Porphyrin Derivatives Mediated Sonodynamic Therapy on Malignant Gliomas in Vitro

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ABSTRACT

Due to its highly infiltrative property, malignant glioma is a cancer of poor prognosis despite multidisciplinary treatment strategies, such as aggressive surgery and chemoradiotherapy, unveiling a need for new therapeutic approaches to control migrated tumor cells. In our study, we investigated the efficacy of sonodynamic therapy (SDT) using porphyrin derivatives involving 5-aminolevulinic acid, protoporphyrin IX, and talaporfin sodium as sonosensitizers on glioma cells, in vitro. These substances have been known to accumulate in glioma cells and are expected to have cytotoxic effects by sonication. Our study demonstrated that the cytotoxicity of sonication on glioma cells was enhanced by each sonosensitizer and the efficacy of the SDT may depend on the degree of intracellular accumulation of sonosensitizer. Also, induction of apoptosis was suggested to be a major mechanism of cell death. Though further investigations will be necessary, our preliminary result showed a potential of SDT with sonosensitizers for glioma treatment.
KEYWORDS: 5-aminolevulinic acid, in vitro, malignant glioma, porphyrin derivatives, protoporphyrin IX, sonodynamic therapy, talaporfin sodium, ultrasound
INTRODUCTION

Gliomas account for approximately 20% of primary brain tumors in adults (Nakamura et al. 2011). Regarding the malignant subtype of glioma, prognosis is very poor. Despite intensive therapy, the actual 2-year survival rate is only 33.9% (Chinot et al. 2014; Gilbert et al. 2014). Though novel therapeutic trials have been recently carried out, patients’ survival has not been improved remarkably. Thus, it is necessary to develop a different approach to treatment, enhancing conventional therapy. At present, the standard treatment for malignant gliomas is maximal safe resection combined with chemoradiation. Removal of the tumor is still the most important role in a therapeutic strategy, but the boundary of the tumor is often unclear because glioma cells infiltrate into adjacent normal brain tissue. Actually, 90% of tumors relapse within 2 cm of the primary site despite the use of chemoradiotherapy after gross total resection (Wilson 1992). These results suggest a key of successful treatment is to control these infiltrative tumor cells.

Photodynamic therapy (PDT) is one of the expected treatments (Muragaki et al. 2013). The mechanism of this therapy is that photochemically activated photosensitizers
generate reactive oxygen species (ROS), such as singlet oxygen and free radicals, which lead to oxidative damage of target tumor cells. Among various photosensitizers, porphyrin derivatives have been utilized in PDT for gliomas because these substances are selectively accumulated in glioma cells, and they have been already applied for clinical use in PDT and intraoperative fluorescence diagnosis (Stylli et al. 2006a; Stylli et al. 2006b). Particularly, 5-Aminolevulinic acid (ALA) is widely used for intraoperative fluorescence diagnosis. ALA, which is a natural amino acid, is the first compound of the porphyrin biosynthetic pathway. In this pathway, protoporphyrin IX (PpIX) is synthesized from ALA in mitochondria, and PpIX is suggested to be retained in tumor cells when administrating exogenous ALA excessively. This is thought to be due to metabolic dysfunction of PpIX in malignant tumor cell (Ishizuka et al. 2011). Accumulated PpIX in a tumor cell acts as a strong photosensitizer. Furthermore, talaporfin sodium (TS) is a second-generation photosensitizer that has recently been approved in Japan for PDT to treat primary malignant brain tumors (Muragaki et al. 2013). Although PDT is an attractive treatment in glioma therapy, there is a great problem of penetration depth of the light in brain tissue. To overcome this issue,
ultrasound has been expected as alternative energy source to excite the photosensitizers. Rosenthal et al. have shown that various photosensitizers are excited by ultrasound as well (Rosenthal et al. 2004). The aim of this study is to investigate the anti-tumor effects of SDT with porphyrin derivatives such as ALA, PpIX, and TS for gliomas in vitro.

**MATERIAL AND METHODS**

*Reagents*

ALA was purchased from COSMO BIO Co., Ltd (Koto, Tokyo, Japan). PpIX, Hoechst 33342 (bis Benzimide H 33342 trihydrochloride), and Propidium iodide (PI) were purchased from Sigma-Aldrich (St. Louis, MO, USA). TS was purchased from Meiji seika, Ltd. (Chuo, Tokyo, Japan). Calcein AM/Ethidium homodimer (EthD-1) costaining kit (Live/Dead Viability/Cytotoxicity Assay Kit®) was purchased from Life Technologies (Minato, Tokyo, Japan). Annexin V-FLUOS Staining Kit was purchased from Roche Applied Science (Minato, Tokyo, Japan).
**Cell culture**

Rat glioma cell line C6 and human glioblastoma cell line U87MG were obtained from the American Type Culture Collection (Manassas, VA, USA). Cells were maintained as monolayers in Dulbecco’s Modified Eagle Medium (DMEM; Sigma-Aldrich, St. Louis, MO, USA) supplemented with 10 % heat-inactivated fetal bovine serum (FBS; Life Technologies, Minato, Tokyo, Japan), 100 U/mL of penicillin G, and 100 μg/mL of streptomycin (Sigma-Aldrich, St. Louis, MO, USA) at 37 °C in a humidified incubator with 5 % CO₂ in air atmosphere. Unless stated otherwise, the conditions of the incubation were as described above. Cells in the exponential growth phase after more than 5 passages were used for all of the following experiments.

**Ultrasound exposure apparatus**

The ultrasound exposure apparatus was arranged as shown in Fig. 1. A laboratory-assembled planar circular transducer (30 mm in diameter) with air backing was submerged in a water bath (0.4×0.2×0.1 m³) filled with degassed water to avoid attenuation of ultrasound wave, and facing upward with its acoustic surface parallel to
the water surface. A function generator (AFG3022, Tektronix, Minato, Tokyo, Japan) was used to generate burst pulses of 1 MHz in center frequency and 100 Hz in pulse repetition frequency. Burst pulses of 50% in duty cycle contain 5,000 cycles in each burst. The transducer was driven by the signal amplified using a power amplifier (UOD-WB-1000, Tokin, Sendai, Miyagi, Japan). The spatial-average-temporal-average intensity ($I_{SAT}$) of 0.16 W/cm$^2$ was used for all experiments, which was determined by pressure calibration using an apparatus utilized for our previous study. (Yamaguchi et al. 2011) The calibration was carried out in a free water condition using a Polyvinylidene difluoride membrane hydrophone (MHB-500, Onda, Sunnyvale, CA, USA). During ultrasound exposure, the flat bottom of the well plate (MULTIWELL™ 6 well, BD FALCON®, Bedford, MA, USA) was placed 5 mm from the surface of the transducer. In this condition, the estimation of $I_{SAT}$ that cells actually received is technically difficult because strong reflections at the water surface generate a standing wave field inside the well. An acoustic absorbing rubber plate is frequently used to prevent the reflection by placing it on the water surface. However, it was not utilized in the experiments to minimize disturbance in culture condition and the possibility of
contamination because cell incubation should be continued for up to 24 hours after the sonication. The wells measured 35 mm in diameter, 18 mm in depth, and were 1.62 mm thick. Furthermore, the height (volume) of solution in each well was stringently unified at 1.3 mm (4 mL/well) because the sound pressure transferred to the cells from the standing waves varied depending on the fluid surface height (data not shown). The water temperature in the bath was held at 37 °C to eliminate the temperature elevation of treated cells.

SDT treatment protocols

Cells were randomly divided into four treatment groups: control, sonosensitizer exposure, ultrasound exposure, and SDT (ultrasound exposure plus sonosensitizer exposure). Upon reaching 70 % confluence, cells were washed with phosphate-buffered saline (PBS; Nacalai tesque, Nakagyo, Kyoto, Japan) and collected using 0.05 % trypsin with 0.53 mM ethylenediaminetetraacetic acid. Approximately $3.0 \times 10^5$ cells/well were dispersed into the wells of each plate and subsequently incubated for 24 hours.
ALA and TS were stored at 4 °C in the dark, and dissolved in DMEM without phenol-red (Sigma-Aldrich, St. Louis, MO, USA) to a final concentration of 200 and 30 μg/mL respectively. PpIX was dissolved in dimethyl sulfoxide (Sigma-Aldrich, St. Louis, MO, USA) at a concentration of 0.005 g/mL and subsequently stored at -20 °C in the dark, and also dissolved in DMEM without phenol-red to a final concentration of 1.0 μg/mL. For the sonosensitizer and the SDT groups, the culture medium was replaced with an equivalent volume of sonosensitizer solution and incubated thereafter for 4 hours. For the other groups, the culture medium was replaced with fresh DMEM without sonosensitizer solution and phenol-red and subsequently incubated. Consecutively each well in the ultrasound and the SDT groups was subjected to ultrasound treatment at 1.0 MHz and 0.16 W/cm² for 60 seconds respectively.

Fluorescence intensity per cell after co-incubation with each sonosensitizer

The fluorescence intensity per cell after co-incubation with each sonosensitizer was analyzed by flow cytometry to verify differences in the uptake of the sonosensitizer in different cell lines. Following co-incubation with each sonosensitizer, cells were
harvested by trypsin and analyzed by a flow cytometer (Attune® Acoustic Focusing Cytometer, Life Technologies, Minato, Tokyo, Japan) using excitation and emission wavelengths of 488 and 690/50 nm, respectively.

Cytotoxicity of SDT

Following ultrasound exposure, cells were re-incubated for 24 hours. A Calcein AM/EthD-1 costaining kit was used with Hoechst 33342 following incubation to assess cell viability. After washing the cells with 1 mL of PBS, 100 μL of PBS containing 2 μM of Calcein AM and 4 μM of EthD-1, and 0.2 μL of Hoechst 33342 were added to each well and the cells were incubated for 30 minutes at room temperature in the dark. Cells were observed using a fluorescent microscope (IX71, U-LH100HGAP0, OLYMPUS, Shinjuku, Tokyo, Japan). The polyanionic dye Calcein AM is well retained within live cells, producing an intense uniform green fluorescence. EthD-1 enters cells with damaged membranes and binds to nucleic acids, thereby producing a bright red fluorescence in dead cells. Hoechst 33342 is a popular cell-permeation nuclear counterstain that emits blue fluorescence when bound to double-stranded
deoxyribonucleic acid (DNA), regardless of cell survival. The average numbers of green, red, and blue fluorescent cells were calculated in five random microscopic fields with at least 1000 cells/field for each treatment group. Experiments were repeated independently three times. Notably, exfoliated floating cells at 24 hours after treatment were defined as dead. The number of dead cells was calculated as \[ number \ of \ Hoechst \ 33342 \ positive \ cells \ in \ control \ group \ - \ number \ of \ Hoechst \ 33342 \ positive \ cells \ in \ each \ treatment \ group \]. Thus, compensated dead cells were estimated as \[ number \ of \ EthD-1 \ positive \ cells \ in \ each \ treatment \ group + number \ of \ dead \ cells \ in \ each \ treatment \ group \]. The survival rate (%) of the tumor cells was calculated as \[ 100 - \frac{number \ of \ compensated \ dead \ cells \ in \ each \ treatment \ group}{number \ of \ Hoechst \ 33342 \ positive \ cells \ in \ control \ group} \times 100 \].

Mechanism of ALA mediated SDT cytotoxicity on C6 cells

Treated C6 cells were stained with Annexin V-FLUOS and PI to assess the cell damage mechanism following re-incubation for 6 hours. After washing once with 1 mL
of PBS, 100 μL of the incubation buffer (10 mM Hepes, pH 7.4, 140 mM NaCl, 5 mM CaCl₂) containing 2 μL of Annexin V-FITC and 2 μL of PI (50 μg/mL) was added to each well. Cells were then incubated for 15 minutes at room temperature in the dark.

The cells were trypsinized, and then analyzed using a flow cytometer with an excitation wavelength of 488 nm and emission wavelength of 530/30 nm for Annexin V-FLUOS and 574/26 nm for PI respectively. Phosphatidyl serine (PS) is located on the cytoplasmic surface of the cell membrane in live cells; however, it is translocated from the inner to the outer leaflet of the plasma membrane in the acute phase of apoptosis, while Annexin V-FLUOS has a high affinity for PS exposed on the outer leaflet and produces green fluorescence. PI is impermeant to live and apoptotic cells; however, it binds tightly to the nucleic acids in dead cells, producing red fluorescence.

Statistical analysis

Experimental data was analyzed statistically with Statcel 3 software (OMS publishing Inc, Tokorozawa, Saitama, Japan). All values were presented as means ± standard deviation of triplicate experiments. The comparisons among these groups were assessed
by one-way analysis of variance (ANOVA) followed by Tukey-Kramer test and p
values < 0.05 were considered to be statistically significant.

RESULTS

Cytotoxicity of SDT with each sonosensitizer on C6 cells

The survival rate was evaluated by counting the number of live and dead cells with a
fluorescent microscope. The number of exfoliated cells was prominent in the SDT
group (Fig. 2). The cytotoxicity of the ultrasound was enhanced by each sonosensitizer
on C6 cells, while neither each sonosensitizer alone nor ultrasound alone caused any
cytotoxicity. The survival rate of C6 cells treated by ALA mediated SDT decreased
from 85.1±6.8 % to 69.1±10.8 % (Fig. 3a), as well as for PpIX mediated SDT (from
100.0±11.7 % to 78.9±1.0 %, Fig. 3b) and for TS mediated SDT (from 100.0±11.7 % to
59.8±8.1 %, Fig. 3c). Furthermore, for all sonosensitizers, the survival rates of the SDT
groups were lower than those from any of the other groups.

Mechanism of ALA mediated SDT cytotoxicity on C6 cells
Flow cytometry analysis demonstrated marked changes in the cell profile following ultrasound exposure on C6 cells (Fig. 4). The results of flow cytometry analysis were the following: cells in the lower-left quadrant (Annexin-V-/PI-) represent living cells, those in the lower-right quadrant (Annexin-V+/PI-) represent early apoptotic cells, and those in the upper-right quadrant (Annexin-V+/PI+) represent late apoptotic cells or necrotic cells. The early apoptotic cell rate in both ultrasound alone and SDT group were more than twice as high as that in the other groups, respectively (ultrasound alone group: Figure 4c 14.2 % and SDT group: Figure 4d 20.4 % vs. Control group: Figure 4a 5.1 % and ALA alone group: Figure 4d 1.5 %). In addition, late apoptotic and necrotic cell rates in the SDT group were the highest among all groups (Figure 4d).

Comparison of the cytotoxicity of SDT between C6 and U87MG cells

The cytotoxic effect of SDT on U87MG cells (Figure 3d-f) was smaller than that on C6 cells despite the fact that it was applied under the same conditions as for C6 in each sonosensitizer administered (Figure 3a-c). Figure 5 shows the results of the difference in uptake of sonosensitizer between these two cell lines in flow cytometry analysis. For
either sonosensitizer, the fluorescence intensity per cell of U87MG cells (Figure 5d-f) was lower than that of C6 cells (Figure 5a-c).

DISCUSSION

Although various therapeutic strategies for malignant gliomas have been tried clinically, satisfactory results have not been obtained yet (Chinot et al. 2014; Miyatake et al. 2014; Thomas et al. 2014). Japanese national survey reported that 1-, 2-, and 3-years survival rates of patients with malignant gliomas was approximately 60.3 %, 25.4 %, 15.9 %, respectively (Report of brain tumor registry of Japan (2001-2004) 13th edition 2014). Therefore, new methods of treatment should be developed as a breakthrough of this situation.

Since the 80’s, PDT has been explored as a novel option to treat malignant gliomas because this therapy is of low toxicity for normal brain tissue by tumor-selectivity of photosensitizers (Stylli et al. 2006a; Stylli et al. 2006b). However, the most important problem of PDT is the limited penetration depth of the light in brain tissue. It was reported that the effective depth of laser beam penetration in brain tissue is
approximately 0.75–1.5 cm (Muragaki et al. 2013). This issue was recognized from the beginning of the development of PDT. To overcome this issue, basic experiments were conducted to confirm whether ultrasound could be used as an alternative to the light for excitation of photosensitizers (Rosenthal et al. 2004; Chen et al. 2014). Further studies have shown the cytotoxic effect of SDT, which was combined ultrasound and conventional photosensitizers, for various cancers (Canaparo et al. 2006; Song et al. 2011; Yamaguchi et al. 2011; Yumita et al. 2011). Studies regarding ALA and PpIX mediated SDT have also provided encouraging results (Wang et al. 2010; Song et al. 2011; Li et al. 2012; Lv et al. 2012; Su et al. 2014), however, knowledge of ALA mediated SDT for malignant gliomas is limited yet (Ohmura et al. 2011; Jeong et al. 2012).

Ultrasound exposure to biological tissues is categorized into two effects: thermal and non-thermal. Acoustic energy absorbed by a biological tissue produces a temperature elevation and the thermal effect is utilized for hyperthermia. But heat injury sometimes causes crucial problem in treatment of intracranial lesions. Consequently, our concept of this study is that the principle of SDT for brain tumor should be based on non-thermal.
A non-thermal effect is caused by various mechanisms of mechanical and chemical phenomena. Acoustic radiation force produces tissue motion and acoustic streaming, which increase cell membrane permeability and resulting drug delivery efficiency.

Cavitation is an important mechanism of the non-thermal effect and is categorized into non-inertial and inertial cavitation. Non-inertial cavitation is caused by microbubble oscillation under applied pressure and mainly produces mechanical effects, such as microstreaming, excess temperature elevation, and acoustic radiation force. Inertial cavitation occurs under a condition with higher intensity and longer duration than those in the non-inertial condition. Bubble expansion and subsequent rapid adiabatic contraction by inertia of the surrounding liquid causes a temperature rise inside the bubble up to several thousands of degrees, in theory, generating free radicals by pyrolysis of H$_2$O molecules. Generation of the free radicals is essential to induce SDT effects, a series of chemical reactions of a sonosensitizer such as porphyrin derivatives that was originally used as light activated chemicals. Porphyrin derivatives themselves have no anti-tumor effects, and are less cytotoxic for normal tissues; however, inertial
cavitation can be followed by energy transfer to oxygen to generate ROS in the presence of porphyrin derivatives, (Canaparo et al. 2006) and they are expected to provide an anti-tumor effect without temperature rising.

Apoptosis is suggested as the mechanism of cell death by PDT. Apoptosis can be induced by low intensity ultrasound exposure even below the threshold for temperature elevation of the culture medium (Lagneaux et al. 2002; Feng et al. 2010; Cheng et al. 2013). In addition, Zhang et al. showed that low intensity ultrasound exposure can induce apoptosis related to caspase-3, Bcl-2, and survivin in C6 cells (Zhang et al. 2012). In our study, flow cytometry analysis confirmed that ultrasound exposure induced cell apoptosis at 0.16 W/cm², even in the absence of a sonosensitizer. On the other hand, cell survival analysis using immunofluorescences did not indicate the presence of apoptosis in the ultrasound alone group. This discrepancy may be caused by tumor-doubling time because measurement was performed 24 hours after treatment. However, an important consideration is that survival rate of SDT group is still suppressed. The sonication conditions are not fully determined, although they may have been well reproduced between experiments, and will require further research.
Our findings also demonstrated a relationship between sonodynamic efficacy and the relative fluorescence intensity of sonosensitizer per cell, referring to the degree of intracellular accumulation of sonosensitizer per cell. We confirmed that the cytotoxicity of SDT was distinct between C6 and U87MG cells, and the efficacy of SDT could depend on the degree of intracellular accumulation of sonosensitizer for each cell line. Yumita et al. also reported that the efficacy of SDT closely correlate with the concentration of sonosensitizer in medium (Yumita et al. 2010). Moreover, results of clinical research showed that PpIX accumulates in tumor cells in a histopathological grade-specific manner after administration of ALA for intraoperative fluorescence diagnosis of malignant gliomas (Ishizuka et al. 2011). This data may support that the peak of the fluorescence intensity of the photo-/sonosensitizer is distinct among different cell lines. Therefore, therapeutic strategies to avoid the regulatory feedback in heme biosynthesis for accumulation of PpIX in tumor cells should be considered to obtain an efficient anti-tumor effect of ALA mediated SDT. For example, Xu et al. reported the efficacy of SDT for the glioma stem-like cells was improved by the inhibition of ATP-binding cassette sub-family G member 2 (ABCG2) transporter.
function, which regulates cellular accumulation of porphyrin derivatives, with a specific inhibitor (Xu et al. 2013).

When considering the development of SDT as clinical application, if the sonication could be given through the skull, SDT will be an ideal noninvasive treatment for malignant brain tumors. This may be possible in the near future given the research conducted nowadays on brain ultrasound imaging and therapy. Because magnetic resonance imaging guided focused ultrasound surgery (ExAblate® Neuro; InSightec, Haifa, Israel) has already been applied in global clinical trial to treat neurological disorders. This device induces focused tissue coagulation via ultrasound exposure generating the thermal effect through intact skull, which has been attracted as an alternative to deep brain stimulation (Lipsman et al. 2014). Clinical trials for transcranial focal ablation of deep-seated tumors, especially glioblastoma, have started (Ram et al. 2006; McDannold et al. 2010). However, there might be a limit to control infiltrative tumor cells by “thermal” effect when considering the injurious effect upon adjacent normal brain tissues. Hence, SDT based on “non-thermal” effect should be established for treatment of deep-seated brain tumors.
CONCLUSIONS

In this study, SDT with porphyrin derivatives such as ALA, PpIX, and TS demonstrated the cytotoxic effect on glioma cells via apoptosis. Although further investigation will be necessary, the “non-thermal” SDT has the potential to be a new therapeutic modality for malignant gliomas.
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FIGURE LEGENDS

Figure 1
Schematic diagram of experimental ultrasound irradiation apparatus set up.

Figure 2
Calcein AM (upper line), Ethidium homodimer (EthD-1) (middle line), and Hoechst 33342 (lower line) were stained on C6 cells 24 hours after treatment. Control represents cells without any treatments, 5-Aminolevulinic acid (ALA) alone represents cells cultured with 200 μg/mL of ALA for 4 hours, Ultrasound (US) alone represents cells treated only with ultrasound at 1.0 MHz and 0.16 W/cm² for 60 seconds, and Sonodynamic therapy (SDT) represents cells treated by ultrasound as described above in the presence of 200 μg/mL of ALA. A significant number of exfoliated cells were observed in the SDT group.

Figure 3
Survival rates of C6 cells and U87MG cells after treatment (a, d: 5-Aminolevulinic acid (ALA) mediated sonodynamic therapy (SDT), b, e: Protoporphyrin IX (PpIX) mediated SDT, c, f: Talaporfin sodium (TS) mediated SDT, upper line: C6 cells, lower line: U87MG cells, n=3). Data are presented as the mean ± standard deviation and statistical analysis was performed using one-way analyzed of variance (ANOVA) followed by the Tukey-Kramer test. * shows a significant difference (p<0.05) between groups. ALA alone; cells cultured with 200 μg/mL of ALA for 4 hours, PpIX alone; cells cultured with 1.0 μg/mL of PpIX for 4 hours, TS alone; cells cultured with 30 μg/mL of TS for 4 hours, Ultrasound (US) alone; cells treated only with ultrasound at 1.0 MHz and 0.16 W/cm² for 60 seconds, and SDT; cells treated by US as described above in the presence of each sonosensitizer.

Figure 4

Apoptosis and necrosis rates on C6 cells were analyzed via flow cytometry with Annexin V and Propidium iodide (PI) costaining to assess the mechanism of cell damage in 5-Aminolevulinic acid (ALA) mediated sonodynamic therapy (SDT) (a:
control group, b: ALA alone group, c: ultrasound alone group, d: SDT group). The results were interpreted in the following fashion: cells in the lower-left quadrant (Annexin-V-/PI-) represent living cells, those in the lower-right quadrant (Annexin-V+/PI-) represent early apoptotic cells, and those in the upper-right quadrant (Annexin-V+/PI+) represent late apoptotic or necrotic cells.

Figure 5

Relative fluorescence intensities of accumulated sonosensitizer in C6 and U87MG cells were analyzed via flow cytometry. For each sonosensitizer, the fluorescence intensity per cell of the U87MG cells was lower than that of the C6 cells. (a, d: 5-Aminolevulinic acid (ALA), b, e: Protoporphyrin IX (PpIX), c, f: Talaporfin sodium (TS), upper line: C6 cells, lower line: U87MG cells)
Figure 1

- Cell plate
- Gas removal equipment
- Water temperature gauge
- Water bath (keep at 37°C)
- Transducer
- Oscilloscope (DPO2012, Tektronix, Tokyo, Japan)
- Amplifier (UOD-WB-1000, Tokin, Miyagi, Japan)
- Function generator (AFG3022, Tektronix, Tokyo, Japan)
Figure 2

Calcein AM

EthD-1

Hoechst 33342
Figure 3

[Bar charts showing survival rates for ALA, PpIX, and TS in C6 and U87MG cells under different conditions: Control, ALA alone, US alone, and SDT.]
Figure 4
Figure 5

AL

PpIX

TS

C6

a

b

c

U87MG

d

e

f

counts

counts

counts

counts

PpIX

PpIX

PpIX

PpIX

TS