

TRIM47 Promotes the Development of Glioma by Ubiquitination and Degradation of FOXO1

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Objective: To investigate the effect of *TRIM47* on glioma cells and further explore its underlying molecular mechanisms.

Methods: Mouse xenograft model was used in this study. The mRNA expression of *TRIM47* was detected by qRT-PCR. The cell viability and proliferation activity was detected by MTT assay and colony formation assay. The migration and invasion of glioma cells were determined by Transwell assay. The protein levels of *TRIM47*, FOXO1, CyclinD1, C-myc, MMP-2 and TIMP-1 were assessed by Western-blotting. The interaction between TRIM47 and FOXO1 was measured by Co-immunoprecipitation (Co-IP) assay.

Results: In glioma tissues and cells, *TRIM47* was significantly up-regulated. Silencing the expression of *TRIM47* inhibited the cell viability and proliferation of cells A172 and U251, as well as their ability to invade and migrate. Among them, the expression levels of C-myc and CyclinD1 also decreased, and MMP-2 was down-regulated and TIMP-1 was up-regulated. Similarly, in vivo model, tumor volume and weight also decreased after *TRIM47* knockout. Further research showed that TRIM47 inhibited FOXO1 expression by ubiquitination and degradation of FOXO1, thereby promoting glioma growth and progression.

Conclusion: In our study, we confirmed functional role of the *TRIM47-FOXO1* axis in the progression of gliomas and provided a potential target for glioma treatment.

Keywords: glioma, tripartite motif 47, forkhead box O1, proliferation, migration, invasion

Introduction

Malignant glioma is a primary tumor of the central nervous system, accounting for about 80% of the intracranial malignancies, with the highest morbidity and mortality.^{1,2} Malignant glioma exhibited increased levels of invasive growth, and is easy to invasion and metastasis.³ At present, the main clinical treatment methods for gliomas are surgical resection plus radiotherapy and chemotherapy. However, because glioma stem cells can resist radiotherapy and chemotherapy, their recurrence rate is still high, and very few patients survive >1 year. In addition, glioma cannot be completely cured, and its prognosis is poor.⁴ Therefore, it is urgent to study the potential molecular mechanisms and explore new potential diagnostic or therapeutic targets for the treatment of glioma.

A growing body of clinical evidence indicated that ubiquitin-mediated degradation of oncogene products or tumor suppressors may be related to the cause of cancer.⁵ In the ubiquitin-proteasome system (UPS) component, the E3 ubiquitin ligase, which recognized the most specific substrates, was considered a potential diagnostic and therapeutic target for cancer.⁶ Tripartite motif (TRIM) family proteins are evolutionarily conserved proteins consisting of circular finger domains

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with one or two b-box domains and a related coiled spiral domain in the amino-terminal region.⁷ It is involved in a wide range of cell processes, including cell proliferation,⁸ differentiation,⁹ apoptosis,¹⁰ cell cycle regulation, carcinogenesis,⁵ etc. Most TRIM proteins confer ubiquitin E3 ligase activity and promote posttranslational modification. Among them, *TRIM47* was first identified as overexpression in astrocytoma. In addition, other studies have shown that the high expression of *TRIM47* was closely related to the occurrence of human prostate cancer¹¹ and non-small cell lung cancer.¹² However, so far, the pathological and clinical role of *TRIM47* in glioma has not been revealed.

Forkhead box O1 (*FOXO1*) belongs to the FOXO transcription factor family and is located on chromosome 13q14, characterized by a conserved winged helix DNA binding domain.¹³ In recent years, the broad role of FOXO in physiological processes has been extensively studied, including cell cycle arrest, angiogenesis, apoptosis, stem cell differentiation and stress resistance.¹⁴ Previous studies have shown that *FOXO1* has an anticancer effect and has been identified as an anticancer gene. In addition, there is increasing evidence that *FOXO1* is downregulated in many cancer types, such as Hodgkin's lymphoma,¹⁵ breast cancer,¹⁶ and alveolar rhabdomyosarcoma.¹⁷ Notably, although many studies have linked *FOXO1* expression to glioma, the exact pattern and role of *FOXO1* in glioma remains elusive.

More importantly, no one has studied the complex functions of *FOXO1* and *TRIM47* in the development and progression of gliomas. Hence, our current study aims to investigate the correlation between *TRIM47* and *FOXO1* in glioma and to explore its potential mechanisms.

Materials and Methods

Clinical Sample Collection

Seventy-nine glioma patients were selected to obtain paired glioma tissue samples and corresponding adjacent tissue samples. The glioma tissue samples were classified as either low-grade glioma (WHO I–II) or high-grade glioma (WHO III–IV) according to the clinicopathologist's diagnosis. Each patient signed a written informed consent and the ethical committee of Lianshui County People's Hospital (LSXRM2019-04).

Cell Culture

Human normal glial cell line (HEB) and five glioma cell lines (U251, Hs683, U87, SHG-44 and A172) were

obtained from Mibio (Shanghai, China). All cells were cultured in RPMI 1640 medium supplemented with 10% FBS in a humidified environment. Subsequent transfection experiments could be performed when cells reached 70% ~ 80% confluences.

Cell Transfection

A172 and U251 with a confluence of 70%~80% were collected and then inoculated into 6-well plates, respectively. They were divided into 5 groups: Mock group (without any treatment), si-NC group (transfected with siRNA NC), si-*TRIM47*#1 (transfected with siRNA1), si-*TRIM47*#2 (transfected with siRNA2) and si-*TRIM47*#3 (transfected with siRNA3), and then follow the instructions of Lipofectamine 2000 liposome transfection kit (Invitrogen, USA). After 48 h transfection, the cells were collected and then for further experiments.

Real-Time PCR Analysis

Total RNA was isolated from glioma tissues and cell lines using TRIzol reagent (Invitrogen, USA). The extracted RNA was reversely transcribed to cDNA by using a Prime Script™ RT Reagent Kit (TaKaRa, Japan). Subsequently, RT-qPCR was performed. The primer sequences were as follows: *TRIM47* sense, 5'-GCTTCAGGAGGCTGAGCAGT-3' and antisense, 5'-TCTGCTACGGCTGCACTCTT-3'; *FOXO1* sense, 5'-TACGAGTGGATGGTCAAGAG-3' and antisense, 5'-ATGAACTTGCTGTGTAGGGAC-3'; *GAPDH* sense, 5'-GCATTGCCCTCAACGACCAC-3' and antisense, 5'-CCACCACCCTGTTGCTGTAG-3'.

Western Blotting

According to the manufacture's instruction, the proteins were extracted and its concentration was measured. Protein samples were separated by polyacrylamide-SDS gels and electro blotted onto nitrocellulose membranes. After blocking with 5% nonfat dry milk for 2.5 h, the PVDF membrane incubated with primary antibodies: cyclinD1 (1:1000, Sigma Aldrich, SAB4502603), C-myc (1:1000, Thermo Fisher Scientific, 13–2500), MMP-2 (1:1000, Sigma Aldrich, SAB4501891), TIMP-1 (1:1000, Sigma Aldrich, AB770), *TRIM47* (1:1000, Sigma Aldrich, SAB2108331), *FOXO1* (1:1000, Thermo Fisher Scientific, MA5-14,846). On the following day, the protein samples were incubated with the secondary antibody at room temperature for 45 min, after which blots were visualized by enhanced chemiluminescence method. The intensity of

bands was quantified by ImageJ software (Biorad, Richmond, CA).

Co-Immunoprecipitation (Co-IP) Assay

For co-immunoprecipitation assay, IgG or IP antibody, suspended IP Matrix and PBS were incubated 1 hour at 4 °C, following by centrifugation and washing by PBS containing protease inhibitor, phosphatase inhibitor and PMSF for 3 times, and then discard the supernatant. Subsequently, the transfected cells were lysed and transferred to the matrix, and then cells were incubated at 4 °C overnight. The matrix was then centrifuged and washed for 5 times. The SDS-PAGE sample loading buffer was added to the immunoprecipitates, and then was boiled for 10 min at 100 °C. The IP and Input proteins were detected by Western blot.

Cell Viability Assay

Cell viability in each group was assessment by MTT and the transfection time was 0 h. The cells were seeded in a 96-well plate and cultured in a 37°C incubator containing 5% CO₂. First, 20 µL of MTT solution (5mg/mL, Sigma) was added to each well, and then continued to incubate in a 5% CO₂ incubator at 37°C. After 4h, the culture was terminated and 150 µL of DMSO was added. Gently evenly shake the mixture for 10 min to promote crystallization dissolution. The absorbance values of each hole at 570 nm wavelength were measured on a microplate and data were expressed as absorbance.

Colony Formation Assay

The same quantities of A172 and U251 cells of each group were collected and cultured in 6-well plates. The medium was changed every 2 to 3 days and cultured continuously for 14 days. After washing the cells twice with PBS, the cells were fixed with 4% formaldehyde for 15 min at 37°C. Subsequently, residual formaldehyde was removed, and cell clones were stained with crystal violet for 10–20 min. Cells in each hole were observed under the microscope and the number of clones was counted. Each group ≥ 50 cells were recorded as effective clones, and cameras took photos directly.

Transwell Assay

A172 and U251 cells were trypsinized and re-suspended in serum-free medium to adjust cell density. Subsequently, a total of 1×10^5 cells were added to the Transwell chamber coated with Matrigel (BD Biosciences, New Jersey, USA), while a medium containing 15% fetal bovine serum was added to the lower chamber. After 24 h in the incubator,

the cells were rinsed with PBS, fixed with 4% paraformaldehyde 37 °C for 16 min and stained with 0.1% crystal violet at 37 °C for 25 min. The cell invasion assay was performed in a similar manner, but 20 µL Matrigel (BD Biosciences) was added to the upper chamber. Five fields were randomly selected under an inverted microscope to represent the invasion ability and migration ability of cells in each group.

Mouse Xenograft Model

To establish the xenograft model, 1×10^6 U251 cells were subcutaneously injected into BALB/c mice (4-week-old). When tumors were measurable, the mice were randomly divided into 3 groups, and PBS, si-NC and si-TRIM47 viruses were injected into the tumor every 2 days for 14d. After 30 days, tumor volume and tumor weight were recorded.

Immunohistochemistry (IHC) Staining

The xenograft tumors were formalin-fixed and paraffin-embedded, sliced into 4µm-thick sections. The xenograft tumors were formalin-fixed and paraffin-embedded, sliced into 4 µm-thick sections. Graded ethanol was used to rehydrate the sections after deparaffinization in xylene at room temperature. After washing with PBS, the sections were placed in 3% hydrogen peroxide for 20 min to inhibit endogenous peroxidase, followed by antigen retrieval by heating for 30 min in a microwave. After being blocked with 10% goat sera, the tissue sections were incubated with primary anti-Ki67 antibody (1:500, Cell Signaling Technology, MA, USA) and at 4°C overnight. After being washed with PBS and incubated with secondary antibody for 30 min at 37°C, 3,3'-Diaminobenzidine tetrahydrochloride (DAB) was applied as a chromogen, and the sections were counterstained with hematoxylin.

Statistical Analysis

The data were presented as the mean \pm standard deviation (SD) and analyzed by the Student's *t*-test and one-way analysis of variance (ANOVA) using the SPSS version 19.0 software. Survival rate was analyzed by Kaplan-Meier analysis. $P < 0.05$ was considered to be statistically significant.

Results

TRIM47 Was Highly Expressed in Glioma Tissues and Glioma Cell Lines

As presented in Figure 1A, the mRNA expression of *TRIM47* in glioma tissues were obviously higher than that in the adjacent normal tissue ($P < 0.05$). Notably,

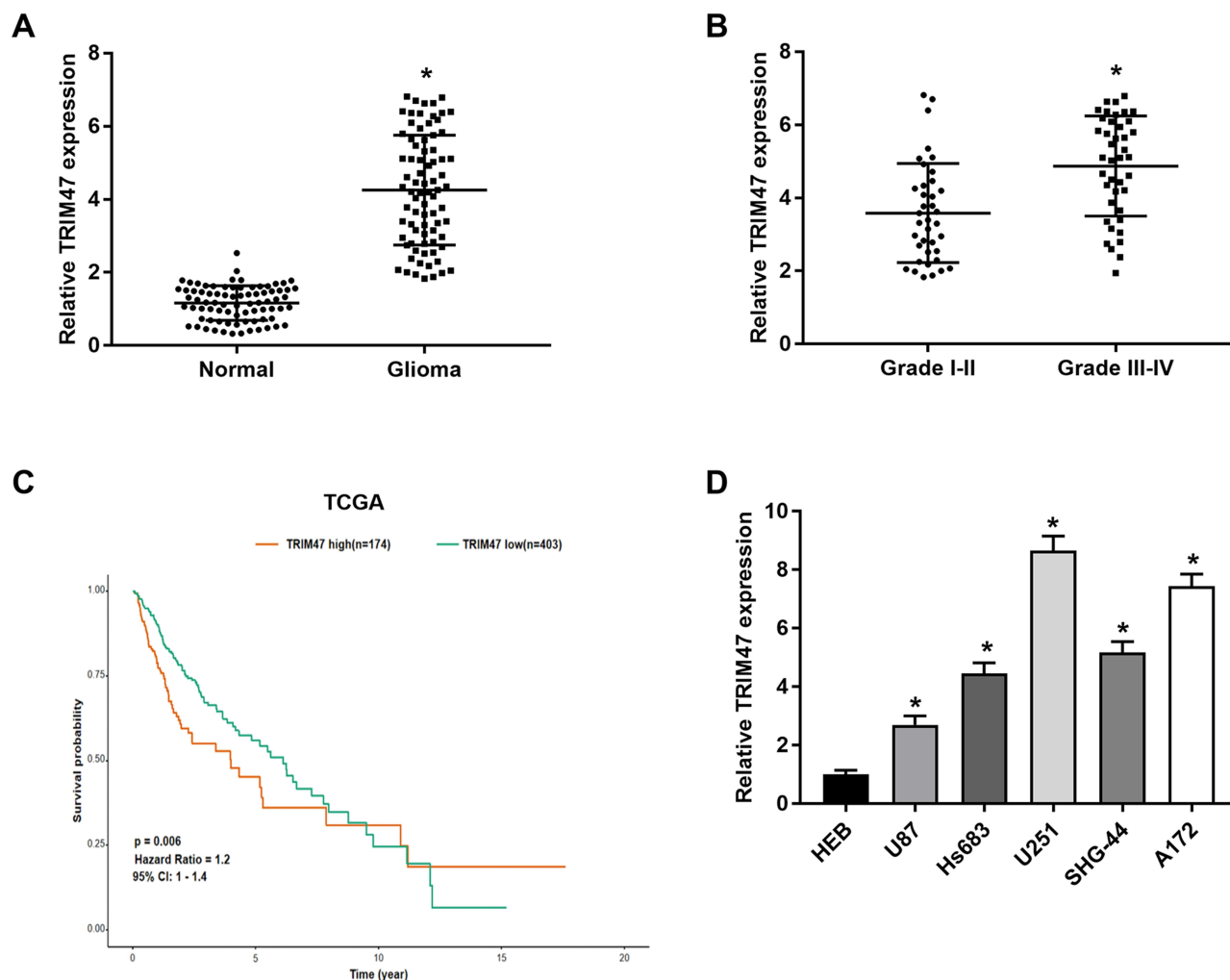


Figure 1 TRIM47 was highly expressed in glioma tissues and glioma cell lines. **(A)** The expression of TRIM47 in glioma tissues and corresponding paracancer tissues were detected by qRT-PCR; **(B)** The expression of TRIM47 in glioma tissues at different stages were detected by qRT-PCR; **(C)** Kaplan-Meier was used to analyze overall survival in patients with low expression of TRIM47 and patients with high expression of TRIM47; **(D)** The expression of TRIM47 in normal glial cells (HEB) and glioma cells (U251, Hs683, U87, SHG-44 and A172) were detected by qRT-PCR. * $P < 0.05$.

compared with grades I–II, the level of *TRIM47* was significantly increased in grades III–IV, indicating that the level of *TRIM47* expression was closely related to the clinical grade of glioma (Figure 1B; Table 1). Subsequently, we analyzed the overall survival based on the TCGA database, and found that the glioma patients with high expression of *TRIM47* had a worse overall survival (Figure 1C). At the same time, we also tested the expression of *TRIM47* in different glioma cell lines (U251, Hs683, U87, SHG-44 and A172). Compared to human normal glial cell line (HEB), *TRIM47* was significantly up-regulated in the above cell lines, with U251 and A172 being the most pronounced (Figure 1D). Therefore, U251 and A172 were selected for follow-up experiments.

TRIM47 Promoted the Proliferation of Glioma Cell in vitro and in vivo

To further investigate the role of *TRIM47* on glioma, U251 and A172 cells were then transfected with si-*TRIM47* or si-NC. From Figure 2A, we can see that compared with group si-NC, the silencing efficiency of *TRIM47* gene in si-*TRIM47* groups were very significant, especially in group si-*TRIM47*#3. Next, we selected group si-*TRIM47*#3 for MTT assay to evaluate the cell viability of A172 and U251 cells. The MTT results indicated that the above cell viability had been significantly reduced in a time-dependent manner (Figure 2B). Similarly, the colony formation assay also indicated that the low expression of *TRIM47* resulted in decreased cell proliferation and

Table I The Correlation Between TRIM47 mRNA Expression and Clinicopathological Factors

variables	Number of Cases	TRIM47 Level		P value
		Low (n)	High (n)	
Overall	79	40	39	
Age (years)				
≤50	37	21	16	0.3069
>50	42	19	23	
Gender				
Male	42	21	21	0.9046
Female	37	19	18	
Pathological grade				
Grade I-II	44	27	11	0.0005*
Grade III-IV	35	13	28	

Note: *P<0.05.

colony number, suggesting that si-*TRIM47* inhibited the proliferation of A172 and U251 cells (Figure 2C).

C-myc and Cyclin D1 were all proliferation-related proteins, to further detect the effect of TRIM47 in the progression of glioma, Western blotting assay was conducted. As expected, the protein expression of C-myc and Cyclin D1 were decreased in *TRIM47* knockdown cells when in comparison to Mock or si-NC groups (Figure 2D). Consistently, down-regulation of *TRIM47* sharply reduced the tumor volume (Figure 2E) and tumor weight (Figure 2F). The result of IHC staining showed that there were less positive staining for Ki67 in si-*TRIM47* group compared with that in the Mock or si-NC groups (Figure 2G).

TRIM47 Promoted the Migration and Invasion of Glioma Cell

To investigate the role of *TRIM47* in migration and invasion of glioma cell, Transwell assay was conducted after transfecting si-NC or si-*TRIM47*. As shown in Figure 3A and B, si-*TRIM47* markedly reduced the number of cell migration and invasion when compared with the Mock or si-NC groups. These results indicate that si-*TRIM47* may inhibit migration and invasion of A172 and U251 cells.

MMP-2 and TIMP-1 were all migration-related proteins, and their functions are opposite. Among them, MMP-2 can promote the migration and invasion of tumor cells, while TIMP-1, on the contrary, can inhibit

the infiltration and migration of tumor cells. To further clarify the changes of the above proteins in the *TRIM47* down-regulated group, we conducted Western blotting. As indicated in Figure 3C, MMP-2 was sharply reduced in si-*TRIM47* group, while TIMP-1 protein expression increased significantly in comparison to Mock or si-NC groups.

TRIM47 Interacted with FOXO1 and Enhanced Ubiquitylation and Degradation of FOXO1

To further understand the potential molecular mechanism of *TRIM47* in glioma progression, we first detected the expression of TRIM47 and FOXO1 in glioma cells A172 and U251. As presented in Figure 4A, compared with Mock or si-NC groups, the protein expression of TRIM47 was decreased in A172 and U251 cells, whereas the protein expression of FOXO1 was increased with *TRIM47* knockdown ($P<0.05$). In addition, up-regulating the expression of TRIM47 remarkably inhibited the protein expression of FOXO1 (Figure 4B). Next, the molecular mechanism by which *TRIM47* regulated the expression level of FOXO1 protein was further explored.

Previously, *TRIM47* has been reported to have an E3 ubiquitin ligase effect, so we speculated whether *TRIM47* might down-regulate *FOXO1* expression by promoting ubiquitination and degradation of FOXO1 protein. To prove this, MG132 (proteasome inhibitor, 10 μ L/mL) or DMSO was added to treat glioma cells for 2h. The western-blotting results showed that, compared with DMSO group, *TRIM47*-induced down-regulation of FOXO1 was counteracted by MG132 (Figure 4C, $P<0.05$). Furthermore, the Co-IP assay showed that *TRIM47* and *FOXO1* interacted with each other in A172 and U251 cells (Figure 4D and E).

TRIM47 Promoted Glioma Development Through FOXO1 Ubiquitination

To further investigate the relationship between *TRIM47* and *FOXO1* in the progression of glioma, MTT assay, Colony formation assay, Transwell assay and Mouse xenograft model were next conducted. As exhibited in Figure 5A, overexpression of TRIM47 significantly increased the cell viability of glioma cells, while overexpression of FOXO1 sharply inhibited the cell viability of glioma cells. It was noted that when pc-*TRIM47* and pc-*FOXO1* were co-transfected with glioma cells, their cell viability

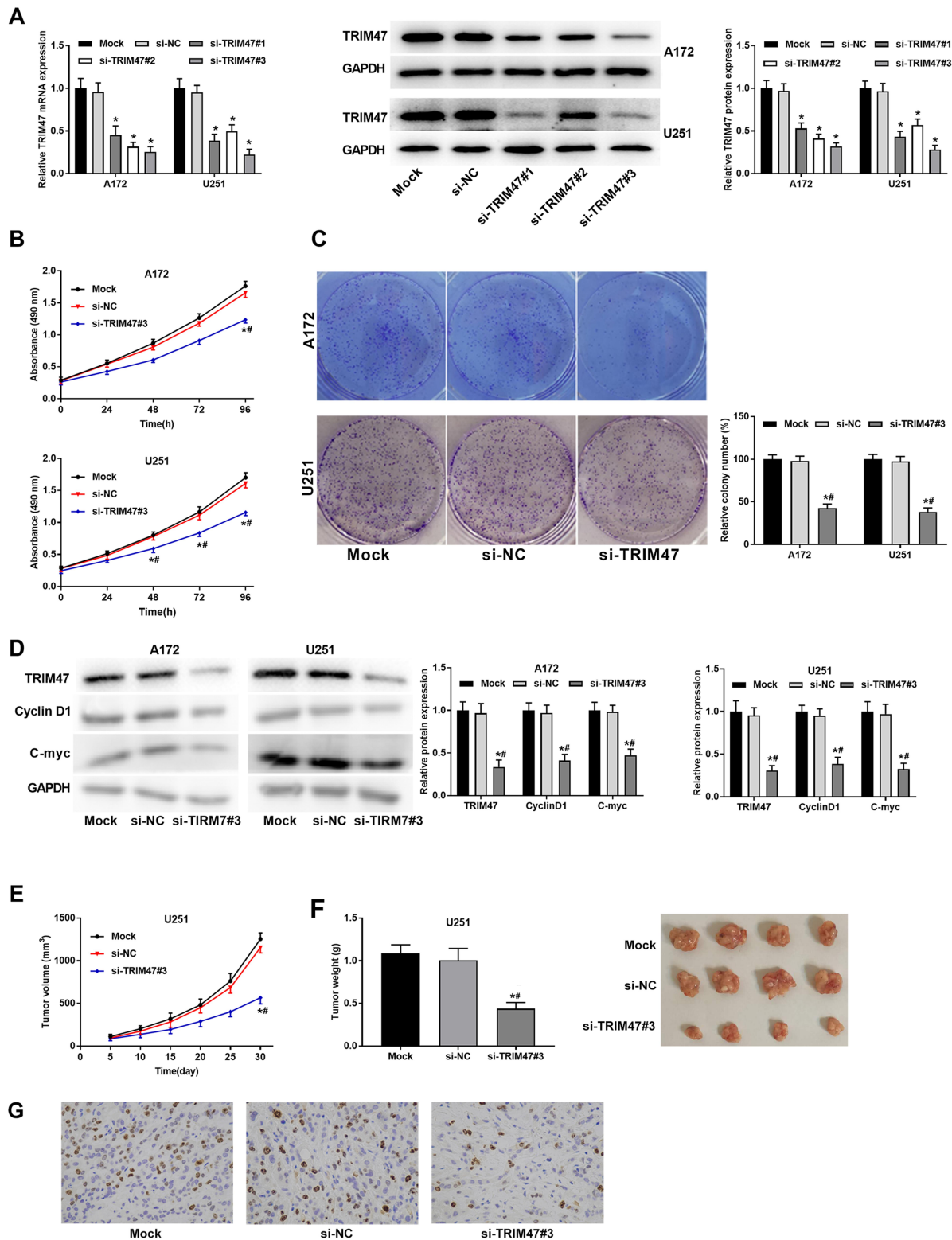


Figure 2 TRIM47 promoted the proliferation of glioma cell in vitro and in vivo. **(A)** The silencing efficiency of TRIM47 was detected by qRT-PCR and Western blotting; **(B)** MTT was used to detect glioma cell proliferation after silencing TRIM47; **(C)** Colony formation assay was used to detect glioma cell proliferation after silencing TRIM47; **(D)** The expression changes of TRIM47 and proliferation-related proteins C-myc and CyclinD1 were detected by Western blotting; **(E)** After silencing TRIM47 expression, the xenograft tumor volume of glioma mice became smaller; **(F)** After silencing TRIM47 expression, tumor weight of xenograft was reduced in glioma mice. **(G)** After silencing TRIM47 expression, the positive staining for Ki67 of glioma mice tumor was remarkably decreased. *P<0.05 vs Mock group, #P<0.05 vs si-NC group.

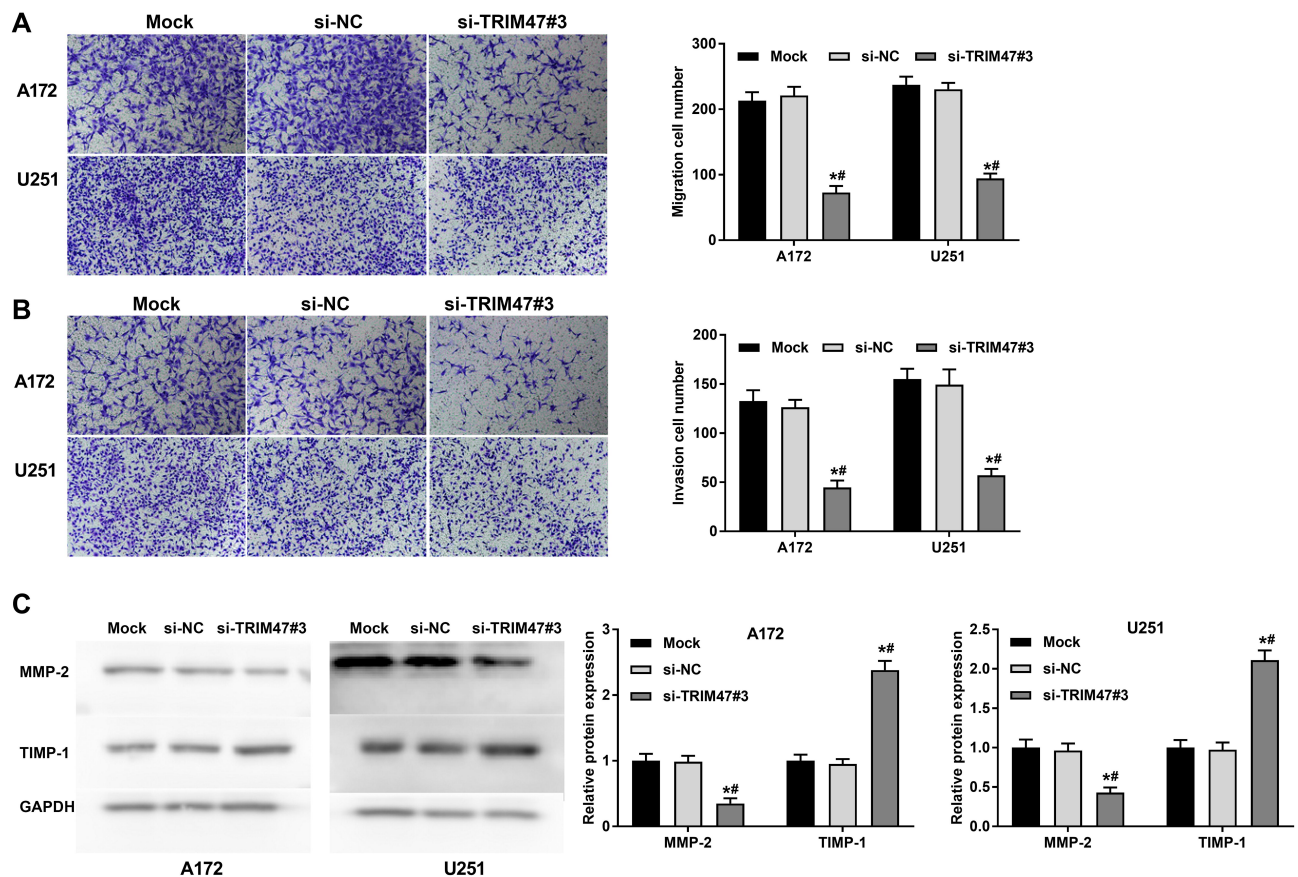


Figure 3 TRIM47 promoted the migration and invasion of glioma cell. **(A)** Transwell assay was used to detect glioma cell migration after TRIM47 silencing; **(B)** Transwell was used to detect the invasion ability of glioma cells after TRIM47 silencing; **(C)** The expression changes of Migration-related proteins MMP-2 and TIMP-1 were detected by Western blotting. * $P < 0.05$ vs Mock group, ** $P < 0.05$ vs si-NC group.

was also significantly reduced, which was still higher than that of pc-FOXO1 group. Consistently, pc-TRIM47 group had the largest number of cell clones, followed by pc-TRIM47+pc-FOXO1 group, and pc-FOXO1 group had the least number of clones (Figure 5B). In Figure 5C, we can clearly see that co-transfection pc-TRIM47 and pc-FOXO1 could to some extent offset the migration of glioma cells induced by overexpression *TRIM47*. In addition, evidence that the *TRIM47* and *FOXO1* interactions affected tumor progression can be confirmed in mouse xenograft model. Overexpression of *TRIM47* increased tumor volume and tumor weight, while pc-FOXO1 has the opposite effect, it would inhibit tumor development. When the two interact with glioma cells, the effects of the two are partially offset (Figure 5D and E). The result of HIC staining showed that *TRIM47* overexpression increased the positive staining for Ki67 of tumor tissues, while overexpression of *FOXO1* revealed the opposite effect (Figure 5F, $P < 0.05$). *FOXO1* overexpression could

eliminated the effect of *TRIM47* overexpression on the xenograft tumor (Figure 5F, $P < 0.05$).

Discussion

Worldwide, glioma is a life-threatening primary brain cancer. Its uncontrollable growth and expansion are the main reasons for its destruction of surrounding normal brain tissue and damage to nerve function.¹⁸ Despite advances in chemotherapy, radiation, and surgery for gliomas over the past few decades, there is currently no effective treatment to improve long-term survival of glioma patients. Therefore, there is an urgent need to elucidate the biological and molecular mechanisms of the development of gliomas, and to develop more effective treatments on this basis.

TRIM47 is an E3 ubiquitin ligase, located at 17q24-25, which is frequently acquired or amplified in many other types of tumors.⁶ Recent pathological analysis showed that changes in the expression of certain TRIM proteins are

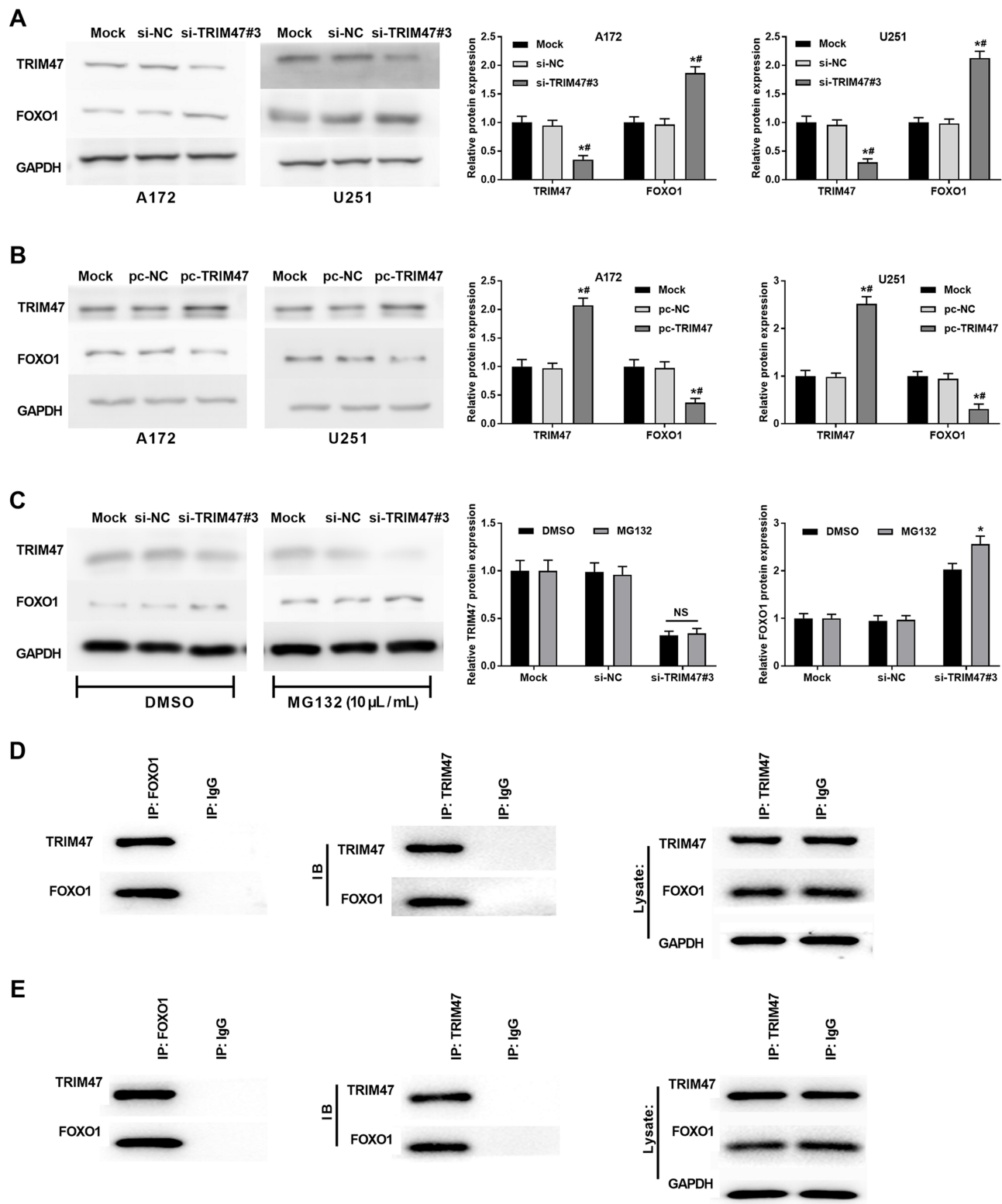


Figure 4 TRIM47 interacted with FOXO1 and enhanced ubiquitylation and degradation of FOXO1. **(A)** The protein levels of TRIM47 and FOXO1 were detected in A172 and U251 cells transfected with si-TRIM47 by Western blotting; **(B)** The protein levels of TRIM47 and FOXO1 were detected in A172 and U251 cells transfected with pc-TRIM47 by Western blotting; * $P < 0.05$ vs Mock group, [#] $P < 0.05$ vs pc-NC group. **(C)** Western blotting was performed in A172 and U251 cells transfected with si-TRIM47 and treated with DMSO or MG132 (10 μ L/mL). * $P < 0.05$ vs DMSO group. **(D)** Coimmunoprecipitation (Co-IP) detected the interaction of TRIM47 and FOXO1 in A172 cells. **(E)** Co-IP detected the interaction of TRIM47 and FOXO1 in U251 cells.

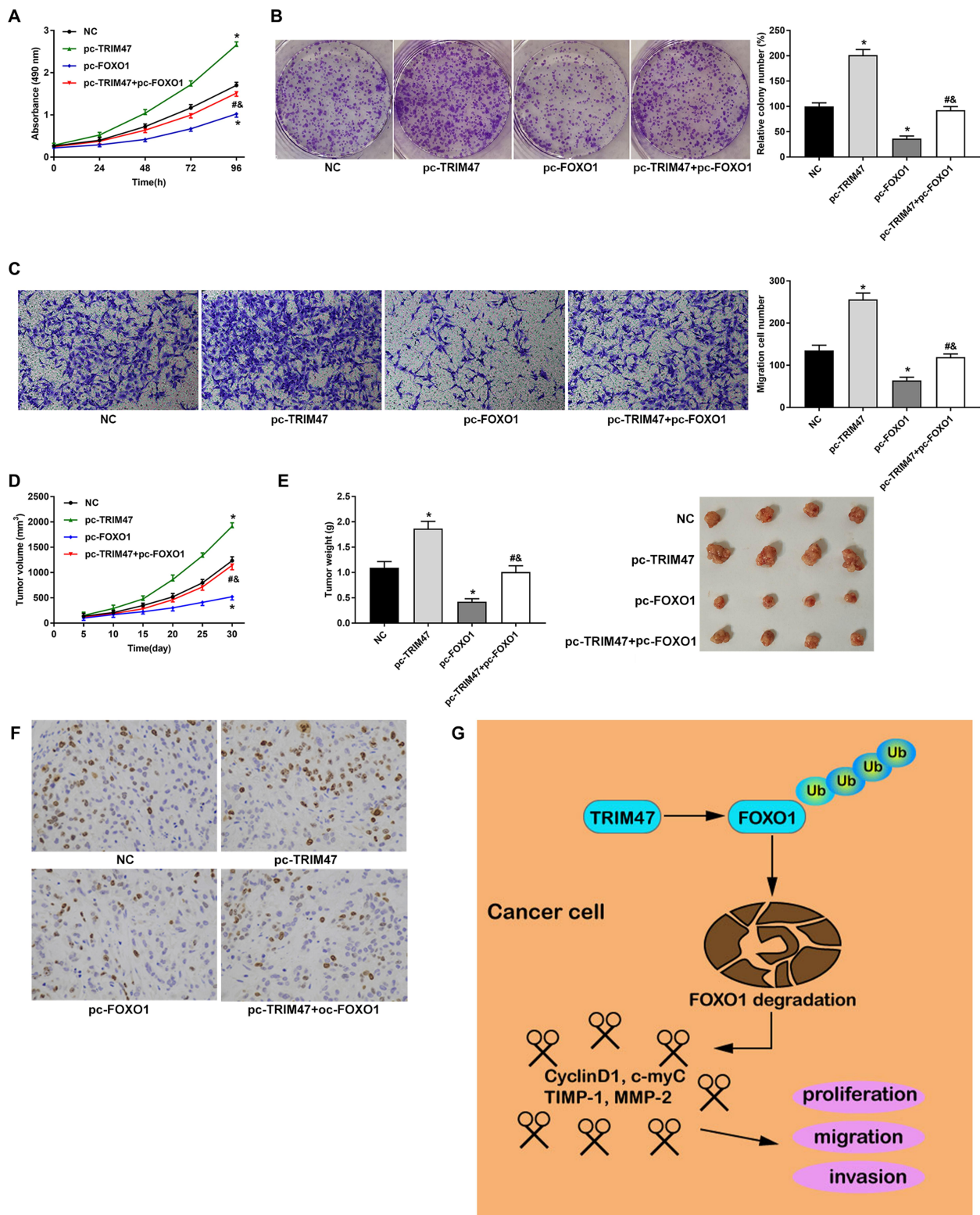


Figure 5 TRIM47 promoted glioma development through FOXO1 ubiquitination. **(A)** MTT was used to detect the proliferation of glioma cells; **(B)** Colony formation assay was used to detect glioma cell proliferation; **(C)** Transwell assay was used to detect glioma cell migration ability after TRIM47 silencing; **(D)** The changes of tumor volume in xenograft glioma mice; **(E)** The changes of xenograft tumor weight in glioma mice. **(F)** Immunohistochemistry (IHC) staining was performed to analyze the positive expression of proliferation marker Ki67 in glioma mice tumor. **(G)** A schematic model of TRIM47 function in glioma. TRIM47 interacts with FOXO1 and promotes the ubiquitination and degradation of FOXO1, ultimately promoting glioma cells proliferation, migration and invasion. *P<0.05 vs NC group, #P<0.05 vs pc-TRIM47 group, &P<0.05 vs pc-FOXO1 group.

closely related to the malignancy of the tumor and the prognosis.¹⁹ In our study, we observed that *TRIM47* was highly expressed in glioma tissue and cells, and its expression level was significantly correlated with pathological stage. In addition, compared with patients with low *TRIM47* expression, the overall survival rate of patients with high *TRIM47* expression was significantly reduced. Therefore, we speculate that *TRIM47* may play an important role in glioma development. To verify this hypothesis, we then constructed *TRIM47* low expression vectors, and evaluated the effect on the proliferation ability and migration ability of glioma cells by silencing *TRIM47* expression. As expected, in glioma cells cultured in vitro, silencing the expression of *TRIM47* significantly inhibited the proliferation and migration of glioma cells. Consistent with this, tumor volume and tumor weight were also significantly reduced in xenograft nude mouse models after silencing *TRIM47* expression. C-myc²⁰ and Cyclin D1²¹ were all proliferation-related proteins, and changes in the expression of these proteins can affect tumorigenesis and progression. MMP-2 and TIMP-1 were both migration-related proteins, but they play the opposite role in tumor progression. It has been reported that MMP-2 can promote the development of a variety of tumors,^{22,23} and its expression level increased with the increase of malignant potential; while TIMP-1 was to inhibit tumor invasion and metastasis.²⁴ Since the expression change of *TRIM47* affected the proliferation, migration and invasion of glioma, does it play a role through the above proteins? With this in mind, we examined changes in the expression levels of the aforementioned proteins. Our results showed that after *TRIM47* knockout, the protein levels of C-Myc, Cyclin D1 and MMP-2 significantly decreased, whereas the protein levels of TIMP-1 increased, which was consistent with previous reports. Collectively, the above data suggested that *TRIM47* may be an oncogene in human glioma.

TRIM proteins are typically involved in cancer development through ubiquitination and degradation of target proteins.²⁵ However, the potential molecular mechanisms of *TRIM47* in glioma cell growth, migration and invasion have been rarely investigated. *FOXO1* is a transcription factor that belongs to the FOX family. The gene encoding *FOXO1* is located on chromosome 13q14, and methylation, mutation, and allele loss are common in cancer. These characteristics suggest that there are potential tumor-related genes involved in the development of human malignancies in this region.²⁶ A growing body of

research reported that *FOXO1* as a cancer inhibitor can inhibit the development of different types of cancer, and the inactivation of *FOXO1* is related to the poor prognosis of patients.^{26,27} In addition, *FOXO1* protein levels and transcriptional activation are tightly regulated by a variety of post-translational modifications, including phosphorylation, acetylation, ubiquitination, and methylation.²⁸ In the presented study, we concluded that *TRIM47* can interact with *FOXO1* and degrade *FOXO1*. First, we found that knocking down *TRIM47* increased the expression of *FOXO1* in glioma cells, whereas overexpression of *TRIM47* decreased the expression of *FOXO1*. Secondly, treatment with proteasome inhibitor MG132 disrupted *TRIM47*-induced down-regulation of *FOXO1* in glioma cells. To further explore the role of the relationship between *TRIM47* and *FOXO1* in glioma progression, pc-*TRIM47* and pc-*FOXO1* were used for subsequent MTT, colony formation assay, Transwell assay experiments. The above data consistently demonstrated that *TRIM47* interacted with *FOXO1* to degrade *FOXO1* to promote the development and progression of glioma.

Conclusion

In our study, *TRIM47* was up-regulated in glioma cells. *TRIM47* inhibited *FOXO1* by ubiquitination and degradation of *FOXO1*, thereby promoting glioma growth and progression. This indicated that *TRIM47* can be considered as a biomarker to guide the diagnosis and treatment of glioma patients, and the measurement of *TRIM47* expression may become an effective method for predicting the prognosis of patients.

Statement of Ethics

This study was authorized by the ethical committee of Lianshui County People's Hospital (LSXRM2019-04) and all patients provided written informed consent.

Ethical and legal approval was obtained prior to the commencement of the study by the ethical committee of Lianshui County People's Hospital (LSXRM2019-04). All experiments were performed following the ethical committee of Lianshui County People's Hospital and national guidelines and regulations. All patients provided written informed consent.

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Disclosure

The authors indicate no potential conflicts of interest.

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