Abstract. Photodynamic therapy (PdT) is a relatively novel type of tumor therapy method with low toxicity and limited side-effects. The aim of the present study was to investigate the underlying mechanism and potential microRNAs (miRNAs) involved in the treatment of glioma by PdT with hematoporphyrin, a clinical photosensitizer. The photodynamic activity of hematoporphyrin on the cell viability and apoptosis of gliomas was investigated by MTT, and flow cytometry and fluorescence microscopy, respectively. Alterations in singlet oxygen and mitochondrial membrane potential were detected. The differentially expressed miRNAs and proteins were evaluated by miRNA gene chip and apoptosis-associated protein chip, respectively. The results demonstrated that cell viability significantly decreased with hematoporphyrin concentration. PDT with hematoporphyrin significantly increased cell apoptosis at a later stage, induced the content of reactive oxygen species (ROS) and decreased the mitochondrial membrane potential, indicating that PdT with hematoporphyrin inhibited cell growth via induction of radical oxygen, decreased the mitochondrial membrane potential and induced apoptosis. The upregulated miRNAs, including hsa-miR-7641, hsa-miR-9500, hsa-miR-4459, hsa-miR-21-5p, hsa-miR-663a and hsa-miR-205-5p may be important in PdT-induced cell apoptosis in glioma. Transporter 1, ATP binding cassette subfamily B member- and nuclear factor-κB-mediated apoptosis signaling pathways were the most significant pathways. Thus, the current study presents PdT as a potential therapeutic approach for the treatment of malignant glioma, and identified miRNAs for the molecular design and development of a third-generation photosensitizer (PS).

Introduction
Glioma is the most common type of malignant tumor of the central nervous system, accounting for ~46% of brain tumors (1,2). Glioma is a neuroectodermal tumor with the highest mortality rate of primary intracranial tumors (3). Currently, craniotomy is the most efficient type of therapy to treat gliomas (4). However, due to the invasive growth of gliomas and its unclear demarcation with normal brain tissues, it is difficult to completely remove tumor tissues by surgery, resulting in a high recurrence rate (5,6), and subsequent treatment with radiotherapy and chemotherapy (which damage the immune system) are generally not satisfactory (7). Methods to decrease the high recurrence rates and serious side-effects of glioma treatment are being investigated to develop specific drugs or therapeutic strategies with high efficiency and sensitivity, and low toxicity for the treatment of gliomas (8).

Photodynamic therapy (PDT) is a novel therapeutic method for tumor treatment (9). PDT was first discovered by Dougherty et al (10) in 1900, and was first used in medicine to ablate and destroy unwanted tissue in 1975 (10). PDT utilizes the activation of a photosensitizer (PS) by light irradiation with a specific wavelength to generate oxygen radicals and further inhibit growth of tumor cells (11,12). The advantages of PDT are its minimally invasive characteristic, repeatability, low toxicity accumulation when compared with the traditional treatments, in addition to its low recurrence rate and more definitive outcome (9). Low toxicity to normal tissue is a significant advantage of PDT, compared with traditional treatment. In recent years, PDT has shown a particularly good therapeutic effect on bladder cancer, early lung cancer, Barrett's esophagus, head and neck cancer and skin cancer, amongst others (13-15). For example, when using 2-(1-hexyloxethyl)-2-devinyl pyropheophorbide-a as a PS in...
the clinical setting, PDT demonstrated a particularly effective outcome in 16 cases of patients with lung cancer in a treatment cycle, with a very slight photosensitive side-effect; after 3 days of PDT treatment, patients with oral cancer recovered (16). Furthermore, PDT treatment for esophageal cancer and Barrett's esophagus resulted in a good outcome (17). However, fundamental difficulties encountered in the treatment of PDT, such as targeting, light source and light dose have not been addressed. Advances in chemistry, materials, optics, clinical treatment and other disciplines contribute to overcoming these difficulties, by elucidating the underlying mechanism of PDT.

Typically, tumor treatment methods involve enhancing the antitumor immune response of the body via immunology. In recent years, PDT was identified as able to inhibit the growth of tumor cells and regulate the conditional body immune system; thus, has gradually been adopted as an auxiliary treatment to reduce the recurrence rate of tumors (18). In previous studies, PDT was identified as a promising treatment strategy for glioma (19-22). Using 5-aminolevulinic acid (5-ALA) as a PS, the expression level of proteasomes was increased in cultured U251 cells and the growth of glioma cells was inhibited under light irradiation (19). Using hematoporphyrin as a PS, PDT induced the transscriptional level of transporter 1, ATP binding cassette subfamily B member (TAPI) and the expression level of HLA-1 in U251 cells under light irradiation, inhibiting the growth of glioma cells in a dose- and time-dependent manner (20). In addition, PDT enhanced the immunogenicity of glioma cells, indicating that a PDT-generated glioma vaccine from dendritic cells presents as a promising novel method for tumor treatment (21). However, the molecular mechanisms underlying PDT-inhibited growth of glioma cells remain unclear. In the current study, based on previous studies, the role of PDT in the induction of cell apoptosis of glioma cells was investigated using hematoporphyrin as the PS, as well as microRNAs (miRNAs) and their corresponding signaling pathways, such as target molecule networks. The current results provide a potential therapeutic strategy involving PDT for the treatment of gliomas.

Materials and methods

Cell culture. Human glioma cell lines U87 and U251 cells were obtained from the American Type Culture Collection (ATCC; Manassas, VA, USA). Cells were cultured in 5% CO2 and 95% humidified atmosphere in DM 1640 medium (Invitrogen, Thermo Fisher Scientific, Inc., Waltham, MA, USA) containing 10% fetal bovine serum (Invitrogen; Thermo Fisher Scientific Inc.) with 100 U/ml penicillin and 100 µg/ml streptomycin. Cells were passaged subsequent to reaching 80% confluence.

MTT assay. Cell viability was determined using MTT. Briefly, 5x10^4 cells per well were seeded in 96-well plates. Following incubation for 12 h at 37°C, the cells were treated with phosphate-buffered saline (PBS; control) or various concentrations of hematoporphyrin (0-120 nM; MedChemExpress, Princeton, NJ, USA) for 60 min. Following light irradiation using an M8 Spectrum Power Energy meter (San Diego, Taizhong, Taiwan, China) at 625 nm, 5 mW/cm² for 60 min (23) and culturing for 22 h at 37°C, 100 µl MTT solution (5.0 mg/ml) was then added to each well and incubated for 4 h at 37°C, followed by removal of culture medium and the addition of dimethyl sulfoxide (100 µl/well) to dissolve the formed formazan crystals. The absorbance value was determined using a microplate reader (Varioskan Flash, Thermo, USA) at a wavelength of 520 nm. The relative cell viability was calculated by normalization to the control. The experiments were independently performed three times.

Cell apoptosis assays. Cell apoptosis was detected using Hoechst 33342 and flow cytometry. Briefly, 2x10^5 cells/well were seeded in 6-well plates. Subsequent to incubation for 12 h at 37°C, the cells were treated with hematoporphyrin (85 nM, the IC50) under light irradiation (625 nm, 5 mW/cm²) for 60 min. The untreated cells served as the control. Cells were subsequently cultured for 23 h at 37°C, then collected with trypsin-ethylenediaminetetraacetic acid (EDTA) and resuspended in PBS. Cell apoptosis was measured by flow cytometry (BD FACSCalibur; BD Biosciences, Franklin Lakes, NJ, USA) using the Annexin V-fluorescein isothiocyanate (Beyotime Institute of Biotechnology, Haimen, China) according to the manufacturer's instructions. The flow cytometry data was analyzed using FlowJo version 10.0 software (FlowJo LLC, Ashland, OR, USA). For the Hoechst 33342 assay, 10 µl Hoechst 33342 (5 µg/ml) was added to each well and incubated for 30 min at 37°C in the dark, followed by removal of the culture medium and two washes with PBS. The apoptotic nuclei were detected by inverted fluorescence microscopy (SR GSD; Leica Microsystems, Inc., Buffalo Grove, IL, USA).

TUNEL assay. Cells (2x10^5/well) were seeded in 12-well plates. After adherence, cells were treated with hematoporphyrin (85 nM) under light irradiation (625 nm, 5 mW/cm²) for 60 min. The untreated cells served as the control. Subsequently, cells were cultured for 23 h at 37°C, and fixed with 4% paraformaldehyde for 30 min, washed with PBS, incubated with 0.3% Triton X-100 (Solarbio, Shanghai, China) in PBS for 5 min, and incubated with 0.3% hydrogen peroxide in PBS for 20 min at 37°C to inactivate endogenous peroxidase. Cells were then incubated in TUNEL detection solution (50 µl/well; Beyotime Institute of Biotechnology) for 60 min at 37°C and 50 µl streptavidin-horseradish peroxidase solution for 30 min. Cells were visualized by 3,3'-diaminobenzidine tetrahydrochloride (DAB) solution (Solarbio). The nucleus was stained with hematoxylin in 20 min at 37°C.

Singlet oxygen detection. Alterations in singlet oxygen were detected using 2'-7'-dichlorodihydrofluorescein diacetate (DCFH-DA; Beyotime Institute of Biotechnology) according to the manufacturer's instruction. Briefly, 2x10^5 cells per well were seeded in 6-well plates. After incubation for 12 h at 37°C, the cells were treated with hematoporphyrin (85 nM, the IC50) under light irradiation (625 nm, 5 mW/cm²) for 60 min. The untreated cells served as the control. Subsequently, cells were cultured for 6 h and incubated with 10 µM DCFH-DA for 30 min in the dark. The cells were then washed with PBS-free Dulbecco's modified Eagle's medium (Life, Thermo Fisher Scientific, Inc., Waltham, MA, USA), collected with trypsin-EDTA and resuspended in PBS. The absorbance values were detected using a microplate reader (using an
emission wavelength of 488 nm and an excitation wavelength of 525 nm. The experiments were independently performed three times.

Mitochondrial membrane potential analysis. Alterations in mitochondrial membrane potential were detected using JC-1 kit (C2005; Beyotime Institute of Biotechnology, Beijing, China) according to the manufacturer’s instructions. Briefly, 2x10^5 cells per well were seeded in 6-well plates. Following incubation for 12 h at 37˚C, the cells were treated with hematoporphyrin (85 nM, the IC_{50} under light irradiation (625 nm, 5 mW/cm^2) for 60 min. The untreated cells served as the control. Cells were incubated with 1 ml work solution of JC-1 for 20 min at 37˚C in the dark, washed three times with JC-1 buffer, collected with trypsin-EDTA, resuspended in PBS and detected by inverted fluorescence microscopy. JC-1 dye exhibits potential-dependent accumulation in mitochondria, which is indicated by a fluorescence emission shift from green (~529 nm) to red (~590 nm). Consequently, mitochondrial depolarization is indicated by a decrease in the red/green fluorescence intensity ratio. Carboxyl cyanide m-chlorophenylhydrazone (CCCP) served as positive controls (in the JC-1 kit, Beyotime Institute of Biotechnology). The ratios of the absorbance values at 525 and 590 nm were determined with a microplate reader. The experiments were independently performed three times.

Mature miRNA microarray analysis. The four groups, which included untreated glioma cells (in the dark; n=3; named Control-dark group), glioma cells treated with hematoporphyrin and without light irradiation (n=3; named Hep-dark group), glioma cells that underwent light irradiation (n=3 named Control-light group), and glioma cells treated with hematoporphyrin and light irradiation (n=3; named Hep-light group), were analyzed using Agilent Human miRNA microarray chips (8x60K) V21.0 (ShanghaiBio Corp. (SBC), Shanghai, China). The differentially expressed miRNAs that exhibited a two-fold or greater change were screened.

Reverse transcription-quantitative polymerase chain reaction (RT-qPCR). Glioma cells were treated with or without PDT and their total RNA was separately extracted using an RNA extraction kit (cat. no. DP431; Tiangen Biotech Co., Ltd., Beijing, China) according to the manufacturer’s instructions. The RNA concentration was quantified using an ultraviolet spectrophotometer (NanoDrop 2000; Thermo Fisher Scientific, Inc.) and was reverse transcribed into cDNA using a FastQuant first strand cDNA synthesis kit (cat. no. KR104; Tiangen Biotech co., Ltd., Beijing, China) according to the manufacturer’s instructions. The sequences of specific primers for cDNA synthesis kit (cat. no. FP411; Biotech co., Ltd.) and run for 40 cycles at 95˚C for 20 sec and 60˚C for 34 sec. The relative miRNA levels were normalized to that of U6 using the 2^-ΔΔcq method (24). The experiment was performed three times.

Gene Ontology (GO) and pathway enrichment analyses. The differentially expressed miRNAs were further analyzed for predicted gene targets simultaneously using at least two of the following five databases: TargetMiner, miRDB, microRNA, TarBase and RNA22 via the SBc analysis system (http://sas.ebioservice.com). GO enrichment analyses and Kyoto Encyclopedia of Genes and Genomes (KEGG) pathway enrichment analyses were performed by the SBC analysis system, which uses clusterProfiler data from R and Bioconductor software (http://www.r-project.org and http://www.bioconductor.org/) with public databases that include NcBI Entrez Gene (http://www.ncbi.nlm.nih.gov/gene), GO (http://www.geneontology.org), KEGG (http://www.genome.jp/kegg) and BioCarta (http://www.biocarta.com). The hypergeometric test was employed with a Benjamini-Hochberg-False Discovery Rate-based multiple testing correction (correction, P<0.01) (25).

Protein array detection assay. A Human Apoptosis Antibody Array (cat. no. AAH-APO-1; RayBiotech, Inc., Norcross, GA, USA) was used according to the manufacturer's instructions. Following development, films were scanned using ImageQuant LAS 4000 (GE Healthcare, Chicago, IL, USA) and the images were processed and quantified using the AAH-APO-1 software.

Statistical analysis. Statistical analysis was performed using GraphPad Prism Software (version 6.0; GraphPad Software, Inc., La Jolla, CA, USA) and Student's t-test. Each experiment was performed at least three times and all figures represent means ± standard deviation. P<0.05 was considered to indicate a statistically significant difference.

Results

PDT with PS, hematoporphyrin inhibits cell viability in a dose-dependent manner. The cell viability in U87 and U251 glioma cells following PDT was detected with PS, hematoporphyrin (Fig. 1). In the U87 and U251 cells (Fig. 1A), cell viability was significantly decreased along with the hematoporphyrin concentration. The level of cell viability was reduced to a greater extent in the U87 cells than in the U251
cells (IC$_{50}$: 85 vs. 166 Nm for U87 vs. U251), indicating that the inhibition of PDT in U87 is better than in U251. The viability of cells that irradiated with white or red light in the U87 cells was compared (Fig. 1B) and the results demonstrated that the inhibition of hematoporphyrin under red light was better than under white light. Thus, red light-irradiated U87 cells were used in the following experiments.

Photodynamic activity of hematoporphyrin (85 nM) induced changes in cell morphology in the U87 cells (Fig. 1C and D). The U87 cells demonstrated morphological changes following PDT-hematoporphyrin treatment, including shrinking, fragmentation, which was not observed in cells without hematoporphyrin treatment (in the light or dark; Fig. 1C and D).

**PDT with PS, hematoporphyrin induces cell apoptosis.** Cell apoptosis in U87 cells following PDT with PS, hematoporphyrin (85 nM) were detected by Hoechst 33342 and flow cytometry (Fig. 2). As presented in Fig. 2A, photodynamic activity of hematoporphyrin induced apoptotic nuclei in U87 cells with low cell density. Similarly, the flow cytometric analysis revealed that photodynamic activity of hematoporphyrin increased cell apoptosis in early (6.1 vs. 5.91% for PDT vs. control) and later (26.35 vs. 0.44% for PDT vs. control; P<0.05) stages (Fig. 2B). Similar to the Hoechst 33342 and flow cytometry data, the TUNEL assay indicated that the photodynamic activity of hematoporphyrin increased cell apoptosis (Fig. 1D). Thus, photodynamic activity of hematoporphyrin inhibited cell growth via induction of cell apoptosis.

**PDT with PS, hematoporphyrin induces cell apoptosis via induction of ROS.** As previous studies demonstrated that, under light irradiation, hematoporphyrin exerts its action via production of singlet oxygen and superoxide anions (26,27), the current study investigated the alterations in singlet oxygen and mitochondrial membrane potential in cells following PDT with PS, hematoporphyrin treatment (Fig. 3). In healthy cells, the mitochondrial membrane potential was not markedly changed, while obvious green fluorescence was observed in the U87 cells following PDT treatment (Fig. 3A). A similar result was obtained using a microplate reader; the mitochondrial membrane potential was significantly decreased following PDT treatment, which was lower than the CCCP group (Fig. 3B). The content of reactive oxygen species (ROS) was significantly increased by PDT, which was greater than that in the Rosup group (Fig. 3C). Thus, PDT with hematoporphyrin may result in the induction of ROS and the decrease in mitochondrial membrane potential, which are important in cell apoptosis and mitochondrial disorders (28).

**Screening and validation of miRNA involved in PDT with PS, hematoporphyrin-induced cell apoptosis.** The potential target miRNA was analyzed by miRNA microarray, and the differentially expressed miRNAs between Hep-light (cells with hematoporphyrin and light irradiation) and Control-dark (cells without treatment), Hep-dark (cells with hematoporphyrin) or Hep-light (cells with light irradiation) groups were screened. There were 185 miRNAs differentially expressed in the PDT treatment groups compared with the other groups.
Figure 2. Effect of photodynamic therapy with photosensitizer, hematoporphyrin in cell apoptosis of U87 cells. (A) Effect of photodynamic activity of hematoporphyrin on apoptotic nuclei in U87 cells; magnification, x200. (B) Effect of photodynamic activity of hematoporphyrin on U87 cell apoptosis detected by flow cytometry and analysis by FlowJo software. Hep-light, cells treated with hematoporphyrin and light irradiation; Control-dark, untreated cells; Hep-dark, cells treated with hematoporphyrin; Control-light, cells treated by light irradiation. **P<0.01, significantly different compared with the control by t-test, n=3.

Figure 3. Mitochondrial membrane potential and reactive oxygen species content in U87 cells treated with PDT and hematoporphyrin. (A and B) JC-1 dye with emissions of 590 nm (red) and 525 nm (green). (C) Mitochondrial depolarization is indicated by a decrease in the red/green fluorescence intensity ratio. (D) Singlet oxygen was detected using 2,7'-dichlorodihydrofluorescein diacetate. CCCP and Rosup served as positive controls. Hep-light, cells treated with hematoporphyrin and light irradiation; Control-dark, untreated cells; Hep-dark, cells treated with hematoporphyrin; Control-light, cells treated by light irradiation. *P<0.05, **P<0.01 and ***P>0.01 vs. the control (untreated cells) (Student's t-test; n=3). CCCP, carbonyl cyanide m-chlorophenylhydrazone.
The top six upregulated miRNAs included hsa-miR-7641, hsa-miR-9500, hsa-miR-4459, hsa-miR-21-5p, hsa-miR-663a and hsa-miR-205-5p. The top eight downregulated miRNAs included hsa-miR-1273e, hsa-miR-642a-3p, hsa-miR-6893-5p, hsa-miR-7641, hsa-miR-135b-5p, hsa-miR-210-3p, hsa-miR-30b-5p and hsa-miR-31-5p. The expression levels...
of the six upregulated miRNAs, including hsa-miR-7641, hsa-miR-9500, hsa-miR-4459, hsa-miR-21-5p, hsa-miR-663a and hsa-miR-205-5p were confirmed by RT-qPCR (Fig. 4A). In addition, the expression levels of the top eight downregulated miRNAs were detected by RT-qPCR (Fig. 4B). The expression levels of hsa-miR-1273e, hsa-miR-642a-3p, hsa-miR-6893-5p, hsa-miR-210-3p and hsa-miR-30b-5p were significantly downregulated in the PDT treatment groups, compared with in the other groups.

GO and KEGG enrichment analysis demonstrated that the most significant cellular functions were cell apoptosis, cell necrosis and ROS-mediated functions, and the most significant signaling pathways were the chemokine signaling pathway, osteoclast differentiation and the insulin signaling pathway, indicating that PDT inhibited glioma cell growth via induction of cell apoptosis and necrosis (Fig. 5). In the process, the upregulated miRNAs, including hsa-miR-1273e, hsa-miR-642a-3p, hsa-miR-6893-5p, hsa-miR-7641, hsa-miR-9500, hsa-miR-135b-5p, hsa-miR-205-5p, hsa-miR-21-5p, hsa-miR-210-3p, hsa-miR-30b-5p, hsa-miR-31-5p, hsa-miR-4459 and hsa-miR-663a are significant and represent important targets for therapy of PDT-induced cell apoptosis in glioma.

Detection of cell apoptosis-associated proteins. The proteins associated with cell apoptosis were detected by apoptosis antibody array (Fig. 6). Quantitative analysis revealed that TAPI- and nuclear factor (NF)-xB-mediated apoptosis...
pathways were the most important signaling pathways involved in PDT with hematoporphyrin-induced apoptosis observed in glioma U87 cells (Fig. 6A). Following PDT with hematoporphyrin treatment in U87 cells, the expression levels of NF-κB, TAPI, Bcl-2-associated X protein and tumor necrosis factor-α were significantly upregulated, and the expression level of Bid was significantly downregulated (P<0.05; Fig. 6B).

**Discussion**

The current study examined the possibility of using PDT for the treatment of gliomas, and evaluated the molecular mechanism of PDT using hematoporphyrin as a PS for the inhibition of glioma cell growth by screening target miRNAs and apoptosis-associated proteins. The aim of the study was to provide a theoretical reference for the design and preparation of novel PSs for PDT.

Using the first generation of PS for PDT, hematoporphyrin on U251 and U87 glioma cells, the inhibition of PDT in U87 cells was identified to be better than that in U251 cells. Under the same conditions, the IC₅₀ of hematoporphyrin in the U87 cells was 80 nM, while that in the U251 cells was 120 nM. Physical and chemical factors act on cells together (29), and PS and light source are two key factors of PDT. Therefore, the effect of single wavelength red light and non-single wavelength white light on photodynamic activity of hematoporphyrin were detected by irradiation of U87 cells. The results demonstrated that the inhibition of hematoporphyrin under red light was better than that under white light. Thus, red light-irradiated U87 cells were used to investigate the molecular mechanism.

The inhibition of cell growth by drugs is generally caused by induction of apoptosis or cell cycle arrest (23). The photodynamic activity of hematoporphyrin was found to inhibit cell growth via induction of cell apoptosis at the later stage. This is consistent with the results reported in previous studies (28). For example, the glucose-modified porphyrin derivatives, synthesized by Králová et al (30) under light irradiation significantly inhibited cell growth of MDA-MB-231 cells via induction of cell apoptosis, which was also verified in a mouse model. Thus, the PDT activity of hematoporphyrin on glioma U87 cells appears to be due to the induction of apoptosis.

It was demonstrated that hematoporphyrin under light irradiation exerts its action via production of singlet oxygen and superoxide anions (26,27). The current study found that PDT treatment with hematoporphyrin induced ROS and decreased mitochondrial membrane potential, which contributed to mitochondrial disorder.

Previous studies demonstrate that apoptosis is closely regulated by miRNAs (31,32). The potential target miRNA was analyzed by miRNA microarray and differentially expressed miRNAs, including six upregulated and eight downregulated miRNAs between Hep-light (cells with hematoporphyrin and light irradiation) and Control-dark (cells without treatment), Hep-dark (cells with hematoporphyrin), or Hep-light (cells with light irradiation) groups, were screened. There were 185 miRNAs differentially expressed in the PDT treatment groups when compared with other groups. The expression of six upregulated miRNAs, including hsa-miR-7641, hsa-miR-9500, hsa-miR-4459, hsa-miR-21-5p, hsa-miR-663a, and hsa-miR-205-5p were confirmed by RT-qPCR. It was reported that miR-21 regulated proliferation and apoptosis of human glioma cells by downregulating the phosphatase and tensin homolog protein (33). In a previous study, overexpression of miR-663 inhibited proliferation, migration and invasion in A172 and U87 glioblastoma cells (34). Furthermore, the miR-205 serum level was identified as an individual diagnostic marker for glioma (35). However, the roles of hsa-miR-7641, hsa-miR-9500, hsa-miR-4459, hsa-miR-21-5p, hsa-miR-663a and hsa-miR-205-5p in PDT-induced cell apoptosis have not yet been confirmed using a cell model.

GO and KEGG enrichment analysis identified that the most significant cellular functions were cell apoptosis, cell necrosis and ROS-mediated functions, and the most significant signaling pathways were the chemokine signaling pathway, osteoclast differentiation and the insulin signaling pathway, indicating that PDT inhibited glioma cell growth via induction of cell apoptosis and necrosis. The proteins associated with cell apoptosis were detected by apoptosis antibody array, and quantitative analysis revealed that TAPI and NF-κB-mediated apoptosis signaling pathway was the most important pathway involved in PDT, with hematoporphyrin-induced apoptosis observed in glioma U87 cells. A recent study reported that miR-21 significantly contributes to the inflammatory response via the sirtuin 1-NF-κB signaling pathway (36). It implied that the miRNAs that specifically targeted TAPI and NF-κB are important therapeutic targets (Fig. 7). However, whether changes of Ap1- and NF-κB are associated with those miRNAs remains unclear.

TAPI and NF-KB proteins may be important in the apoptotic process. Furthermore, 5-ALA may significantly inhibit the growth of glioma U251 cells under light irradiation, and the expression levels of TAPI-associated proteasomes has been demonstrated to be significantly increased (20). In addition, PDT was demonstrated to induce phosphorylation of proteins in the Bcl-2 family, enhancing the intracellular oxidative stress response and thereby accelerating apoptosis of U87 cells (37). According to the results of a previous study and previous clinical experience (38,39), PDT was hypothesized to be a potential treatment strategy for glioma via apoptosis induction and immune enhancement (19-22), which was confirmed in the current study. PDT represents a potential treatment strategy for glioma, and the treatment effect appears to be more obvious under a red light source. PDT inhibited cell growth via induction of cell apoptosis and immune enhancement. PDT treatment with hematoporphyrin inhibits cell growth of gliomas, which is regulated by miRNAs; however, the target genes for miRNAs and the molecular network remain unclear. Therefore, further studies are required in future. In the current study, the experiments, such as cell apoptosis, were only performed at one time point (23 h post-PDT treatment); thus, further research is required to verify the mechanism of action at additional time points. Furthermore, in this study, the human glioma cell lines, U87 and U251, were used; in future studies, we also aim to use normal control primary glioma cells from rats or mice to further confirm our findings.

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Competing interests

The authors declare that they have no competing interests.

References


2296