



Inhibition of Sonic Hedgehog Signaling Suppresses Glioma Stem-Like Cells Likely Through Inducing Autophagic Cell Death

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Glioblastoma (GBM) often recurs after radio- and chemotherapies leading to poor prognosis. Glioma stem-like cells (GSCs) contribute to drug resistance and recurrence. Thus, understanding cellular mechanism underlying the growth of GSCs is critical for the treatment of GBM. Here GSCs were isolated from human U87 GBM cells with magnetic-activated cell sorting (MACS) using CD133 as a marker. The CD133⁺ cells highly expressed sonic hedgehog (Shh) and were capable of forming tumor spheroids *in vitro* and tumor *in vivo*. Athymic mice received intracranial injection of luciferase transduced parental and CD133⁺ GBM cells was utilized as orthotopic GBM model. Inhibited Shh by LDE225 delayed GBM growth *in vivo*, and downregulated Ptch1 and Gli1. CD133⁺ cell proliferation was more sensitive to inhibition by LDE225 than that of CD133⁻ cells. Treatment with LDE225 significantly reduced CD133⁺-derived tumor spheroid formation. Large membranous vacuoles appeared in the LDE225-treated cells concomitant with the conversion of LC3-I to LC3-II. In addition, LDE225-induced cell death was mitigated in the presence of autophagy inhibitor 3-methyladenine (3-MA). Tumor growth was much slower in *Shh* shRNA-knockdown mice than in control RNA-transfected mice. Conversely, tumor growth was faster in Shh overexpressed mice. Furthermore, combination of LDE225 and rapamycin treatment resulted in additive effect on LC3-I to LC3-II conversion and reduction in cell viability. However, LDE225 did not affect the phosphorylated level of mTOR. Similarly, amiodarone, an mTOR-independent autophagy enhancer, reduced CD133⁺ cell viability and tumor spheroid formation *in vitro* and exhibited anti-tumor activity *in vivo*. These results suggest that Shh inhibitor induces autophagy of CD133⁺ cells likely through mTOR independent pathway. Targeting Shh signal pathway may overcome chemoresistance and provide a therapeutic strategy for patients with malignant gliomas.

Keywords: sonic hedgehog, glioblastoma, stem-like cells, autophagy, amiodarone

INTRODUCTION

Glioblastoma (GBM) is one of the most common and malignant subtype of brain tumors in adults (1). Although the incident rate is relatively low, GBM is very invasive that leads to rapid neurological destruction and a disproportionately high mortality. Despite the improvement of new surgical and radiation techniques, the median survival rate of patients with GBM is rather low (2, 3). A blood-brain-barrier permeable DNA alkylating agent temozolomide (4) is currently the first line chemotherapy for treating GBM. In clinics, combination of radiotherapy with temozolomide significantly prolongs the survival rate for GBM patients (5, 6). Unfortunately, tumor often regrows after radio- and chemotherapies (7, 8) resulting in poor prognosis (9, 10). The recurrence is caused at least in part, by the resistance of GBM to conventional chemo- and radio-therapies (11–14), highlighting an urgent necessity of designing new strategies for the treatment of GBM.

Sonic hedgehog (Shh) is a member of the hedgehog (Hh) family which functions as a chemical signal in transmitting information to the embryonic cells required for normal development. Shh plays a critical role in the regulation of vertebrate organogenesis and the development of brain and spinal cord including midbrain and ventral forebrain neuronal differentiation and proliferation, and many other parts of the body (15–18). Because of its role in embryonic development, aberrant or dysregulation of Shh signaling has been implicated in the initiation and/or maintenance of different types of tumor (19–21) including GBM (22). Consistent with these reports, Shh antagonists have been shown to possess anti-tumor activity in patients with basal cell carcinoma and medulloblastoma (21, 23, 24).

Cancer stem cells (CSCs) are cancer cells that possess the ability to replenish tumors through the self-renewal and differentiation into multiple cell types (25, 26). CSCs have been identified in the breast, colon, brain and other areas and can differentiate into all the cell phenotypes of the parental tumor (27, 28). CSCs are hypothesized to be associated with chemo- and radio-resistance that lead to recurrence of tumor formation. In this theory, conventional radio- and chemotherapies kill differentiated or differentiating cells which form the bulk of the tumor, while sparing CSCs. Therefore, targeting CSCs offers a promising approach to improve cancer treatment or even cure cancer (29). In the present study, we isolated glioma stem-like cells (GSCs) from human GBM cell line U87MG (U87) using CD133 as a marker. We found that Shh expression is higher in the CD133⁺ cells than in the CD133[−] cells. LDE225, a smoothened antagonist (30, 31), delayed GBM growth *in vivo* and significantly reduced the number of tumor spheroids derived from CD133⁺ cells. Furthermore, tumor growth was much slower in *Shh* knockdown mice suggesting that glioma growth may be dependent on a small population of CD133⁺ cells that are regulated by the Shh pathway.

MATERIALS AND METHODS

Animals

The BALB/cAnN.Cg-Foxn1^{nu}/CrJNarl mice were purchased from the National Laboratory Animal Center (NLAC). Five mice were housed in a cage with controlled temperature (22 ± 2°C) and humidity (55 ± 5%), kept on a 12 h light/dark cycle, and were given free access to water and food. Care and use of laboratory animals were in accordance with National Institutes of Health (NIH) guidelines. All the procedures were approved by the Institutional Animal Care and Use Committee of the College of Medicine, NCKU, with project approval number (#104064 and #107106).

Cell Culture

The human glioblastoma (GBM) cell lines U87MG (U87) was provided by Dr. Michael Hsiao (Genomics Research Center, Academia Sinica, Taiwan). GBM patient-derived cell line P#5 was developed by Dr. Jian Ying Chuang. Both cells were cultured in Dulbecco's Modified Eagle medium-high glucose (DMEM-high glucose, Caisson) supplemented with 10% fetal bovine serum (FBS, Sigma-Aldrich), 100 U/ml penicillin, and 0.1 mg/ml streptomycin (Caisson). All cells were maintained in a humidified incubator with 5% CO₂ at 37°C.

Isolation and Characterization of Cancer Stem Cells From Glioblastoma Cell Line

For magnetic-activated cell sorting (MACS) purification, fresh GBM and spheroids were dissociated, washed, and incubated either with PE conjugated CD133/2 or IgG2b (Miltenyi Biotec, 1:11) at a concentration of 10⁸ nucleated cells per ml at room temperature for 15 min. EasySep² PE selection cocktail at 100 µl/ml cells was added and mixed down for more than 5 min. Magnetic cell separation was performed using manual FalconTM polystyrene round-bottom tubes and an EasySep² Magnet machine. Tube was removed from magnet and cells resuspended in an appropriate amount of desired medium. The CD133⁺ cells were incubated in neural stem cells selection medium (NeuroCultTM NS-A Basal Medium, NeuroCultTM NS-A Proliferation Supplement, bFGF 10 ng/mL, EGF 10 ng/mL, 20 µg/mL Heparin; STEMCELL, Canada) and gave rise to non-adherent spheres on Ultra Low Attachment Multiple Well Plates (CORNING). CD133⁺ cells were initially allowed to form tumor spheroids in suspension culture, dissociated using Accutase (BD Biosciences) at 37°C for 30 min, and then split 1:3 to 1:5. The number of tumor spheroids formed by CD133⁺ cells treated with or without LDE225 (25 µM) for 7 days in culture were determined with an Olympus DP72 image analysis system and Inverted fluorescence microscope Olympus IX71.

Cell Proliferation Assay

Cell proliferation was measured using WST1 [2-(4-iodophenyl)-3-(4-nitrophenyl)-5-(2,4-disulphophenyl)-2H-tetrazolium] assay (Clontech Laboratories, California, USA). The CD133⁺ cells were seeded at a concentration of 8,000 cells/well in 200 µl culture medium containing various concentrations of LDE225 or

cyclopamine (e.g., final concentration of 1–100 μM) into 96-well plates. The cells were incubated for 48 h at 37°C and 5% CO_2 . WST-1 reagent (10 μl /well) was added, and cells were incubated for 1 h at 37°C and 5% CO_2 . The absorbance of the product was measured at 440 nm with a microplate (ELISA) reader. The cell counts were determined by the percentage of the absorption relative to the vehicle-treated control culture.

Western Blotting Assay

Cell pellets were collected, centrifuged at 4,000 rpm and stored at -80°C . Drugs- or vehicle-treated cell pellets were lysed in a RIPA lysis buffer containing 50 mM Tris-HCl, pH 7.4, 150 mM NaCl, 1% Nonidet P-40, 0.25% sodium deoxycholate, 0.1% sodium dodecyl sulfate (SDS), protease inhibitor (Roche) and phosphatase inhibitor (Roche). Lysates were shaken at 40 rpm on ice for 1 h and then centrifuged at 12,000 rpm for 30 min at 4°C. The mouse brain tissues were homogenized in a lysis buffer (50 mM Tris-HCl, pH 7.5, 0.3 M sucrose, 5 mM EDTA and protease/phosphatase inhibitor cocktail (Roche, Nutley, USA). Supernatants were collected and then protein concentration was measured by Bradford assay. The protein was resuspended in 5X sample buffer (12.5 mM Tris, 25% glycerol, 4% SDS, 1.54% DTT, and 0.02% Bromophenol blue). The total protein was denatured at 95°C for 5 min. Protein electrophoresis on 15 or 8% SDS-polyacrylamide gel under 120 volt, and the separated protein was transferred to a PVDF membrane (Immunobilon transfer membranes, Millipore) by semi-dry transfer system (BIO-RAD) under 350 mA, 20 volt for 1.5 h. Membranes were incubated with blocking buffer (3% bovine serum albumin in PBS) for 1 h. Membranes were incubated with the following primary antibodies: rabbit anti-Shh (1:3,000, Millipore, Darmstadt, Germany), rabbit anti-P62 (1:2,000, Cell Signaling Technology, Danvers, USA), mouse anti-LC3 (1:2,000, MBL, Japan), mouse anti-CD133 (1:5,000, Millipore, Darmstadt, Germany), rabbit anti-Ptch1 (Protintech, Illinois, USA), rat anti-Gli1 (Sigma, Darmstadt, Germany) antibody overnight at 4°C, followed by treatment with HRP-conjugated secondary antibodies (Jackson ImmunoResearch Lab., West Grove, USA) for 1 h at room temperature. After three rinses with 0.3% Triton X-100 in PBS for 10 min each, the ECL-plus chemical reagents (PerkinElmer) were added to the membrane. Films (Fuji, Japan) were exposed at different time points to ensure the optimum density. The band intensities were analyzed for densitometry by using Windows Image-J version 1.29. The protein levels in all groups were expressed as a percentage of those in controls.

Immunofluorescent Staining

For immunostaining of tumor spheroids, CD133⁺ cells were initially allowed to form spheroids in suspension culture for 7 days. The cells were then fixed with 4% paraformaldehyde in PBS pH 7.4 for 15 min at room temperature. The cells were then incubated for 1 h in blocking solution (1% bovine serum albumin), and then incubated in primary antibodies at 4°C overnight. The following primary antibodies were used: mouse anti-CD133 (1:100, Millipore, Darmstadt, Germany), rabbit anti-SOX-2 (1:1,000, Abcam, Cambridge, UK) and

rabbit anti-Shh (1:200, Millipore, Darmstadt, Germany), rabbit anti-ALDH1A1 (Protintech, Illinois, USA), rat anti-ABCG2 (Santa Cruz, Texas, US), rabbit anti-LC3 (Novus, Colorado, US). After three rinses with PBS containing 0.1% Triton X-100 for 10 min each, the cells were incubated in secondary antibodies at room temperature for 1 h. The following secondary antibodies were used: Alexa Fluor®594-conjugated Goat-anti-rabbit IgG (Jackson ImmunoResearch Lab., West Grove, USA) and Alexa Fluor®488-conjugated Sheep-anti-mouse IgG and Alexa Fluor®488-conjugated Goat-anti-rat IgG (Jackson ImmunoResearch Lab., West Grove, USA). DAPI (4',6-diamidino-2-phenylindole) (1:1,000, Sigma-Aldrich, St. Louis, USA) dye was used as nuclear counterstain. The cells are washed in PBS containing 0.1% Triton X-100 three times for 10 min. The sections were cover-slipped with Prolong® gold anti-fade reagent (Life Technologies, Carlsbad, USA). Fluorescence signals were detected with a Leica DM 2500 microscope and MetaMorph image software used for counting the cells number.

JC-1 Mitochondrial Membrane Potential Assay

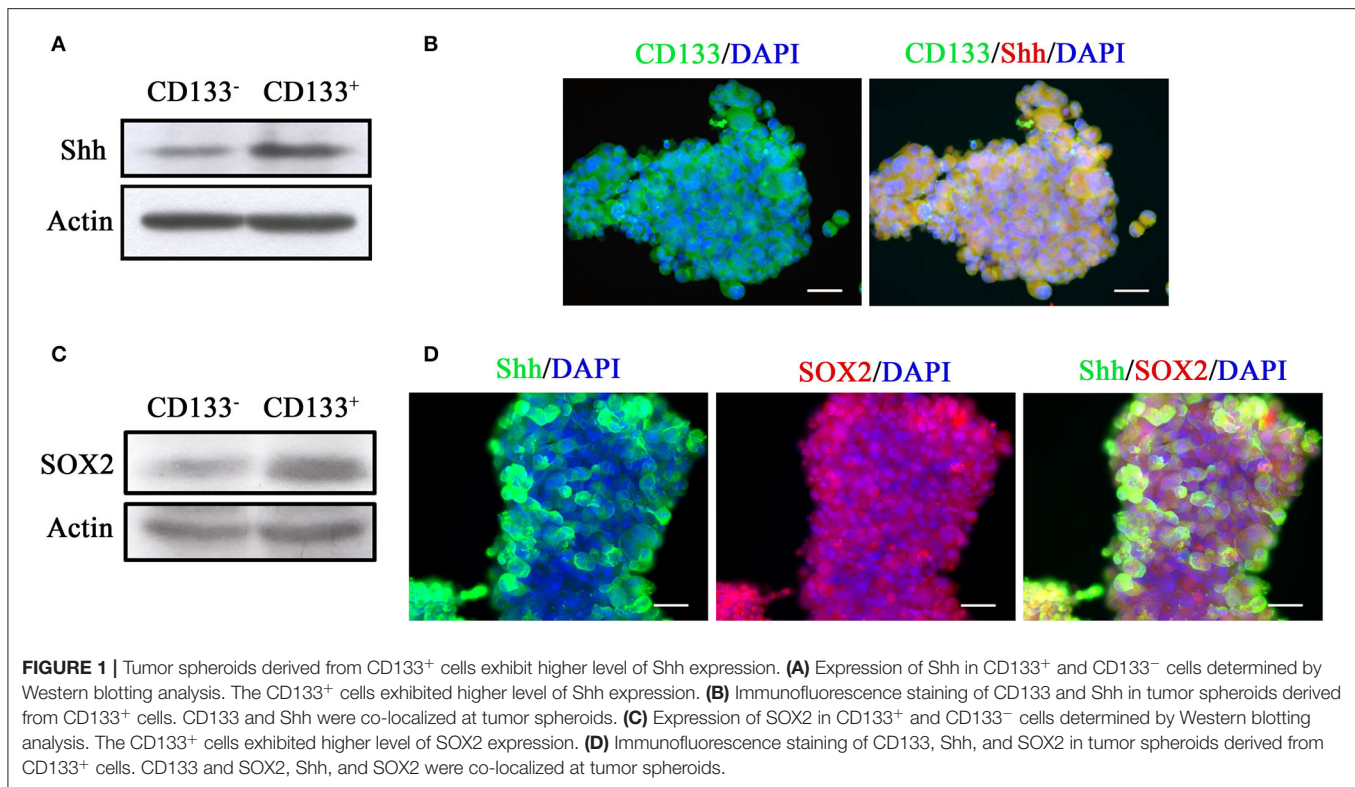
Early apoptosis was measured by using JC-1 Mitochondrial Membrane Potential assay (Cayman Chemical, Michigan, USA). The CD133⁺ cells were seeded at a concentration of 5,000 cells/well in 100 μl culture medium containing 25 μM LDE225 or 0.1% DMSO as vehicle control into black 96-well plates. The cells were incubated for 48 h at 37°C and 5% CO_2 . JC-1 Staining solution (10 μl /well) was added, and cells were incubated for 30 min at 37°C and 5% CO_2 . Plates were centrifuged at 300 $\times$ g at room temperature. The supernatant was removed, and JC-1 buffer was used to suspend JC-1 stained cells. The fluorescence intensity of J-aggregate and JC-1 monomer was measured with excitation and emission at 535/595 nm and 485/535 nm with a microplate (ELISA) reader. The ratio of J-aggregate and JC-1 was determined as early apoptosis.

Caspase-Glo® 3/7 Assay

The activity of caspase 3 and 7 was measured by using Caspase-Glo® 3/7 Assay (Promega, Wisconsin, US). The CD133⁺ cells were seeded at a concentration of 1,000 cells/well in 100 μl culture medium containing 25 μM LDE225 or 0.1% DMSO as vehicle control into white 96-well plates. The cells were incubated for 48 h at 37°C and 5% CO_2 . Caspase-Glo® 3/7 Reagent (10 μl /well) was added, and cells were incubated for 1 h at 37°C and 5% CO_2 . The luminescence was recorded with a microplate (ELISA) reader. The activity of caspase 3 and 7 was determined by the percentage of the luminescence signal relative to the control culture.

In vivo Intracranial Xenograft Animal Model and Bioluminescence Imaging

U87 GBM cells were transduced with lentiviral vector expressing GFP and firefly luciferase. GFP/Luc expressing cells were sorted out for further passages (FACS-Aria, BD Biosciences). For tumorigenesis, luciferase-expressing GBM cells were inoculated intracranially into the 8- to 10-week-old male nude mice



(BALB/cAnN-Foxnlnu/CrlNarl mice, National Laboratory Animal Center). Nude mice were anesthetized with chloral hydrate and placed on a stereotaxic device. Subsequently, a hamilton syringe with 30-gauge needle was mounted on a stereotaxic device, and luciferase-expressing GBM cells were injected into the left side of the brains, 1.5 mm caudal and lateral to the bregma, and at a depth of 3.5 to 4 mm. LDE225 (Cayman) was injected intraperitoneally injected at a dose of 20 mg/kg twice weekly. Tumor growth was monitored by IVIS spectrum Live Imaging System (IVIS-200, Xenogen) twice weekly. Before monitoring, mice were injected with 150 mg/kg D-luciferin (PerkinElmer), and simultaneously anesthetized with isoflurane. The results of luciferase radiance were quantitated by Live Imaging Software (Xenogen) and the results were analyzed by using GraphPad Prism software.

Shh shRNA Lentivirus Production

Production of lentivirus was initiated by triple transfection of HEK293T cells by a Lipofectamine[®] LTX Reagent (Life Technologies, Carlsbad, USA) method using small hairpin interfering RNA (shRNA) together with pCMV-dR8.91 and pMD2.G. The *Shh*-shRNA or scrambled shRNA conjugated on the vector of pLKO.1 with puromycin-resistant region was provided by National RNAi Core Facility (Institute of Molecular Biology, Academia Sinica, Taiwan). Cells were harvested 48 h later, medium containing lentiviruses filtered with 0.45 μ m filters and viral particles were concentrated from the supernatant by Lenti-X[™] Concentrator (Clontech Laboratories, Mountain View, USA) and purified to yield 1×10^8 transducing units/ml

storing at -80°C until use. The target sequence of *Shh*-shRNA is described as follows: *Shh*-shRNA: 5'-GCGGAAGGTATGAAGGGAAGA-3'.

Shh-Over-Expression

To create lentiviruses over-expressing *Shh*, full-length *Shh* open reading frames (ORFs) (NM_000193; GenScript, New Jersey, USA) was amplified by PCR and was inserted into pLVX-IRES-ZsGreen1 expression vector (Clontech Laboratories, California, USA). The pLVX-NES1-IRES-ZsGreens1 vector encoding *Shh* (or empty vector) and the two packaging plasmids (pCMV-dR8.91 and pMD2.G) were co-transfected into HEK293T cells by lipofectamine[®] LTX Reagent (Life Technologies, Carlsbad, USA). Lentiviruses were harvested at 48 h after transfection, filter lentivirus supernatant through a 0.45 μ m PVDF membrane filters, concentrated by Lenti-X[™] Concentrator (Clontech Laboratories, Mountain View, USA), purified to yield 1×10^8 transducing units/ml and stored at -80°C until use.

Statistical Analysis

Experiments were performed at least in triplicate. All results were presented as mean $\pm$ standard error of the mean (SEM). Independent experiments were analyzed by unpaired *t*-test. Two-way ANOVA was used to analyze the differences in tumor spheroids numbers *in vitro* and intracranial tumor growth *in vivo* at different times of treatment. Levels of $p < 0.05$ were considered to be of statistical significance.

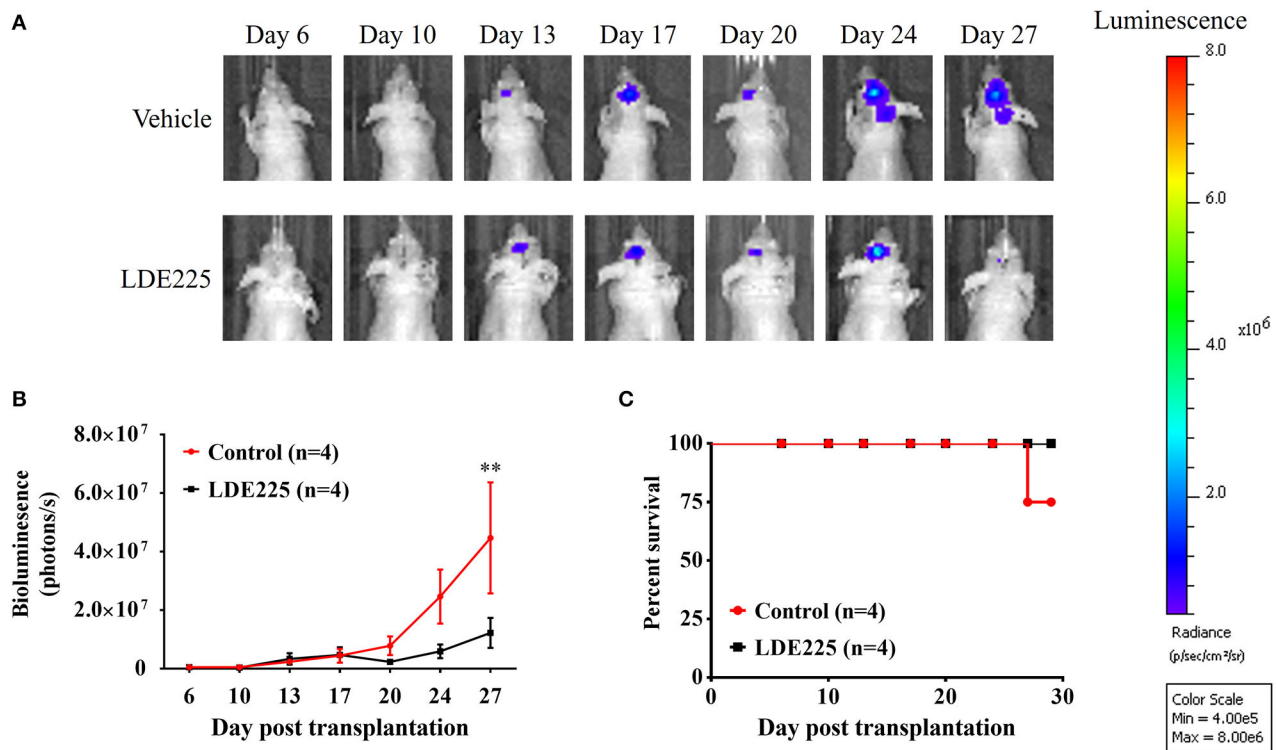


FIGURE 2 | LDE225 delays tumor growth in an intracranial tumor model. **(A)** Luc-expressing parental U87 cells (3×10^5 cells) were injected intracranially into athymic mice and tumor growth was monitored using the IVIS-200 imaging system. **(B)** Tumor growth was slower in LDE225 treated mice than in vehicle control mice. ** $p < 0.001$ vs. vehicle control. **(C)** Kaplan–Meier analysis of LDE225 treatment on survival.

RESULTS

CD133 has been reported as a marker of human neural and brain tumor stem cells (32, 33). We previously have isolated cancer stem-like cells with CD133 from glioblastoma (GBM) cell lines using magnetic bead cell sorting (34). Then CD133⁺ and CD133[−] cell populations were collected and cultured separately.

Shh Expression Is Higher in CD133⁺ Cells

We examined Shh expression in CD133⁺ and CD133[−] cells. Western blotting analysis showed that Shh expression was higher in the CD133⁺ cells than in the CD133[−] cells (**Figure 1A**). Immunofluorescence staining revealed that most of CD133⁺ cells were positive and co-localized with Shh (**Figure 1B**). SRY (sex determining region Y)-box 2 (SOX2) is a transcription factor that plays an important role in maintaining embryonic and neural stem cells (35, 36). **Figure 1C** showed that SOX2 expression was higher in the CD133⁺ cells than in the CD133[−] cells. These Shh-positive cells were also co-immunolabeled with SOX2 (**Figure 1D**). ABCG2 is a member of the ATP-binding cassette (ABC) transporter superfamily. The expression level of ABCG2 has been implicated in multidrug resistance (MDR) in cancer chemotherapy and the ability of self-renewal which correlates with CD133 (37, 38). ALDH1A1 is a member of the highly

conserved ALDH family that is observed in several cancer stem cells, and is often used to isolate and functionally characterize cancer stem cells (39). We also examined the expression of ABCG2 and ALDH1A1 by immunofluorescence in parental and CD133⁺ cells. The expression of ABCG2 and ALDH1A1 is higher in CD133⁺ cells than in parental cells (**Supplemental Figure 1**).

We used a potent and selective smoothened antagonist LDE225 (30) to determine the requirement of Shh signaling in the proliferation and tumor growth of GBM cell lines. We first confirmed the effect of LDE225 on tumor growth by using intracranial injection model. Luc-expressing parental U87 cells were injected intracranially into athymic mice and tumor growth was monitored using IVIS-200 imaging system. At day 7, LDE225 or DMSO as vehicle was injected intraperitoneally twice per week (20 mg/kg) into the mice and tumor growth was observed for 22 more days. Comparing with vehicle control, **Figure 2A** showed that LDE225 was able to delay tumor growth. A two-way ANOVA revealed a main effect of group (LDE225 vs. control) [$F_{(1,42)} = 6.126$, $p < 0.05$], interaction [$F_{(6,42)} = 2.297$, $p = 0.0524$] and days after application [$F_{(6,42)} = 5.849$, $p < 0.001$] (**Figure 2B**). Kaplan–Meier analysis of the survival data of vehicle control and LDE225-treated mice displayed in **Figure 2C**. We also confirmed whether LDE225 inhibits Shh signal pathway. CD133⁺ cells were treated with LDE225 (25 μ M)

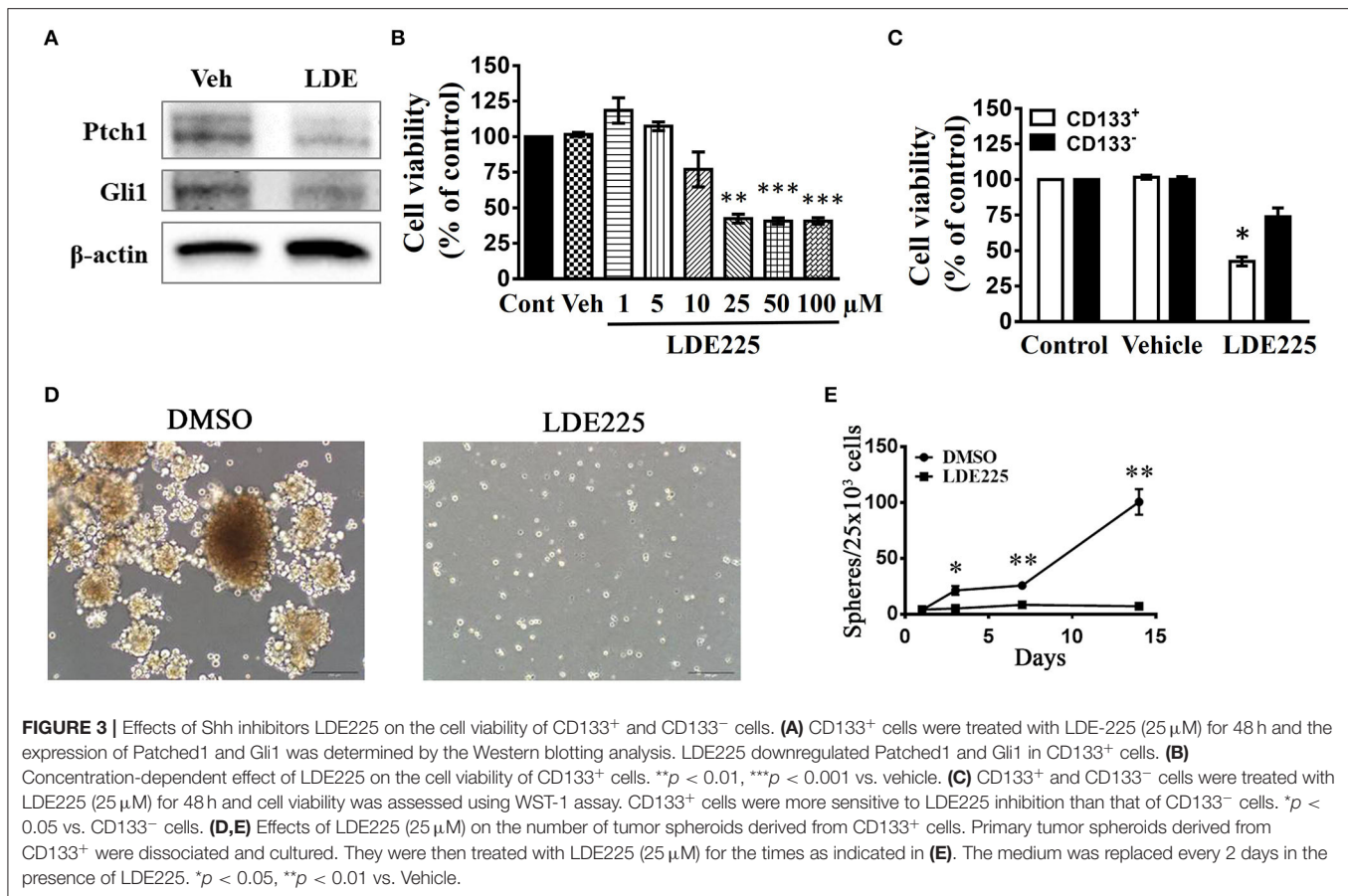


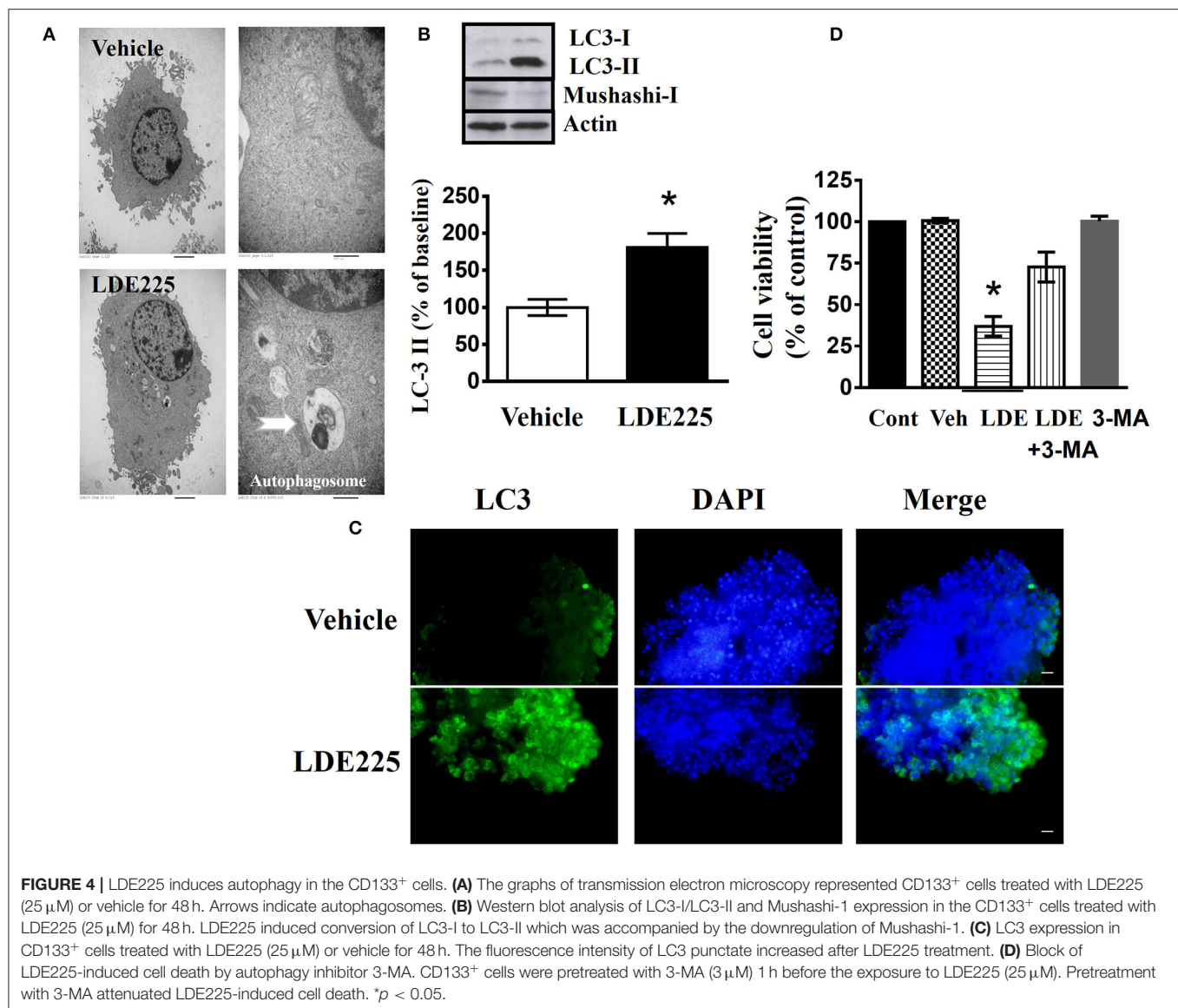
FIGURE 3 | Effects of Shh inhibitors LDE225 on the cell viability of CD133⁺ and CD133⁻ cells. **(A)** CD133⁺ cells were treated with LDE-225 (25 μM) for 48 h and the expression of Patched1 and Gli1 was determined by the Western blotting analysis. LDE225 downregulated Patched1 and Gli1 in CD133⁺ cells. **(B)** Concentration-dependent effect of LDE225 on the cell viability of CD133⁺ cells. **(C)** CD133⁺ and CD133⁻ cells were treated with LDE225 (25 μM) for 48 h and cell viability was assessed using WST-1 assay. CD133⁺ cells were more sensitive to LDE225 inhibition than that of CD133⁻ cells. **(D,E)** Effects of LDE225 (25 μM) on the number of tumor spheroids derived from CD133⁺ cells. Primary tumor spheroids derived from CD133⁺ were dissociated and cultured. They were then treated with LDE225 (25 μM) for the times as indicated in **(E)**. The medium was replaced every 2 days in the presence of LDE225. **(A)** $p < 0.05$, **(B)** $p < 0.01$, **(C)** $p < 0.05$ vs. CD133⁻ cells. **(D,E)** $p < 0.05$, $p < 0.01$ vs. Vehicle.

or vehicle for 48 h and the expression of Patch1 and Gli1 were determined by Western blotting analysis. As shown in **Figure 3A**, LDE225 downregulated the expression of PATCH1 and GLI1. Next, CD133⁺ and CD133⁻ cells were treated with LDE225 (25 μM) for 48 h and cell viability was assessed using WST-1 assay. Concentration dependent relationship estimated the IC₅₀ of about 20 μM for LDE225 to reduce CD133⁺ cell viability (**Figure 3B**), while IC₅₀ of LDE225 was about 50 μM for parental cells (**Supplemental Figure 2A**). As shown in **Figure 3C**, CD133⁺ cells were more sensitive to LDE225 inhibition than that of CD133⁻ cells. At the concentration of 25 μM, LDE225 inhibited the proliferation of CD133⁺ and CD133⁻ cells by $57.57 \pm 3.17\%$ ($n = 3$) and $26.18 \pm 6.22\%$ ($n = 4$), respectively [$t_{(5)} = 4.013$, $p < 0.05$]. We also determined the effect of LDE225 on patient-derived cell line P#5 which displays various characteristics of GBM (40, 41). As illustrated in **Supplemental Figure 2B**, LDE225 inhibited cell viability with IC₅₀ ~423 μM. Similar inhibition of CD133⁺ was observed by another Shh inhibitor cyclopamine, which binds to and inactivates smoothened protein (42) (IC₅₀ ≈ 100 μM, **Supplemental Figure 2C**). We determined the effect of Shh inhibition on self-renew capacity of CD133⁺ cells (**Figure 3D**). CD133⁺ cells were treated with LDE225 (25 μM) or vehicle and CD133⁺-derived tumor spheroids were counted at 1, 3, 7, and 14 days. Treatment with LDE225 significantly

reduced the number of tumor spheroids. A two-way ANOVA revealed a main effect of drug (LDE225 vs. vehicle) [$F_{(1,16)} = 100.9$, $p < 0.001$], interaction [$F_{(3,16)} = 44.27$, $p < 0.001$] and days after application [$F_{(3,16)} = 44.25$, $p < 0.001$] (**Figure 3E**).

Inhibition of Shh Induces Autophagy in CD133⁺ Cells

In LDE225-treated cells, the observation by transmission electronic microscope showed the appearance of large membranous vacuoles in the cytoplasm which is a characteristic feature of cells undergoing autophagy (**Figure 4A**). In addition, LC3 is distributed in the autophagosome membrane (43). The conversion of LC3-I to LC3-II is a common biomarker for autophagy activation (44, 45). We determined the conversion of LC3-I to LC3-II with anti-LC3 antibody. Immunoblotting using lysates from LDE225 (25 μM)-treated CD133⁺ cells revealed a significant increase in processed LC3-II [$t_{(5)} = 3.39$, $p < 0.05$] (**Figure 4B**). This was accompanied by the reduced expression of Mushashi-1, a RNA-binding protein selectively expressed in neural progenitor cells (46). The intensity of LC3 fluorescence punctate also increased after LDE225 treatment (**Figure 4C**). Furthermore, LDE225-induced cell death was rescued by autophagy inhibitor 3-methyladenine (3-MA) (47) (**Figure 4D**) suggesting that



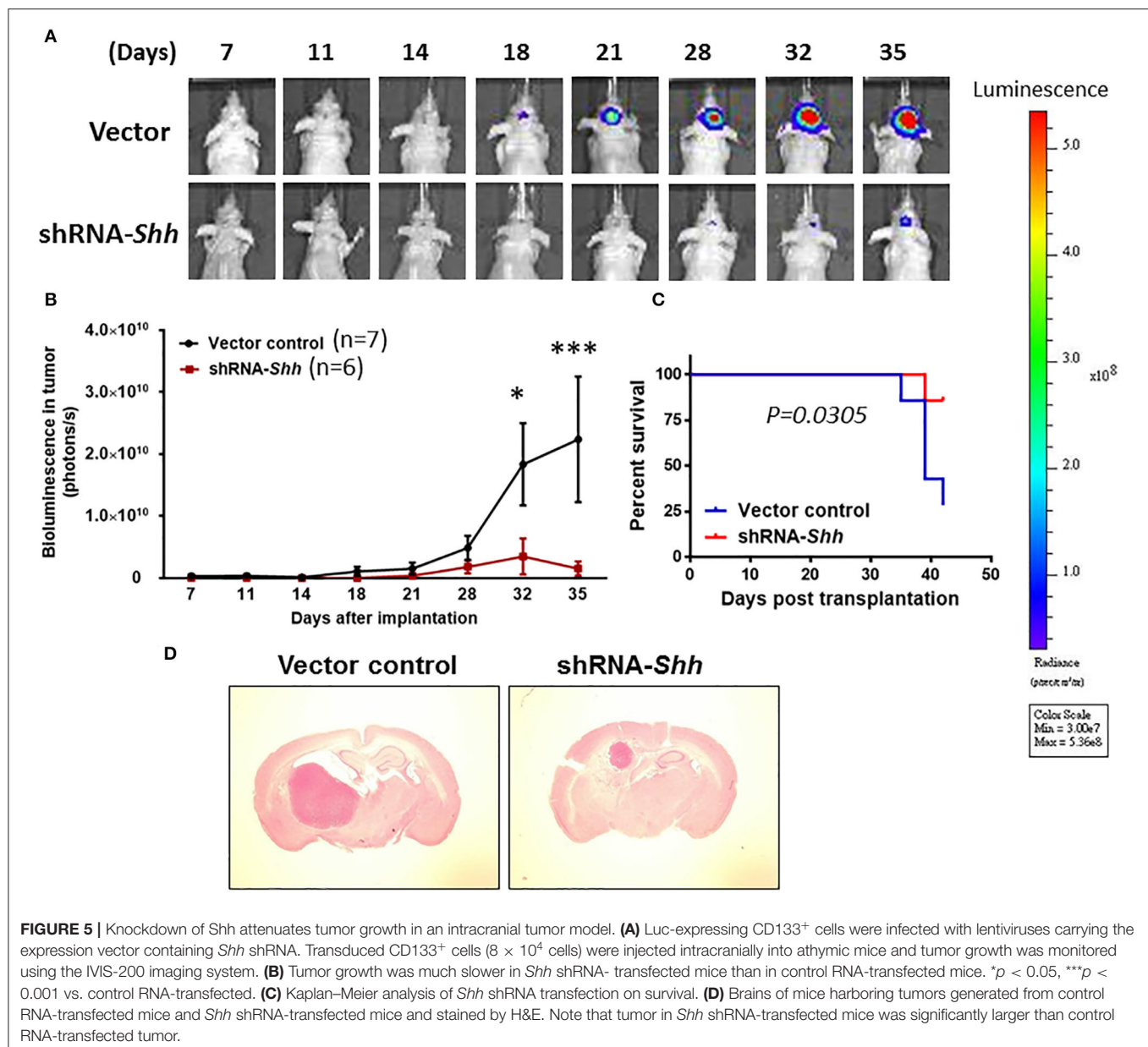
autophagy plays a critical role in LDE225-induced cytotoxicity in CD133⁺ cells.

We performed different apoptosis assays to evaluate whether LDE-induced cell death involves apoptosis. We used JC-1 mitochondrial membrane potential assay which measures the mitochondrial membrane potential as the indicator of cell health. Change in fluorescent property of JC-1 dye can be utilized to evaluate early apoptosis. CD133⁺ cells were treated with LDE225 (25 μ M) or 0.1% DMSO as vehicle control for 48 h. **Supplemental Figure 3A** shows that vehicle led to mitochondrial depolarization and decreased the ratio of J-aggregate/JC-1 monomer while that was not different between vehicle and LDE225 (25 μ M) treatment. Caspase-Glo assay also revealed that Caspase 3/7 activity was not different between LDE225 and vehicle (**Supplemental Figure 3B**). These results suggest that

LDE225-induced cell death likely was not mediated primarily through apoptosis.

Knockdown of Shh Slows Tumor Growth in an Intracranial Tumor Model

We clarified whether Shh plays a role in tumor growth under *in vivo* conditions using an orthotopic GBM model. To monitor intracranial tumor growth, we infected Luc-expressing CD133⁺ cells with lentiviruses carrying the expression vector containing *Shh* shRNA. Transduced CD133⁺ cells (8×10^4 cells) were injected intracranially into athymic mice and tumor growth was monitored using IVIS-200 imaging system. **Figure 5A** shows that tumor growth was much slower in *Shh* shRNA-knockdown mice than in control RNA-transfected mice. A two-way ANOVA revealed a main effect of group (*Shh* shRNA vs.



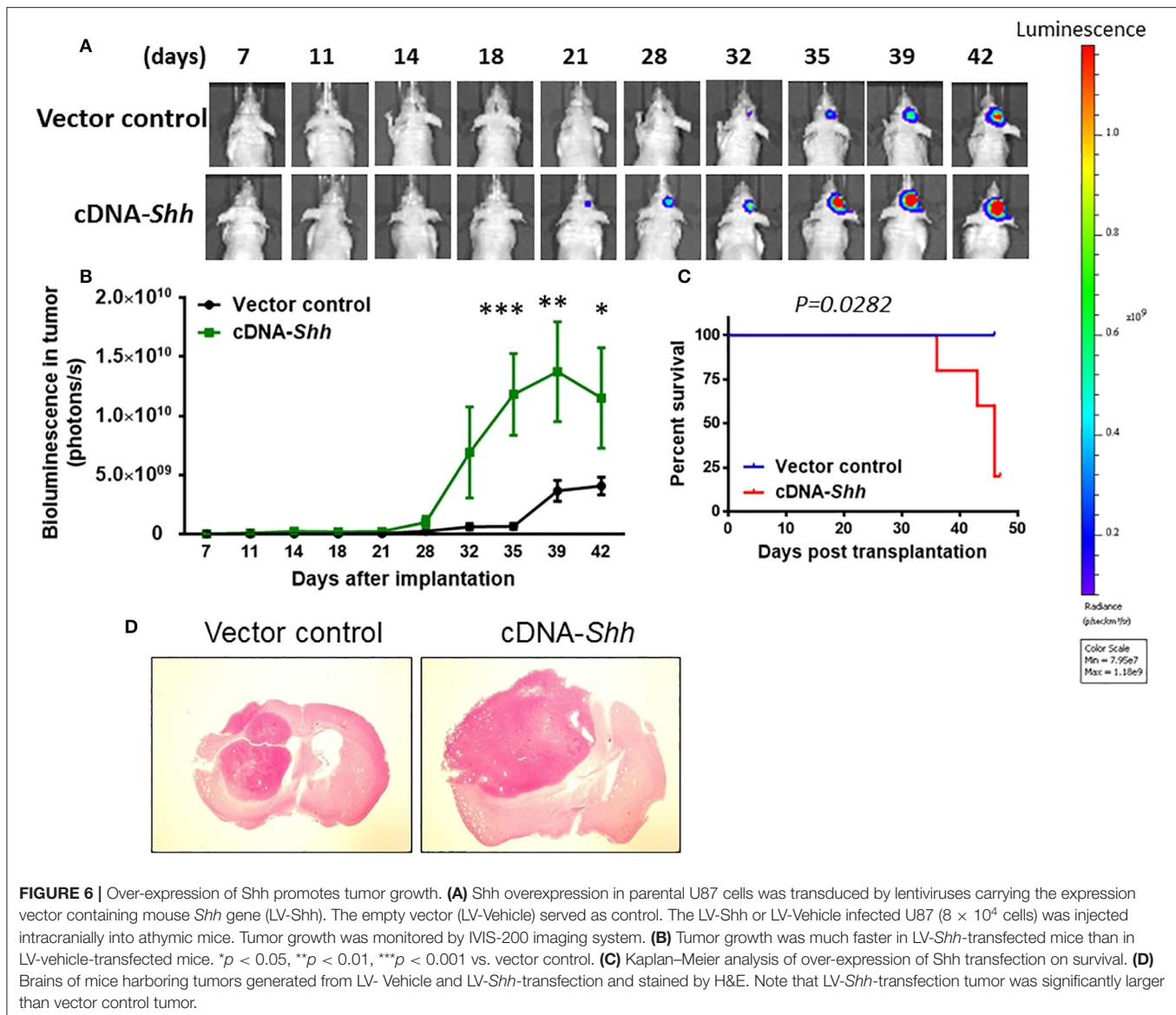
control) [$F_{(1,11)} = 4.980$, $p < 0.05$], interaction [$F_{(7,77)} = 3.110$, $p < 0.01$] and days after application [$F_{(7,77)} = 4.853$, $p < 0.001$] (**Figure 5B**). Kaplan–Meier analysis of the survival data demonstrated a statistically significant difference ($p < 0.05$) in the median survival between vector control and *Shh* shRNA-treated mice (**Figure 5C**). Brain tumor growth was analyzed in coronal brain slices by H&E staining. As shown in **Figure 5D**, in comparison with vector control, shRNA-*Shh*-transfection generated significantly smaller intracranial tumor.

To examine whether knockdown of *Shh* affected autophagy and stemness *in vivo*, we examined the expression of LC3-II in *Shh* shRNA-treated mice. **Supplemental Figure 4** shows that the levels of *Shh*, CD133, *mushashi-1* and *SOX2* were lower whereas the conversion of LC3-I to LC3-II was higher in mice

of *Shh* knockdown CD133⁺ cells compared to CD133⁺ cells of control mice. These results suggest that knockdown of *Shh* reduces stemness and GBM tumor growth.

Over-Expression of *Shh* Promotes Tumor Growth

To over-express *Shh*, we cloned the mouse *Shh* gene into parental U87 cells with lentiviruses carrying the expression vector containing *Shh* gene. The empty vector (LV-Vehicle) served as control. The LV-*Shh* or LV-vehicle transfected parental U87 GBM cells (8×10^4 cells) was injected intracranially into athymic mice. As shown in **Figures 6A,B**, tumor growth was much faster in LV-*Shh*-transfected mice than in LV-vehicle-transfected mice. A two-way ANOVA revealed a main effect



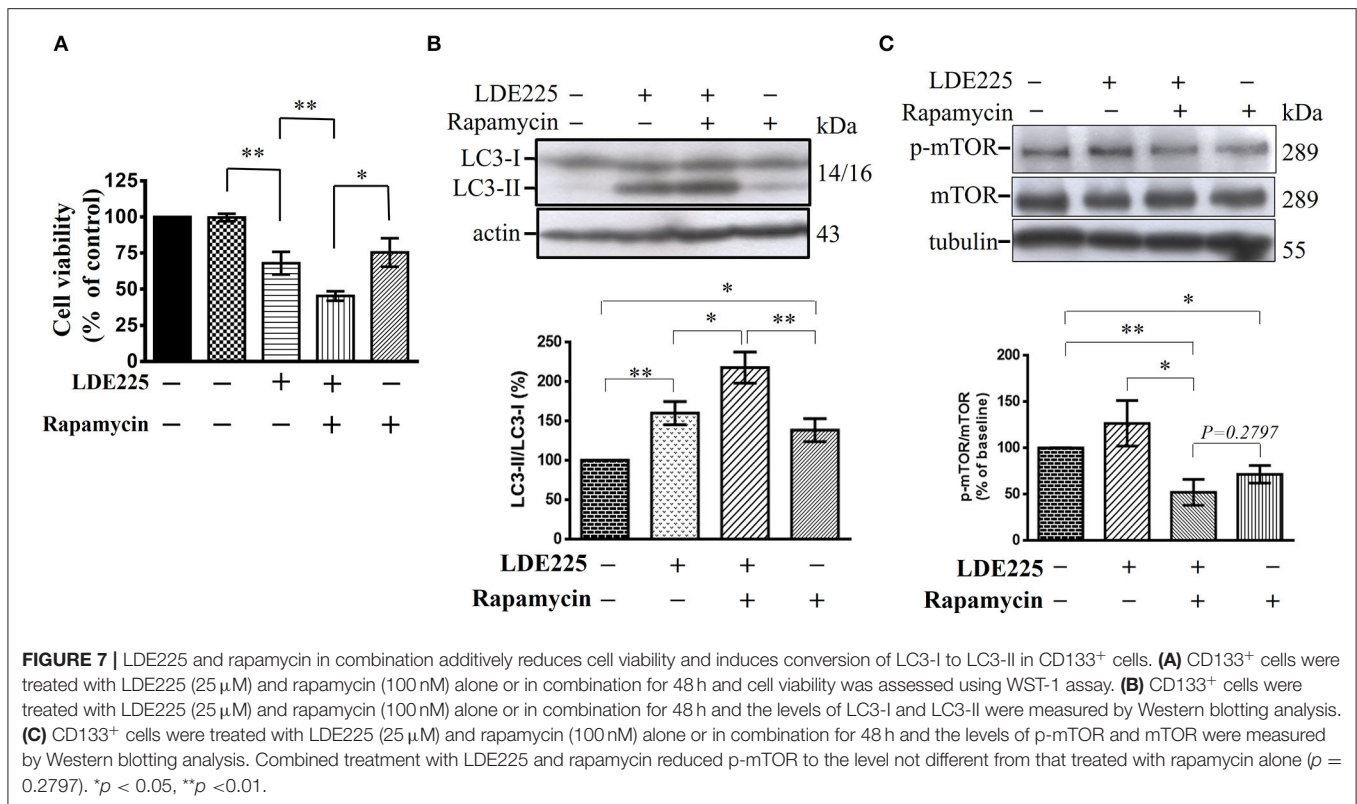
of group (LV-*Shh* vs. control) [$F_{(1,6)} = 7.665$, $p < 0.05$], interaction [$F_{(9,54)} = 4.022$, $p < 0.001$] and days after application [$F_{(9,54)} = 9.691$, $p < 0.001$]. Kaplan-Meier analysis of the survival data demonstrated a statistically significant difference ($p < 0.05$) in median survival between control and LV-*Shh*-treated mice (Figure 6C). Brain tumor size was analyzed in coronal brain slices. As shown in Figure 6D, in comparison with vector control, LV-*Shh*-transfection generated significantly larger intracranial tumor.

To examine whether over-expression of Shh affected autophagy and stemness *in vivo*, we examined the expression of LC3-II in LV-*Shh*-treated mice. Supplemental Figure 5 shows that the levels of Shh, CD133, mushashi-1 and SOX2 were higher whereas the conversion of LC3-I to LC3-II was lower in Shh over-expression GBM compared

to that of control. These results suggest that over-expression of *Shh* promotes cancer stemness and GBM tumor growth.

Additive Effect of Rapamycin and LDE225 on The Viability of CD133⁺ Cells

Autophagy is negatively regulated by the mammalian target of rapamycin (mTOR) and can be induced by the mTOR inhibitor rapamycin (48–50). To confirm our hypothesis, we tested the effect of rapamycin on CD133⁺ cells. Single treatment of CD133⁺ cells with LDE225 (25 μ M) or rapamycin (100 nM) inhibited the cell viability by $32.0 \pm 7.9\%$ ($n = 6$) and $24.6 \pm 9.9\%$ ($n = 6$), respectively. As shown in Figure 7A, combination of LDE225 and rapamycin resulted in $54.6 \pm 3.3\%$ ($n =$



6) inhibition ($p < 0.05$ vs. single treatment). Interestingly, combined treatment resulted in additive conversion of LC3-I to LC3-II. The conversion of LC3-I to LC3-II after LDE225 or rapamycin treatment was $159.8 \pm 14.7\%$ ($n = 7$) and $138.8 \pm 14.5\%$ ($n = 7$) of control, respectively. Combined treatment resulted in $217.7 \pm 19.7\%$ ($n = 7$) of control ($p < 0.05$ vs. LDE225, $p < 0.01$ vs. rapamycin) (Figure 7B).

Autophagy can also be induced through mTOR-independent pathway (50). We determined the effect of LDE225 (25 μ M) on mTOR phosphorylation. LDE225 (25 μ M) did not significantly influence the phosphorylated level of mTOR (p-mTOR) whereas rapamycin (100 nM) reduced it. Combined treatment with LDE225 and rapamycin reduced p-mTOR to the level not significantly different from treatment with rapamycin alone ($p = 0.2797$) (Figure 7C). Thus, it is likely that LDE225 induced autophagy through mTOR-independent pathway.

Amiodarone Reduces Stem-Like Cell Viability and Inhibits Tumor Formation

Amiodarone, a clinically used anti-arrhythmic drug, could induce autophagy via mTOR-independent signaling (51). We determined whether amiodarone reduced cancer stem-like cell viability *in vitro* and exhibited anti-tumor activity *in vivo*. CD133⁺ cells were treated with amiodarone for 48 h and cell viability was assessed using WST-1 assay. As

shown in Figure 8A, amiodarone dose-dependently reduced cell viability [$F_{(7,24)} = 16.71$, $p < 0.001$]. Immunoblotting using lysates from amiodarone (5 μ M)-treated CD133⁺ cells revealed a significant increase in processed LC3-II [$t_{(4)} = 4.3976$, $p < 0.05$] (Figure 8B). Furthermore, amiodarone-induced cell death was reversed by autophagy inhibitor 3-methyladenine (3-MA) (Figure 8C) suggesting that autophagy plays a critical role in amiodarone-induced cell death of CD133⁺ cells. CD133⁺ cells were treated with amiodarone (5 μ M) or vehicle and CD133⁺-derived tumor spheroids were counted at 1, 3, 7, and 14 days. Treatment with amiodarone significantly reduced the number of tumor spheroids [$t_{(4)} = 10.09$, $p < 0.001$, at 14 days culture] (Figure 8D).

We determined whether amiodarone inhibited tumor formation. Transduced CD133⁺ cells (6×10^4 cells) were injected intracranially into athymic mice and tumor growth was monitored using IVIS-200 imaging system. At day 11, amiodarone was injected intraperitoneally once per day for 8 days (80 mg/kg) into the mice and tumor growth was observed for 17 more days. Figure 9A shows that tumor growth at day 35 was much slower in amiodarone-treated mice than in control vehicle-treated mice [$t_{(8)} = 2.758$, $p < 0.05$]. Brain tumor size was analyzed in coronal brain slices. As shown in Figure 9B, in comparison with vehicle control, amiodarone-treated mice generated significantly smaller intracranial tumor.

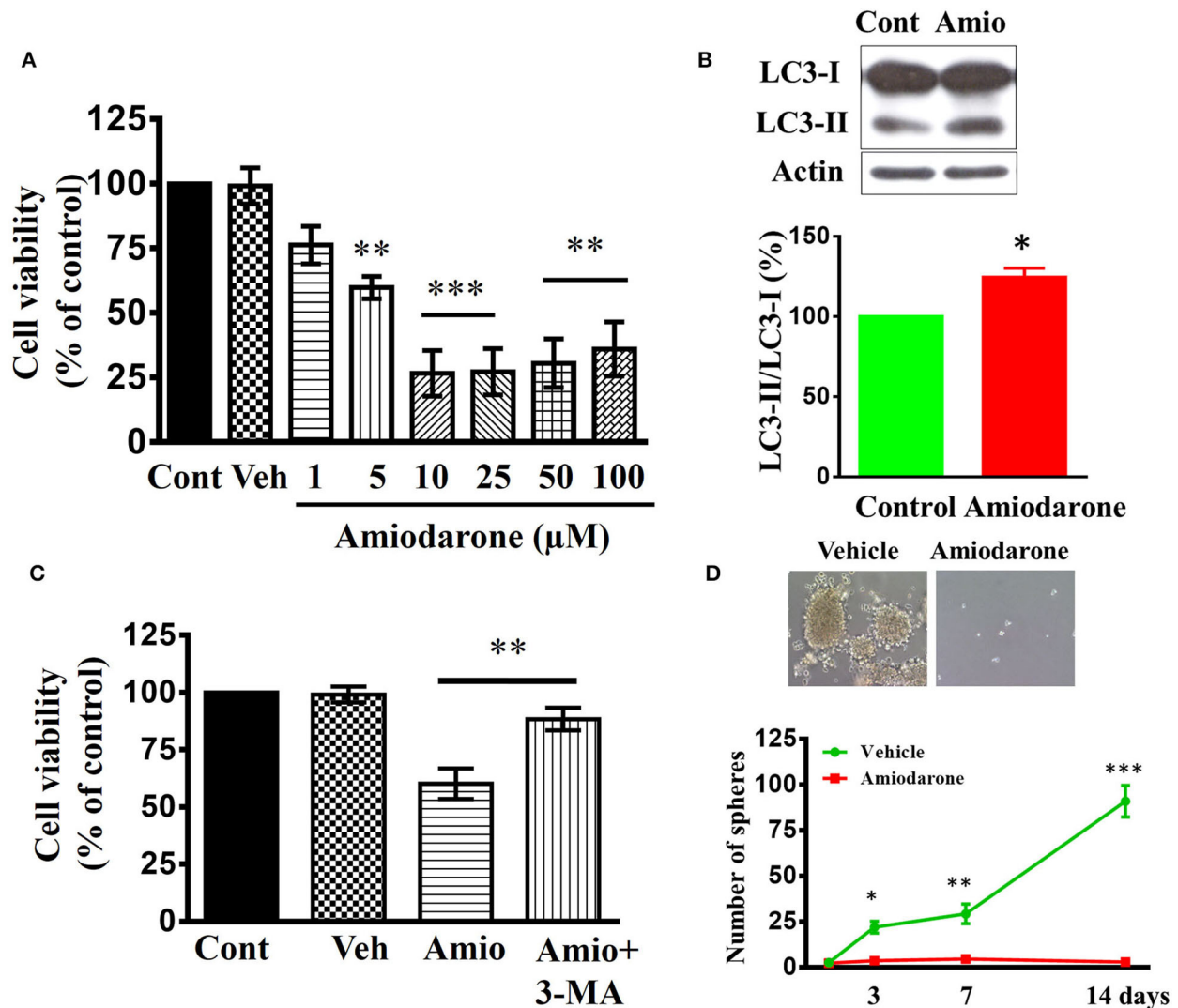


FIGURE 8 | Effects of amiodarone on the cell viability and tumor spheroid formation of CD133⁺ cells. **(A)** Concentration-dependent effect of amiodarone on the cell viability of CD133⁺ cells. ** $p < 0.01$, *** $p < 0.001$ vs. vehicle. **(B)** Western blot analysis of LC3-I and LC3-II expression in the CD133⁺ cells treated with amiodarone (5 μ M) for 48 h. **(C)** Block of amiodarone-induced cell death by autophagy inhibitor 3-MA. CD133⁺ cells were pretreated with 3-MA (3 μ M) 1 h before the exposure to amiodarone (5 μ M). Pretreatment with 3-MA attenuated amiodarone-induced cell death. ** $p < 0.01$. **(D)** Effects of amiodarone (5 μ M) on the number of tumor spheroids derived from CD133⁺ cells. Primary tumor spheroids derived from CD133⁺ were dissociated and cultured. They were then treated with amiodarone (5 μ M) for the times as indicated. The medium was replaced every 2 days in the presence of amiodarone. * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$ vs. Vehicle.

DISCUSSION

Roles of Shh Pathway in CD133⁺ Cells-Derived Tumor Spheroid Formation and Tumorigenesis

We have isolated CD133-positive cells from human U87MG (U87) GBM cells using magnetic bead cell sorting (34). CD133⁺-derived cell clones were able to grow *in vitro* in tumor spheroids and generate a tumor *in vivo* by intracranial cell injection in immunocompromised mice. The developmental signal transduction pathway mediated by Shh plays an important role during embryogenesis by regulating

body patterning, cell proliferation and differentiation (52, 53). Activation of Shh has been causatively correlated with initiation and/or maintenance of cancer (54). In the present study, using Western blotting analysis and immunofluorescent staining, we showed that CD133⁺ cells exhibited higher Shh expression compared to CD133⁻ cells. Treatment with smoothened antagonist LDE225 significantly reduced the number of CD133⁺ cells-derived tumor spheroids suggesting Shh signaling likely contributes to tumor spheroids formation *in vitro*. Silencing *Shh* gene with small hairpin interfering RNA inhibited tumor growth in an intracranial mouse model. Conversely, over-expression of *Shh* gene facilitated tumor

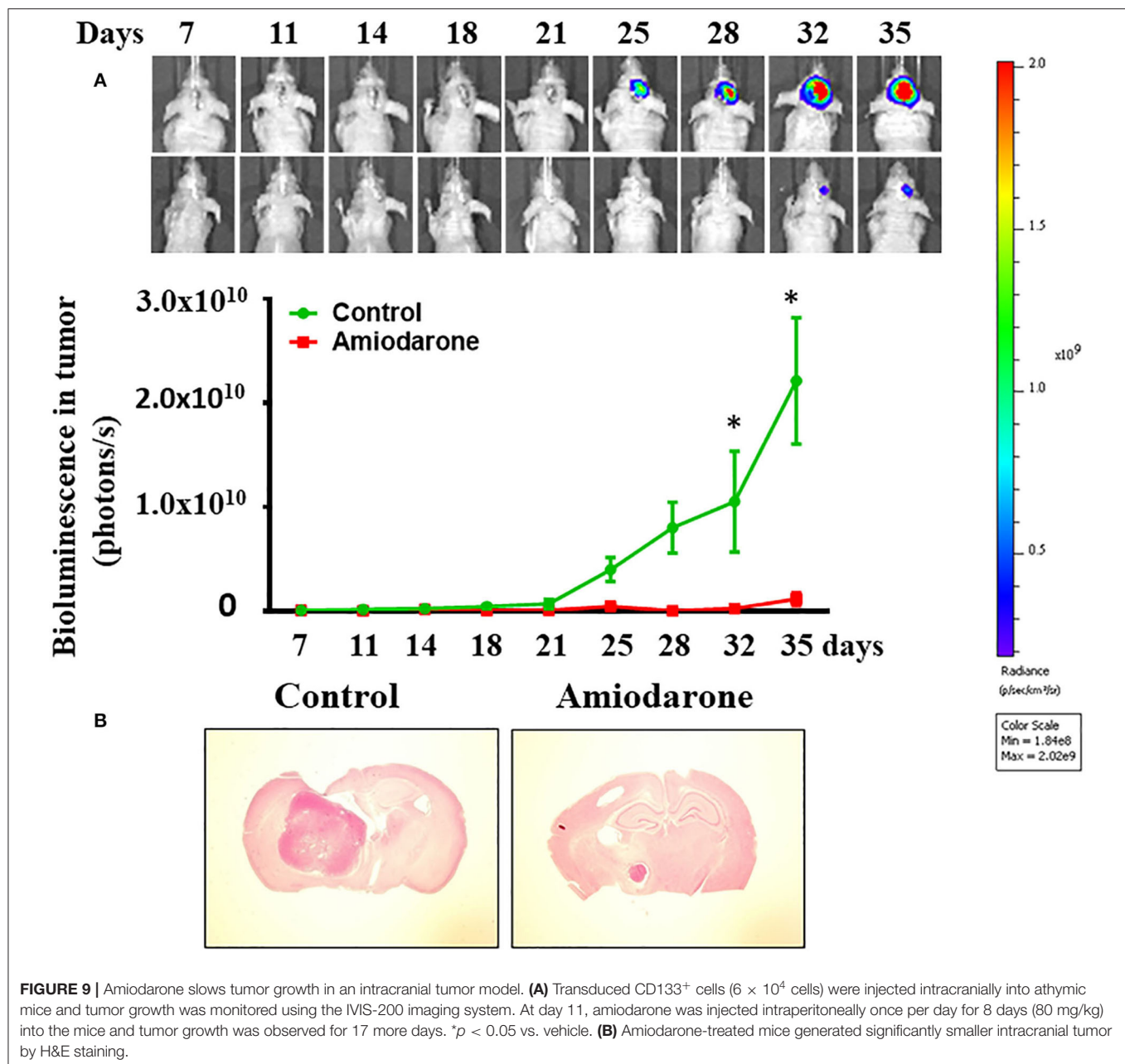


FIGURE 9 | Amiodarone slows tumor growth in an intracranial tumor model. **(A)** Transduced CD133⁺ cells (6×10^4 cells) were injected intracranially into athymic mice and tumor growth was monitored using the IVIS-200 imaging system. At day 11, amiodarone was injected intraperitoneally once per day for 8 days (80 mg/kg) into the mice and tumor growth was observed for 17 more days. * $p < 0.05$ vs. vehicle. **(B)** Amiodarone-treated mice generated significantly smaller intracranial tumor by H&E staining.

growth. Taken together, these results suggest that Shh signaling is involved in *in vitro* tumor spheroid formation and *in vivo* tumor growth.

Shh Inhibitor Reduces the Number of CD133⁺-Derived Tumor Spheroids by Inducing Autophagy

The reports about the relationship between Shh and autophagy pathway in GBM are limited. One study in SHSY5Y cells showed that cyclopamine reduced serum starvation-induced LC3 positive cells but increased the levels of cleaved caspase

3 (54). They suggested that Shh inhibitor induced cell death by reducing autophagic processes concomitant with facilitating apoptotic cell death. In the present study, we demonstrated that LDE225 induced cell death in CD133⁺ cells in a concentration-dependent manner with IC₅₀ of $\sim 20 \mu\text{M}$. Similar inhibition of cell survival was observed after application of another Shh inhibitor cyclopamine. Electron microscopy examination revealed that, after exposure to LDE225, CD133⁺ cells exhibited large membranous vacuoles in the cytoplasm that is a characteristic feature of cells undergoing autophagy. Further experiments of the conversion of LC3-I to LC3-II and attenuation of cell death by autophagy inhibitor 3-MA confirmed that

LDE225-induced autophagy in the CD133⁺ cells. This study, in contrast to previous results, suggests that Shh inhibitor produces cytotoxicity in CD133⁺ cells by inducing autophagic cell death.

To further test this hypothesis, we used rapamycin which is a mTOR inhibitor and is capable of inducing autophagy (49, 51). We found that rapamycin reduced CD133⁺ cell viability. Interestingly, the effects of LDE225 and rapamycin were additive. Furthermore, LDE225 at the concentration that induced autophagy and reduced cell viability of CD133⁺ cells did not affect the phosphorylated level of mTOR. Combined LDE225 and rapamycin failed to show further reduction of p-mTOR compared with rapamycin alone. These results suggest that LDE225 induces autophagy through mTOR-independent pathway. These findings also provide additional support that Shh regulation of autophagy plays an important role in the survival of GSCs. Thus, Shh pathway inhibitor could act as a sensitizer to increase efficiency of conventional chemotherapeutic agents in GBM by inducing cancer stem cell autophagic death.

Anti-tumorigenesis of Amiodarone

Amiodarone, a frequently prescribed anti-arrhythmic drugs in clinics, has been identified as a mTOR-independent autophagy enhancer (55, 56). If LDE225-induced cell death was mediated by the mTOR-independent autophagy, then amiodarone should produce similar effects as LDE225. Indeed, this was the case. Amiodarone reduced CD133⁺ cell viability and tumor spheroid formation *in vitro* and exhibited anti-tumor efficacy *in vivo*.

In summary, there was a small percentage of human U87 GBM cells which were CD133-positive, exhibited cancer stem cell markers and were capable of forming tumor spheroids *in vitro* and tumor *in vivo*. These CD133⁺ cells showed higher Shh expression and inhibition of Shh pathway with LDE225 reduced their survival and self-renew property. In LDE225-treated CD133⁺ cells, there appeared large membranous vacuoles in the cytoplasm and the conversion of LC3-I to LC3-II. LDE225-induced cell death was reversed by autophagy inhibitor 3-MA, suggesting that LDE225-induced autophagic cell death in CD133⁺ cells. Further, LDE225 did not affect the level of p-mTOR and combined LDE225 and rapamycin failed to show further reduction of p-mTOR when compared with rapamycin alone. Taken together, these results suggest that targeting Shh signal pathway may overcome chemoresistance and provide a therapeutic strategy for the treatment of malignant gliomas.

DATA AVAILABILITY STATEMENT

All datasets generated for this study are included in the article/**Supplementary Material**.

ETHICS STATEMENT

The animal study was reviewed and approved by the Institutional Animal Care and Use Committee of the College of Medicine, NCKU.

AUTHOR CONTRIBUTIONS

J-YC, C-LS, and P-WG contributed to the conception and design of the experiments. H-CH and C-CL performed the experiments and statistical analysis. C-LS and P-WG wrote the paper. All authors contributed to the article and approved the submitted version.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: <https://www.frontiersin.org/articles/10.3389/fonc.2020.01233/full#supplementary-material>

Supplemental Figure 1 | The differential expression of CD133 and cancer stem cell markers in parental GBM cells and tumor spheroids derived from CD133⁺ cells. **(A)** Immunofluorescence staining of CD133, SOX2, ABCG2, and AIDH1A1 in parental U87 cells. **(B)** Immunofluorescence staining of ABCG2 and AIDH1A1 in tumor spheroids derived from CD133⁺ cells.

Supplemental Figure 2 | The inhibitory effects of shh inhibition on cell proliferation. **(A)** Concentration-dependent effect of LDE225 on the cell viability of parental U87 cells. **p* < 0.05, ****p* < 0.001 vs. vehicle. **(B)** Concentration-dependent effect of LDE225 on the cell viability of patient-derived P#5 cell line. **p* < 0.05, ****p* < 0.001 vs. vehicle. **(C)** Concentration-dependent effect of cyclopamine on the cell viability of CD133⁺ cells. **p* < 0.05, ****p* < 0.001 vs. vehicle.

Supplemental Figure 3 | LDE225-induced cell death is not mediated primarily through apoptosis. CD133⁺ cells were treated with LDE225 (25 μM) or vehicle for 48 h. **(A)** JC-1 assay detect the changes of mitochondria potential which is able to measure early apoptosis. Vehicle (0.1% DMSO) led to mitochondrial depolarization with the decreased ratio of J-aggregate/JC-1 monomer. There was not different from LDE225 (25 μM) treatment. **(B)** Caspase/Glo assay revealed the activity of Caspase 3/7. There was not different between LDE225 and vehicle treatment.

Supplemental Figure 4 | The conversion of LC3-I to LC3-II was enhanced in *Shh* shRNA transfection CD133⁺-bearing mice. Tumor tissues were collected from *Shh* shRNA or vector-control transfection CD133⁺-bearing mice. **(A)** The efficiency of *Shh* shRNA-mediated knockdown of Shh was confirmed by western blot analysis. **(B–D)** The levels of CD133 **(B)**, mushashi-1 **(C)**, and SOX2 **(D)** were lower in *Shh* shRNA transfection CD133⁺-bearing mice. **(E)** The conversion of LC3-I to LC3-II was enhanced by *Shh* shRNA transfection. **p* < 0.05 vs. control.

Supplemental Figure 5 | The conversion of LC3-I to LC3-II was lower in *Shh* over-expression GBM-bearing mice. Tumor tissues were collected from LV-*Shh* or vector-control transfection GBM-bearing mice. **(A)** The efficiency of LV-*Shh*-mediated over-expression of Shh was confirmed by western blot analysis. **(B–D)** The levels of CD133 **(B)**, mushashi-1 **(C)**, and SOX2 **(D)** were higher in LV-*Shh* transfection GBM-bearing mice. **(E)** The conversion of LC3-I to LC3-II was reduced by LV-*Shh* transfection. **p* < 0.05, ***p* < 0.01 vs. control.

REFERENCES

- Jansen M, Yip S, Louis DN. Molecular pathology in adult gliomas: diagnostic, prognostic, and predictive markers. *Lancet Neurol.* (2010) 9:717–26. doi: 10.1016/S1474-4422(10)70105-8
- Soderberg-Naucler C, Rahbar A, Stragiotto G. Survival in patients with glioblastoma receiving valganciclovir. *N Engl J Med.* (2013) 369:985–6. doi: 10.1056/NEJMc1302145
- Wen PY, Kesari S. Malignant gliomas in adults. *N Engl J Med.* (2008) 359:492–507. doi: 10.1056/NEJMra0708126
- Stevens MF, Hickman JA, Langdon SP, Chubb D, Vickers L, Stone R, et al. Antitumor activity and pharmacokinetics in mice of 8-carbamoyl-3-methylimidazo[5,1-d]-1,2,3,5-tetrazin-4(3H)-one (CCRG 81045; M & B 39831), a novel drug with potential as an alternative to dacarbazine. *Cancer Res.* (1987) 47:5846–52.
- Stupp R, Mason WP, van den Bent MJ, Weller M, Fisher B, Taphoorn MJ, et al. Radiotherapy plus concomitant and adjuvant temozolomide for glioblastoma. *N Engl J Med.* (2005) 352:987–96. doi: 10.1056/NEJMoa043330
- Stupp R, Hegi ME, Mason WP, van den Bent MJ, Taphoorn MJ, Janzer RC, et al. Effects of radiotherapy with concomitant and adjuvant temozolomide versus radiotherapy alone on survival in glioblastoma in a randomised phase III study: 5-year analysis of the EORTC-NCIC trial. *Lancet Oncol.* (2009) 10:459–66. doi: 10.1016/S1470-2045(09)70025-7
- Hou LC, Veeravagu A, Hsu AR, Tse VC. Recurrent glioblastoma multiforme: a review of natural history and management options. *Neurosurg Focus.* (2006) 20:E5. doi: 10.3171/foc.2006.20.4.2
- Kioi M, Vogel H, Schultz G, Hoffman RM, Harsh GR, Brown JM. Inhibition of vasculogenesis, but not angiogenesis, prevents the recurrence of glioblastoma after irradiation in mice. *J Clin Invest.* (2010) 120:694–705. doi: 10.1172/JCI40283
- Wong ET, Hess KR, Gleason MJ, Jaeckle KA, Kyrtsis AP, Prados MD, et al. Outcomes and prognostic factors in recurrent glioma patients enrolled onto phase II clinical trials. *J Clin Oncol.* (1999) 17:2572–8. doi: 10.1200/JCO.1999.17.8.2572
- Yung WK, Albright RE, Olson J, Fredericks R, Fink K, Prados MD, et al. A phase II study of temozolomide vs. procarbazine in patients with glioblastoma multiforme at first relapse. *Br J Cancer.* (2000) 83:588–93. doi: 10.1054/bjoc.2000.1316
- Bao S, Wu Q, McLendon RE, Hao Y, Shi Q, Hjelmeland AB, et al. Glioma stem cells promote radioresistance by preferential activation of the DNA damage response. *Nature.* (2006) 444:756–60. doi: 10.1038/nature05236
- Brada M, Hoang-Xuan K, Rampling R, Dietrich PY, Dirix LY, Macdonald D, et al. Multicenter phase II trial of temozolomide in patients with glioblastoma multiforme at first relapse. *Annals Oncol.* (2001) 12:259–66. doi: 10.1023/A:1008382516636
- Frosina G. DNA repair and resistance of gliomas to chemotherapy and radiotherapy. *Mol Cancer Res.* (2009) 7:989–99. doi: 10.1158/1541-7786.MCR-09-0030
- Schonberg DL, Lubelski D, Miller TE, Rich JN. Brain tumor stem cells: molecular characteristics and their impact on therapy. *Mol Aspects Med.* (2014) 39:82–101. doi: 10.1016/j.mam.2013.06.004
- Ericson J, Morton S, Kawakami A, Roelink H, Jessell TM. Two critical periods of sonic hedgehog signaling required for the specification of motor neuron identity. *Cell.* (1996) 87:661–73. doi: 10.1016/S0092-86740081386-0
- Chiang C, Litingtung Y, Lee E, Young KE, Corden JL, Westphal H, et al. Cyclopia and defective axial patterning in mice lacking sonic hedgehog gene function. *Nature.* (1996) 383:407–13. doi: 10.1038/383407a0
- Wechsler-Reya RJ, Scott MP. Control of neuronal precursor proliferation in the cerebellum by sonic hedgehog. *Neuron.* (1999) 22:103–14. doi: 10.1016/S0896-6273(00)80682-0
- Ruiz i Altaba A. Gli proteins and hedgehog signaling: development and cancer. *Trends Genet.* (1999) 15:418–25. doi: 10.1016/S0168-9525(99)01840-5
- Kubo M, Nakamura M, Tasaki A, Yamanaka N, Nakashima H, Nomura M, et al. Hedgehog signaling pathway is a new therapeutic target for patients with breast cancer. *Cancer Res.* (2004) 64:6071–4. doi: 10.1158/0008-5472.CAN-04-0416
- Katoh Y, Katoh M. Hedgehog target genes: mechanisms of carcinogenesis induced by aberrant hedgehog signaling activation. *Curr Mol Med.* (2009) 9:873–86. doi: 10.2174/156652409789105570
- Gupta S, Takebe N, LoRusso P. Targeting the hedgehog pathway in cancer. *Ther Adv Med Oncol.* (2010) 2:237–50. doi: 10.1177/1758834010366430
- Bar EE, Chaudhry A, Lin A, Fan X, Schreck K, Matsui W, et al. Cyclopamine-mediated hedgehog pathway inhibition depletes stem-like cancer cells in glioblastoma. *Stem Cells.* (2007) 25:2524–33. doi: 10.1634/stemcells.2007-0166
- Rudin CM, Hann CL, Laterra J, Yauch RL, Callahan CA, Fu L, et al. Treatment of medulloblastoma with Hedgehog pathway inhibitor GDC-0449. *N Engl J Med.* (2009) 361:1202–5. doi: 10.1056/NEJMoa0902903
- Von Hoff DD, Lorusso PM, Rudin CM, Reddy JC, Yauch RL, Tibes R, et al. Inhibition of the hedgehog pathway in advanced basal-cell carcinoma. *N Engl J Med.* (2009) 361:1164–72. doi: 10.1056/NEJMoa0905360
- Chen J, Li Y, Yu TS, McKay RM, Burns DK, Kernie SG, et al. A restricted cell population propagates glioblastoma growth after chemotherapy. *Nature.* (2012) 488:522–6. doi: 10.1038/nature11287
- Reya T, Morrison SJ, Clarke MF, Weissman IL. Stem cells, cancer, and cancer stem cells. *Nature.* (2001) 414:105–11. doi: 10.1038/35102167
- O'Brien CA, Pollett A, Gallinger S, Dick JE. A human colon cancer cell capable of initiating tumour growth in immunodeficient mice. *Nature.* (2007) 445:106–10. doi: 10.1038/nature05372
- Hemmati HD, Nakano I, Lazareff JA, Masterman-Smith M, Geschwind DH, Bronner-Fraser M, et al. Cancerous stem cells can arise from pediatric brain tumors. *Proc Natl Acad Sci USA.* (2003) 100:15178–83. doi: 10.1073/pnas.2036535100
- Korkaya H, Wicha MS. Selective targeting of cancer stem cells: a new concept in cancer therapeutics. *BioDrugs.* (2007) 21:299–310. doi: 10.2165/00063030-200721050-00002
- Jalili A, Mertz KD, Romanov J, Wagner C, Kalthoff F, Stuetz A, et al. NVP-LDE225, a potent and selective SMOOTHENED antagonist reduces melanoma growth *in vitro* and *in vivo*. *PLoS ONE.* (2013) 8:e69064. doi: 10.1371/journal.pone.0069064
- Miller-Moslin K, Peukert S, Jain RK, McEwan MA, Karki R, Iamas LL, et al. 1-amino-4-benzylphthalazines as orally bioavailable smoothened antagonists with antitumor activity. *J Med Chem.* (2009) 52:3954–68. doi: 10.1021/jm900309j
- Singh SK, Clarke ID, Terasaki M, Bonn VE, Hawkins C, Squire J, et al. Identification of a cancer stem cell in human brain tumors. *Cancer Res.* (2003) 63:5821–8.
- Pallini R, Ricci-Vitiani L, Montano N, Mollinari C, Biffoni M, Cenci T, et al. Expression of the stem cell marker CD133 in recurrent glioblastoma and its value for prognosis. *Cancer.* (2011) 117:162–74. doi: 10.1002/cncr.25581
- Chang CH, Liu WT, Hung HC, Gean CY, Tsai HM, Su CL, et al. Synergistic inhibition of tumor growth by combination treatment with drugs against different subpopulations of glioblastoma cells. *BMC Cancer.* (2017) 17:905. doi: 10.1186/s12885-017-3924-y
- Graham V, Khudyakov J, Ellis P, Pevny L. SOX2 functions to maintain neural progenitor identity. *Neuron.* (2003) 39:749–65. doi: 10.1016/S0896-6273(03)00497-5
- Suh H, Consiglio A, Ray J, Sawai T, D'Amour KA, Gage FH. *In vivo* fate analysis reveals the multipotent and self-renewal capacities of Sox2+ neural stem cells in the adult hippocampus. *Cell Stem Cell.* (2007) 1:515–28. doi: 10.1016/j.stem.2007.09.002
- Xu J, Peng H, Zhang JT. Human multidrug transporter ABCG2, a target for sensitizing drug resistance in cancer chemotherapy. *Curr Med Chem.* (2007) 14:689–701. doi: 10.2174/092986707780059580
- Ma SA, Liu T, Jin Y, Wei J, Yang Y, Zhang H. ABCG2 is required for self-renewal and chemoresistance of CD133-positive human colorectal cancer cells. *Tumor Biol.* (2016) 37:12889–96. doi: 10.1007/s13277-016-5209-5
- Silva IA, Bai S, McLean K, Yang K, Griffith K, Thomas D, et al. Aldehyde dehydrogenase in combination with CD133 defines angiogenic ovarian cancer stem cells that portend poor patient survival. *Cancer Res.* (2011) 71:3991–4001. doi: 10.1158/0008-5472.CAN-10-3175
- Chang KY, Hsu TI, Hsu CC, Tsai SY, Liu JJ, Chou SW, et al. Specificity protein 1-modulated superoxide dismutase 2 enhances temozolomide resistance in glioblastoma, which is independent of

- O(6)-methylguanine-DNA methyltransferase. *Redox Biol.* (2017) 13:655–64. doi: 10.1016/j.redox.2017.08.005
41. Chang KY, Huang CT, Hsu TI, Hsu CC, Liu JJ, Chuang CK, et al. Stress stimuli induce cancer-stemness gene expression via Sp1 activation leading to therapeutic resistance in glioblastoma. *Biochem Biophys Res Commun.* (2017) 493:14–9. doi: 10.1016/j.bbrc.2017.09.095
 42. Chen JK, Taipale J, Cooper MK, Beachy PA. Inhibition of hedgehog signaling by direct binding of cyclopamine to smoothened. *Genes Dev.* (2002) 16:2743–8. doi: 10.1101/gad.1025302
 43. Kabeya Y, Mizushima N, Ueno T, Yamamoto A, Kirisako T, Noda T, et al. LC3, a mammalian homologue of yeast Apg8p, is localized in autophagosomal membranes after processing. *EMBO J.* (2000) 19:5720–8. doi: 10.1093/emboj/19.21.5720
 44. Klionsky DJ, Cuervo AM, Seglen PO. Methods for monitoring autophagy from yeast to human. *Autophagy.* (2007) 3:181–206. doi: 10.4161/auto.3678
 45. Li P, Du Q, Cao Z, Guo Z, Evankovich J, Yan W, et al. Interferon- γ induces autophagy with growth inhibition and cell death in human hepatocellular carcinoma (HCC) cells through interferon-regulatory factor-1 (IRF-1). *Cancer Lett.* (2012) 314:213–22. doi: 10.1016/j.canlet.2011.09.031
 46. Kaneko Y, Sakakibara S, Imai T, Suzuki A, Nakamura Y, Sawamoto K, et al. Musashi 1: an evolutionarily conserved marker for CNS progenitor cells including neural stem cells. *Dev Neurosci.* (2000) 22:139–53. doi: 10.1159/000017435
 47. Klionsky DJ, Abeliovich H, Agostinis P, Agrawal DK, Aliev G, Askew DS, et al. Guidelines for the use and interpretation of assays for monitoring autophagy in higher eukaryotes. *Autophagy.* (2008) 4:151–75. doi: 10.4161/auto.5338
 48. Fleming A, Noda T, Yoshimori T, Rubinsztein DC. Chemical modulators of autophagy as biological probes and potential therapeutics. *Nat Chem Biol.* (2011) 7:9–17. doi: 10.1038/nchembio.500
 49. Kim YC, Guan KL. mTOR: a pharmacologic target for autophagy regulation. *J Clin Invest.* (2015) 125:25–32. doi: 10.1172/JCI73939
 50. Sarkar S, Ravikumar B, Floto RA, Rubinsztein DC. Rapamycin and mTOR-independent autophagy inducers ameliorate toxicity of polyglutamine-expanded huntingtin and related proteinopathies. *Cell Death Differ.* (2009) 16:46–56. doi: 10.1038/cdd.2008.110
 51. Renna M, Jimenez-Sanchez J, Sarkar S, Rubinsztein DC. Chemical inducers of autophagy that enhance the clearance of mutant proteins in neurodegenerative diseases. *J Biol Chem.* (2010) 285:11061–7. doi: 10.1074/jbc.R109.072181
 52. Ingham PW, Nakano Y, Seger C. Mechanisms and functions of Hedgehog signalling across the metazoa. *Nat Rev Genet.* (2011) 12:393–406. doi: 10.1038/nrg2984
 53. Robbins DJ, Fei DL, Riobo NA. The hedgehog signal transduction network. *Sci Signal.* (2012) 5:re6. doi: 10.1126/scisignal.2002906
 54. Milla LA, Gonzalez-Ramirez CN, Palma V. Sonic hedgehog in cancer stem cells: a novel link with autophagy. *Biol Res.* (2012) 45:223–30. doi: 10.4067/S0716-97602012000300004
 55. Zhang L, Yu J, Pan H, Hu P, Hao Y, Cai W, et al. Small molecule regulators of autophagy identified by an image-based high-throughput screen. *Proc Natl Acad Sci USA.* (2007) 104:19023–8. doi: 10.1073/pnas.0709695104
 56. Williams A, Sarkar S, Cuddon P, Ttofi EK, Saiki S, Siddiqi FH, et al. Novel targets for Huntington's disease in an mTOR-independent autophagy pathway. *Nat Chem Biol.* (2008) 4:295–305. doi: 10.1038/nchembio.79

Conflict of Interest: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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