

# Identification of abnormally high expression of POGZ as a new biomarker associated with a poor prognosis in osteosarcoma

Sikuan Zheng,<sup>1,2,3,4\*</sup> Yue Liu,<sup>5\*</sup> Haohe Sun,<sup>2</sup> Jingyu Jia,<sup>1,3,4</sup> Tianlong Wu,<sup>1,3,4</sup> Rui Ding,<sup>1,2,3,4</sup> Xigao Cheng<sup>1,3,4</sup>

<sup>1</sup>The Second Affiliated Hospital of Nanchang University, Jiangxi

<sup>2</sup>The Second Clinical Medical College of Nanchang University, Jiangxi

<sup>3</sup>Institute of Orthopedics of Jiangxi Province, Nanchang, Jiangxi

<sup>4</sup>Institute of Minimally Invasive Orthopedics of Nanchang University, Jiangxi

<sup>5</sup>Queen Mary School, Medical College of Nanchang University, Nanchang, Jiangxi, China

\*These authors contributed equally

## ABSTRACT

Osteosarcoma (OS) is the most prevalent malignant bone tumor in children and young adults. There is an urgent need for a novel biomarker related to the prognosis of OS. We performed a meta-analysis incorporating six independent datasets and performed a survival analysis with one independent dataset GSE21257 in the GEO database for gene screening. The results revealed that one potential biomarker related to OS survival, *POGZ*, was the most significantly upregulated gene. We also verified that the *POGZ* was overexpressed in clinical samples. The survival analysis revealed that *POGZ* is associated with a poor prognosis in OS. Moreover, flow cytometry analysis of isolated OS cells demonstrated that OS cells were arrested in the G<sub>1</sub> phase after *POGZ* knockdown. The RNA-seq results indicated that *POGZ* was co-expressed with *CCNE1* and *CCNB1*. Pathway analysis showed that genes associated with high expression levels of *POGZ* were related to the cell cycle pathway. A cell model was constructed to detect the effects of *POGZ*. After *POGZ* knockdown, OS cell proliferation, invasion and migration were all decreased. Therefore, *POGZ* is an important gene for evaluating the prognosis of OS patients and is a potential therapeutic target.

**Key words:** Prognosis; osteosarcoma; biomarkers; *POGZ*; cell cycle.

**Correspondence:** Prof. Xigao Cheng, The Second Affiliated Hospital of Nanchang University, No.1 Minde Road, EastLaker District, Nanchang, Jiangxi, China. Tel. +86.791.8633-1753. E-mail: 228206846@qq.com

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## Introduction

Osteosarcoma (OS) accounts for approximately 60% of malignant bone tumors in children and young people, making it the most prevalent tumor of this type; it seriously affects patients' quality of life and results in death due to rapid development of the disease.<sup>1,2</sup> Treatment options for OS include surgery, chemotherapy and radiotherapy. Despite the fact that advances in treatment methods have increased the five-year survival rate of OS patients to approximately 60-70%, approximately 30%-40% of patients still develop lung metastases or die.<sup>3,4</sup> It is worth noting that the prognosis of patients with metastatic OS is extremely poor.<sup>5</sup> Prognostic evaluation and early diagnosis are crucial and can effectively improve the survival of OS patients.<sup>6</sup> However, currently available biomarkers cannot effectively predict the prognosis of OS patients.<sup>7</sup> Therefore, there is a pressing clinical need to discover new biomarkers correlated with the prognosis of OS.<sup>6-8</sup> In medical oncology, high-throughput sequencing and gene chip technology are promising tools used to identify potential molecular targets. They provide new methods to explore OS-related genes and predict new tumor molecular interactions, key regulatory molecules, and therapeutic drug targets.<sup>9</sup> Recently, these technologies have been successfully utilized to discover new biomarkers in tumors, including the *MYC*<sup>10,11</sup> and *RAS*,<sup>12</sup> which both promote OS metastasis. In addition, *VEGF*<sup>13</sup> was found to be positively associated with metastasis and a poor prognosis of in OS patients. Nonetheless, these aforementioned methods do not offer many potential biomarkers related to the prognosis of OS.<sup>14</sup> Hence, it is of tremendous clinical significance to discover new biomarkers related to the prognosis of OS and explore their molecular mechanisms. To expand the research on molecules related to the prognosis of OS, we identified a gene known as *POGZ* that is considerably linked to OS survival using bioinformatics. To further evaluate the relationship between the expression level of *POGZ* and OS, we constructed a cytological model verifying the effects of *POGZ* inhibition on OS cell proliferation and metastasis. *POGZ* might be regarded as a potential therapeutic target for OS.

## Materials and Methods

### Clinical specimens

The tissue microarray chip was provided by Typos Biotechnology Company (Xi'an, China) and included 41 OS samples and 19 normal control samples. Table 1 displays the clinicopathological characteristics of the corresponding OS patients.

### Data extraction and analysis of screened genes

In the Gene Expression Omnibus (GEO) database (<http://www.ncbi.nlm.nih.gov/geo/>), we selected 6 expression profiles and downloaded the original (.CEL file) and platform files. Background correction, quantile normalization, professional summary, log<sub>2</sub> conversion and missing values supplementation of the matrix data for each GEO dataset were performed using the "affy" software package and R/Bioconductor software (version 3.5.3) "limma" package. Subsequently, the "mama" package was used to group OS patients and normal controls, perform a meta-analysis, and obtain the z-score value. We used the "meta" package in the R software to download the pan genome project containing multi-tumor expression data to draw a forest map.

The Kaplan-Meier survival curve was plotted using the GEO expression profile and the survival data through the R software "survival" package for single factor Cox survival analysis.

## Immunohistochemical staining

After deparaffinization, the tissue microarray chip was subjected to antigen retrieval and endogenous peroxidase blocking. After serum blocking, polyclonal rabbit anti-human *POGZ* (1:100, Absin, abs132798) was added and incubated overnight at 4°C. Next, the secondary antibody (1:100, Absin, abs20044) of the corresponding species was added, and the DAB staining kit was used to (Vector Laboratories, USA) to detect the signal. The histochemical score was calculated using the Quant Center analysis tool:  $H\ score = \sum(P_i \times I_i) = (\text{percentage of weak intensity cells} \times 1) + (\text{percentage of medium intensity cells} \times 2) + (\text{percentage of high intensity cells} \times 3)$ .<sup>23</sup>

## Receiver operating characteristic (ROC) curve and logistic regression analysis

ROC curve analysis was performed to evaluate the sensitivity (true positive rate) and specificity (true negative rate) of *POGZ* for OS diagnosis. We also investigated the size of the area under the curve (AUC) by using the "pROC" package of R software.

## Biological function and pathway enrichment analysis

Gene set enrichment analysis (GSEA) is a computational method that determines whether an a priori defined set of genes shows statistically significant, concordant differences between two biological states. It can be used to evaluate microarray data at the gene set level. The Kyoto Encyclopedia of Genes and Genomes (KEGG) is a bioinformatics resource. We can use it to study the relationship between genes. Samples from the GSE42352 datasets were divided into two groups based on the expression levels of *POGZ* (median value) and the gene set enrichment analysis (GSEA) software (<http://software.broadinstitute.org/gsea/index.jsp>) was applied for both groups.

## Cell culture and transient transfection

OS cell lines including Saos-2 and U-2OS were supplied by the China Center Type Culture Collection (CCTCC, Shanghai, China). Saos-2 and U-2OS cells were cultured in a DMEM media (Gibco, Waltham, MA) supplemented with 10% FBS (Gibco, Waltham, MA). All the cells were grown at 37°C in 5% CO<sub>2</sub>. Following the manufacturer's recommendations, Turbofect transfection reagent was used (Thermo Fisher Scientific Inc, Shanghai, China) to transfect OS cells with negative control (NC) and *POGZ* siRNA (GenePharma, Shanghai, China) into OS cells. The siRNA sequences targeting *POGZ* were as follows: si*POGZ*-1: 5'-GCCACGAACUGUCCUGUATT-3', 5'-UACAGGAACAGUUCGUGGCTT-3'; si*POGZ*-2: 5'-CCUAAUUAUUCCCUACUUTT-3', 5'-AAGUAGGAAAUGAUUAGGTT-3'; and si*POGZ*-3: 5'-CCACAUGAUCAACAAUCAUTT-3', 5'-

**Table 1.** Clinicopathological characteristics of OS patients.

	Character	Number of case (%)
Grade	G1~G2	17 (0.41)
	G3~G4	24 (0.59)
Sex	F	14 (0.34)
	M	27 (0.66)
Stage	I~II	38 (0.93)
	III~IV	3 (0.07)
Age	<20	13 (0.32)
	>20	28 (0.68)
Location	Lower limb bone	28 (0.68)
	Upper limb bone	13 (0.32)

AUGAUUGUUGAUGAUGUGGTT-3'. The negative control siRNA sequence was as follows: 5'-UUCUCCGAACGUGU-CACGUTT-3', 5'-ACGUGACACGUUCGGAGAATT-3'.

### RNA extraction and real-time quantitative RT-PCR

The total amount of RNA from the OS cells was extracted with the AxyPrep Multisource Total RNA Miniprep Kit (Axygen Scientific, Union City, CA, USA) according to the manufacturer's protocol. A cDNA volume of 20  $\mu$ L was synthesized utilizing the Takara PrimeScript™ RT Reagent Kit and gDNA Eraser (Cat# RR047A, Lot# AK2802) using 1  $\mu$ g total RNA following the measurements of total RNA concentration using the software Quantity One software (PDI Inc., New York, New York). Afterwards, the quantitative real-time PCR (RT-qPCR) primers were synthesized by Xiangyin Biotechnology China (Hangzhou China). RT-qPCR was performed using TB Green™ Premix Ex Taq™ II (TakaRa Code: DRR820A) following the standard protocol with the 7900HT Fast real-time System (Applied Biosystems, Foster City, CA, USA). The thermal cycling program consisted of 1 cycle at 95°C for 1 min, followed by 40 cycles at 95°C for 5 s, and then 60°C for 30 s. *GAPDH* was used as the reference gene. The relative gene expression levels were determined according to the critical threshold (Ct) number and calculated using the  $2^{-\Delta\Delta C_t}$  method. The primers for *POGZ* used for RT-qPCR were as follows: *POGZ* F1 ACCCAGTTTGTAAAGCCGACA, *POGZ* R1 CTGGAGACTGAACAGCTAGTTG, *POGZ* F2 GTGAAGCGACCTGGTGTACA and *POGZ* R2 ACATCGTG-GACATATTTCCGTC. The primers for the housekeeping gene were as follows: 5'-TGACTTCAACAGCGACACCCA-3' (*GAPDH* forward primer) and 5'-CACCTGTTGCTGTAGC-CAAA-3' (*GAPDH* reverse primer).

### Sample collection and high-throughput sequencing

After the U-2OS and Saos-2 cells were transfected with si*POGZ*-2 and si*POGZ*-3 and NC was used as a control. Next, RNA was extracted separately for high-throughput sequencing 48 h later. We used high-throughput sequencing technology to purify mRNA from total RNA and amplified it with PCR technology to build our RNA database. The concentration of the RNA library was detected and diluted to 1 ng/ $\mu$ L. After accurate quantification, the insertion size was qualified and used to cluster with the coded samples to form clusters. Finally, these clusters were sequenced to determine the expression of the desired gene expression.

### Western blotting

Cells were lysed with RAPI buffer 48 h following siRNA transfection, and then centrifuged at 12,000 g for 10 min, and total protein samples were collected. Protein concentration was determined in each sample using the Pierce™ BCA Protein Assay Kit (Thermo Fisher Scientific). Subsequently, 25  $\mu$ g of protein was separated by SDS-PAGE, and then transferred to polyvinylidene difluoride membranes. After being blocked in Tris-buffered saline (TBS) containing 5% non-fat milk for 1 h to saturate additional protein binding sites, the blots were incubated with the following primary antibodies: anti-*POGZ* (1:1,000, Absin, abs132798), anti-E-cadherin (1:5,000, Proteintech, cat20874-1-AP), anti-N-cadherin (1:2,000, Proteintech, cat22018-1-AP), anti-vimentin (1:2,000, Proteintech, cat10366-AP), anti-cyclinE1 (1:1,000, Proteintech, cat11554-1-AP), and cyclinB1 (1:1,000, Proteintech, cat55004-1-AP). The antibodies were maintained at 4°C for 12 h, followed by incubation with horseradish peroxidase-conjugated secondary (anti-mouse or anti-rabbit) IgGs at room temperature for 2 h or 4 h. The proteins were visualized by using a BM Chemiluminescence Western blotting kit (Roche Diagnostics GmbH). To ensure the equal loading and the accuracy of changes

in protein abundance, the level of each protein was normalized to that of GAPDH as a housekeeping control.

### Colony formation assay

Cells were plated on six-well tissue culture plates at a density of 50 cells/cm<sup>2</sup>. Fourteen days later, the colonies were fixed with ethanol and stained with 2% crystal violet, then washed with water to remove the excess dye, and imaged using a scanner. Changes in clonogenicity were quantified by counting the number of colonies, using the ImageJ software.

### Cell proliferation by EdU and Cell Counting Kit-8 (CCK8) assay

To assess the degree of cell proliferation, 5x10<sup>3</sup> U-2OS and Saos-2 cells were plated on 24-well plates, and then the cells were incubated under standard conditions in a complete media. After 48 h following siRNA transfection, cell proliferation was detected based on the incorporation of 5-ethynyl-2'-deoxyuridine (EdU) with the EdU Cell Proliferation Assay Kit. Images were captured using a fluorescence microscope (OLYMPUS, BX53, China). The level of cell proliferation was evaluated using the CCK-8 assay (DOJINDO, Kumamoto, Japan, Cat# CK04). For this purpose, approximately 3,000 cells in 100  $\mu$ L medium were seeded in each well of a 96-well plate, and three independent parallel experiments were set up. The cells were incubated at 37°C in 5% CO<sub>2</sub>, and 10  $\mu$ L CCK-8 reagent was added to the wells at 1, 2, 3, 4, 5 and 6 days, and incubated for 2 h. Finally, the absorbance was measured at a wavelength of 450 nm.

### Cell migration and invasion assay

OS cells in 200  $\mu$ L of serum-free DMEM were added to upper chamber of prepared Transwell plates for the migration and invasion assays, and media containing 10% FBS was added to the lower chamber. The plates were incubated in 5% CO<sub>2</sub> at 37°C overnight. The cells on the upper surface were removed using a cotton bud. The remaining invading cells were fixed and stained with 2% crystal violet. Five representative fields of view for each membrane were selected, then images were taken *via* microscopy, and the number of migrating cells was counted using the ImageJ software.

### Cell cycle analysis by flow cytometry

Fixed cells were stained with propidium iodide (PI) (50  $\mu$ g/mL, Sigma) 48 h after transfection with siRNA targeting *POGZ*. The tests were performed in triplicates. To guarantee the accuracy of the cell cycle analysis, the cells that interfered after 48 hours were centrifuged at 1,500 rpm for 5 min and then resuspended in 500  $\mu$ L of PBS, and 1.5 mL of a 95% ethanol solution (-20°C pre-cooled) was added to fix the cells at -20°C for 10 min. After centrifugation of the cells centrifuged at 1,500 rpm for 10 min, the supernatant was discarded, and 500  $\mu$ L PBS was added to resuspend the hydrated cells for 10 min. Subsequently, RNase A was added at 37°C for 10 min. Ultimately, we added PI (50  $\mu$ g/ml) to DNA and the content was stained for 15 min. Then, the EdU test was performed. EdU analysis and cell cycle determination were performed using BD CellQuest Pro™ on a BD FACSCalibur flow cytometer (BD Biosciences, New Jersey, USA) by obtaining at least 20,000 mononuclear cells.

### Statistical analyses

Statistical analyses were performed using the SPSS software version 22.0 and GraphPad Prism 7 software. All the data are expressed as the mean  $\pm$  standard deviation. One-way analysis of variance (ANOVA) was conducted to compare the data from multiple groups simultaneously. Student's *t*-test was used for data



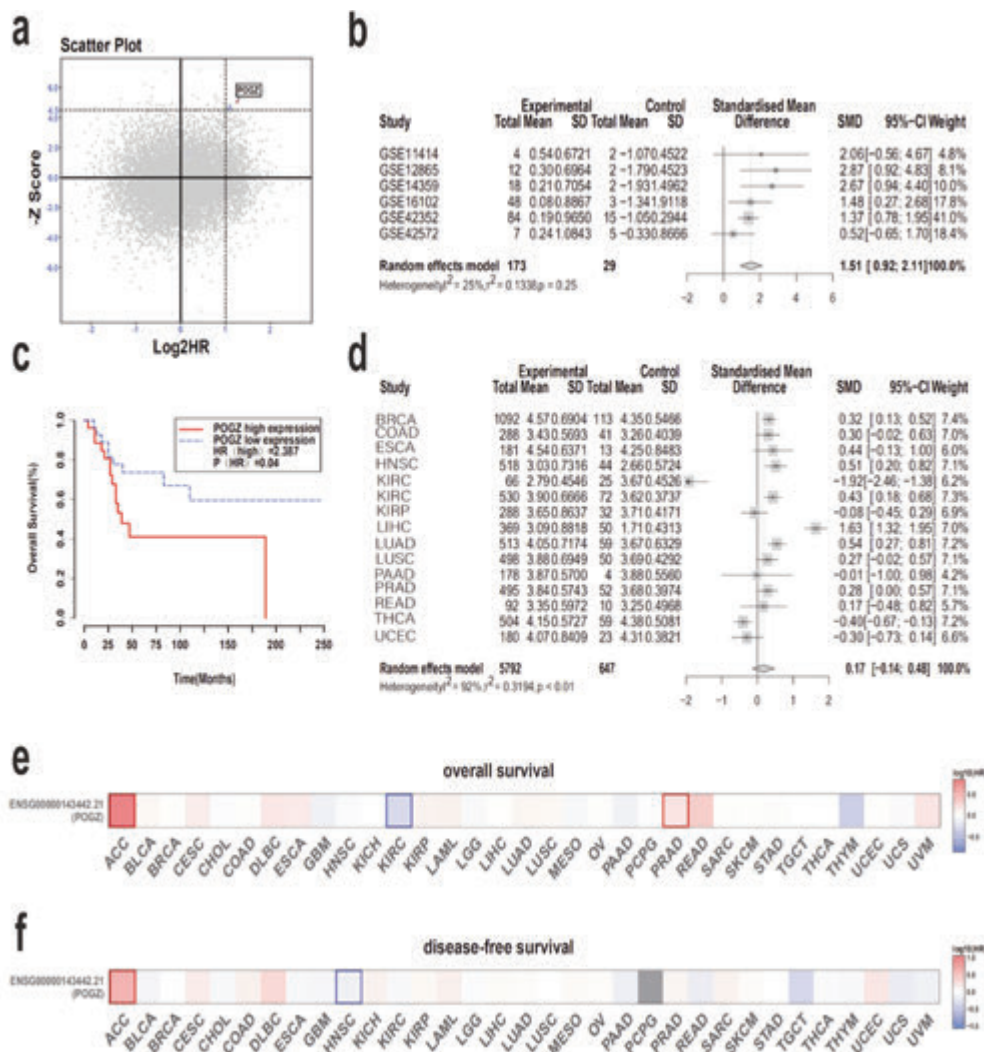
comparison between two groups. For all analyses, two-tailed p-values below 0.05 were considered statistically significant.

## Results

### Identification of genes that are significantly related to the prognosis of OS

After excluding 244 studies involving cell lines and 31 non-conforming studies in the GEO, we selected 6 studies, whose data were included in the GSE11414,<sup>15</sup> GSE12865,<sup>16</sup> GSE14359,<sup>17</sup> GSE16102,<sup>18</sup> GSE42352,<sup>19,20</sup> GSE42572<sup>21</sup> datasets, for inclusion in

the meta-analysis (Supplementary Figure 1). We selected 10,390 genes from the 6 datasets for meta-analysis and obtained the corresponding negative Z score. There were no survival data in the six datasets for meta-analysis, so the GSE21257 dataset was used for survival analysis to obtain the log<sub>2</sub> HR value corresponding to each gene. Based on the cut-off criteria of a negative Z score greater than 4.5 and a log<sub>2</sub> hazard ratio (HR) value greater than 1, we screened out 4 genes that met the criteria (*POGZ*, *CTSE*, *GALNT14*, *HSD11B2*, Figure 1a). Among these 4 genes, we selected the most significantly upregulated gene (with the most negative Z score), *POGZ* (negative Z score=-5.06, log<sub>2</sub> HR=1.26, p=0.04). Additionally, the forest plot revealed that within the 6 datasets, *POGZ* was steadily upregulated and had no heterogeneity among



**Figure 1.** Identify genes that are unfavorable upregulated and associated with a poor prognosis of OS. A) Identifying candidate genes according to the criteria: negative Z score >4.5 and log<sub>2</sub> HR >1; among these 4 genes, *POGZ* had the most significantly upregulated gene (the most negative Z score). b) Forest plot of *POGZ* expression across meta-analysis; the plot revealed that within the 6 datasets, *POGZ* was steadily upregulated and had no heterogeneity ( $I^2 = 25%$ ,  $T^2=0.1338$ ,  $p=0.25$ ). c) KM analysis of overall survival was performed to indicate higher expression of *POGZ* were correlated with the poor survival of OS patients in GSE21257. d) Forest plot of *POGZ* expression in different types of tumors; BRCA, breast invasive carcinoma; COAD, colon adenocarcinoma; ESCA, esophageal carcinoma; HNSC, head and neck squamous cell carcinoma; kidney chromophobe-primary tumor; KIRC kidney renal clear cell carcinoma; KIRP, kidney renal papillary cell carcinoma; LIHC, liver hepatocellular carcinoma; LUAD, lung adenocarcinoma; LUSC, lung squamous cell carcinoma; PAAD, pancreatic adenocarcinoma; PRAD, prostate adenocarcinoma; READ, rectum adenocarcinoma; THCA, thyroid carcinoma; UCEC, uterine corpus endometrial carcinoma. e-f) The relationship between high expression of *POGZ* and prognosis of multiple tumors was drawn.

the datasets ( $I^2 = 25\%$ ,  $T^2 = 0.1338$ ,  $p=0.25$ , Figure 1b).

As illustrated in Figure 1c, the survival analysis of 53 OS patients from the GSE21257 dataset suggested that the high expression level of *POGZ* was significantly related to the poor prognosis of OS ( $p<0.05$ , HR = 2.387). In addition, we performed the survival analysis using the TARGET\_OS and GSE39055 datasets. The results also showed a consistent trend (Supplementary Figures 2 and 3). The data of GSE21257, GSE39055 and TARGET\_OS data were used for meta-analysis. The results showed that *POGZ* was steadily upregulated without heterogeneity among the datasets ( $I^2 = 0\%$ ,  $Z=2.66$ ,  $p=0.008$ , Supplementary Figure 4).

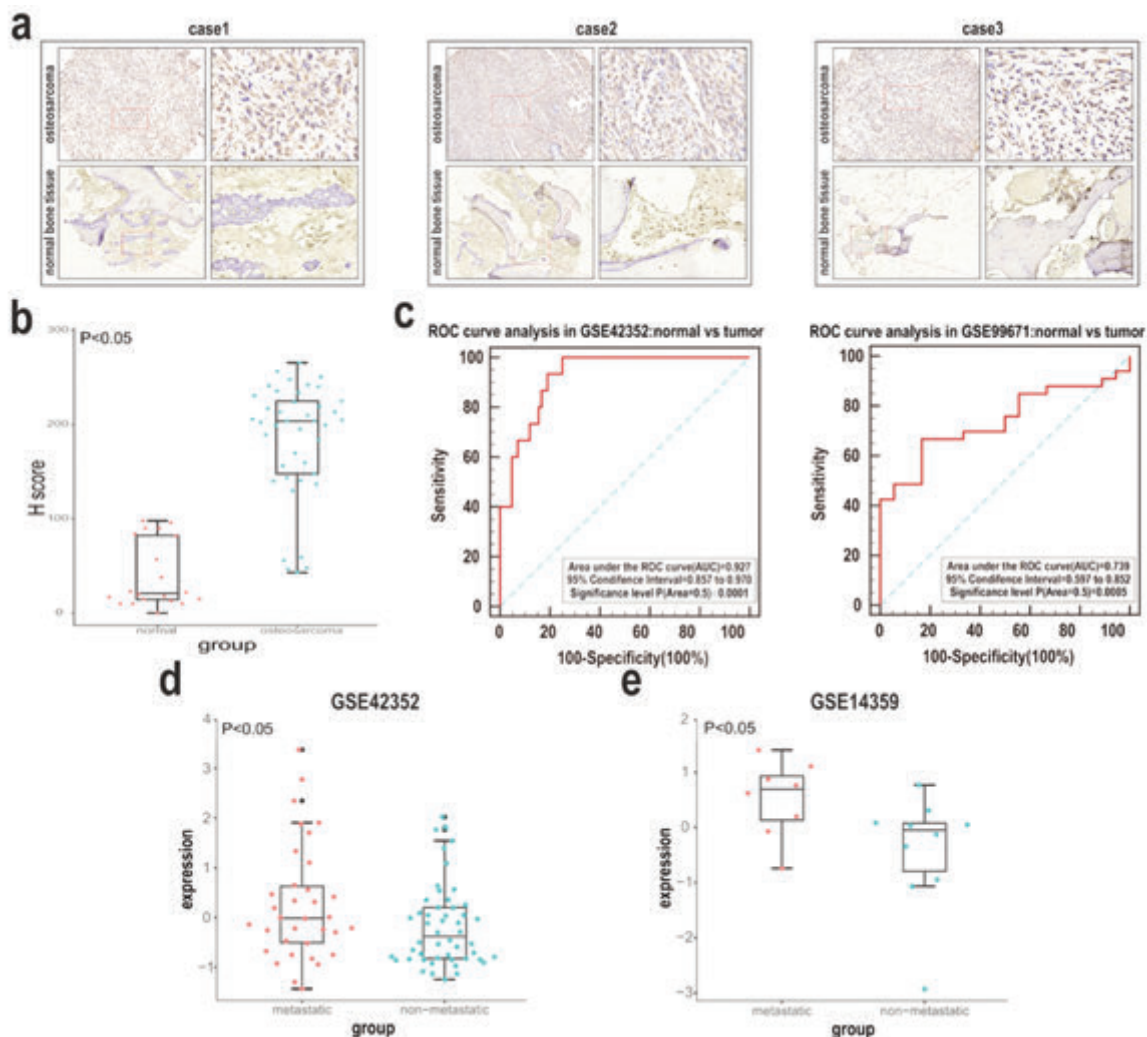
To explore the effects of *POGZ* on a broad range of tumors, we analyzed tumor tissue versus adjacent normal tissue gene expression in 15 tumor types (a total of 7,316 samples), based on the data containing the tumor and controls in The Cancer Genome Atlas (TCGA) database. The results indicated that the expression level of *POGZ* in each tumor was unstable (Figure 1d). Compared with that in the control group, *POGZ* expression was upregulated in liver

cancer ( $P<0.05$ ) and downregulated in renal chromophobe cell carcinoma ( $p<0.05$ ). Moreover, we analyzed the correlation between *POGZ* expression levels and multi-tumor overall survival and disease-free survival, and the results confirmed that the upregulation of *POGZ* in liver cancer was not related to the prognosis ( $p>0.05$ , Figure 1 e-f). Generally, we believe that *POGZ* is an unfavorable risk factor for OS.

### Verification that *POGZ* is upregulated and related to OS metastasis

In order to verify the upregulation of *POGZ* in OS tissue, we performed IHC staining on 41 OS tissues and 19 control tissues. *POGZ* was overexpressed in OS tissues versus control tissues (Figure 2a). Subsequently, we conducted ROC curve analysis to evaluate the sensitivity and specificity of *POGZ* for the diagnosis of OS. The sensitivity and specificity values of the ROC curve were 0.927 and 0.739, respectively based on the GSE42352 and GSE99671 datasets (Figure 2 b,c).

Moreover, based on 6 datasets including metastatic and non-

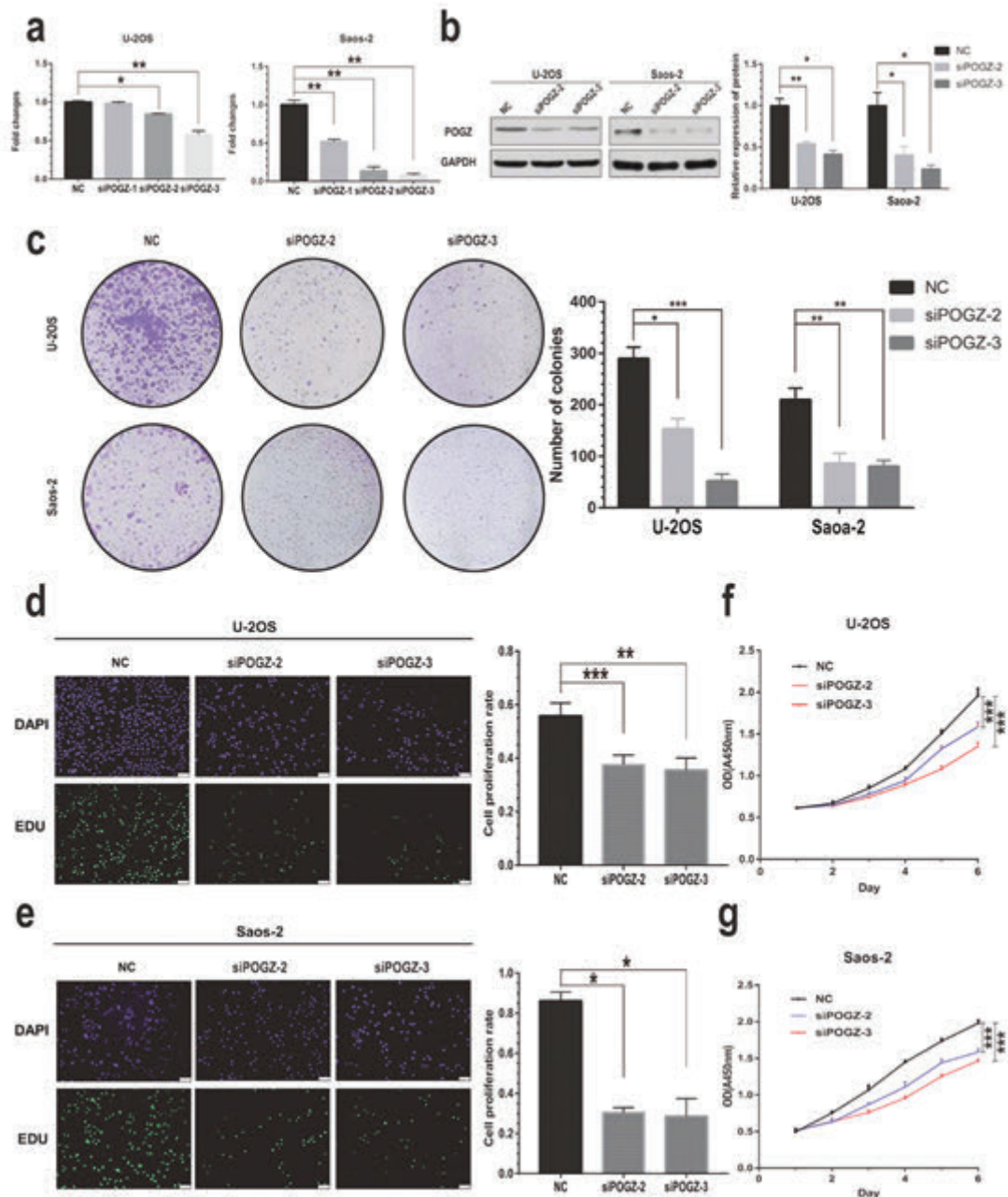


**Figure 2.** *POGZ* expression is upregulated in clinical OS samples and related to metastasis. a) Representative images of IHC staining for *POGZ* in normal bone tissues and OS tissues. b) Quantification of *POGZ* IHC staining in OS (n=41) and normal tissues (n=19). c,d) The ROC curve from GSE42352 and shows that *POGZ* is a potential marker enabling to distinguish OS tissues from normal tissues. e,f) High expression of *POGZ* is associated with metastasis of osteosarcoma; \* $p<0.05$ , \*\* $p<0.01$ , \*\*\* $p<0.001$ .

metastatic OS samples, we developed a forest map of *POGZ* expression level analysis (Supplementary Figure 5). After excluding 4 datasets with small sample sizes, we selected the gene expression profiles and corresponding clinical information in the GSE42352 and GSE14359 datasets for the further independent analysis. Next, we utilized *POGZ* expression levels to perform a *t*-test and found that *POGZ* expression levels were increased in the metastatic groups within the GSE42352 ( $t = 1.90$   $p=0.031$ ) and GSE14359 cohorts ( $t = 2.29$   $p=0.018$ ) were all increased (Figure 2 d,e). Ultimately, high expression levels of *POGZ* play a vital role in the progression of OS.

### *POGZ* knockdown inhibited OS cell proliferation

To determine the impact of *POGZ* on OS proliferation, two siRNA sequences were used to knock down the expression of *POGZ* in U-2OS and Saos-2 cells. RT-qPCR and Western blotting were carried out to gauge the knockout efficiency (Figure 3 a,b). A clonogenic assay proved that compared with that in the NC group, the number of colonies formed by OS cells transfected with si*POGZ*-2 and si*POGZ*-3 was greatly reduced (Figure 3c). The EdU detection method was implemented to evaluate the cell proliferation. This method is an immunochemical detection method that measures the incorporation of nucleotide analogs into newly



**Figure 3.** Cell proliferation assay in *POGZ* inhibited OS cell lines. a) The capability of different siRNAs in terms of downregulating *POGZ* expression. b) The protein levels of *POGZ* by western blot after different siRNAs knock down. c) Representative clonogenic assay of U-2OS cells and Saos-2 cells expressing the indicated plasmids. Quantitative analysis was performed using the ImageJ software. d,e) EdU assays for proliferation rates. f,g) CCK8 assays for proliferation rates; \* $p<0.05$ , \*\* $p<0.01$ , \*\*\* $p<0.001$ .



copied DNA. In U-2OS and Saos-2 cells, the percentage of EdU-positive cells treated with siPOGZ-2 and siPOGZ-3 was significantly reduced (Figure 3 d,e). Additionally, we assessed the proliferation of OS cells using the CCK-8 assay. The growth rate of OS cells with *POGZ* knockdown was decreased compared to that of control cells (Figure 3 f,g). These results indicate that interference with *POGZ* expression inhibits OS cell proliferation.

### Interference with *POGZ* inhibits the migration and invasion of OS cells

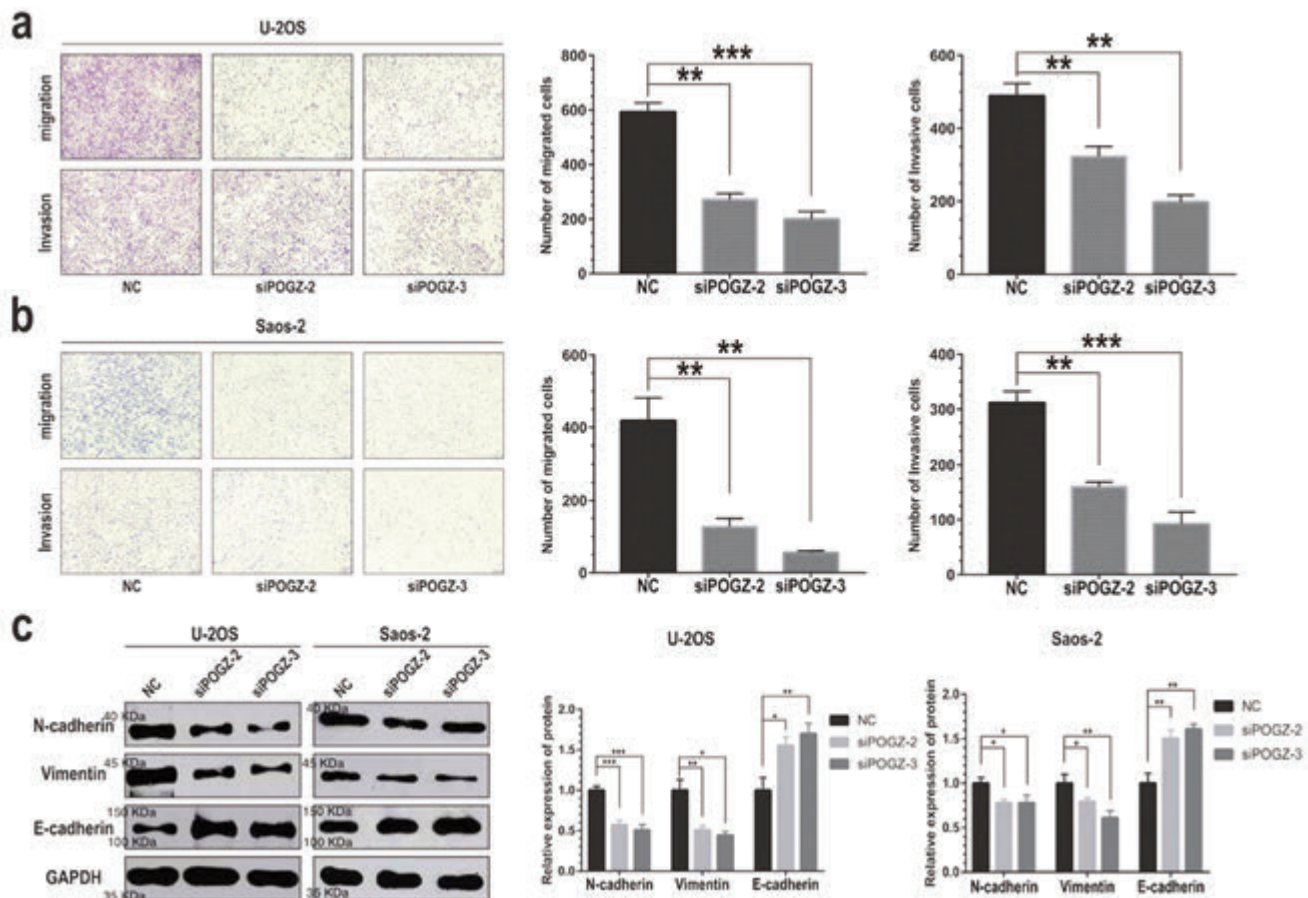
Migration and invasion are key steps in the development of tumors and metastasis of tumors. *In vitro* transwell analysis was used to study the effect of *POGZ* on the migration and invasion of OS cells. Compared with NC cells, U-2OS and Saos-2 cells in the siPOGZ-2 and siPOGZ-3 groups exhibited remarkably reduced migration and invasion (Figure 4 a,b). Furthermore, Western blotting was performed to assess the levels of epithelial-mesenchymal transition (EMT)-related proteins, and the results indicated that vimentin and N-cadherin were both downregulated, while E-cadherin was more upregulated in the U-2OS and Saos-2 cells of the siPOGZ groups than in those of the NC group (Figure 4c). These findings suggest that interference with *POGZ* expression inhibits OS cell metastasis.

### Genes associated with *POGZ* are enriched in cell cycle related processes

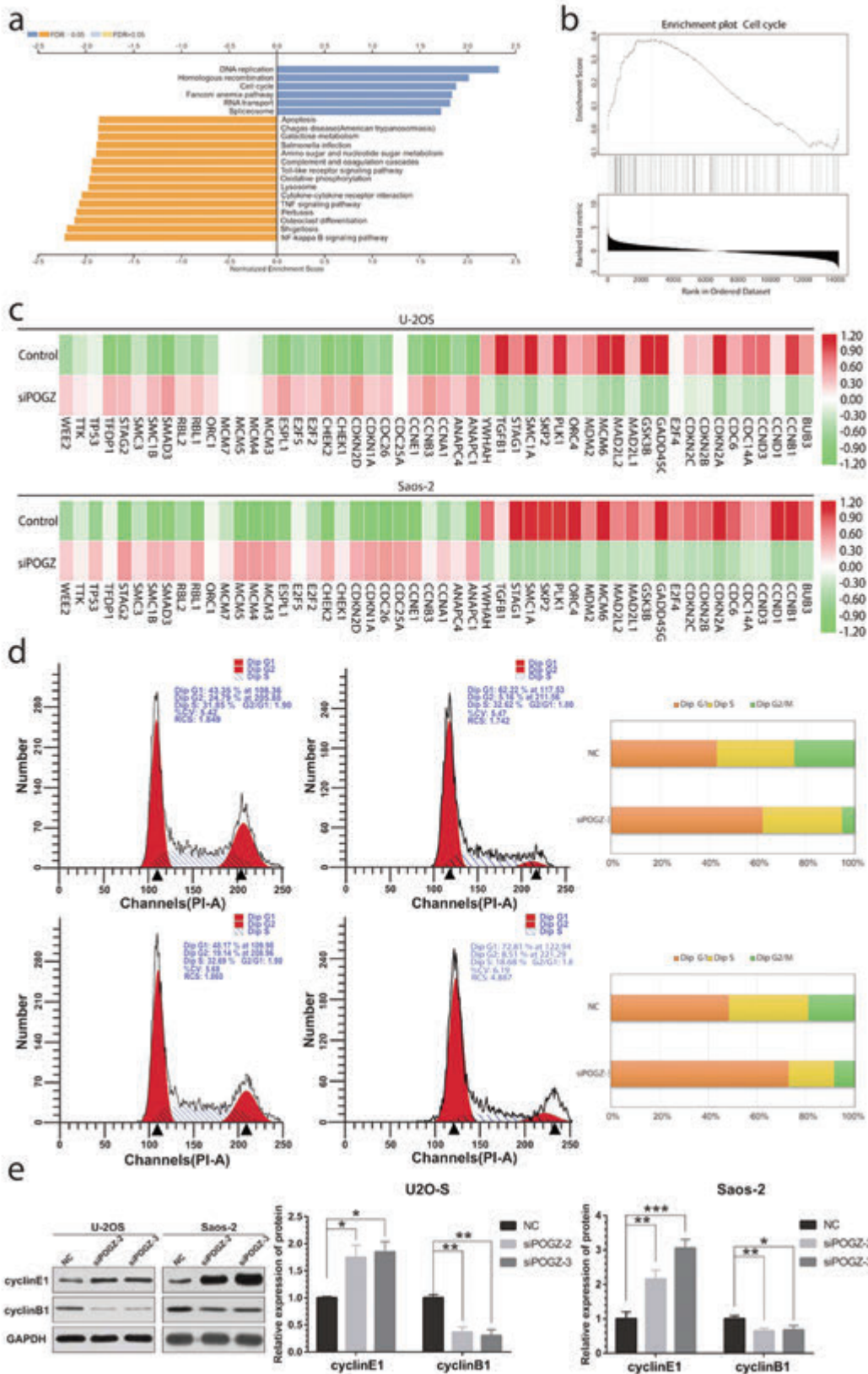
Based on the RNA-seq, the KEGG pathway analysis results revealed that the following pathways were positively related to *POGZ* expression: the DNA replication, homologous recombination, the cell cycle, the Fanconi anemia, RNA transport, spliceosomes pathways. Most of these pathway terms (cell cycle, DNA replication, homologous recombination, and Fanconi anemia) are related to the cell cycle (Figure 5 a,b). To determine which genes in the *POGZ* and cell cycle pathways were co-expressed to affect the growth of OS cells, we used the high-throughput sequencing results to perform a differential analysis, which demonstrated that *CCNE1* was upregulated and *CCNB1* was downregulated (Figure 5c). The above results confirm that *POGZ* has a regulatory effect on cell cycle pathways and also affects the expression of downstream *CCNE1* and *CCNB1*.

### *POGZ* controls the cell cycle progression *in vitro*

To verify the results of our biological analysis and high-throughput sequencing, we performed a flow cytometric analysis of isolated OS cell cycle. OS cells with *POGZ* knockdown exhibited cell cycle arrest in the G<sub>1</sub> phase, confirming the role of *POGZ* in regulating cell cycle progression (Figure 5d). Western



**Figure 4.** *POGZ* inhibitions prevent the metastasis and invasion of OS cells. a,b) Transwell assays showed that *POGZ* inhibitions significantly reduced the migration (upper) and invasion (lower) of OS cells. c,d) EMT markers were detected by Western blot; \* $p < 0.05$ , \*\* $p < 0.01$ , \*\*\* $p < 0.001$ .



**Figure 5.** Genes associated with *POGZ* were enriched in cell cycle related processes and *POGZ* controls the cell cycle progression *in vitro*. a) KEGG pathway enrichment analysis of genes positively and negatively associated with *POGZ*. b) *POGZ* is associated with cell cycle progression. c) Heat map of core-DEGs related to cell cycle process in *POGZ* inhibited U-2OS and Saos-2 cells; the heat map reveals that *CCNE1* was upregulated and *CCNB1* was downregulated in OS cells after *POGZ* inhibition. d) Representative images (left) and quantification (right) of negative control (si*POGZ*-3-transfected U-2OS and Saos-2 cells) were analyzed in the cell cycle assay. e) Cell cycle related makers were detected by Western blot; \* $p < 0.05$ , \*\* $p < 0.01$ , \*\*\* $p < 0.001$ .



blot detection of cycle-related proteins revealed that *CCNE1* was upregulated and *CCNB1* was downregulated in knockdown cells compared to control cells (Figure 5e). Overall, *POGZ* was shown to affect the cell cycle of OS cells by participating in the regulation of the G1 phase.

## Discussion

*POGZ* is an important gene for evaluating the prognosis of OS patients and is highly expressed in OS cells. Inhibiting the expression of *POGZ* can reduce OS cell proliferation and tumor metastasis. *POGZ* regulates OS cell cycle-related processes, but the mechanism by which OS cells become arrested in the G1 phase after *POGZ* inhibition requires more in-depth investigation.

In this study, four candidate genes (*POGZ*, *CTSE*, *GALNT14*, *HSD11B2*) met our inclusion criteria: highly expressed in OS and related to a poor prognosis. *POGZ* contains at least one short region called the zinc finger domain and also contains a centromere protein (CENP)-B-like DNA binding domain and a DDE domain derived from a transposase encoded by a pogo-like DNA transposon. *POGZ* is involved in normal kinetochore assembly, cohesion of mitotic sister chromatids, and separation of mitotic chromosomes.<sup>26</sup> Previous genetic findings have shown that *POGZ* is responsible for various neurodevelopmental disorders, including autism spectrum disorders and intellectual disabilities, and is also considered to be related to 'synaptic disorders'.<sup>27</sup> Since *POGZ* is the most upregulated of the four target genes, as well as the most harmful to the survival of OS patients, and given that there is no literature report revealing its impact on tumorigenesis, we decided to conduct an in-depth research on it.

Based on the TCGA database, we used Gene Expression Profiling Interactive Analysis (GEPIA) to analyze the relationship between *POGZ* expression and the prognosis of multiple tumors. Supplementary Figures 6 and 7 show that *POGZ* acts as a survival protective factor in other solid tumors (disease-free survival HR = 0.91, p=0.017; overall survival HR = 0.9, p=0.0065). Interestingly, these results indicate that high expression of *POGZ* is a specific risk factor in OS. Furthermore, based on the gene expression profiles and clinical information of the two datasets, we divided the dataset samples into metastatic and non-metastatic OS groups, and found that the expression of *POGZ* was significantly higher in the metastatic OS group.

In terms of molecular phenotype, we established a cell model with *POGZ* downregulation to explore the role of *POGZ* in OS. The proliferation and colony formation of OS cells decreased after *POGZ* knockdown. Transwell experiments demonstrated that the migration and invasion abilities of OS cells were weakened after *POGZ* knockdown. EMT is a process involving the invasion and metastasis of various tumors.<sup>28</sup> In this study, we found that the expression of EMT-related proteins vimentin, N-cadherin and E-cadherin in OS cells was positively correlated with the expression of *POGZ*. This finding indicates that *POGZ* could induce the metastasis of OS *via* EMT.

The function of many oncogenes is to either stimulate cell division or counteract cell cycle arrest.<sup>29</sup> Western blot analysis of cell cycle-related gene levels by Tanak *et al.* showed that OS cells that were not treated with a gamma-secretase inhibitor (GSI) had higher expression levels of cyclin E1 and a higher number of cells in the G<sub>1</sub> phase than those treated with a GSI treatment.<sup>30</sup> In an experiment to study the effect of TCTP expression on the growth of OS cells, Shen *et al.* found that cyclinB1 protein levels in Saos-2 and U2O-S cells decreased after Lv-shTCTP infection and the proportion of cells in the G<sub>2</sub>/M phase was significantly reduced.<sup>31</sup> In our study, the KEGG pathway analysis in Figure 5a indicates that the

OS cell cycle pathway is activated, and the first four pathways with the most significant enrichment are all cell cycle related. According to *POGZ* and OS cell cycle pathway gene co-expression annotation analysis, we found that after knocking down *POGZ*, *CCNE1* was upregulated and *CCNB1* was downregulated. Western blotting was used to verify the above results. *CCNB1* (cyclin B) is necessary for adequate control of the G<sub>2</sub>/M transition phase of the cell cycle.<sup>32</sup> *CCNE1* (cyclin E) is a positive regulator of the cell cycle. It promotes the G<sub>1</sub>/S phase transition by binding to *CDK2* and activating *CDK2*.<sup>33</sup> Thus, we believe that *POGZ* regulates the OS cells, thereby affecting cell proliferation. However, the limitation of our research was the underlying molecular mechanism was not investigated.

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