miR-342 inhibits glioma cell proliferation by targeting GPRC5A

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Abstract. Accumulating evidence suggests that microRNAs (miRNAs) play a key role in the biological behaviors and progression of glioma. However, the function and bio-molecular mechanisms of miR-342 in glioma remain largely unclear. In the present study, reverse transcription quantitative-polymerase chain reaction and western blotting were performed to determine the mRNA and protein expression levels of the factors investigated. MTT assay was performed to examine the proliferation rates. Luciferase reporter assay was performed to test the binding between miRNA-342 and its putative target. Data indicated that mir-342 expression was markedly decreased in human glioma tissues and cell line U87, and reduced mir-342 expression significantly promoted cell proliferation. In order to explore the mechanisms, G-protein coupled receptor family C group 5 member A (GPRC5A) was identified as a target of mir-342 and depletion of GPRC5A suppressed cell proliferation. Our findings demonstrated that miR‑342 regulates the cell proliferation of glioma by targeting GPRC5A, which indicates that mir-342 is a target of interest regarding the treatment of refractory glioma, and it may provide a promising prognostic and therapeutic strategy for glioma treatment.

Introduction

Gliomas are the most common malignant tumors of the central nervous system, accounting for approximately 40% of all brain tumors (1). Despite recent advances including adjuvant chemotherapy, radiotherapy and extensive tumor resection, the prognosis of glioma patients and the 5-year survival rate remain unfavorable (2). Thus, there is an urgent need to identify the mechanisms involved in glioma progression and metastasis to elucidate novel diagnostic and therapeutic targets for this malignancy.

MicroRNAs (miRNAs), single-stranded RNAs with a length of approximately 19-24 nucleotides, play significant roles in a series of biological cellular processes such as cell proliferation, migration, invasion and tumorigenesis (3). miR-342 serves a critical role in numerous physiological and pathological processes. It has been reported to be involved in the pathogenesis of many types of cancers, such as gastric cancer (4), hepatocellular carcinoma (5) and non-small cell lung cancer (6). Although these studies have demonstrated the important role of miR-342 in cancer progression, the modes of action of miR-342 in glioma have not been fully characterized.

G-protein-coupled receptors are the largest protein superfamily with more than 700 genes in the human genome (7). They play an important role in a variety of biological processes (8). It has been demonstrated that GPRC5A, one member of the GPCR family, is upregulated in many cancer tissues and cell lines (9-11). Yet, the relationship between GPRC5A and miR-342 remains unclear. We hypothesized that mir-342 directly targets GPRC5A, and the present study was designed to explore this hypothesis.

In the present study, the expression level of miR-342 was measured to assist the investigation of its regulatory roles in glioma. miR-342 was found to be markedly downregulated in glioma tissues and cell lines, and to exert tumor-suppressing functions in glioma. Moreover, miR-342 was found to regulate cell proliferation by targeting GPRC5A.

Materials and methods

Clinical samples and cell culture. Human glioma cell lines U-87MG (ATCC HTB-14 (RRID: CVCL_0022, glioblastoma of unknown origin), U251 (The Cell Bank of Type Culture Collection of Chinese Academy of Sciences; cat. no. TCHu 58), T98G [American Type Culture Collection (ATCC); cat. no. CRL-1690] and SNB19 (ATCC; cat. no. CRL-2219) and normal human astrocytes (NHAs; ScienCell Research Laboratories, Inc.; cat. no. 1820) were cultured in Dulbecco’s modified Eagle’s medium (DMEM;
Gibco™, 10564011) supplemented with 10% fetal bovine serum (FBS, Gibco™, 10099141) (both from Thermo Fisher Scientific, Inc., Waltham, MA, USA), 100 IU/ml penicillin, and 100 µg/ml streptomycin at 37°C under a 5% CO₂ atmosphere. Glioma specimens were collected from 39 patients following prior approval and written informed consent. Ten normal brain tissue samples used as controls were collected by donations from individuals who died in traffic accidents. All clinical samples were collected and histologically examined by pathologists from July 2016 to May 2018 at The Second Affiliated Hospital of Harbin Medical University. The present study was approved by The Ethics Committee of The Second Affiliated Hospital of Harbin Medical University. Written informed consent was obtained from all enrolled subjects.

Cell transfection. The miR-342 mimics, inhibitors and their corresponding miRNA negative control (miR-NC) were chemically synthesized by GenePharma (Shanghai, China) and transfected using Lipofectamine 2000 (Invitrogen; Thermo Fisher Scientific, Inc.). Overexpression or knockdown plasmids were transfected using Lipofectamine 2000 following the manufacturer’s instructions.

GPRC5A overexpression and shRNA-mediated knockdown plasmids. The longest transcript human genes of GPRC5A (NCBI Reference Sequence: NM_003979.3) were cloned into pcDNA 3.1 plasmids and then sequenced for validation. The siRNA duplexes targeting GPRC5A were obtained from Invitrogen; Thermo Fisher Scientific, Inc. In order to knock down GPRC5A expression in U87 cells, subconfluently cultured U87 cells were transfected with GPRC5A shRNA, or negative control shRNA using the RNAiMAX transfection reagent (Invitrogen; Thermo Fisher Scientific, Inc.), according to the manufacturer’s protocol. GPRC5A expression was assessed following 3 days. The shRNAs were designed by Invitrogen (Invitrogen; Thermo Fisher Scientific, Inc.) and cloned using Lipofectamine 2000 (Invitrogen; Thermo Fisher Scientific, Inc.), according to the manufacturer’s protocol. Stable population transfection was obtained following selection with 1 µg/ml G418 (Amresco, LLC, Solon, OH, USA) for 2 weeks. All shRNA oligos were obtained from Invitrogen; Thermo Fisher Scientific, Inc.

Quantitative real-time PCR (qPCR). Trizol reagent (Invitrogen; Thermo Fisher Scientific, Inc.) was used for total RNA extraction. Six microliters of the extracted RNA was reverse transcribed using the PrimeScript™ RT reagent kit with gDNA Eraser (Takara) in the StepOnePlus Real-Time System (Applied Biosystems™ ABI Prism 7500 Fast; Thermo Fisher Scientific, Inc.). The sequences of primers used were: GPRC5A forward 5'-CGC CACAAAGCACCAGGA-3' and reverse primer 5'-ATAGAG CGTGTCCCCGTGTCT-3'; GAPDH forward 5'-GAAAGC CGTCGGGTGTACTA-3' and reverse primer 5'-GCATCA CCCGGAGAGAAAT-3'; U6 small nuclear RNA forward 5'-CTCGCTTCGCGCCACA-3' and reverse primer 5'-AAC GCTTCACGAATTTCGCCG-3'; and 342 forward 5'-AGGTGA GGGTGCTATCTGT-3' and reverse primer 5'-GGGTGC GATTTCGTGTGAG-3'. All the samples were amplified in triplicate and each experiment was repeated three times. The conditions for the real-time fluorescent quantitative PCR were: 1 cycle at 95°C for 5 min in the holding stage; 40 cycles at 95°C for 15 sec and 60°C for 60 sec in the cycling stage; 1 cycle at 95°C for 15 sec; 60°C for 1 min and 95°C for 15 sec in the melt curve stage. Thermal cycling and real-time detection were conducted using the StepOnePlus Real-Time PCR Systems (ABI, Thermo Fisher Scientific, Inc.). The quantities of each mRNA were calculated using the comparative \(2^{-\Delta\Delta Cq}\) method (12).

Western blot analysis. Protein was isolated from the cells and tissues with RIPA lysis buffer containing 1% protease inhibitor cocktails (Pierce Biotechnology, Inc.; Thermo Fisher Scientific, Inc.). After sample buffer was added to the proteins (each well, 30 µg per sample), proteins were boiled at 95°C for 10 min. Then, the proteins were separated using 10% polyacrylamide gel electrophoresis. After electrophoresis, proteins were transferred to polyvinylidene fluoride (PVDF) membranes with 100 V transfer-molded voltage lasting for 45 to 70 min. After determination of the protein concentration, primary antibodies for western blotting were applied which included anti-GPRC5A (dilution, 1:1,000; PAB14597; Abnova, Taipei, Taiwan) and anti-GAPDH (dilution, 1:2,000; ab8245; Abcam, Cambridge, UK). HRP-conjugated IgG (1:5,000) antibody was used as the secondary antibody. After which membranes were washed 3 times (5 min/time). Development was completed with chemiluminescence reagents. GAPDH was used as an internal reference. Bands were visualized with a Bio-Rad Gel Doc EZ imager (Bio-Rad Laboratories, Inc., Hercules, CA, USA). The specific bands were visualized with a chemiluminescence system (Millipore), and then visualized with Quantity One software 4.6.2 (Bio-Rad Laboratories, Inc.).

Cell Counting Kit-8 (CCK-8) assay. Cells in the logarithmic growth phase were digested with trypsin and seeded on 96-well plates at 100 µl of medium containing 1x10⁴ cells per well. Then we measured the cell proliferation rate at 0, 24, 48, and 72 h after transfection. Ten microliters of CCK-8 reagent was added into each well following another 2-h incubation at 37°C. The absorbance value was determined by using the XT-96DJ ELISA analyzer at a wavelength of 490 nm.

Luciferase reporter assay. Wild-type and mutated GPRC5A 3’-UTR containing the putative binding site of miR-342 were synthesized and sequenced. Cells were seeded in 24-well plates and transfected with reporter vectors together with miR-342 mimics, miR-342 inhibitor or the corresponding miR-NC. The firefly luciferase activity was measured and normalized to Renilla signals at 48 h post-transfection.

Tumorigenicity assay. In total, 16 BALB/c male nude mice (specific pathogen-free grade; weight, 16-18 g; age, 4-6 weeks) were purchased from the Laboratory Animal Center of Harbin Medical University. Lentiviral-mediated stable GAPC5A, GAPC5A-miR-342, miR-342 mimic cells and miR-NC cells were resuspended in Hank’s buffer and mixed with an equal...
volume of Matrigel (BD Biosciences) at a concentration of 5x10^6 cells/ml. The cells were subcutaneously injected into the flanks of nude mice (n=4 in each group). Subsequently, the mice were maintained in a specific pathogen-free grade laboratory, under the following conditions: Controlled temperature, 23±2°C; humidity, 40-70%; 12-h light/dark cycle) at the Laboratory Animal Center in our hospital with ad libitum access to food and water for 4 weeks. The volume of xenograft tumors was monitored every 3 days by measuring the length and width (Volume=length x width x width/2). The animal study was conducted in accordance with the Institutional Animal Care and Use Committee (IACUC) guidelines, and was approved by the Experimental Animal Ethics Committee of The Second Affiliated Hospital of Harbin Medical University.

Statistical analysis. All values are expressed as mean ± SEM and were analyzed by one-way analysis of variance (ANOVA) followed by Tukey's post hoc test among groups using Statistical Product and Service Solutions (SPSS) (version 17.0) (SPSS, Inc., Chicago, IL, USA). Pearson's correlation analysis was performed to study the correlation between the expression of miR-342 and GPRC5A in cancer tissues. A P-value <0.05 was considered to indicate a statistically significant difference between groups.

Results

miR-342 is downregulated and GPRC5A is upregulated in glioma tissues and cell lines. Downregulation of miR-342 was observed in the glioma tissues (P<0.01; Fig. 1A) and cell lines (P<0.05; Fig. 1B) when compared with that noted in the normal human prostate tissues and the normal human astrocytes (NHAs). Meanwhile, the western blot results showed that the protein expression of GPRC5A was significantly upregulated in human glioma tissues (Fig. 1C) and cell lines (Fig. 1D). According to the results of RT-qPCR, GPRC5A expression in the U87 cell line was the highest, therefore, we chose this cell line for further experiments. The correlation analysis confirmed that the expression of miR-342 and GPRC5A was significantly negatively correlated (r=-0.4559, P=0.0035; Fig. 1E).

miR-342 inhibits the proliferation of glioma cells. Next, we explored the potential role of miR-342 in glioma. The transfection efficiency was determined according to the level of miR-342 was achieved after miR-342 mimic transfection and a significantly decreased expression of miR-342 was achieved after miR-342 inhibitor transfection. Moreover, upregulation of miR-342 resulted in greater suppression of cell proliferation than the control (mimics NC), whereas downregulation of miR-342 promoted cell proliferation (Fig. 2B) as determined using the CCK-8 assay. These results indicated that miR-342 inhibited the proliferation of U87 and U251 cells and downregulation of miR-342 promoted the proliferation of cells.

GPRC5A is a direct target of miR-342 in glioma. In order to determine the mechanism of miR-342 in cell proliferation, G-protein coupled receptor family C group 5 member A (GPRC5A) was found to be a putative target of miR-342 (Fig. 3A). At the protein and mRNA levels, overexpression or knockdown of miR-342 resulted in a significant decrease or increase in the expression level of GPRC5A, respectively (Fig. 3B and C). In addition, luciferase reporter assays were performed to ascertain whether miR-342 targets GPRC5A by binding to its 3'UTR. The results from the luciferase reporter assay indicated that upregulated expression of miR-342 significantly inhibited the activity of the reporter gene, whereas miR-342 inhibitor significantly increased it (Fig. 3D). The results indicate that GPRC5A is a direct target of miR-342.

GPRC5A promotes cell proliferation of U87. In order to investigate the cellular function of GPRC5A, the expression level of GPRC5A in U87 cells was manipulated by transfection with an overexpression (OE) or shRNA-mediated knockdown plasmids. The mRNA and protein levels of GPRC5A were determined in the transfected cells for which the expression levels of GPRC5A were markedly increased in the presence of overexpression plasmids or decreased in the absence of plasmids or silenced by shRNA. The inhibitory effect of each shRNA (sh1, sh2, sh3) was not significantly different while sh2 had the highest inhibition rate of GPRC5A, thus this shRNA was selected for further experiments (Fig. 4A and B). Moreover, CCK-8 assay results showed that overexpression of GPRC5A significantly promoted cell growth, while knockdown of GPRC5A suppressed it (Fig. 4C). The observation indicated GPRC5A expression associated with cell proliferation in vitro.

Restoration of GPRC5A reverses the effects of miR-342 in vitro and in vivo. Based on the above results, it was proposed that miR-342 suppresses cell proliferation via up-regulation of GPRC5A. Considering the low expression level of miR-342, rescue experiments were performed by co-transfecting the miR-342 mimics with or without GPRC5A followed by determination of the cell proliferation of U87 cells. Growth curves obtained by CCK-8 assay showed that upregulation of the expression of miR-342 alone significantly inhibited cell growth whereas overexpression of GPRC5A alone significantly promoted cell growth; co-expression of GPRC5A with miR-342 abrogated the inhibitory effects of miR-342 mimics on the cell proliferation (Fig. 5A). Based on the in vitro data, the effect of miR-342 and GPRC5A on tumor growth was further evaluated in vivo. Lentivirus of miR-342 and GPRC5A were either used to infect cells alone or simultaneously. We found that tumor xenografts derived from cells infected with miR-342 alone were significantly smaller than the negative control while those infected with GPRC5A showed an opposite trend; co-infection of GPRC5A with miR-342 abrogated the inhibitory effects of miR-342 (Fig. 5B). The expression level of GPRC5A in tumor xenografts was also assessed. The expression level of GPRC5A was much lower in the miR-342 mimic group and higher in the GPRC5A overexpression or co-infected groups, compared with their counterparts in the negative control (NC) group (Fig. 5C). The in vitro and in vivo observations suggest that miR-342 targets the 3'UTR of GPRC5A directly and inhibits U87 cell proliferation via GPRC5A up-regulation.
Figure 1. miR-342 is downregulated in human glioma cancer tissues and glioma cell lines. (A) Expression of miR-342 in human glioma cancer tissues and adjacent non-tumor tissues were detected by quantitative real-time PCR. (B) Expression of miR-342 in several glioma cell lines and normal human astrocytes (NHAs) was detected by quantitative real-time PCR. (C) Protein levels of GPRC5A in human glioma cancer tissues and adjacent non-tumor tissues were detected by western blot analysis. T, tumor tissue; N, normal tissue. (D) Protein levels of GPRC5A in several glioma cell lines and normal human astrocytes (NHAs) were detected by western blot analysis. (E) Correlation analysis between expression of miR-342 and GPRC5A in cancer tissues. *P<0.05, **P<0.01, compared to normal non-tumor tissues or NHAs. GPRC5A, G-protein coupled receptor family C group 5 member A.
Gliomas are the most common primary brain tumors, which show an extremely high proliferation and invasive capacity (13,14). Therefore, effective diagnostic and therapeutic strategies for glioma are urgently needed. Recently, studies have confirmed that miRNAs play important roles in tumorigenesis and development, and are involved in the regulation of tumor growth, apoptosis, differentiation, invasion, angiogenesis, and metastasis (15,16). In addition, miRNAs can also act as oncogenes or tumor-suppressor genes, and their role in the development of gliomas acts on the mechanisms of glioma (17). Downregulation of miR-342 has been found in a number of cancer types, such as breast cancer (18), prostate cancer (19) and nasopharyngeal carcinoma (20). Thus, it has been proposed that miR-342 could be used as a diagnostic and prognostic biomarker. In the present study, a significantly downregulated miR-342 level was observed in glioma tissues and cell lines and the increase in the expression level of miR-342 was found to suppress the proliferation of glioma cells in vitro, suggesting that miR-342 functions as an anti-oncogene.

G-protein-coupled receptors are the largest protein superfamily with more than 700 genes in the human genome (7) playing an important role in a variety of biological processes (8). This protein superfamily also acts as drug targets in many different diseases and more than 40% of FDA (Food and Drug Administration)-approved drugs target
GPCRs (G protein-coupled receptors) or GPCR-associated pathways (21). GPCRs also play an integral role in regulating and activating cancer-associated signaling pathways; therefore, they may be used as biomarkers for the early diagnosis of various types of cancer (22). Further investigation of the pharmacological potential of GPCRs and their downstream regulators is required in order to develop therapies that can efficiently target cell signaling pathways in cancer (23,24).

GPC5A, also termed RA13 (retinoic acid-induced protein 3), located on chromosome 12p13-p12.3, has been found
to play significant roles in various biological processes, such as cell proliferation and the cell cycle. However, the influences of GPRC5A on different cancers vary. GPRC5A was reported to be strongly expressed in the lung (25) and is regarded as a tumor suppressor in lung cancer as well as in head and neck squamous cell carcinoma (26). However, there are many studies that have reported that high expression of GPRC5A is correlated with a worse survival rate in colon, breast and gastric cancer (27). Liu and colleagues found high expression of GPRC5A in pancreatic cancer and it was found to suppress the activity of phosphorylated GSK-3β at Ser9 (26). Moreover, Zhou and Rigoutsos reported that miR-103a-3p also targets the 5′UTR of GPRC5A and reduced GPRC5A protein expression in pancreatic cells. It also may indirectly regulate many

Figure 4. GPRC5A promotes U87 cell proliferation in vitro. (A) mRNA levels of GPRC5A in the cells transfected with the GPRC5A overexpression (OE) plasmids or shRNA-mediated knockdown plasmids were determined by quantitative real-time PCR. (B) Proteins levels were also confirmed by western blot analysis. Representative images of the western blots are shown; bands were quantitated by densitometry and normalized against GAPDH. (C) Growth curves of each group were obtained by CCK-8 assay in U87 cells transfected with OE GPRC5A or shRNA of GPRC5A plasmids. **P<0.01 vs. CV or SC; ***P<0.01 vs. Scramble. GPRC5A, G-protein coupled receptor family C group 5 member A.
cell processes, such as DNA repair, metabolism and the cell cycle (28). In the present study, we confirmed that GPRC5A promoted the proliferation of U87 cells and confirmed that GPRC5A is a direct target of miR-342. Furthermore, the tumor-suppressive effect of miR-342 was reduced by enforced expression of GPRC5A in vivo and in vitro. These results suggest that miR-342 functions as an anti-oncogene via multiple gene targeting, such as on GPRC5A.

In conclusion, the study availed a better understanding of the function of miR-342 and GPRC5A in glioma. We confirmed the downregulated level of miR-342 in glioma and revealed the role of miR-342 in glioma cell proliferation and invasion. Our data indicated the suppressive role of miR-342 in glioma development and miR-342 may serve as a potential therapeutic target in glioma. However, our results were based on one single cell line, U87, and thus additional cell lines are needed to be included in further research. Meanwhile, although the role of miR-342 in cellular invasiveness and cancer progression is clear, its mechanisms remain to be investigated.

Figure 5. Upregulation of GPRC5A reverses the inhibitory effects of miR-342 mimics on U87 cells. (A) Growth curves were obtained by CCK-8 assay in the cells transfected with miR-342 mimic, GPRC5A overexpression plasmid alone or co-transfected with miR-342 mimic and GPRC5A overexpression plasmid or the negative control (NC). (B) Representative images of tumor xenografts and tumor growth curves are shown. (C) Protein levels of GPRC5A9 in each group were determined by western blot analysis. *P<0.05, **P<0.01, ***P<0.001 vs. mir-nc; ##P<0.01 vs. GPRC5A+mir-342.
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Availability of data and materials

The datasets used during the present study are available from the corresponding author upon reasonable request.

Authors' contributions

JW, YY, YC and XT interpreted and analyzed the data, and drafted the manuscript. JW analyzed the data. XT designed the study. All authors interpreted the results and wrote the manuscript.

Ethics approval and consent to participate

The animal study was conducted in accordance with the Institutional Animal Care and Use Committee (IACUC) guidelines, and was approved by the Experimental Animal Ethics Committee of The Second Affiliated Hospital of Harbin Medical University. The present study was approved by the Ethics Committee of The Second Affiliated Hospital of Harbin Medical University. Written informed consent was obtained from all enrolled subjects.

Patient consent for publication

Not applicable.

Competing interests

The authors state that they have no competing interests.

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