TRIM59 Promotes Retinoblastoma Progression by Activating the p38–MAPK Signaling Pathway

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Purpose. Retinoblastoma is a malignant tumor of the developing retina that mostly occurs in children. Our study aimed to investigate the effect of tripartite motif-containing protein 59 (TRIM59) on retinoblastoma growth and the underlying mechanisms.

Methods. We performed bioinformatic analysis of three datasets (GSE24673, GSE97508, and GSE110811) from the Gene Expression Omnibus database. Quantitative reverse-transcription PCR and immunoblotting of three retinoblastoma cell lines were conducted to verify TRIM59 as a differentially expressed gene. Specific siRNAs were used to inhibit TRIM59 expression in the HXO-Rb44 cell line. A lentiviral vector was transfected into the Y79 cell line to overexpress TRIM59. The effects of TRIM59 on retinoblastoma cell proliferation, cell cycling, and apoptosis were explored in vitro using the abovementioned cell lines. The effect of TRIM59 expression on retinoblastoma cell proliferation was evaluated in a mouse xenograft tumor model.

Results. TRIM59 expression in three retinoblastoma cell lines was remarkably elevated compared with normal control. Knocking down TRIM59 expression remarkably suppressed cell proliferation and growth and promoted cell apoptosis in HXO-Rb44 cells, whereas TRIM59 overexpression promoted tumor progression in Y79 cells. Silencing TRIM59 also markedly inhibited in vivo tumor growth in the xenograft model. Mechanistic studies revealed that TRIM59 upregulated phosphorylated p38, p-JNK1/2, p-ERK1/2, and p-c-JUN expression in retinoblastoma cells. Notably, the p38 inhibitor SB203580 attenuated the effects of TRIM59 on cell proliferation, apoptosis, and the G1/S phase transition.

Conclusions. TRIM59 plays an oncogenic role in retinoblastoma and exerts its tumor-promotive function by activating the p38–mitogen-activated protein kinase pathway.

Keywords: Retinoblastoma, TRIM59, p38–MAPK signaling pathway, biomarker
METHODS

Data Sources

Three microarray, gene expression datasets (GSE24673, GSE97508, and GSE110811) were retrieved from the Gene Expression Omnibus (GEO) database (https://www.ncbi.nlm.nih.gov/geo/). Among these datasets, GSE24673 was based on the GPL244 platform ([HuGene-1.0-st] Affymetrix Human Gene 1.0 ST Array); GSE97508 was based on the GPL15207 platform ([Affymetrix Human Gene Expression Array], and GSE110811 was based on the GPL16868 platform ([Affymetrix Human Gene 2.0 ST Array]). Nine retinoblastoma samples (GSM607948–GSM607946) and two normal healthy adult retinas (GSM607947 and GSM607948) were obtained from the GSE24673 dataset. Six retinoblastoma samples (GSM2570469–GSM2570474) and three control tissues (GSM2570466–GSM2570468) were obtained from the GSE97508 dataset. A total of 28 retinoblastoma samples (GSM3017123–GSM3017150) and three control tissues (GSM3017151–GSM3017153) were obtained from GSE110811.

After careful review, all samples in the GSE24673 dataset were considered to be satisfactory; however, three retinoblastoma samples (GSM2570472, GSM2570473, and GSM2570474) were eliminated from GSE97508, and one normal sample (GSM3017153) was removed from GSE110811 due to low sample quality.

Bioinformatic Analysis

Original .CEL files were analyzed and underwent background correction, quality control, and data standardization. Affy packages were used for the GSE24673 and GSE97508 datasets, and Oligo packages were used for the GSE110811 dataset. Principal component analysis was analyzed using the built-in princomp function in R 3.6.3 (R Foundation for Statistical Computing, Vienna, Austria), and plots were generated using the scatterplot3d package. Differentially expressed genes (DEGs) were screened using the limma package in R. The cutoff criteria for statistical significance were |logFC| (an absolute log2 value in the fold change of the expression of the genes) > 1 and P < 0.05. Screened DEGs were used to create a heatmap using the built-in heatmap function and a Venn diagram using the VennDiagram package in R.

Cell Culture

The human retinoblastoma cell lines Weri-Rb1, Y79, and HXO-Rb44 and the normal retinal epithelial cell line ARPE-19 were purchased from ATCC (Rockville, MD, USA). All of the cells were cultured in Dulbecco's Modified Eagle Medium/Nutrient Mixture F-12 (DMEM/F-12; Thermo Fisher Scientific, Waltham, MA, USA) supplemented with penicillin G, streptomycin, and 10% fetal bovine serum (Thermo Fisher Scientific). The cells were incubated in a humidified atmosphere with 5% CO2 at 37°C.

Lentiviral Transduction

For downregulation of TRIM59 expression, three human siRNA sequences (RNa1, GGAAGCTGTTCCTCAGATAT [small interfering RNA of TRIM59-1, siTRIM59-1]; RNa2, GAAGAGTTCCTCAGTAAT [siTRIM59-2]; and RNa3, GAATGGACGACAGAGAA [siTRIM59-3]) synthesized by Geneway Company (Shanghai, China) were cloned into the pLKO.1 plasmid (Addgene, Cambridge, MA, USA). The coding DNA sequence region of TRIM59 (NM_173084.2) was cotransfected with the viral packaging plasmids pSPAX2 and pMD2.G (Addgene) into 293T cells with Lipofectamine 2000 Transfection Reagent (Thermo Fisher Scientific). Forty-eight hours after infection, the lentiviral particles in the supernatants were collected. The HXO-Rb44 cell line was infected with the siRNA viral supernatants, and the Y79 cell line was infected with the overexpression viral supernatants. TRIM59 expression was examined by reverse-transcription quantitative polymerase chain reaction (qRT-PCR). TRIM59-overexpressing Y79 cells were treated in the absence or presence of the p38 inhibitor SB203580 (Cell Signaling Technology, Danvers, MA, USA) for 24 hours in subsequent experiments.

qRT-PCR

Total RNA was extracted from cells using TRIzol reagent (Thermo Fisher Scientific) according to the manufacturer's protocol. The extracted RNA was quantitated by spectrophotometry at 260 nm using a NanoDrop ND-1000 Spectrophotometer (NanoDrop Technologies, Wilmington, DE, USA). First-strand cDNA was synthesized using the Quantitect Reverse Transcription Kit (Qiagen, Hilden, Germany). Later, RT-PCR with cDNA as the template was performed on the ABI PRISM 7000 Sequence Detection System (Thermo Fisher Scientific) using the Quantifast SYBR Green PCR Kit (Qiagen). The thermal cycling conditions were 95°C for 10 minutes followed by an initial denaturation step at 95°C for 15 seconds, 40 cycles at 60°C for 45 seconds, and 60°C for 30 seconds. The experiments were carried out in triplicate for each data point. The primers for TRIM59 and β-actin were as follows: TRIM59 forward, 5'-TTGTCACCTGCCCCTGAAC-3'; TRIM59 reverse, 5'TCTTTATGCGCTTGGATC-3'; β-actin forward, 5'-TTGGATCCAGAAGTACTG-3'; and β-actin reverse, 5'-CTTGATCTTCATGGTGCTG-3'. Quantifications were normalized using β-actin as an internal reference and calculated by using the 2^-ΔΔCt method.

Immunoblotting

Cell lysates were prepared from cell lines using a RIPA Lysis and Extraction Buffer kit (Thermo Fisher Scientific), and the procedure was performed as described previously.22 Primary antibodies against target proteins, such as TRIM59, Bcl-2, cleaved caspase-3 (Abcam, Cambridge, UK), cyclin D1, phosphorylated p38 (p-p38), p38, p-JNK1/2, JNK1/2, p-ERK1/2, ERK1/2, p-c-JUN, c-JUN, and β-actin (Cell Signaling Technology) were diluted between 1:500 and 1:2000, and secondary antibodies (Beyotime Biotechnology, Shanghai, China) were diluted 1:1000.

Cell Proliferation

A Cell Counting Kit-8 (CCK-8; Dojindo Molecular Technologies, Rockville, MD, USA) assay was applied to assess the
Figure 1. Bioinformatic analysis of TRIM59 expression in retinoblastoma samples from the GEO database. Violin plot of the number of genes in retinoblastoma patient and normal control samples from the GSE24673 (A), GSE97508 (B), and GSE110811 (C) datasets. \textit{Black dots} represent mean values of gene expression after sample normalization. Volcano plot of the distribution of DEGs in the GSE24673 (D), GSE97508 (E), and GSE110811 (F) datasets, mapping upregulated genes (\textit{red dots}) and downregulated genes (\textit{blue dots}). Genes showing no significant changes are marked as \textit{gray dots}. Heatmap using hierarchical clustering of DEGs in the GSE24673 (G), GSE97508 (H), and GSE110811 (I) datasets. J) Venn diagram DEGs were selected with a fold change $> 2$ and $P < 0.01$ in the GSE24673, GSE97508, and GSE110811 datasets. All three datasets showed overlap of 243 genes. Scatterplot of TRIM59 expression among retinoblastoma patient samples and normal control samples in the GSE24673 (K), GSE97508 (L), and GSE110811 (M) datasets.
Figure 2. TRIM59 expression is upregulated in three retinoblastoma cell lines. Evaluation of mRNA expression using qRT-PCR (A) and protein expression using immunoblotting (B, C) of TRIM59 in the three human retinoblastoma cell lines, Weri-Rb1, Y79, and HXO-Rb44, as well as the normal retinal epithelial cell line ARPE-19. **P < 0.05, ***P < 0.01 compared with ARPE-19 cells. Confirmation of knockdown performed by siRNAs targeting TRIM59 in HXO-Rb44 cells at the mRNA (D) and protein (E, F) level. **P < 0.01 compared with the siNC group. Validation of upregulated mRNA (G) and protein expression (H, I) of TRIM59 in lentivirus transfected Y79 cells. **P < 0.01 compared with vector-transfected cells.

cell proliferation rate. Approximately 3 \times 10^3 HXO-Rb44 or Y79 cells were seeded in a 96-well plate and transfected with siRNA negative control (siNC), siTRIM59, vector, overexpression (oe)TRIM59, vehicle + vector, vehicle + oeTRIM59, SB203580 + vector, or SB203580 + oeTRIM59. At 0, 12, 24, and 72 hours after transfection, CCK-8 reagent (10 μL) was added to each well and incubated for 2 hours at 37°C. The absorbance at 450 nm was measured using an ELISA microplate reader (Perlong, Beijing, China).

Cell-Cycle Analysis
HXO-Rb44 and Y79 cells were harvested and washed with ice-cold PBS. After fixation with 70% ethanol overnight at 4°C, the cells were washed twice with PBS and then treated with 1 mg/mL RNase A in 500 mL of PBS at 37°C for 30 minutes. Next, the cells were stained with 50 μg/mL propidium iodide (PI; Thermo Fisher Scientific) in the dark for 30 minutes. Flow cytometric analysis was performed using a FACScan flow cytometer (BD Biosciences, San Jose, CA, USA), and the results were analyzed using FlowJo 7.6 (FlowJo LLC, San Diego, CA, USA).

Apoptosis Assays
Cellular apoptosis was evaluated using an Annexin V-FITC Apoptosis Detection Kit (Beyotime Biotechnology). Cells were harvested using trypsin/EDTA and washed with PBS. Binding buffer was subsequently added to resuspend the cells. Next, 5 μL of the annexin V-FITC and PI solution were added, and samples were incubated for 15 minutes at room temperature. Finally, stained cells were analyzed using a CytoFLEX Flow Cytometer (Beckman Coulter, Indianapolis, IN, USA). Data were analyzed using the FlowJo 7.6 software. Apoptotic cells were defined by positive annexin V staining and negative PI staining. Normal, living cells were not stained by either annexin V-FITC or PI (cells in the third quadrant); early apoptotic cells were only stained by annexin V-FITC and were stained negative by PI (cells in the second quadrant); late apoptotic cells were simultaneously stained by both annexin V-FITC and PI (cells in the first quadrant); and necrotic cells were stained only by PI (cells in the fourth quadrant).

In Vivo Tumorigenicity Assay
Twelve 6-week-old male BALB/c nude mice weighing 18 to 20 g were obtained from Shanghai Experimental
Animal Center (Shanghai, China) and bred in a specific pathogen-free animal facility. The animal protocol adhered to the ARVO Statement for the Use of Animals in Ophthalmic and Vision Research and was approved by the Ethics Committee of the Second Affiliated Hospital of Nanchang University. Prior to injection, the mice were randomly divided into siTRIM59 and siNC groups (n = 6 for each group). A total of 5 × 10^5 HXO-Rb44 cells were subcutaneously injected into the left upper flank region. Tumor length and width were measured every three days. Tumor size was calculated as follows: volume (m^3) = (length × width^2) × 0.5. All mice were sacrificed 33 days later, and the tumor nodules were removed and weighed, and immunohistochemistry was then performed. The expression of TRIM59, cyclin D1, Bcl-2, and cleaved caspase-3 was determined by immunoblotting.

**Immunohistochemistry**

Paraffin-embedded tumor samples were sectioned into 4- to 7-μm-thick sections. Subsequent immunohistochemistry was performed according to the protocol in a previous report. An anti-TRIM59 primary antibody (Abcam) was used at a 1:600 dilution, and a secondary antibody (Abcam) was used at a 1:400 dilution. The slides were then stained with 3,3'-diaminobenzidine and counterstained with hematoxylin (Baso Diagnostics, Inc., Zhuhai, China) following the manufacturer's protocol. Immunohistochemical images were acquired using a Eclipse Ni microscope (Nikon Instruments, Inc., Melville, NY, USA). The sizes of positive areas were calculated using Nikon NIS-Elements, version 5.11.00.

**Statistical Analysis**

SPSS Statistics 23.0 (IBM, Chicago, IL, USA) was used for statistical analyses. In vitro experiments were performed in triplicate, and in vivo tests were repeated six times. Data are presented as the mean ± SD. Differences among multiple groups were analyzed by one-way ANOVA with Tukey’s post hoc test, and the means of two groups were compared by Student’s t-test. Two-way, repeated-measures ANOVA was used to analyze CCK-8 assay results. P < 0.05 was considered to indicate a statistically significant difference.

**RESULTS**

**Analysis of the GEO Database**

The mean values of gene expression for each sample in the GSE24673, GSE97508, and GSE110811 datasets were fundamentally the same after normalization, suggesting that the sample data sources were reliable and consistent for further downstream analyses (Figs. 1A–1C; Supplementary Fig. S1). After standardizing microarray results, 2758 DEGs were identified. Among these, 386 upregulated genes and 770 downregulated genes were screened in the GSE24673 dataset (Fig. 1D; Supplementary Table S1), 894 upregulated genes and 1007 downregulated genes were screened in the
FIGURE 4. Effects of TRIM59 on the cell-cycle distribution. (A, B) G1 phase arrest in HXO-Rb44 cells induced by TRIM59 expression downregulation was detected by flow cytometry. **P < 0.01 compared with the siNC group. (C, D) TRIM59 significantly promoted the G1/S phase transition in Y79 cells **P < 0.01 compared with vector-transfected cells. The cell-cycle-related protein cyclin D1 was detected in HXO-Rb44 cells (E, F) and Y79 cells (G, H) by immunoblotting. **P < 0.01 compared with the siNC and vector groups, respectively.

GSE97508 dataset (Fig. 1E; Supplementary Table S2), and 229 upregulated genes and 580 downregulated genes were screened in the GSE110811 dataset (Fig. 1F; Supplementary Table S3). The relative expressions of DEGs in each sample showed significant differences (Figs. 1G–1I).

Furthermore, a Venn diagram revealed 108 overlapping upregulated genes in the GSE24673, GSE97508, and GSE110811 datasets (Fig. 1J). Compared with the normal control samples, the expression levels of TRIM59 in retinoblastoma samples were significantly increased in all three datasets (Figs. 1K–1M).

Upregulation of TRIM59 Expression in Retinoblastoma Cells

The expression levels of TRIM59 in the human retinoblastoma cell lines HXO-Rb44, Y79, and Weri-Rb1 were significantly higher than that in ARPE-19 cells (Fig. 2A). Meanwhile, the TRIM59 expression rate was highest in HXO-Rb44 cells and lowest in Y79 cells. Moreover, TRIM59 upregulation was further confirmed using immunoblotting, and the trend was consistent with the qRT-PCR data (Figs. 2B, 2C). To explore the biological function of TRIM59 in retinoblastoma cells, we used specific siRNAs to knock down the expression of TRIM59 in HXO-Rb44 cells, which had relatively high expression of TRIM59. In addition, Y79 cells with relatively low TRIM59 expression were used to generate a cell line with stable TRIM59 overexpression by infection with a lentivirus. As shown in Figures 2D to 2F, the mRNA and protein levels of TRIM59 in HXO-Rb44 cells were both significantly lower than those in control cells. Additionally, upregulated expression of TRIM59 in Y79 cells was observed by qRT-PCR (Fig. 2G) and immunoblotting (Fig. 2H, 2I).

TRIM59 Accelerates the Cell Proliferation of Human Retinoblastoma Cells

CCK-8 assays revealed that knocking down TRIM59 expression significantly suppressed cell proliferation in HXO-Rb44 cells compared with corresponding control cells (incubation time of 24 hours, P < 0.05; incubation times of 48 and 72 hours, P < 0.01) (Fig. 3A). There was no significant
TRIM59 Promotes Retinoblastoma Cell-Cycle Progression

Flow cytometry analysis revealed that TRIM59 silencing significantly increased the proportion of cells in the G1 phase but reduced the proportion of cells in the S phase in HXO-Rb44 cells (Figs. 4A, 4B). In contrast, the opposite effects were observed in Y79 cells with TRIM59 overexpression (Figs. 4C, 4D). Furthermore, we examined the expression level of cyclin D1, a key cell-cycle-regulating protein, and found that the cyclin D1 expression levels in TRIM59-knockdown HXO-Rb44 cells were lower than those in control cells (Figs. 4E, 4F). Additionally, TRIM59-overexpressing cells had higher expression levels of the cyclin D1 protein than vector-transfected cells (Figs. 4G, 4H). All of these results suggest a positive correlation between the TRIM59 expression level and cell-cycle progression in retinoblastoma.

Downregulation of TRIM59 Expression Induces Retinoblastoma Cell Apoptosis

Compared with transfection of a negative control siRNA, TRIM59 depletion increased the percentage of apoptotic HXO-Rb44 cells (Figs. 5A, 5B). In contrast, compared with vector-transfected cells, overexpression of TRIM59 produced a remarkable decrease in the apoptotic cell percentage in Y79 cells (Figs. 5C, 5D). In addition, immunoblotting was used to detect the expression levels of two apoptosis-related biomarkers: Bcl-2 and cleaved caspase-3. As shown in Figures 5E and 5F, the results demonstrate that the expres-
TRIM59 Promotes RB Progression by p38 Pathway

TRIM59 promotes retinoblastoma tumor growth in a xenograft model. (A, B) Representative images of subcutaneous tumors in mice receiving siTRIM59 or the negative control (siNC). (C) Quantification of tumor weight. *P < 0.05 compared with the siNC group. (D) Tumor growth curves comparing the group of mice receiving siTRIM59 with the control group. *P < 0.05 compared with the siNC group. (E) TRIM59 knockdown confirmed by immunohistochemical staining. Magnification, 200×. (F, G) Immunoblotting results indicating that, compared with control treatment, silencing TRIM59 led to reduced expression of TRIM59, cyclin D1, and Bcl-2 and increased expression of cleaved caspase-3. **P < 0.01 compared with the siNC group.

Silencing TRIM59 Suppresses the Tumorigenesis of Retinoblastoma Cells In Vivo

As shown in Figure 6A, all nude mice developed xenograft tumors at the injection site. However, tumor volume and weight were significantly lower in the mice injected with TRIM59-silenced cells than in those in the control group by approximately 69.96% and 63.73%, respectively (Figs. 6B–6D). TRIM59 knockdown was confirmed by immunohistochemical staining (Fig. 6E). Moreover, we also measured the protein levels of TRIM59, cyclin D1, Bcl-2, and cleaved caspase-3 and found that TRIM59-knockdown xenograft tumors had relatively low expression of TRIM59, cyclin D1, and Bcl-2 and relatively high expression of cleaved caspase-3 (Figs. 6F, 6G). Therefore, these results suggest that silencing TRIM59 can attenuate tumor growth and promote cell apoptosis.

TRIM59 Regulates Retinoblastoma Cells Via the p38–Mitogen-Activated Protein Kinase Pathway

To elucidate the mechanism underlying the effects of TRIM59 on retinoblastoma cells, we evaluated the
TRIM59 Promotes RB Progression by p38 Pathway

**DISCUSSION**

Retinoblastoma is the most common intraocular neoplasm and a serious threat to vision and life in infants and young children. Although multidisciplinary treatments that include surgery, chemotherapy, and radiotherapy are currently available, the mortality rate of retinoblastoma patients is still approximately 70%, especially in the low- and middle-income countries where most affected children live. Even if patients receive enucleation or orbital exenteration, intracranial invasion, metastasis, or second nonocular neoplasm can lead to death. Therefore, investigating the molecular mechanism and promising biomarkers of intraocular retinoblastoma progression should be beneficial for early diagnosis and effective therapeutic target exploration.

Bioinformatic analysis is a well-established method that is widely used to help researchers identify potential biomarkers. Thus, we performed bioinformatic analysis of the GEO database to reveal that TRIM59 was one of the most upregulated genes in retinoblastoma samples, which was consistent among the three datasets analyzed. Through qRT-PCR and immunoblotting, we verified that the level of TRIM59 in retinoblastoma cells differed significantly from that in control cells. Previous studies have shown that TRIM59 appears to be an oncoprotein involved in various types of cancer. Elevated TRIM59 expression is correlated with a relatively poor prognosis and progression in breast cancer. Knocking down TRIM59 expression remarkably inhibits cell proliferation in neuroblastoma, hepatocellular carcinoma, and cholangiocarcinoma. However, to date, the role of TRIM59 in retinoblastoma has not been reported. Therefore, TRIM59 was selected for the present research. We have identified TRIM59 as a promising prognostic biomarker or therapeutic target in retinoblastoma. Moreover, gain- and loss-of-TRIM59 experiments with retinoblastoma cell lines suggested that TRIM59 promoted cell proliferation and growth and suppressed cell apoptosis. An in vivo assay further confirmed the tumorigenic function of TRIM59. Our findings suggest that TRIM59 plays an oncogenic role in the progression of retinoblastoma, which is consistent with the results previously observed in other cancers. According to the literature, TRIM59 can regulate colorectal cancer through the Akt pathway, regulate neuroblastoma through the Wnt pathway, and regulate bladder cancer through the TGF-β pathway. The intrinsic mechanism of TRIM59 involved in retinoblastoma needed to be identified.

The p38–MAPK signaling pathway has been implicated in the inflammatory response, cell cycle regulation, cell death, differentiation, and tumor progression. Goldsmith et al. reported that inhibition of the p38–MAPK pathway led to decreased glioma cell invasion. Another study found that miR-3188 promoted cell proliferation and migra-
FIGURE 8. p38 inhibition alleviates TRIM59-mediated retinoblastoma progression. The p38-specific inhibitor SB203580 effectively suppressed cell proliferation (A, B) and the G1/S phase transition (C, D) and promoted cell apoptosis (E, F) in TRIM59-overexpressing Y79 cells using the methods described above. *P < 0.01, "SB203580 + oeTRIM59" group compared with "vehicle + oeTRIM59" group; ††P < 0.01, "SB203580 + vector" group compared with "vehicle + vector" group. (G, H) Immunoblotting was performed to evaluate the expression of p-p38, cyclin D1, Bcl-2, and cleaved caspase-3. *P < 0.05, †P < 0.01, "SB203580 + oeTRIM59" group compared with "vehicle + oeTRIM59" group; ††P < 0.01, "SB203580 + vector" group compared with "vehicle + vector" group.
tion via the p38–MAPK pathway in breast cancer. Furthermore, miRNA-655 regulates retinoblastoma cell proliferation, invasion, and apoptosis via the p38–MAPK pathway. Our results showed that downregulation of TRIM59 expression significantly decreased the expression of the p38–MAPK pathway components p-p38, p-JNK1/2, p-ERK1/2, and p-c-JUN. The opposite results were observed in TRIM59-overexpressing Y79 cells. Moreover, inhibition of p38 with SB203580 abrogated the effects of TRIM59 on cell proliferation, apoptosis, and the G1/S phase transition. These results suggest involvement of the p38–MAPK signaling pathway in the functions of TRIM59 in retinoblastoma (Fig. 9).

These findings may be somewhat limited by the rarity of retinoblastoma patients, and we have not examined the level of TRIM59 in the metastatic samples. We also did not analyze the correlation between TRIM59 expression and survival in retinoblastoma patients. Therefore, the clinical significance of TRIM59 is worthy of further investigation.

In summary, to the best of our knowledge, our study shows for the first time that upregulated TRIM59 expression exerts a crucial role in the progression of retinoblastoma both in vitro and in vivo by activating the p38–MAPK pathway. Our findings offer insights into manipulating TRIM59 as a novel biomarker and target gene for retinoblastoma early diagnosis and treatment.

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Figure 9. Schematic diagram indicates the proposed role of TRIM59 in retinoblastoma. Upregulation of TRIM59 promotes retinoblastoma cell proliferation and cell-cycle transition and suppresses cell apoptosis via p38–MAPK pathway activation.


