Overexpression of microRNA-129-5p in glioblastoma inhibits cell proliferation, migration, and colony-forming ability by targeting ZFP36L1

Xu Guo, Haozhe Piao*, Ye Zhang, Peixin Sun, Bing Yao

ABSTRACT

Glioblastoma multiforme (GBM) is a highly invasive cancer with a high recurrence rate. The prognosis of GBM patients remains poor, even after standard surgical resection combined with chemoradiotherapy. Thus, there is an urgent need for new therapeutic targets in GBM. In recent years, microRNAs have received considerable attention due to their important role in tumor development and progression. In this study, we investigated the role of miR-129-5p and miR-129-5p/ZFP36L1 axis in GBM tumorigenesis. Analysis of GSE103228 microarray data from the GEO database showed that miR-129-5p was significantly downregulated in GBM vs. normal brain tissues. Quantitative reverse transcription PCR analysis of miR-129-5p expression in seven GBM cell lines (LN229, A172, U87, T98G, U251, H4, and LN118) vs. normal human astrocytes (NHA) showed miR-129-5p was significantly downregulated in GBM cells. Overexpression of miR-129-5p in LN229 and A172 cells significantly suppressed cell proliferation, migration, invasion, and colony-forming ability. Target Scan analysis identified ZFP36L1 as the target of miR-129-5p. UALCAN dataset analysis found that ZFP36L1 was significantly upregulated in GBM vs. normal brain tissues, and high ZFP36L1 expression was positively associated with poor survival of GBM patients. Western blot analysis demonstrated that ZFP36L1 was significantly upregulated in seven GBM cell lines vs. NHA. Overexpression of miR-129-5p in LN229 and A172 cells significantly inhibited ZFP36L1 mRNA and protein expression, while overexpression of ZFP36L1 in LN229 and A172 cells reversed miR-129-5p-mediated inhibition on GBM tumorigenesis. Our results revealed an important role of miR-129-5p in the negative regulation of ZFP36L1 expression in GBM, suggesting new candidates for targeted therapy in GBM patients.

KEYWORDS: Glioblastoma multiforme; GBM; microRNA; miR-129-5p; ZFP36L1; migration; proliferation; colony-forming ability; tumor-suppressor
MicroRNAs are also involved in glioma development [13-17]. Overexpression of miR-148a and miR-378-3p suppressed GBM cell proliferation, migration, and invasion by targeting integrin alpha 9 (ITGA9) and transmembrane 4 superfamily member 17 (tetraspanin 17, TSPAN17), respectively [18,19]. In addition, the expression of microRNAs was found to be associated with chemoresistance of glioblastoma and non-small cell lung cancer [20-22]. Upregulation of miR-145 enhanced the sensitivity of GBM stem cells to demethoxycurcumin, and overexpression of miR-29a significantly inhibited the proliferation of GBM stem cells.

In recent years, miR-129-5p has received considerable attention as a driver of Alzheimer’s disease [23], osteogenesis [24], and intervertebral disk degeneration [25]. Furthermore, miR-129-5p was reported to be involved in the tumorigenesis of gastric [26-28], pancreatic [29], laryngeal [30], prostate [31], and lung cancer [32] as well as of hepatocellular carcinoma [33]. Recent reports demonstrated that miR-129-5p was able to regulate the proliferation and invasion of GBM cell lines [16,34]. However, despite these studies, the specific regulatory mechanisms of miR-129-5p in glioma remain unclear and require further investigation.

ZFP36 ring finger protein-like 1 (ZFP36L1) is a member of the ZFP36 family of RNA-binding proteins [35], which contains two standard repeating zinc finger motifs (CCCH). The CCCH motif binds to the AU-rich elements (AREs) in the 3’-UTR of the target mRNA resulting in mRNA instability [36]. It was reported that ZFP36L1 was functionally involved in post-transcriptional regulation of bile acid metabolism in hepatocytes, myogenesis, and lymphocyte development [37-40]. Disruption of ZFP36L1 expression in mice led to death on embryonic day 11 [41]. Increased expression of ZFP36L1 inhibited cell growth in lung and colorectal cancer cell lines [42,43]. However, little is known about the biological role of ZFP36L1 and its regulatory factors (such as microRNAs) in GBM tumorigenesis.

In this study, we investigated the role of miR-129-5p and miR-129-5p/ZFP36L1 axis in GBM tumorigenesis. First, we used the GSE103228 microarray data from the gene expression omnibus (GEO) database to analyze differentially expressed microRNAs between GBM and normal tissues and found that miR-129-5p was significantly downregulated in GBM tissues. Subsequently, we confirmed that overexpression of miR-129-5p inhibited the proliferation, migration, invasion, and colony-forming ability of LN229 and A172 GBM cells. Furthermore, we identified that ZFP36L1 was the downstream target gene of miR-129-5p. Overexpression of miR-129-5p in LN229 and A172 cells inhibited the expression of ZFP36L1, while forced expression of ZFP36L1 reversed miR-129-5p-mediated inhibition on GBM tumorigenesis, including tumor proliferation, migration, and colony-forming ability. Our results reveal an important role of the miR-129-5p/ZFP36L1 axis in GBM tumorigenesis and shed a new light on targeted therapy in GBM.

MATERIALS AND METHODS

Analysis of microarray data

The microRNA and mRNA expression profile GSE103228 was selected from the GEO database (https://www.ncbi.nlm.nih.gov/geo/). The profile included mRNA and microRNA expression of five human GBM and five normal brain tissues. Analysis of microRNA expression was performed on the GPL18058 platform (Exiqon miRCURY LNA microRNA array, 7th generation). The differentially expressed microRNAs in GBM samples were further analyzed by the GEO’s R language analysis tool (http://www.ncbi.nlm.nih.gov/geo/geo2r/). Adjusted p values were applied based on the Benjamini-Hochberg method to minimize the false discovery rate. To determine dysregulated microRNAs in GBM samples, the cut-off values were defined as the fold change >2 and adjusted p < 0.05 was used. After removing unknown microRNAs and missing information, we found that miR-129-5p was among the top three dysregulated microRNAs and was selected for further analysis.

Identification of microRNA target genes

The candidate target genes of miR-129-5p were identified using TargetScan (http://www.targetscan.org/vert_72/), which is dedicated to the analysis of mammalian microRNA target genes. Based on the cumulative weighted context++ score, ZFP36L1 was among the top 25 target candidates for miR-129-5p. After literature search, pre-analysis of TCGA, and removal of proteins with unknown function, ZFP36L1 remained as the promising candidate gene and was selected for further analysis.

UALCAN dataset analysis

The UALCAN (http://ualcan.path.uab.edu/index.html) database provides an easy access to publicly available cancer transcriptomics data (namely, TCGA and MET500 transcriptome sequencing). In this study, UALCAN was used to identify biomarkers and to validate in silico potential genes of interest. In addition, ZFP36L1 gene expression and patient survival information based on the gene expression were analyzed with the UALCAN dataset to explore ZFP36L1 impact on the prognosis of patients with GBM.
Cell culture

Normal human astrocyte (NHA) cells and human GBM cell lines (LN229, A172, U87, T98G, U251, H4, and H118) were obtained from the Institute of Biochemistry and Cell Biology, Chinese Academy of Sciences (Shangh hai, China). These cells were cultured in RPMI-1640 medium (Invitrogen, Carlsbad, CA, USA) containing 10% fetal bovine serum (FBS, Invitrogen) in a humidified atmosphere of 5% CO₂ at 37°C.

RNA isolation and qRT-PCR

Total RNAs were extracted from cultured cells as described previously [44]. In brief, total RNAs were extracted by TRIZOL reagent (Gibco Invitrogen), and the quality and integrity of the extracted RNAs were determined by agarose gel electrophoresis. Poly(A) tails were added according to the protocol of the Poly(A) Tailing Kit (Tiangen, Beijing, China). Complementary DNA (cDNA) was synthesized using the PrimeScript™ RT reagent Kit with gDNA Eraser (Takara, Dalian, China) and gene-specific primers or random primers. qRT-PCR was performed on a LightCycler 480 II Real-Time PCR system (Roche Diagnostics, Basel, Switzerland) using SYBR Green (Takara). Glyceraldehyde 3-phosphate dehydrogenase (GAPDH) and U6 small nuclear RNA (snRNA) were employed as endogenous controls for mRNA and microRNA, respectively. The comparative Ct method was used to calculate the relative expression of RNA. The primers used in this study were as follows:

<table>
<thead>
<tr>
<th>miR-129-5p primers</th>
<th>miR-NC 5'-UUGUACUACACAAAAGUACUG-3'</th>
</tr>
</thead>
<tbody>
<tr>
<td>ZFP36L1: 5'-ACCACCACCCTCGTGTCTG-3'</td>
<td>U6: 5'-CGGATTTGGTCGTATTGGG-3'</td>
</tr>
<tr>
<td>GAPDH: 5'-TTCACGAATTTGCGTGTCAT-3'</td>
<td>U6: 5'-CGCTTCGGCAGCACATATAC-3'</td>
</tr>
<tr>
<td>CTGGAAGATGGTGATGGGATT-3'</td>
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Western blot analysis

Western blot analysis was performed as described previously [45]. In brief, cells were lysed with lysis buffer (Beyotime, Shanghai, China) containing 1 mmol/L phenylmethylsulfonyl fluoride (PMSF). After centrifugation at 14,000 × g for 10 min at 4°C, the extracted proteins were resuspended, and the concentrations were measured using the bicinchoninic acid (BCA) protein assay kit (Beyotime). Equal amounts (30 μg) of protein were resolved by SDS-PAGE and electo-transferred onto the PVDF membrane (Millipore). The immunoblots were then blocked with 5% fat-free milk for 1 h, and further incubated overnight with primary antibodies against ZFP36L1 (1:1000 dilution, ab42473, Abcam, USA) and GAPDH (1:3000 dilution, ab82602, Abcam, USA) at 4°C. After incubation with peroxidase-conjugated secondary antibodies for 1 h, the blots were washed and then visualized using an electrochemiluminescence (ECL) detection kit (ThermoBiotech Inc, Rockford, IL, USA). The chemiluminescent signal on the blots was determined and quantified using densitometric software (Bio-Rad, California, USA).

Transfection

Transfections were performed using the Lipofectamine 3000 Reagent (Invitrogen) according to the manufacturer’s instruction. For the transfection of microRNAs, 50 nmol/L of miR-129-5p mimic (5'-CUUUUUGCGGCUUGCGCUACG-3') and/or negative control miR-NC (5'-UUGUACUCACAAAAGUACUG-3') were added to each transfection well (6-well culture plate) containing 2 ml culture medium. After 48 h of transfection, total RNAs and proteins were collected for further analysis. For the overexpression of ZFP36L1, 2 × 10⁵ of LN229 and A172 cells were seeded onto 6-well plates. After overnight culture, the cells were transfected with 0.75 μg/mL of pcDNA3.1 (control) or pcDNA3.1-ZFP36L1 for 48 h. The transfection efficiency of ZFP36L1 was evaluated by qRT-PCR and Western blot analysis. In order to visualize and count the numbers of cells, cells were fixed with 4% paraformaldehyde at 4°C for 60 min. After washing twice with ice-cold PBS, cells were stained with impurity-free crystal violet for 2 min and imaged under a microscope.

Cell proliferation assay

Cell proliferation was determined using the Cell Counting Kit-8 (CCK-8) kit (Dojindo, Japan) according to the manufacturer’s instructions. In brief, cells (3000 cells/well) were transfected with NC (transfection reagent control), miR-NC (miRNA sponge-negative scramble control), or miR-129-5p mimics and cultured in 96-well plates. After 0, 24, 48, and 72 h of incubation, 100 μL of culture supernatant from each well was collected and transferred to another 96-well plate and then incubated with 10 μL of CCK-8 solution at 37°C for 4 h. The absorbance was measured at 450 nm wavelength using spectrophotometry (BioTek, USA).

Wound healing assay

Cells were seeded in 6-well plates and then treated with NC, miR-NC, or miR-129-5p mimics. Linear scratch wounds were created on cell monolayers using a sterile 200-μL pipette tip, and the scratched areas were imaged at ×100 magnification using a Leica DMI3000B computer-assisted microscope (Leica, Buffalo Grove, IL, USA). Images were captured at 0, 24, 48, and 72 h after scratching and analyzed using Image-Pro Plus v6.0 image analysis software (Media Cybernetics, Rockville, MD, USA). The remaining wound area was calculated to estimate the migration distance of the cells.
Matrigel invasion assay

Matrigel invasion assay was performed using Transwell plates (BD, Biosciences, CA, USA). Transwell membrane inserts were precoated with Matrigel (BD), and the analyzed cells were appropriately seeded onto the upper compartment of the chamber. The bottom chamber contained complete RPMI-1640 medium, and the upper chamber contained serum-free medium. After 24 h of incubation at 37°C, the cells on the Matrigel-coated upper surface were wiped off with a cotton swab, and the invaded cells throughout the filter were fixed and stained with Giemsa at 37°C for 3 h. The images of invaded cells were photographed with a microscope (Leica) at ×100 magnification. For each membrane isolated from triplicate chambers, the number of invaded cells was counted in three randomly selected fields. The results were presented as bar graphs.

Statistical analysis

All statistical analyses were performed using IBM SPSS Statistics for Windows, Version 20.0 (IBM Corp., Armonk, NY, USA). Data were expressed as mean ± SD. The χ² test, Student’s t-test, and one-way ANOVA analysis were used for comparisons. A value of \( p < 0.05 \) was considered statistically significant.

RESULTS

Downregulated miR-129-5p expression in GBM tissues and cell lines

The microarray dataset GSE103228 was used to examine whether miR129-5p expression was associated with GBM tumorigenesis. After analyzing 3557 differentially expressed microRNAs, we found that the expression of miR-129-5p was significantly downregulated in GBM tissues vs. normal brain tissues (\( p < 0.005 \), Figure 1A). The decreased expression of miR-129-5p in GBM was further validated using NHAs and seven GBM cell lines, including LN229, A172, U87, T98G, U251, H4, and LN118. As shown in Figure 1B, miR-129-5p was significantly downregulated in all GBM cell lines with a reduction of approximately 60–90% compared to NHAs, suggesting that miR-129-5p may play an important role in the tumorigenesis of GBM.

MiR-129-5p inhibited GBM tumorigenesis

To explore the tumorigenic role of miR-129-5p in GBM, the two GBM cell lines LN229 and A172 with the least expression of miR-129-5p were selected. We first overexpressed miR-129-5p in LN229 and A172 cells and then examined whether miR-129-5p could affect GBM cell proliferation, migration, invasion, and colony-forming abilities. Compared to NC group, transfection with miR-NC did not alter the expression of miR-129-5p in LN229 and A172 cells (Figure 2A). Compared to miR-NC group, the level of miR-129-5p was significantly increased in miR-129-5p mimics groups, to about 11.5-fold and 9.5-fold in LN229 and A172 cells, respectively.

Next, CCK-8 assay was performed to examine whether miR-129-5p affects the cell proliferation ability. The overexpression of miR-129-5p significantly inhibited the proliferation of LN229 and A172 cells after 72, 96, and 120 h of culture (Figure 2B), indicating that miR-129-5p was functionally involved in the tumor cell proliferation.

Further, miR-129-5p-mediated proliferation and migration inhibition in GBM was examined by wound-healing assay, a classic method for a comprehensive analysis of cell proliferation and migration. As shown in Figure 3A, the two control groups (NC and miR-NC) of LN229 and A172 cells exhibited
high motility, while the overexpression of miR-129-5p in mimics group reduced the wound healing ability after 72 h of culture. The wound closure rates of LN229 and A172 cells transfected with miR-129-5p mimics were significantly lower at 24, 48, and 72 h compared to those in miR-NC-transfected cells \( (p < 0.05) \) (Figure 3B).

Using colony forming assay, we next examined whether miR-129-5p could affect the ability of GBM cells to form colonies. The results showed that the overexpression of miR-129-5p significantly inhibited colony formation in both LN229 and A172 cells (Figure 3C).

Taken together, the results of CCK-8, wound-healing, and colony-forming assay suggested that miR-129-5p can suppress GBM cell proliferation, migration, and colony-forming ability.

ZFP36L1 as a target gene of miR-129-5p

To further investigate the underlying mechanism of miR-129-5p-mediated inhibition of cell proliferation, migration and colony-forming ability in GBM, TargetScan bioinformatics database was used and ZFP36L1 was identified as a potential target gene of miR-129-5p (Figure 4A). UALCAN was used to examine the expression of ZFP36L1 in GBM samples. As shown in Figure 4B, the expression of ZFP36L1 in GBM samples \( (n = 156) \) was significantly higher than that in normal brain samples \( (n = 5) \) \( (p < 0.0005) \). In addition, survival analysis showed that patients with higher expression of ZFP36L1 had a significantly increased risk of death (Figure 4C).

Next, the expression of ZFP36L1 was analyzed in NHA and GBM cell lines. As shown in Figure 4D, ZFP36L1 was significantly upregulated to 2–6.5 fold in the seven GBM cell lines, suggesting that miR-129-5p was negatively correlated with ZFP36L1 in GBM. To further validate this hypothesis, miR-129-5p mimics was overexpressed in LN229 and A172 cells, and the expression of ZFP36L1 was determined by qRT-PCR. As shown in Figure 4E, the overexpression of miR-129-5p significantly inhibited ZFP36L1 mRNA expression in both LN229 and A172 cells \( (p < 0.05) \). The results of Western blot analysis (Figure 4F) further supported the hypothesis that miR-129-5p overexpression suppresses ZFP36L1 protein expression. Taken together, these results indicated that miR-129-5p negatively regulates the expression of ZFP36L1 in GBM.
Overexpression of ZFP36L1 reverses the inhibitory effect of miR-129-5p in GBM

Next, we examined whether miR-129-5p-regulated ZFP36L1 expression is involved in GBM proliferation, migration, and colony forming ability. LN229 and A172 cells with overexpressed miR-129-5p were transfected with pcDNA3.1-ZFP36L1 to increase the level of ZFP36L1, and the cells in the control group were transfected with pcDNA3.1 vector. The expression of ZFP36L1 was confirmed by qRT-PCR and Western blot (Figure 5A). Compared to pcDNA3.1 control group, transfection with pcDNA3.1-ZFP36L1 successfully increased the mRNA and protein levels of ZFP36L1 in both LN229 and A172 cells (p < 0.05). Afterwards, we examined whether miR-129-5p-mediated inhibition of GBM cell proliferation could be reversed by increased expression of ZFP36L1. As shown in Figure 3B, ZFP36L1 overexpression significantly restored miR-129-5p-mediated proliferation inhibition in LN229 and A172 cells (p < 0.05), suggesting that ZFP36L1 plays a crucial role in GBM proliferation.
We next examined the role of ZFP36L1 in GBM cell migration, invasion, and colony forming ability. The results of wound-healing assay showed that ZFP36L1 overexpression increased tumor migration ability (Figure 6A). Compared to empty-vector transfected cells, the open wound area in pcDNA3.1-ZFP36L1 transfected cells was significantly reduced at 24, 48, and 72 h of culture in LN229 and A172 cells \( (p < 0.05, \text{Figure 6B}) \). Furthermore, miR-129-5p-mediated inhibition of colony-forming ability was restored by ZFP36L1 overexpression (Figure 6C).
Collectively, these experiments indicated that miRNA-129-5p negatively regulates ZFP36L1 expression to affect the proliferation, migration, and colony-forming abilities of LN229 and A172 GBM cells.

**DISCUSSION**

Despite multiple treatment modalities are available for GBM, GBM remains the most common primary malignant brain tumor, with a poor prognosis. In the past few years, increasing attention has been paid to the role of microRNAs in the development, diagnosis, and prognosis of gliomas. In this study, we demonstrated that miR-129-5p and miR-129-5p-regulated target gene ZFP36L1 are involved in GBM. We found that miR-129-5p was markedly downregulated in GBM tissues and seven GBM cell lines. Overexpression of miR-129-5p suppressed GBM cell proliferation, migration, and colony-forming ability. Furthermore, we identified that ZFP36L1 was the downstream target of miR-129-5p and was responsible for miR-129-5p-mediated tumor proliferation, migration, and colony-forming ability. Overexpression of ZFP36L1 in miR-129-5p-overexpressing GBM cells successfully restored all the features of GBM tumorigenesis. Overall, our study suggests that during the development of GBM, miR-129-5p expression is reduced and leads to higher expression of ZFP36L1, which results in increased proliferation, migration, and colony forming ability of GBM cells.

MiR-129-5p has recently received considerable attention as a tumor suppressor. In gastric cancer, miR-129-5p was found to attenuate tumor progression by targeting high mobility group box 1 (HMGB1) [27], WW domain containing E3 ubiquitin protein ligase 1 (WWP1) [28], disintegrin and metalloproteinase domain-containing protein 9 (ADAM9) [46], solute carrier family 2 member 3 (SLC2A3) [47], collagen, type I, alpha 1 (COLA1) [48], and interleukin (IL)-8 [49]. In addition, miR-129-5p was shown to regulate cell invasion and metastasis in lung cancer [32], hepatocellular carcinoma [50], pancreatic [29], and prostate cancer [31]. Gao et al. found that the expression of miR-129-5p was significantly decreased in human prostate cancer tissues, and downregulation of miR-129-5p...
increased ETS variant transcription factor 1 (ETV1) expression. Overexpression of ETV1 reversed miR-129-5p mimics inducing cell growth arrest [31]. On the other hand, Gu et al. found that DNA (cytosine-5)-methyltransferase 3A (DNMT3A), a tumor suppressor in glioma, was negatively regulated by miR-129-5p [51]. Overexpression of DNMT3A partially attenuates miR-129-5p-inhibited cell proliferation, suggesting that other unidentified downstream targets of miR-129-5p may affect GBM proliferation as well as other features of GBM tumorigenesis, such as migration and colony-forming ability.

In this study, miR-129-5p was found to be significantly downregulated in LN229, A172, U87, T98G, U251, H4, and LN118 cells and in GBM tissue samples. To further explore the specific biological role of miR-129-5p in GBM, the two cell lines with the lowest expression of miR-129-5p, LN229 and A172, were selected to restore the expression of miR-129-5p. Notably, miR-129-5p overexpression not only inhibited cell proliferation and migration, but also suppressed the colony-forming ability. These results indicated that miR-129-5p can be considered as a tumor suppressor during GBM tumorigenesis.
development and that miR-129-5p may be used as a potential diagnostic and prognostic biomarker in GBM patients.

In addition to its association with tumorigenesis, miR-129-5p was implicated in other diseases. A study by Yang et al. suggested that dysregulation of miR-129-5p might contribute to the development of intervertebral disc degeneration by targeting bone morphogenetic protein 2 (BMP2) expression [25]. In patients with intervertebral disc degeneration, miR-129-5p was downregulated along with BMP2 upregulation. Overexpression of miR-129-5p apparently increased the survival of neural progenitor cells, whereas downregulation of miR-129-5p increased apoptosis. Moreover, restored BMP2 expression reversed the effect of miR-129-5p. On the other hand, miR-129-5p was also recently suggested to play a crucial role in the protection of cardiac function [52]. In a chronic heart failure rat model, miR-129-5p transfection not only improved heart function and hemodynamic parameters, but also attenuated oxidative stress and inflammation factors. In addition, a study by Zeng et al. demonstrated that miR-129-5p may be a potential candidate for the treatment of Alzheimer’s disease [23]. Overexpression of miR-129-5p promoted neuronal cell proliferation, suppressed apoptosis, and decreased the inflammatory reaction. These studies suggest that miR-129-5p is involved in different biological processes, which deserves further investigation.

It is difficult to develop efficient treatment against GBM, because GBM is characterized by many disregulated pathways. Currently, maximal surgical resection combined with postoperative radiotherapy and chemotherapy are the standard treatment of GBM. However, chemoresistance is the typical hallmark of recurrent GBM. More than 90% of GBM relapsed at the primary tumor site, and 5% of GBM occurred as multiple lesions after treatment [53]. Ma et al. demonstrated that miR-129-5p was able to inhibit the stemness and chemoresistance of non-small cell lung cancer (NSCLC) cells [54]. Namely, overexpression of miR-129-5p suppressed the ability of sphere formation and the expression of stemness makers, including octamer-binding transcription factor 4 (Oct4), SRY-box 2 (Sox2), and Nanog in NSCLC cells [54]. Furthermore, miR-129-5p/HMGB1 axis [55] and NONHSAT101069/miR-129-5p axis [56] were found to regulate the radiosensitivity and epirubicin resistance of breast cancer cells, respectively. In this study, colony-forming assay was conducted to indirectly investigate the stemness ability of GBM cells. Remarkably, our results showed that miR-129-5p significantly suppressed the colony-forming ability of LN229 and A172 GBM cells. This implies that miR-129-5p may affect GBM stemness and warrants further investigation in the future.

The RNA-binding protein ZFP36L1 is able to directly bind to the 3'-UTRs of their target mRNAs and promote mRNA decay. However, a few studies focused on the role of ZFP36L1 in tumorigenesis. Loh et al. suggested that ZFP36L1 may function as a tumor suppressor [57]. Forced expression of ZFP36L1 inhibited cell proliferation in bladder and breast cancer cells. On the other hand, a study by Zindy et al. demonstrated that ZFP36L1 was upregulated in hepatocellular carcinoma, implying that ZFP36L1 may be oncogenic in liver cancer [58]. In this study, we found that ZFP36L1 was highly expressed in GBM cell lines and tissues samples. Remarkably, GBM patients with higher expression of ZFP36L1 were associated with poor survival. In contrast to the study of Loh et al. on bladder and breast cancer cells [57], ZFP36L1 overexpression in GBM cells not only enhanced cell proliferation, but also increased cell migration and colony-forming ability. This suggests that ZFP36L1 may play different biological roles in different tumors and be a potential target for GBM treatment. This deserves further investigation.

The limitation of this study was the small number of GBM patients available in the databases. Although the current study found that miR-129-5p and ZFP36L1 are involved in GBM tumorigenesis, it remains unclear whether the expression of miR-129-5p and ZFP36L1 are associated with the overall survival, disease-free survival, and recurrence-free survival of GBM patients. Thus, further prospective studies should be conducted to investigate the association of miR-129-5p and ZFP36L1 with patient survival taking into consideration important prognostic confounders.

CONCLUSION

The present findings revealed a tumor suppressor role of miR-129-5p/ZFP36L1 axis in GBM cell proliferation, migration, and colony-forming ability. Importantly, ZFP36L1 expression in GBM was negatively regulated by miR-129-5p and associated with poor survival. Altogether, our study not only identified a new tumor suppressor role of miR-129-5p/ZFP36L1 axis in GBM tumorigenesis, but also provided potential targets for the treatment of GBM patients.

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