

# ARID4B Knockdown Suppresses PI3K/AKT Signaling and Induces Apoptosis in Human Glioma Cells

This article was published in the following Dove Press journal:  
*OncoTargets and Therapy*

Siou-Min Luo<sup>1</sup>  
Wen-Chiuan Tsai<sup>2</sup>  
Chia-Kuang Tsai<sup>1,3</sup>  
Ying Chen<sup>4</sup>  
Dueng-Yuan Hueng<sup>1,5,6</sup>

<sup>1</sup>Graduate Institute of Medical Sciences, National Defense Medical Center, Taipei, Taiwan, Republic of China; <sup>2</sup>Department of Pathology, Tri-Service General Hospital, National Defense Medical Center, Taipei, Taiwan, Republic of China; <sup>3</sup>Department of Neurology, Tri-Service General Hospital, National Defense Medical Center, Taipei, Taiwan, Republic of China; <sup>4</sup>Department of Biology and Anatomy, National Defense Medical Center, Taipei, Taiwan, Republic of China; <sup>5</sup>Department of Neurological Surgery, Tri-Service General Hospital, National Defense Medical Center, Taipei, Taiwan, Republic of China; <sup>6</sup>Department of Biochemistry, National Defense Medical Center, Taipei, Taiwan, Republic of China

**Purpose:** Glioblastoma multiforme is a highly malignant primary brain cancer with a poor prognosis. We recently reported that ARID4B could potentially serve as a biomarker associated with poor survival in glioma patients. However, the function of ARID4B in human gliomas remains unclear. The aim of this study is to investigate the molecular cell biology role of ARID4B in human glioma cells.

**Materials and Methods:** Gene Expression Omnibus (GEO) and Human Protein Atlas (HPA) datasets were analyzed for the expression of *ARID4B* in WHO pathological grading, overall survival and immunohistochemical staining. Using quantitative RT-PCR and Western blotting, those findings were confirmed in normal brain tissue and glioma cell lines. *ARID4B* knockdown was conducted via lentivirus-based transfection of small hairpin RNA in human glioma cells to investigate cell proliferation, cell cycle, and apoptosis.

**Results:** In the present study, our analysis of GEO datasets showed that *ARID4B* mRNA expression is higher in WHO grade IV tumors (n = 81) than in non-tumor control tissue (n = 23, P < 0.0001). *ARID4B* knockdown suppressed glioma cell proliferation and induced G1 phase arrest via the PI3K/AKT pathway. It also increased expression of *HDAC1*, leading to higher acetyl-p53 and acetyl-H3 levels and reduced glioma cell migration and invasion. These effects were mediated via down-regulation of AKT pathway components, including p-mTOR, p-PI3K and p-AKT. *ARID4B* knockdown also led to downregulation of Cyclin D1, which increased apoptosis in human glioma cells.

**Conclusion:** These findings that ARID4B expression correlates positively with WHO pathologic grading in glioma. *ARID4B* knockdown suppresses PI3K/AKT signaling and induces apoptosis in human glioma cells. These results suggest that ARID4B acts as an oncogene in human gliomas.

**Keywords:** invasion, migration, cell proliferation, mTOR, Cyclin D1

## Introduction

Glioblastoma multiforme (GBM) is the most common primary brain cancer and is highly aggressive. Despite treatment with chemoradiotherapy and/or surgical resection, GBM patient outcomes are poor.<sup>1</sup> Average survival among GBM patients treated with chemoradiotherapy is 14.6 months.<sup>2</sup>

AT-Rich Interaction Domain 4B (ARID4B) was initially detected in breast cancer tissues and is believed to promote tumor metastasis.<sup>3</sup> Identified as a component of the histone deacetylase (HDAC)-dependent SIN3A chromatin remodeling complex,<sup>4</sup> ARID4B contains both a DNA-binding domain and a tudor domain, and may function as an adapter protein, but in leukemia, ARID4B acts as a tumor suppressor.<sup>5</sup> In

Correspondence: Dueng-Yuan Hueng  
No. 325, Sec. 2, Chenggong Road, Neihu Dist., Taipei City, 114, Taiwan, Republic of China  
Tel +886-2-8792-7177  
Email hony2195@yahoo.com.tw

addition, *ARID4B* functions as a transcriptional activator of *PIK3CA* and *PIK3R2*, which regulate the PTEN-PI3K pathway in PTEN-deficient prostate cancers. *ARID4B* is thus a potential therapeutic target in prostate cancers lacking PTEN, as it is necessary for prostate cancer progression and initiation.<sup>6</sup> Expression of *ARID4B* is also elevated in various other cancer types. For example, it promotes mammary tumor metastasis and proliferation,<sup>7</sup> and it is reportedly a predictive biomarker in gliomas and meningiomas and is highly expressed in high-grade gliomas.<sup>8</sup> But despite these early findings, the role of *ARID4B* in cell biology of gliomas remains unclear. In the present study, therefore, we knocked down *ARID4B* in two glioma cell lines to investigate its function in high-grade gliomas.

## Materials and Methods

### Analysis of GEO Datasets for Human Gliomas

Sample profiles from the Gene Expression Omnibus (GEO) were analyzed. This included 180 files of de-linked data (GDS1962\_214934) on *ARID4B* mRNA expression [https://www.ncbi.nlm.nih.gov/geo/tools/profileGraph.cgi?ID=GDS1815:214943\\_s\\_at](https://www.ncbi.nlm.nih.gov/geo/tools/profileGraph.cgi?ID=GDS1815:214943_s_at). Information on pathological grading and *ARID4B* gene expression from the GDS1962\_214934 dataset included 130 data files from 23 patients without tumors (controls), 7 with grade II gliomas, 18 with grade III gliomas, and 81 with grade IV gliomas. The survival analysis of the datasets from GDS1815/214943\_s\_at ([https://www.ncbi.nlm.nih.gov/geo/tools/profileGraph.cgi?ID=GDS1815:214943\\_s\\_at](https://www.ncbi.nlm.nih.gov/geo/tools/profileGraph.cgi?ID=GDS1815:214943_s_at)) contained 100 high-grade gliomas samples (HG-U133A), which excluded unknown survival of 23 datasets; therefore, 77 patient survival data were analyzed and showed.

### Analysis of The Cancer Genome Atlas (TCGA) Datasets for Human Gliomas from the Human Protein Atlas (HPA)

The survival analysis of the datasets from TCGA in the Human Protein Atlas (<https://www.proteinatlas.org/ENSG00000054267-ARID4B/pathology/glioma>) contained 153 high-grade gliomas samples survival data were analyzed and showed.

### Datasets of Immunohistochemical Staining from the Human Protein Atlas (HPA)

Immunohistochemical staining for *ARID4B* in human gliomas was examined using the Human Protein Atlas

(HPA) (<https://www.proteinatlas.org/ENSG00000054267-ARID4B/pathology/tissue/glioma#>). The dataset included pathological glioma sections from 11 patients that were stained using anti-*ARID4B* antibody (HPA027333). Four of the 11 samples were low-grade gliomas, while the other 7 samples were high-grade. The quantity was defined as 0 (Negative), 1 (<25%), 2 (25–75%), and 3 (>75%). The intensity was scored as 0 (Negative), 1 (Weak), 2 (Moderate), and 3 (Strong). Labeling index was scored based on multiplying quantity by intensity.

### Cell Culture and shRNA Transfection

LN229, U87MG and U118 glioma cell lines purchased from American Type Culture Collection (ATCC). GBM8401 glioma cell line, provided by Dr. Wei-Hwa Lee, was obtained from the patient before the establishment of the Institutional Review Board (IRB) of the National Defense Medical Center (NDMC). The characterization of GBM8401 glioma cell line has published in article entitled as Establishment and characterization of a malignant glioma cell line, GBM8401/TSGH, NDMC.<sup>9</sup> GBM8401, U87MG and U118 human glioma cells were cultured in high-glucose (4.5 g/L) Dulbecco's modified Eagle's medium (DMEM, Corning, USA REF:10–013-CV) containing 10% fetal bovine serum (FBS, Gibco USA REF:10,437–028), penicillin, and streptomycin at 37°C in a 5% CO<sub>2</sub> incubator. LN229 human glioma cells were cultured in low-glucose (1 g/L) DMEM (Corning, USA REF: 10–014-CV) containing 10% FBS (Corning, USA REF: 10,437–028), penicillin, and streptomycin (Corning, USA REF: 30–004-CI) at 37°C in a 5% CO<sub>2</sub> incubator.

For *ARID4B* knockdown experiments, GBM8401 or LN229 cells were infected for 24 h with a shLuc control and *ARID4B*-shRNA-containing lentiviral vector (TRCN0000146977, RNAi core, Academia Sinica, Taiwan, ROC), after which the transfected cells were selected for 14 days in medium containing puromycin (2 g/L InvivoGen CAS:59–58-2). Stable clones were cultured in medium containing puromycin (0.2 mg/L).

### cDNA Synthesis and RNA Isolation

Extraction of total mRNA, reverse transcription and quantitative RT-PCR were performed according to the manufacturer's protocol. For cDNA synthesis, using Oligo dT primer reverse transcribed 1.0 µg RNA into cDNA with Biotin Tetro Reverse Transcriptase (London, UK Cat.No.BIO-65,050). Normal brain cDNA was obtained from Origene Technologies

(Rockville, MD, USA cat.CH1001). The reverse transcripts were amplified and quantified using a Onestep RT-qPCR system. GAPDH served as an internal control to normalize relative gene expression quantified using the  $2^{-\Delta\Delta}$  CT method. The primer pairs used were as follows: for ARID4B, 5'-ATGAGCCTCCCTATTTGACAGT-3' (forward) and 5'-GGCCCTTTATGTGGTCATCCT-3' (reverse); for GADPH, 5'-CTTCATTGACCTCAACTAC-3' (forward) and 5'-GCCATCCACAGTCTTCTG-3' (reverse); for HDAC1, 5'-CGCCCTCACAAAGCCAATG-3' (forward) and 5'-CTGCTTGCTGTACTCCGACA-3' (reverse); for PIK3CA, 5'-CCACGACCATCATCAGGTGAA-3' (forward) and 5'-CCTCACGGAGGCATTCTAAAGT-3' (reverse); for CCND1, 5'-GCTGCGAAGTGAAACCATC-3' (forward) and 5'-CCTCCTTCTGCACACATTGAA-3' (reverse); for PTEN, 5'-TGGATTGACTTAGACTTGACCT-3' (forward) and 5'-GGTGGGTTATGGTCTTCAAAGG-3' (reverse); for SIN3A, 5'-GGTGGAGGATGCGCTATCTTA-3' (forward) and 5'-GGGTGTCGATGCTCTGAGATTT-3' (reverse).

## Cell Lysate Preparation and Western Blotting

GBM8401, LN229 and U87MG cells were lysed in RIPA buffer (100 mM Tris-HCl, 150 mM NaCl, 0.1% SDS, and 1% Triton-X-100) and centrifuged at 13,500 rpm for 10 min, after which the supernatant was collected as the cell lysate. Cell lysates each containing 20  $\mu$ g of protein were analyzed by 8% and 12% sodium dodecyl sulfate polyacrylamide gel electrophoresis. Following electrophoresis, the separated proteins were transferred to PVDF membranes, which were then blocked for 1 h at room temperature in 5% skim milk in TBST buffer. The antibodies used included anti-ARID4B (BETHYL A302-223A, 1:2000), Cyclin D1 (Cell Signaling Technology CST2978, 1:1000), phosphorylated (p)-PI3K (Cell Signaling Technology CST4228, 1:1000), PI3K (Cell Signaling Technology CST4292, 1:2000), p-AKT (Ser473) (Cell Signaling Technology CST9271 1:2000), AKT (Cell Signaling Technology CST4691 1:3000), p-mTOR (Ser2448) (Cell Signaling Technology CST2971, 1:1000), mTOR (Cell Signaling Technology CST2972, 1:1000), acetyl-p53 (Cell Signaling Technology CST2972, 1:2000), p53 (Santa Cruz Biotechnology E2918, 1:3000), acetyl-H3 (Cell Signaling Technology CST9677, 1:1000), GAPDH (Santa Cruz Biotechnology SC-47724, 1:5000) and ACTN (Santa Cruz Biotechnology

SC-17829, 1:5000). Bands were developed using enhanced chemiluminescence and X-ray film (GE Healthcare, RPN2106, Piscataway, NJ, USA).

## Soft Agar Colony Formation Assays

To assay colony formation in soft agar, we seeded LN229 and GBM8401 cells with knockdown *ARID4B*, and control stable clones at a density of  $2 \times 10^3$  cells/well into 6-well plates with medium containing 0.35% SeaPlaque Agarose (Lonza Rockland, Inc. ME, USA cat.50002) layered on top of a base medium containing 0.5% agarose. After 4 weeks, the colonies were stained with 0.1% crystal violet (Merck KGaA DE, Cat.115940), and those greater than 0.05 mm in size were counted using ImageJ software (NIH, Bethesda, MD).

## Cell Migration and Invasion Assay

Cells were seeded into 12-well plates and grown at 37°C in a 5% CO<sub>2</sub> incubator. When the cells reached 90% confluence, we removed the medium, made a wound in the monolayer with a 200  $\mu$ L pipette tip, and washed the plate three times to remove non-adherent cells. The wound area was photographed immediately after wounding (0 hr) and again 16 h later. The migration rates were computed based on the change in wound area measured using ImageJ software (NIH, Bethesda, MD).

Invasion assays were carried out in Transwell chambers (Corning BioCoat cat.354480). A Matrigel matrix layer was coated onto the upper side of the filter separating the upper and lower chambers. Glioma cells in serum-free DMEM were then added to the upper chambers, and medium containing 10% FBS was added to the lower chambers. After incubation for 16 h at 37°C in a CO<sub>2</sub> incubator, the non-migrated cells on the upper side of the filter were removed. Cells that invaded the Matrigel and migrated to the lower side of the filter chamber were stained with 0.1% crystal violet for 10 mins. The glioma cells in upper side of the filter were removed by swab. The invaded glioma cells in lower side of the filter was photographed. Finally, the invading cells were dissolved in 1% SDS, and the optical density at 595 nm was measured.

## Cell Proliferation and Flow Cytometric Cell Cycle Analysis

For cell cycle analysis, glioma cells were fixed in 70% ethanol at 4°C and kept at -20°C overnight. The cells

were then washed twice with cold phosphate-buffered saline (PBS) and stained with propidium iodide (PI, Sigma-Aldrich, MO, USA Cas.25535–16-4) (50 µg/mL PI in PBS, 1% Tween 20 and 10 µg/mL RNase A) for 30 min in the dark. Cell cycle analysis was performed by measuring DNA content using fluorescence activated cell sorting (BD Biosciences, San Jose, CA, USA). Two independent experiments were performed. For the proliferation assays, glioma cells were processed using FITC-BrdU Flow Kits according to the manufacturer's instructions (BD Biosciences, San Jose, CA, USA cat.51–2354AK). The cells were then trypsinized with trypsin EDTA and analyzed immediately using a FACS Calibur flow cytometer (BD Biosciences, San Jose, CA, USA). The data collected were analyzed using FACSDiva software (BD Biosciences, San Jose, CA, USA).

### Protein–Protein Interaction Network Analysis and Genomic Data Commons Data Portal

The distribution of ARID4B in datasets from the Genomic Data Commons Data Portal were determined and analyzed for known and predicted protein–protein interactions using the Search Tool for the Retrieval of Interacting Genes/Proteins (STRING) database (<https://string-db.org/>).

### Apoptosis Assay

To evaluate the incidence of apoptosis, we used the PE Annexin V Apoptosis Detection Kit I according to the manufacturer's instructions (BD Biosciences San Jose, CA, USA, cat.559763). Briefly, glioma cells were seeded into 6-well dishes at a density of  $3 \times 10^5$  cells per well. Stained cells were detected using Annexin V-FITC/7-AAD. The cells were categorized into 3 groups: viable (annexin V<sup>-</sup> and FITC<sup>-</sup>), early apoptotic (annexin V<sup>+</sup> and FITC<sup>-</sup>), and later apoptotic (annexin V<sup>+</sup> and FITC<sup>+</sup>).

### Statistical Analysis

Student's *t*-test or one-way analysis of variance was used to make comparisons among groups. The results are presented as the means  $\pm$  s.d. or as specified. Values of  $*P < 0.05$  were considered significant.

## Results

### ARID4B Expression Correlates Positively with WHO Pathological Grading of Gliomas and with Poor Survival in Patients with High-Grade Gliomas

To correlate WHO pathological grading of gliomas with *ARID4B* expression, we analyzed the association between *ARID4B* expression and pathological grading of human gliomas using a GEO mRNA de-linked databases (GDS1962\_214934). We found that *ARID4B* expression was significantly lower in non-tumor controls ( $n = 23$ ) than in WHO grade III ( $n = 19$ ;  $P = 2 \times 10^{-6}$ ) and grade IV ( $n = 81$ ;  $P = 1.36 \times 10^{-9}$ ) tumors (Figure 1A).

### ARID4B Expression and Survival analyses in High-Grade Gliomas

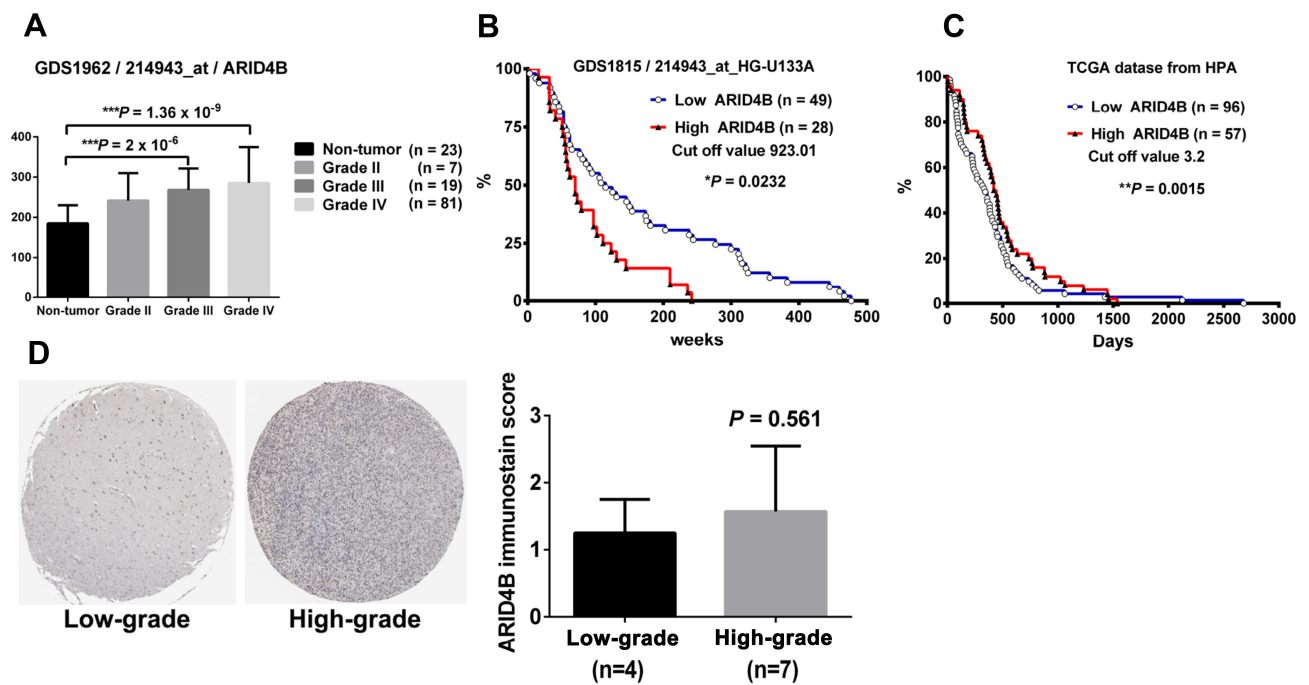
GEO profiles were used to analyze the relationship between *ARID4B* mRNA expression and survival. As shown in Figure 1B, the group from GDS1815/214943\_s\_at HG-U133A showing high *ARID4B* expression had poorer survival than that showing low *ARID4B* expression (median survival: 70 vs 115 weeks,  $*P = 0.0232$ , Gehan-Breslow-Wilcoxon Test; hazard ratio: 0.4210, 95% CI: 0.2398-0.7393, cut-off value set at 923.01). Thus, *ARID4B* expression is higher in high-grade than in low-grade gliomas, which suggests ARID4B overexpression contributes to the poor prognosis of GBM patients. In contrast, the survival curves from TCGA database showed the survival probability in glioma patients with low *ARID4B* expression ( $<3.52$ ) ( $n = 96$ ) was shorter than high *ARID4B* expression,  $**P = 0.0015$  (Figure 1C).

### Immunohistochemical Staining of ARID4B in Human Glioma Patient Samples

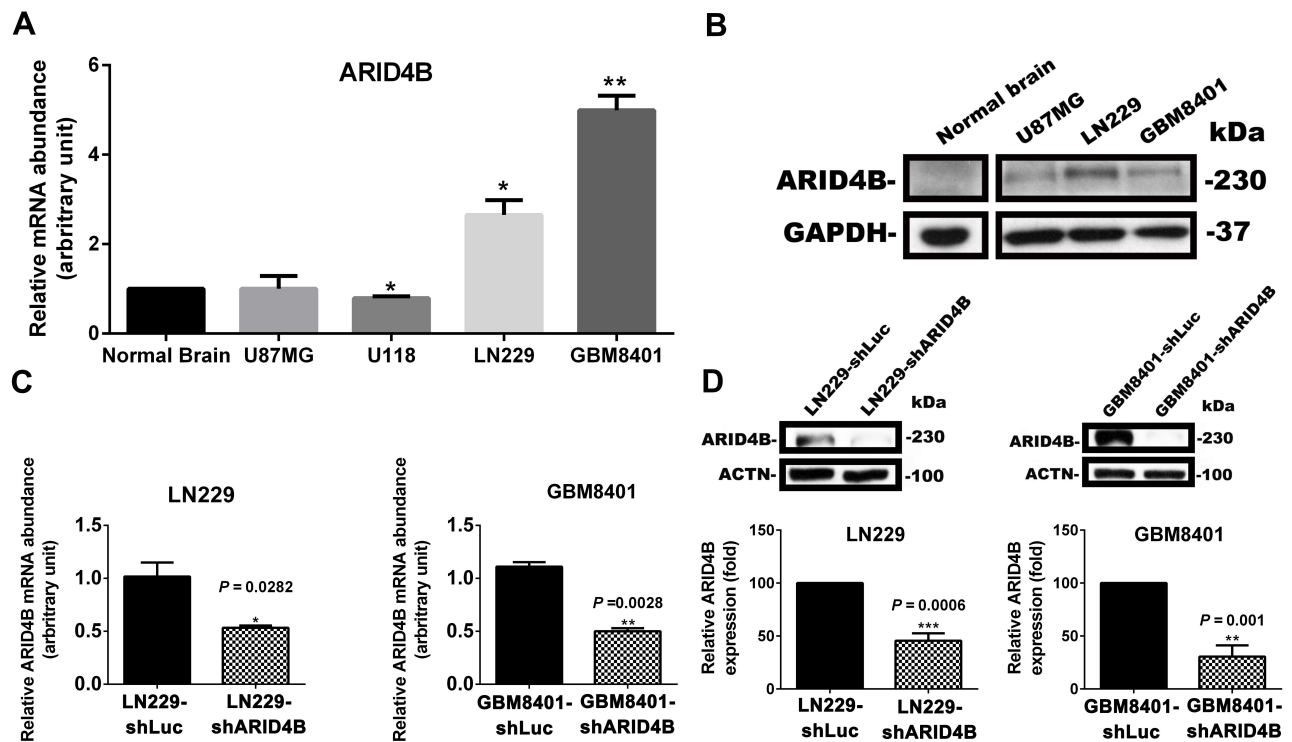
Using HPA database of human proteins, we found that the immunohistochemical staining of ARID4B in high-grade glioma was higher than that in low-grade glioma, but it was not statistically significant (Figure 1D).

### ARID4B Knockdown in Human Glioma Cell Lines

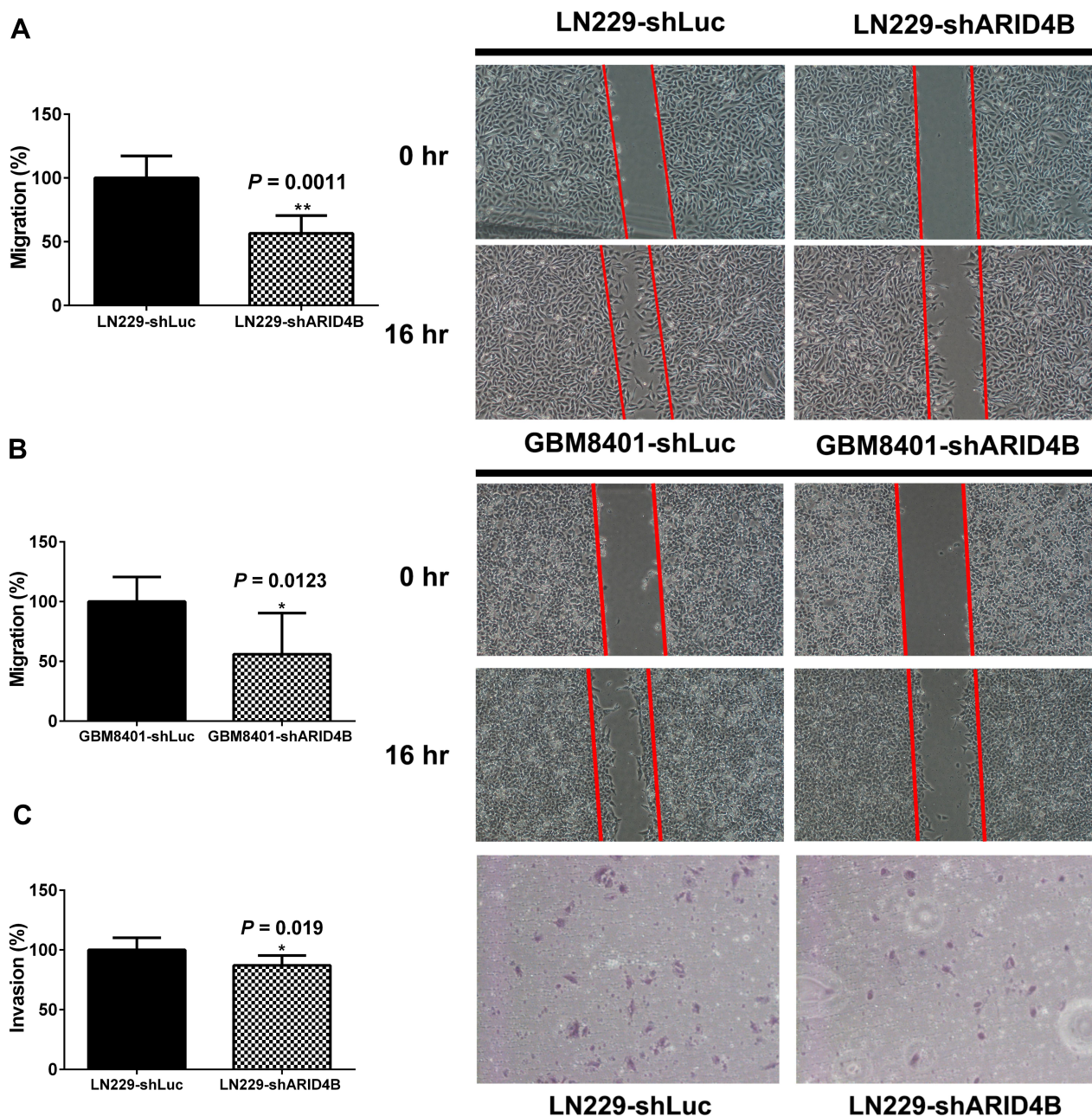
Confirming *ARID4B* expression levels in human glioma cell lines, RT-qPCR showed that both LN229 and GBM8401 human glioma cells expressed higher levels of *ARID4B* mRNA than normal brain cells (Figure 2A).



**Figure 1** Expression of *ARID4B* correlates positively with WHO grading of human gliomas and *ARID4B* staining in human glioma samples from the Human Protein Atlas. (A) Bar graph of *ARID4B* mRNA levels from the GDS1962\_214934 dataset showing a significant correlation between *ARID4B* mRNA expression and WHO tumor grades for human gliomas  $***P < 0.001$ . (B) Kaplan–Meier survival curves showing significantly poorer survival among glioma patients highly expressing *ARID4B* ( $>923.01$ ) ( $n = 28$ ) than among those expressing lower levels of *ARID4B* ( $<923.01$ ) ( $n = 49$ ) expression levels of *ARID4B*.  $*P < 0.05$  (Gehan-Breslow-Wilcoxon Test) from GDS1815/214943\_s\_at contained 100 high-grade gliomas samples (HG-U133A). (C) Kaplan–Meier survival curves showed the survival probability in glioma patients with low *ARID4B* expression ( $<3.52$ ) ( $n = 96$ ) was shorter than high *ARID4B* expression ( $>3.52$ ) ( $n = 57$ )  $**P < 0.01$  (Log rank test) from TCGA database. (D) Immunohistochemical staining showed *ARID4B* expression in high-grade gliomas was higher than that in low-grade gliomas. Statistical analysis of *ARID4B* labeling scores. *ARID4B*-stained areas were defined as being less than 25%, 25–75%, or more than 75% of the sample areas.



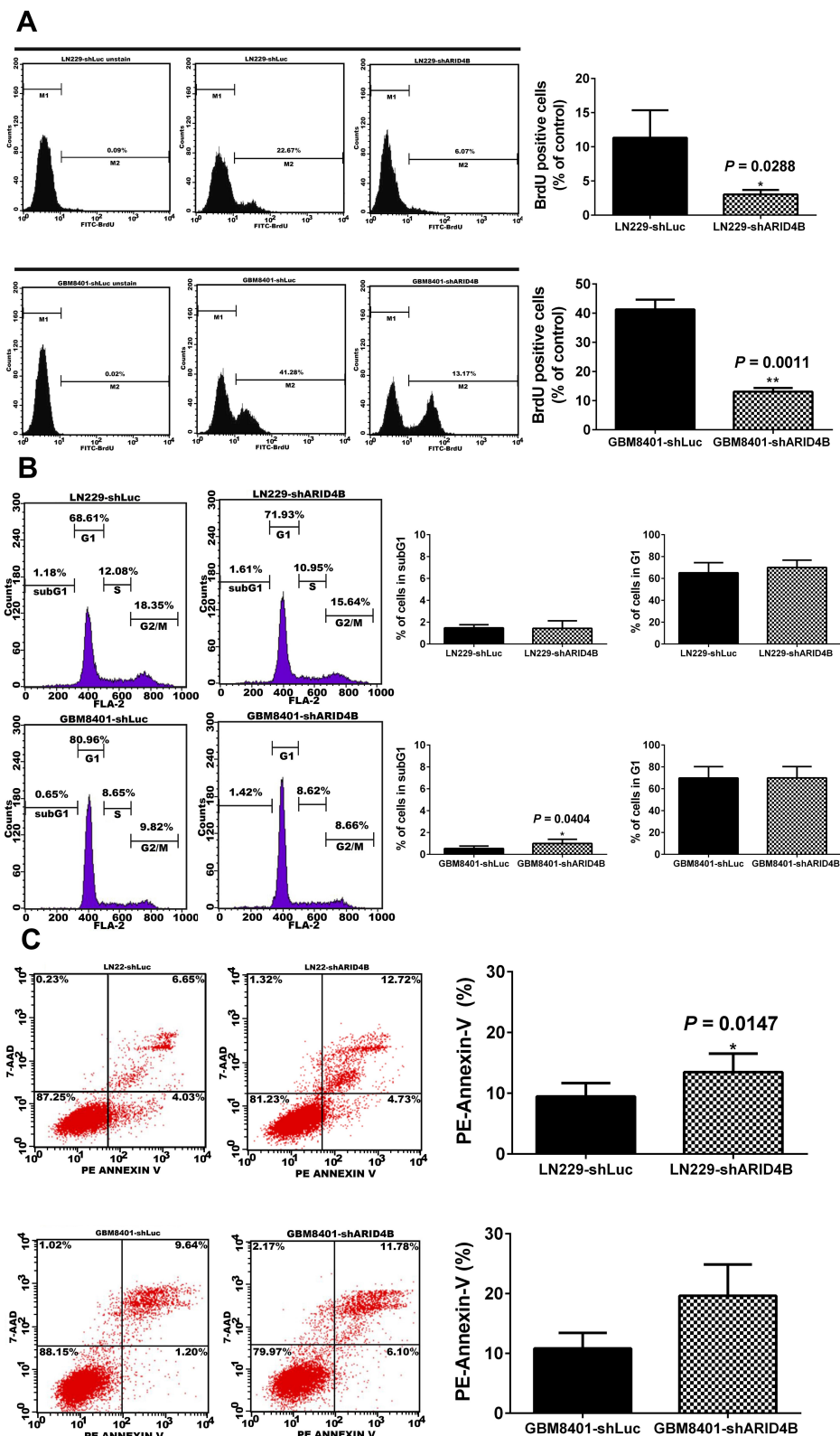
**Figure 2** *ARID4B* expression in gliomas and establishment of *ARID4B* knockdown glioma cells. (A) Levels of *ARID4B* expression in gliomas determined with RT-qPCR.  $*P < 0.05$ ,  $**P < 0.01$  vs the control shLuc group (B) Western blots showing *ARID4B* expression in gliomas. *ARID4B* knockdown using shARID4B in LN229 and GBM8401 validated by (C) RT-qPCR and (D) Western blotting.  $*P < 0.05$ ,  $**P < 0.01$ ,  $***P < 0.001$  vs the control shLuc group.



**Figure 3** Effect of *ARID4B* knockdown on glioma cell migration and invasiveness. (**A** and **B**) Migration assessed based on the decreases in wound area during 16-h wound-healing assays with GBM8401 and LN229 cells expressing shARID4B or shLuc. Wound area at 0 h was assigned a value of 100%. \* $P < 0.05$ , \*\* $P < 0.01$  vs the control shLuc group. (**C**) Invasiveness assayed in 16-h Transwell invasion assays with LN229 cells expressing shARID4B or shLuc. In all panels, the data were from three independent experiments, and expressed as the mean  $\pm$  s.d. \* $P < 0.05$  vs the control shLuc group.

Correspondingly, Western blotting showed higher *ARID4B* protein expression in the two human glioma cell lines (Figure 2B). We therefore chose LN229 and GBM8401 cells for subsequent *ARID4B* knockdown experiments. To determine the mechanism by which *ARID4B* contributes to glioma tumorigenesis, we used *ARID4B* shRNA purchased from the RNAi core at

Academia Sinica. We then established GBM8401 and LN229 glioma cells stably expressing shARID4B, which was confirmed using RT-qPCR. The results showed that the *ARID4B* mRNA expression was significantly suppressed in the *ARID4B* knockdown cells (\* $P = 0.0282$  and \*\* $P = 0.0028$ , respectively) (Figure 2C). Western blotting revealed similar suppression of *ARID4B* protein expression in the



**Figure 4** Effect of *ARID4B* knockdown on cell proliferation, cell cycle, and apoptosis. **(A)** GBM8401 and LN229 cells were labeled with BrdU and then processed for flow cytometric analysis. \* $P < 0.05$ , \*\* $P < 0.01$  vs the control shLuc group. **(B)** Effect of *ARID4B* knockdown on cell cycle regulation in glioma cells assessed using propidium iodide. \* $P < 0.05$  vs the control shLuc group. **(C)** Annexin-V/7-AAD analysis of the effect of *ARID4B* knockdown on apoptosis among glioma cells. In all panels, the data are shown as the means  $\pm$  s.d.;  $n = 3$ ; \* $P < 0.05$  vs the control shLuc group.

knockdown cells ( $***P = 0.0006$  and  $**P = 0.001$ , respectively) (Figure 2D).

## ARID4B Knockdown Suppresses Glioma Cell Migration and Invasion

To assess the contribution of ARID4B to glioma cell migration behavior, we performed wound healing assays with LN229 and GBM8401 glioma knockdown cells. The results demonstrate that ARID4B knockdown significantly reduces glioma cell migration ( $**P = 0.0011$ , and  $*P = 0.0123$ , respectively) (Figure 3A and B). In addition, Matrigel-based invasion assays similarly showed that silencing ARID4B expression in LN229 glioma cells significantly suppressed the invasiveness of the cells ( $*P = 0.019$ ) (Figure 3C).

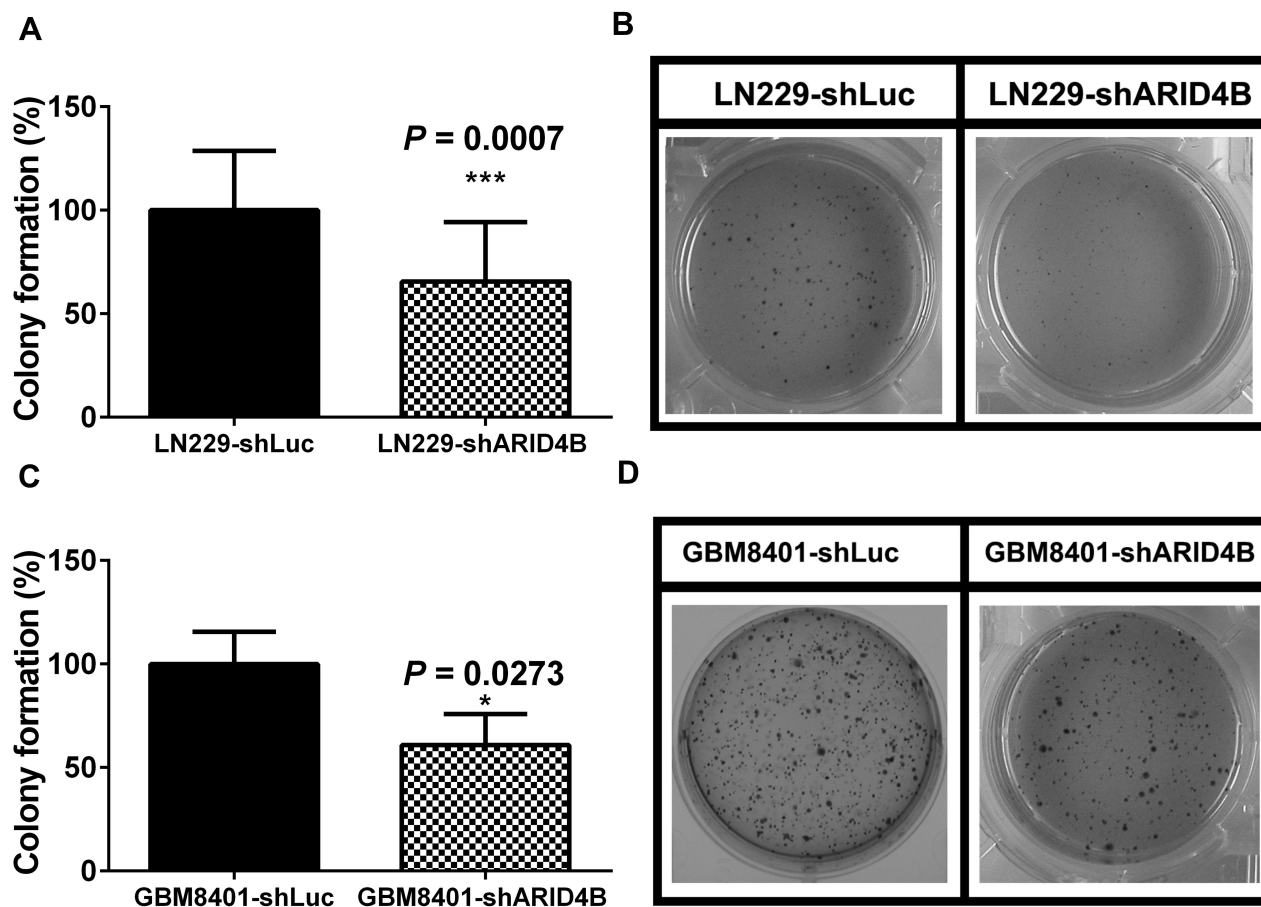
## ARID4B Knockdown Suppresses Cell Proliferation and Increases Apoptosis

We used BrdU assays to investigate LN229 and GBM8401 cell proliferation. Flow cytometric analysis showed that

ARID4B knockdown significantly reduced proliferation of both LN229 ( $*P = 0.0288$ ) and GBM8401 ( $***P = 0.0011$ ) glioma cells (Figure 4A). ARID4B knockdown also reduced expression of Cyclin D1, a pro-survival effector protein, and increased the glioma cell fraction in subG1 phase (Figure 4B). In addition, Annexin-V assays showed that ARID4B knockdown significantly increased Annexin-V/7-AAD ( $*P = 0.0147$ ), indicating that ARID4B knockdown enhanced apoptosis (Figure 4C).

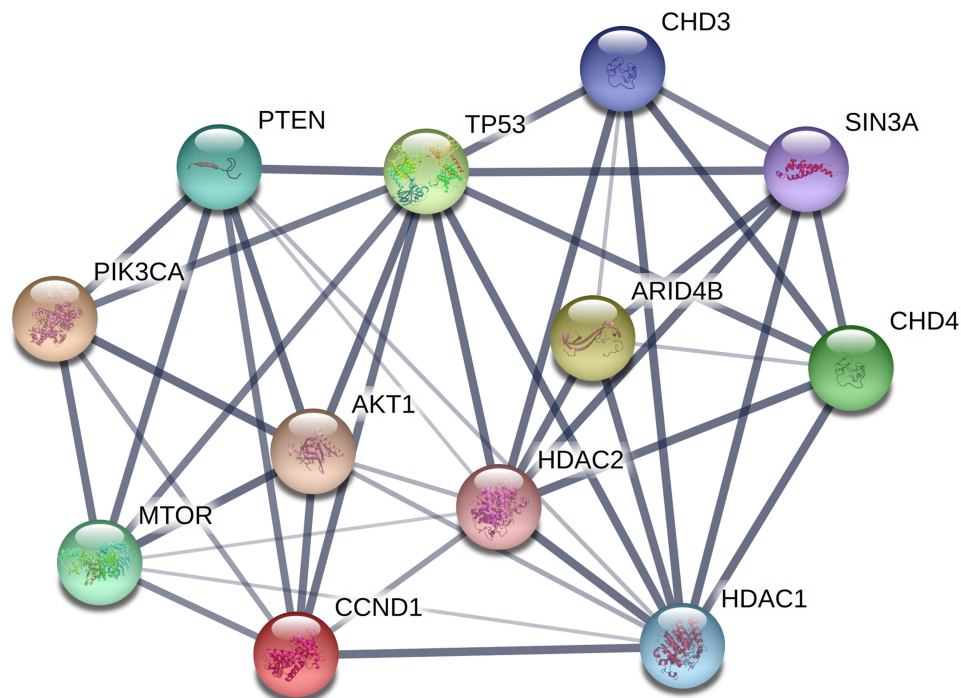
## ARID4B Knockdown Suppresses Colony Formation by Human Glioma Cells

Because ARID4B overexpressed in high-grade gliomas, we investigated the effect of ARID4B knockdown on the tumorigenicity of LN229 and GBM8401 glioma cells by carrying out soft-agar colony formation assays. We found that ARID4B knockdown led to a significant reduction of colony formation by LN229 and GBM8401 glioma cells ( $***P = 0.0007$  and  $*P = 0.0273$ , respectively) (Figure 5).



**Figure 5** Effect of ARID4B knockdown on colony formation by glioma cells. (A and C) Images showing colony formation in soft agar by GBM8401 and LN229 cells expressing shARID4B or shLuc. (B and D) Bar graph showing ARID4B knockdown suppresses colony formation by glioma cells. Data are presented as the mean  $\pm$  s.d.; n = 3;  $*P < 0.05$ ,  $***P < 0.001$  vs the shLuc group.





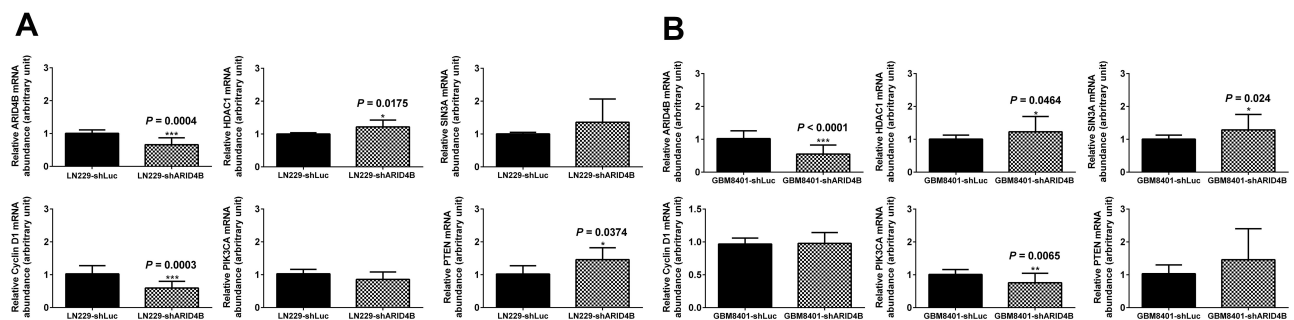
**Figure 6** ARID4B protein-protein interaction (PPI) network. The PPI network was generated using the STRING database.

## ARID4B Inhibits Glioma Growth via the PTEN-ARID4B-PI3K Pathway

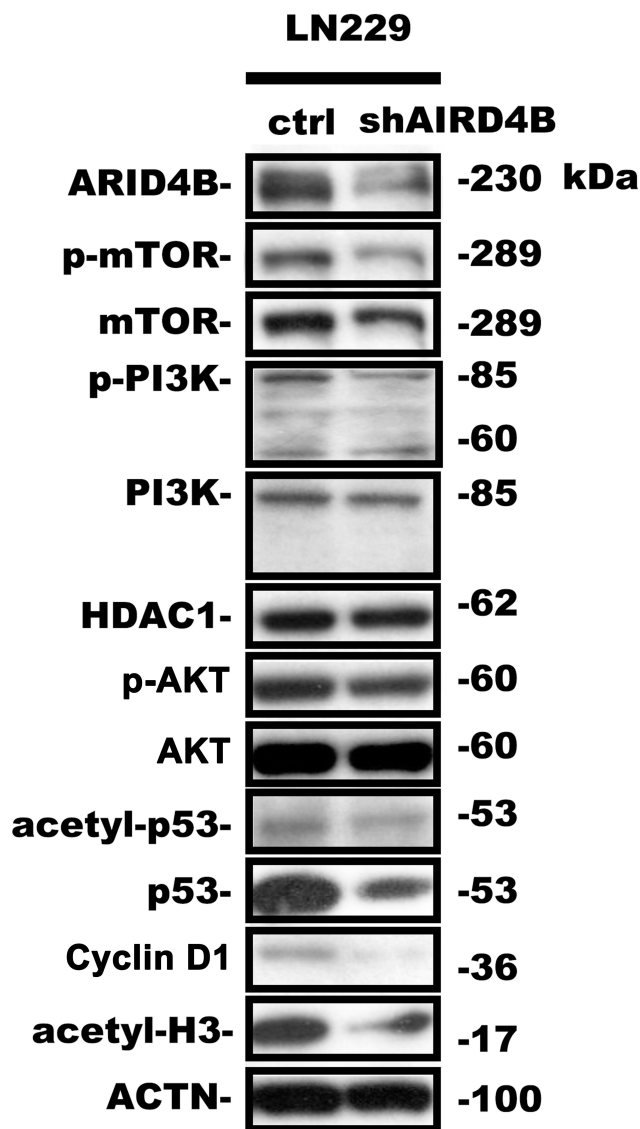
The ARID4B protein-protein interaction (PPI) network obtained with the Search Tool for the Retrieval of Interacting Genes/Proteins (STRING) database showed in [Figure 6](#). We found that ARID4B interacts with HDAC1, HDAC2, chromodomain-helicase-DNA-binding protein 3 (CHD3), chromodomain-helicase-DNA-binding protein 4 (CHD4), TP53, Cyclin D1 (CCND1), phosphatase and tensin homolog (PTEN), SIN3 Transcription Regulator Family Member A (SIN3A), phosphoinositide 3-kinase (PI3K), AKT, and mammalian target of rapamycin (mTOR). AKT, also known as protein kinase B (PKB), is a serine/threonine protein kinase that functions as an effector protein situated downstream of

PI3K. The PI3K/AKT/mTOR signaling pathway is important for mediating cell survival and growth. PTEN suppresses PI3K/AKT signaling by dephosphorylating PIP3. HDAC1 is a key regulator of TP53 activity that acts through deacetylation. ARID4B is involved in the SIN3/HDAC1 complex and recognized to be a transcriptional activator of class I PI3K subunit p110 $\alpha$  (encoded by *PIK3CA*), which is a regulator of the PTEN-PI3K pathway.

ARID4B was recently identified as a component of the HDAC1/SIN3A chromatin remodeling complex and as a regulator of PTEN-PI3K pathway that acts by targeting a *PIK3CA* transcriptional activator. Using real-time qPCR, we found here that *ARID4B* knockdown reduced *PIK3CA* and *Cyclin D1* gene expression and



**Figure 7** Effect of *ARID4B* knockdown on expression of the indicated mRNAs. **(A and B)** RT-qPCR analysis of expression of the indicated mRNAs in LN229 and GBM8401 cells. Data are presented as the mean  $\pm$  s.d.; n = 3; \*P < 0.05, \*\*P < 0.01, \*\*\*P < 0.001 vs the control shLuc group.



**Figure 8** Effect of *ARID4B* knockdown on expression of the indicated proteins. Western blot analysis of expression of the indicated proteins in LN229 cells. Statistical data are presented as the mean  $\pm$  s.d.; n = 3; in [Table 1](#).

increased gene expression of *SIN3A*, *HDAC1* and *PTEN* in LN229 ([Figure 7A](#)) and GBM8401 ([Figure 7B](#)) glioma cells. In addition, Western blotting showed that *ARID4B* knockdown reduced the protein expression of p-AKT, p-mTOR, and the p110 $\alpha$  subunit of PI3K. On the other hand, levels of HDAC1 were increased in *ARID4B* knockdown glioma cells. The elevated HDAC1 levels led in turn to decreases in acetyl-p53 and acetyl-H3 in LN229 ([Figure 8](#) and [Table 1](#)) glioma cells. Taken together, our results demonstrate that *ARID4B* knockdown increases HDAC1 levels, leading to reductions in acetyl-p53 ([Figure 9](#)).

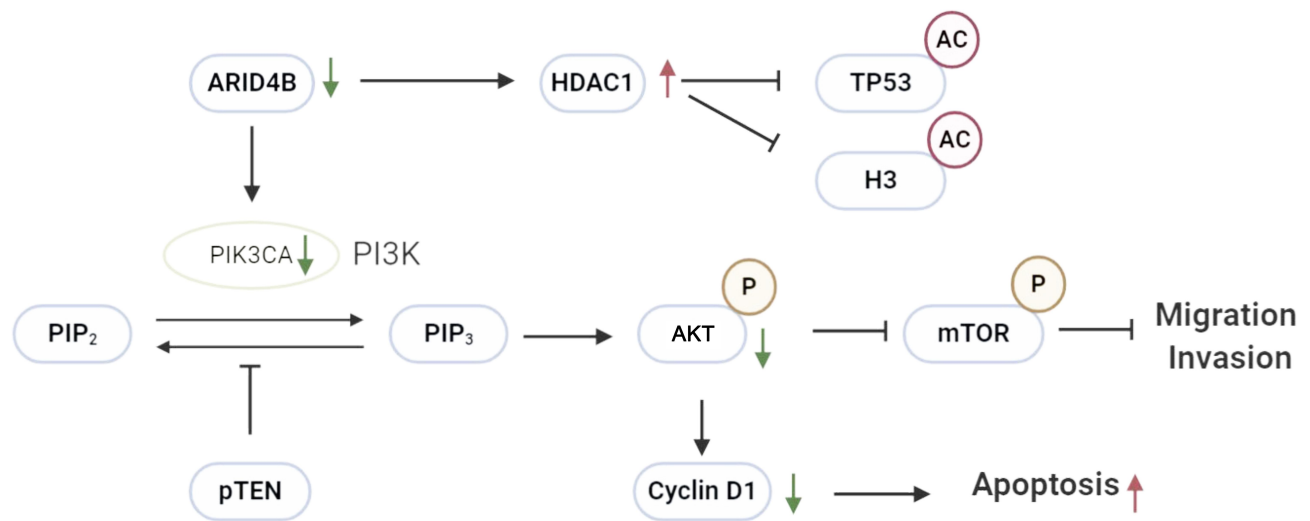
**Table 1** The Proteins are Upregulated and Downregulated in *ARID4B* Knockdown

Protein	Fold Change	P value
ARID4B	-1.36	***p = 0.0003
HDAC1	0.35	*p = 0.0464
Cyclin D1	-2.46	**p = 0.0167
Acetyl-H3	-0.78	P = -0.78
Acetyl-p53	-0.58	***p = 0.0065
Total-p53	0.09	P = 0.09
Acetyl-p53/p53	-0.36	*p = 0.018
p-AKT	-0.96	**p = 0.0016
Total AKT	-0.12	P = -0.12
p-AKT/AKT	-0.98	**p = 0.0028
p-mTOR	-0.88	***p < 0.0001
mTOR	-0.04	P = -0.04
p-mTOR/mTOR	-0.78	**p = 0.0011
p-PI3K(85)	-0.42	P = -0.42
PI3K(85)	0.00	P = 0.00
p-PI3K/PI3K (85)	-0.41	P = -0.41
PI3K(60)	0.22	P = 0.22

**Notes:** Data are presented as the mean  $\pm$  s.d.; n = 3 and shARID4B group vs the control shLuc group; \*P < 0.05, \*\*P < 0.01, \*\*\*P < 0.001 vs the control shLuc group. **Abbreviations:** ctrl, control shLuc; ARID4B, AT-rich interaction domain 4B; HDAC1, histone deacetylase 1; CHD3, chromodomain-helicase-DNA-binding protein 3; CHD4, chromodomain-helicase-DNA-binding protein 4; PTEN, phosphatase and tensin homolog; SIN3A, SIN3 transcription regulator family member A; PI3K, phosphoinositide 3-kinase; AKT, protein kinase B; mTOR, mammalian target of rapamycin.

## Discussion

Our previous study showed that *ARID4B* expression tended to shorten survival rates, but it did not reach statistical significance.<sup>8</sup> We further enrolled more data to analyze, as shown in [Fig1B](#) and [Fig1C](#), these two cohorts showed that the *ARID4B* expression and survival outcome of human gliomas have some differences. The possible explanations included the different cohorts and different analytical methods.<sup>10</sup> Therefore, further enroll large cohort human gliomas are mandatory in the future. In the present study, we showed that *ARID4B* knockdown induces apoptosis via a PI3K/AKT signaling pathway in human glioma cells. To our knowledge, this is the first study investigating the molecular mechanism of action of *ARID4B* in human gliomas. Previous studies showed that *ARID4B* took part in differentiation and cell growth.<sup>11</sup> miR-290-targeted *ARID4B* inhibited the promoting effects in breast cancer.<sup>12</sup> *ARID4B* reported that gastric cancer can be detected through molecular imaging. Furthermore, *ARID4B* might serve as a treatment target in gastric cancer.<sup>3,13-14</sup> *ARID4B* is considered to be an oncogene in hepatocellular carcinoma and can be used as a biomarker for the prognoses of patients.<sup>15</sup>



**Figure 9** Schematic of knockdown *ARID4B* in gliomas. Knockdown *ARID4B* suppresses expression of phosphorylation of the PI3K downstream effectors mTOR and AKT and the downstream target Cyclin D1 has been reduced. Upregulated genes and protein are indicated by red arrows, and down expressed are green arrow.

STRING network analysis showed that *ARID4B* interacts with HDAC1, Cyclin D1, PTEN, SIN3A, PI3K, AKT, TP53, and mTOR, which form important signaling pathways that play key roles in a variety of crucial cellular processes,<sup>16</sup> including cell migration and invasion.<sup>17</sup> Reducing levels of p-PI3K and p-AKT previously showed to suppress glioma cell migration and invasion.<sup>18</sup> Consistent with those findings, we observed that *ARID4B* knockdown reduced expression of both p-AKT and p-mTOR and suppressed cell migration and invasion. Recent studies showed that *ARID4B* is a component of the SIN3A/HDAC1 repressor complex; however, *ARID4B* does not exhibit acetyltransferase or methyltransferase activity,<sup>19</sup> though it does act as a regulator of the PI3K/AKT pathway.<sup>6</sup> Similarly, we found that *ARID4B* knockdown reduces expression of the PI3K p110 $\alpha$  (*PIK3CA*) and p85 subunits, as well as the levels of p-AKT, p-mTOR, and Cyclin D1. Cyclin D1 is an important regulator of cell proliferation that also controls tumorigenesis, cellular senescence, and apoptosis.<sup>20</sup> Through its actions, Cyclin D1 exerts cytoprotective effects against induction of apoptosis.<sup>16</sup> The PI3K/AKT signal transduction pathway is involved in the regulating survival and apoptosis. Inhibiting AKT phosphorylation may decrease the anti-apoptotic effect of AKT. Thus, reducing p-AKT, Cyclin D1 and p-mTOR levels relieved the suppression of apoptosis in glioma cells. It follows then that the reduced expression of p-AKT, p-mTOR and Cyclin D1 following *ARID4B* knockdown leads to enhanced apoptosis in glioma cells.<sup>18</sup>

Our results show that *ARID4B* knockdown increases expression of HDAC1, which is known to deacetylate p53 and H3. Histone deacetylase inhibitors (HDACIs) induce

apoptosis and growth arrest in many cancers.<sup>20</sup> In addition, p53 plays an important role in tumor oncogenic stress and cell death, acting as a crucial tumor suppressor mediating cell cycle arrest and apoptosis.<sup>21</sup> Inhibiting HDAC1 and SIRT1 leads to enhancement of p53 acetylation, which increases p53-dependent senescence and apoptosis in many cancer cell types.<sup>22</sup> This suggests p53 acetylation is an important regulator of apoptosis. The p53 pathway is highly dysregulated in GBM.<sup>23</sup> This is due in part to missense mutations in *p53* that often lead to reduced tumor suppressor activity.<sup>24</sup> In our study, we used LN229 and GBM8401 cells, two *p53*-mutant and *PTEN*-wild type glioblastoma cell lines. LN229 cells, for example, harbor a mutated *p53* with possible homozygous deletions.<sup>11</sup> We suggest that the ability of HDAC1 to suppress LN229 cell apoptosis may be decreased by the *p53* mutation.

## Conclusion

In summary, our data show that the level of *ARID4B* expression correlates positively with WHO pathologic grades of gliomas. *ARID4B* appears to be a potentially useful biomarker of the prognosis in glioma patients, and we suggest this may enable identification of novel therapeutic targets for treatment of human gliomas in the coming future.

## Data Sharing Statement

All data are included in this article.

## Consent for Publication

All participants agreed to publish the study.

## Acknowledgments

We appreciated the help for lentiviruses carrying a ARID4B-specific shRNA (shARID4B) from National RNAi Core Facility, Academia Sinica, Taiwan. We thank Professor Shih-Ming Huang (Graduate Institute of Biochemistry, National Defense Medical Center, Taipei, Taiwan) for providing the experimental help.

## Author Contributions

All authors made a significant contribution to the work reported in the conception, study design, execution, and acquisition of data, analysis and interpretation; took part in drafting, revising, and critically reviewing the article. All authors gave final approval of the version to be published; have agreed on the journal to which the article has been resubmitted; and agree to be accountable for all aspects of the work.

## Funding

This work was supported, in part, by grants from the Ministry of Science and Technology (MOST 106-2314-B-016-012-MY3, and MOST 108-2314-B-016-026 -MY3 to D.-Y.H.), Tri-Service General Hospital (TSGH-C107-008-S05, ATSGH-C107-008-S05, TSGH-C108-007-008-S05, TSGH-C01-109016, TSGH-2-C107-106-2314-B-016-012-MY3, TSGH-1-C108-106-2314-B-016-012-MY3, TSGH-2-C108-106-2314-B-016-012-MY3, TSGH-2-C108-108-2314-B-016-0-MY3 to D.-Y.H.), and Medical Affairs Bureau, Ministry of National Defense (MAB-106-019, MAB-107-009, MAB-108-022, MAB-109-014 to D.-Y.H.).

## Disclosure

The authors report no conflicts of interest in this work.

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