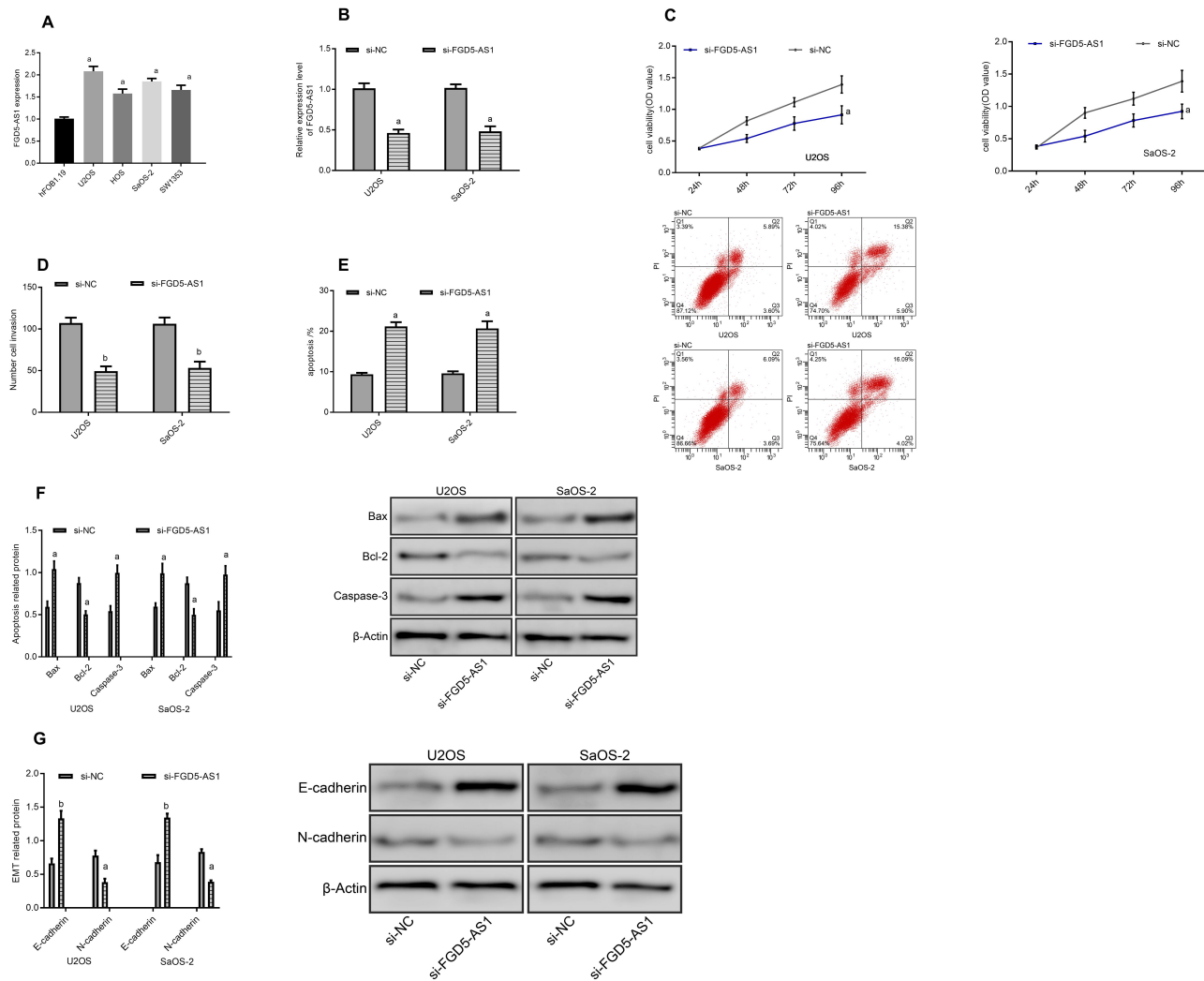
### Effects of miR-320b Inhibition or Overexpression on Osteosarcoma Cells

The miR-320b level of osteosarcoma cell lines U2OS, HOS, SaOS-2 and SW1353 was found to be lower than that of human normal osteoblast cell line hFOB1.19 ( $P<0.05$ ). After constructing miR-320b inhibition/overexpression cell lines, it was found that compared with the CG, miR-320b overexpression significantly reduced the viability and invasion capacity of osteosarcoma cells ( $P<0.05$ ), while increased the apoptosis ability by regulating the apoptotic protein ( $P<0.05$ ). Detection of EMT-related proteins revealed that E-cadherin increased while N-cadherin decreased after miR-320b overexpression ( $P<0.05$ ). However, the inhibition of miR-320b expression showed the opposite effect as that of overexpression (Figure 4).

### Co-Transfection Experiment

To further explore the regulatory relationship between miR-320b and FGD5-AS1, we co-transfected si-FGD5-AS1 and





**Figure 2** Effects of inhibiting FGD5-AS1 on tumor cells. **(A)** Expression of FGD5-AS1 in osteosarcoma cell lines. **(B)** Expression of FGD5-AS1 after transfection. **(C)** Cell proliferation. **(D)** Cell invasion. **(E)** Apoptosis rate and apoptosis graph. **(F)** Apoptosis-related protein diagram. **(G)** EMT-related protein diagram. **Notes:** <sup>a</sup>Represented <sup>a</sup> $P < 0.05$  compared with si-NC; <sup>b</sup>Represented <sup>b</sup> $P < 0.01$  compared with si-NC.

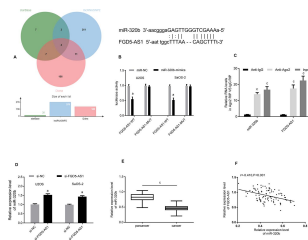
miR-320b-inhibitor into cells. The results showed that there were no significant differences between si-FGD5-AS1 + miR-320b-inhibitor and si-NC in cell viability, invasion, apoptosis and EMT regulation ( $P > 0.05$ ). It can also be seen that si-FGD5-AS1 + miR-320b-inhibitor could reverse the effect of si-FGD5-AS1 on osteosarcoma cells (Figure 5).

## Discussion

lncRNA is a class of non-protein transcript that can regulate various physiological and pathological functions.<sup>17</sup> With the property of regulating signal transduction and protein transformation between protein complex tissues and cells, they are often used as feasible biomarkers for tumors.<sup>18</sup> In osteosarcoma, lncRNAs participate in the occurrence of disease

through epigenetic inheritance, transcription and post-transcription mechanisms.<sup>19</sup> For example, Lu et al<sup>20</sup> proposed that CASC2 could be used as a potential tumor suppressor lncRNA for osteosarcoma and participate in the regulation of malignant behavior of tumor cells. Upon review, it was found that the relationship between lncRNA and osteosarcoma proliferation, prognosis, chemotherapy resistance was introduced in detail, and it was pointed out that the role and regulatory mechanism of lncRNA in human osteosarcoma should be further studied.<sup>21</sup>

FGD5-AS1 is highly expressed in cancers and can function as a therapeutic target in oral squamous cell carcinoma.<sup>22</sup> Since the current report on FGD5-AS1 in osteosarcoma has not been elaborated, we quantitatively detected by PCR to find that FGD5-AS1 was upregulated in osteosarcoma, which was

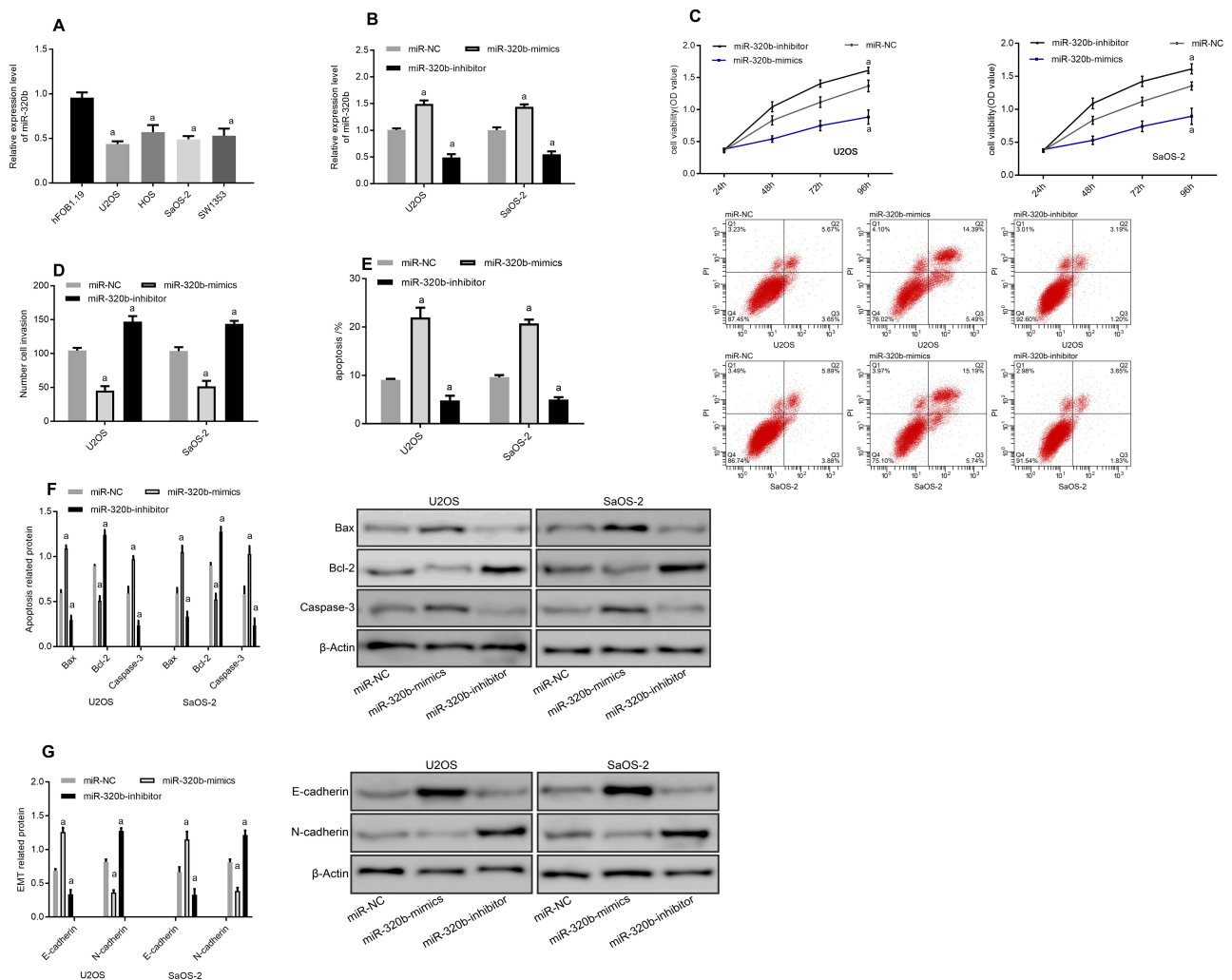


**Figure 3** Target relationship verification. **(A)** Wayne diagram. **(B)** Binding site and luciferase reporter. **(C)** RIP experiment. **(D)** Effects of different transfections on the expression of miR-320b. **(E)** Expression of miR-320b in cancer tissues. **(F)** Correlation between miR-320b and FGD5-AS1. **Notes:** <sup>a</sup>Represented <sup>a</sup>P<0.05 compared with NC; <sup>c</sup>Represented <sup>c</sup>P<0.001 compared between two groups or with Anti-IgG.

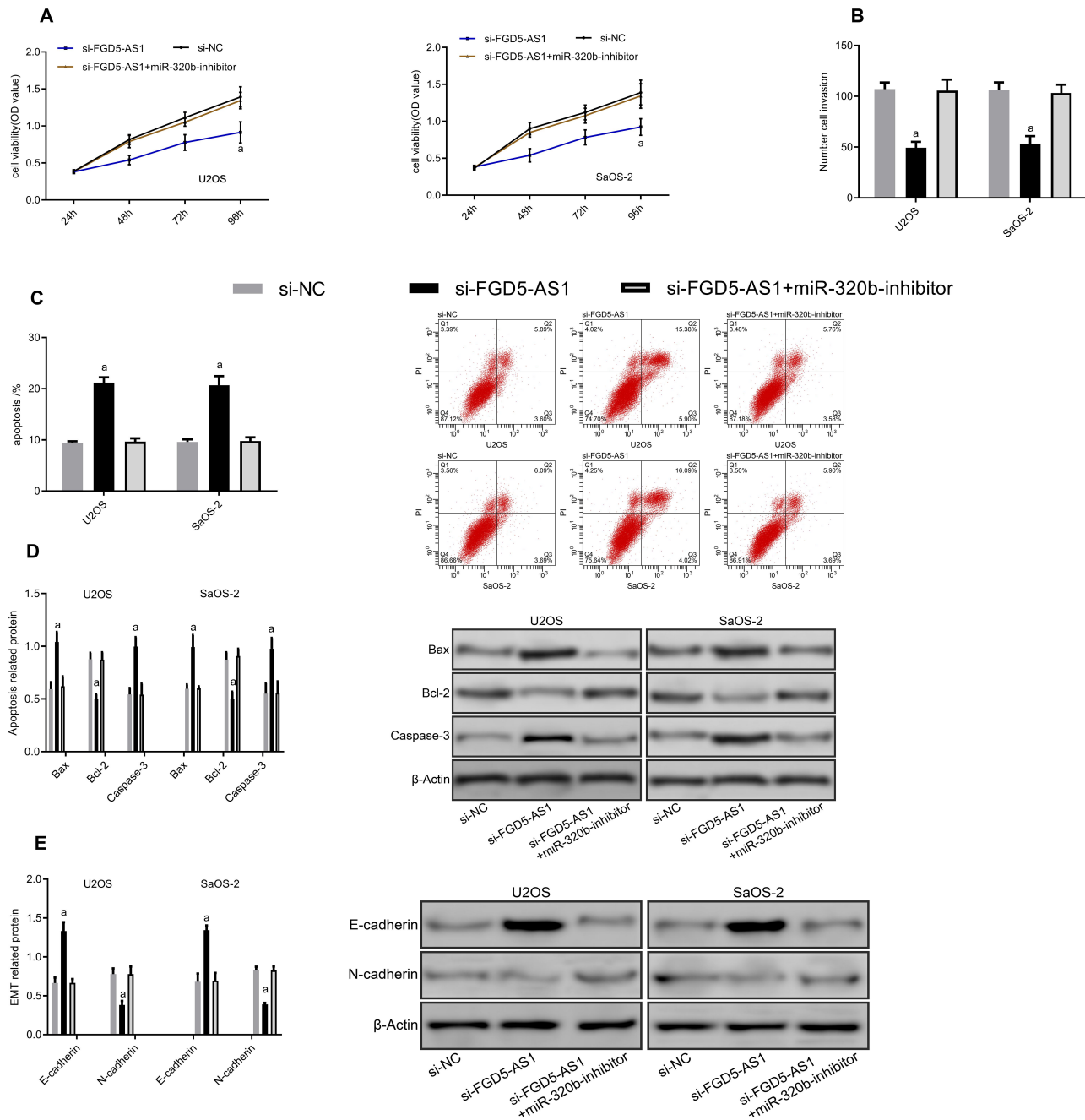
the diagnosis of various diseases due to their easy access. For example, serum lncRNA XIST and HIF1A-AS1 enjoy favorable diagnostic and predictive value in non-small cell lung cancer.<sup>23</sup> In this study, we found a correlation between the expression of serum and tissue through correlation coefficient analysis, and serum FGD5-AS1 can be used for the diagnosis of patients with osteosarcoma. In addition, the survival rate of osteosarcoma patients with high FGD5-AS1 expression was lower. These results suggest that FGD5-AS1 is a prognostic indicator for osteosarcoma, but the role of FGD5-AS1 in osteosarcoma remains unknown.

consistent with the preceding research results, indicating its role as an oncogene. Combined with related literature,<sup>23,24</sup> it is not difficult to find that serum biomarkers are widely used in

The occurrence and development of tumors is a process of gradual proliferation abnormality, while the spread mechanism of cancer cells includes cell invasion, migration and EMT.<sup>25,26</sup> Therefore, we further conducted



**Figure 4** Effects of miR-320b inhibition or overexpression on osteosarcoma cells. **(A)** Expression of miR-320b in osteosarcoma cell lines. **(B)** Expression of miR-320b after transfection. **(C)** Cell proliferation. **(D)** Cell invasion. **(E)** Apoptosis rate and apoptosis graph. **(F)** Apoptosis-related protein diagram. **(G)** EMT-related protein diagram. **Note:** <sup>a</sup>Represented <sup>a</sup>P<0.05 compared with miR-NC.



**Figure 5** Co-transfection experiment. **(A)** Proliferation diagram. **(B)** Cell invasion diagram. **(C)** Apoptosis diagram. **(D)** Apoptosis-related protein diagram. **(E)** EMT-related protein diagram.

**Note:** <sup>a</sup>Represented <sup>a</sup>P<0.05 compared with si-NC.

cytology experiments. First, the selected osteosarcoma cells were transfected. The results showed that the inhibition of FGD5-AS1 statistically reduced the viability and invasion of osteosarcoma cells, and increased the apoptosis ability by regulating apoptotic proteins. Changes in the biological traits of EMT will result in cells acquiring the ability to migrate and invade, while maintaining stem cell characteristics, tumor progression, and metastasis are all

related to EMT.<sup>27,28</sup> By detecting EMT-related proteins, we observed that FGD5-AS1 inhibition led to decreased EMT behavior, increased E-cadherin, and decreased N-cadherin (P<0.05), suggesting that FGD5-AS1 could be used as a therapeutic target for osteosarcoma to regulate the malignant behavior of tumor cells.

Increasing evidence has demonstrated that lncRNAs, highly expressed in the cytoplasm, can act as molecular



sponges, competitively bind or adsorb with other regulatory proteins and miRNAs, thus participating in the biological processes of tumors.<sup>29,30</sup> In this study, we predicted the lncRNA-miRNA binding sites through three biological target prediction sites: starBase, lncRNASNP2, and Diana. A total of 4 shared miRNAs were screened out, of which miR-320b belonged to the miR-320 family, which showed a correlation with osteosarcoma in previous reports.<sup>12</sup> Further, the specific binding between FGD5-AS1 and miR-320b was confirmed by double luciferase report and RIP experiment, and there was a correlation between them. Recently, research has pointed out that miR-320b is involved in the regulation of the progression of a variety of cancers,<sup>31</sup> and the inhibition of miR-320b can promote malignant behaviors such as cell proliferation in nasopharyngeal carcinoma<sup>32</sup> and colorectal cancer.<sup>33</sup> Therefore, we applied cytology to verify the effect of miR-320b on the biological behavior of osteosarcoma. The results showed that after miR-320b overexpression, the viability, invasion and EMT ability of osteosarcoma cells were decreased, and the apoptosis was increased, while the inhibition of miR-320b produced the opposite effects, further confirming the anti-tumor effect of miR-320b in osteosarcoma. Finally, through co-transfection experiments, we found that miR-320b inhibition could reverse the changes in cell behavior caused by silencing FGD5-AS1, suggesting that FGD5-AS1 could participate in the biological process of osteosarcoma by targeting miR-320b.

This study demonstrated the expression of FGD5-AS1 in osteosarcoma and the targeting relationship between miR-320b. However, as the experimental study of nude mice was absent in this study, the effect of FGD5-AS1 on the growth of solid tumors is not yet clear, which is one of the limitations. In addition, there are many network pathways involved in the regulation of osteosarcoma, and more experiments are needed to explore the involvement of FGD5-AS1 in mediating the specific downstream regulatory genes of miR-320b. What is more, the mechanism of drug resistance can also be investigated in future studies to better solve the limitations of clinical treatment. All in all, based on the comprehensive study, we conclude that FGD5-AS1 is involved in the biological process of osteosarcoma by targeting miR-320b.

## Conclusion

This study demonstrated the expression of FGD5-AS1 in osteosarcoma and the targeting relationship between miR-320b.

However, as the experimental study of nude mice was absent in this study, the effect of FGD5-AS1 on the growth of solid tumors is not yet clear, which is one of the limitations. In addition, there are many network pathways involved in the regulation of osteosarcoma, while miRNA is completed by regulating the downstream protein pathway, so the role of lncRNA/miRNA in osteosarcoma related pathways remains to be further investigated. What is more, the mechanism of drug resistance can also be investigated in future studies to better solve the limitations of clinical treatment. All in all, based on the comprehensive study, we conclude that FGD5-AS1 is involved in the biological process of osteosarcoma by targeting miR-320b.

## Funding

This study was financially supported by The mechanism research of HIF-1  $\alpha$  regulating CXCR4 in lung metastasis of osteosarcoma Ningbo Natural Science Foundation, No.2019A610250.

## Disclosure

The authors report no conflicts of interest in this work.

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