***In vitro* demonstration of salinomycin as a novel chemotherapeutic agent for the treatment of SOX2-positive glioblastoma cancer stem cells**

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Abstract. Glioblastoma multiforme (GBM) is the deadliest and most common form of primary brain tumor. Conventional treatments are ineffective at treating GBM due to the heterogeneous cellular makeup of the tumors as well as the existence of drug-resistant cells known as cancer stem cells (CSCs). CSCs have the ability to initiate tumorigenesis and self-renew, which can lead to recurrence. Salinomycin, an antibiotic commonly used in agricultural feed, has been revealed to target CSCs in other cancer types. A few studies have suggested salinomycin can be effective at treating glioblastoma stem cells (GSCs); however, no study has examined the effect of salinomycin treatment on GSC markers. In the present study, flow cytometry, RT-qPCR, and limiting dilution assays were used to further analyze the effects of salinomycin on GSCs. It was revealed that salinomycin decreased the expression of the GSC marker *SOX2* at both the transcriptional and translational level. However, the effect of salinomycin on the GSC markers *Nestin* and *CD133* was inconsistent between GBM subtypes. Additionally, the present findings provide initial evidence of caspase-3-dependent and independent apoptosis as the method by which salinomycin induces cell death in GBM. The present results indicated that salinomycin is an effective candidate as a chemotherapeutic agent that can treat GBM by targeting both bulk tumor cells as well as CSCs.

Introduction

Glioblastoma multiforme (GBM) is the most common and deadliest form of brain cancer (1-3). Although current treatments are rigorous, including surgery and chemotherapy, the prognosis for patients with GBM remains poor. The median survival rate is 14.6 months, and the two-year survival rate is 30% (1,4). This poor prognosis in GBM and other solid tumors is often attributed to the presence of cancer stem cells (CSCs) (5-8). This subpopulation of cells has the unique abilities to self-renew, differentiate, and initiate tumorigenesis. Due to this, CSCs play an important role in recurrence and metastasis. Glioblastoma stem cells (GSCs) have also been revealed to be resistant to conventional therapies (9,10), underscoring the need to develop treatments that account for the cellular heterogeneity found in GBM environments, specifically with regard to GSCs.

While the properties of self-renewal, differentiation, and tumorigenesis define GSCs, practically, it is easier to use GSC markers for their identification. These markers are proteins that are commonly expressed by GSCs and associated with the stem cell phenotype (11). A variety of GSC markers have been proposed and debate continues on which markers are the most accurate (11). Three of the most commonly examined GSC markers are *CD133*, *SOX2* [SRY (sex determining region Y)-box transcription factor 2], and *Nestin*. *CD133* is a cell surface glycoprotein of unknown function and was the first identified GSC marker (6,8,11). Cells that are *CD133*-positive have exhibited an increased ability to form tumors *in vivo*. *SOX2* is a marker of embryonic stem cells used in somatic reprogramming. The knockdown of *SOX2* in GSCs has been revealed to reduce their tumorigenesis and chemoresistance (12). *Nestin* is a class VI intermediate filament also found in non-malignant neural stem cells. *Nestin*-positive cells have demonstrated radioresistance and increased tumor formation (13).

One drug that has shown promise in targeting CSCs is salinomycin, a monocarboxylic polyether antibiotic isolated from *Streptomyces albus* (14). Salinomycin has been used as an anticoccidial in poultry since the 1970s, and it acts as an ionophore with a preference for Na⁺ and K⁺ (15). A high-throughput study in 2009 identified the potential of salinomycin to kill stem cell-like breast cancer cells and significantly reduce

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their tumor seeding ability in mice (14). Several other studies since then have revealed the effect of this drug on other types of CSCs including acute myeloid leukemia, lung cancer, colorectal cancer, and prostate cancer (16-18). Salinomycin has been revealed to overcome ATP-binding cassette transporters in acute myeloid leukemia, inhibit the Wnt signaling in chronic lymphocytic leukemia, and induce apoptosis in both of these cancers (19-21). The mechanism by which salinomycin exerts these capabilities remains unknown.

In a previous study, we hypothesized that salinomycin would specifically target the GSCs (22). Although several studies have suggested that salinomycin is effective against GBM, the effect of the drug on GSCs is understudied. Prior studies demonstrated that salinomycin decreases the viability of GSC-enriched cultures more than non-GSC enriched cultures, and that salinomycin inhibits neurosphere formation, a functional marker for stemness (23-25). In the present study, we corroborated these previous studies on the susceptibility of GSC-enriched cultures and neurosphere formation. Furthermore, novel evidence was provided of the ability of salinomycin to decrease the expression of the GSC marker SOX2 at both the transcriptional and translational level. Finally, preliminary evidence was provided that salinomycin-induced death of GSCs is achieved via the apoptotic pathway.

Materials and methods

Cell culture. Three human glioblastoma cell lines were used: Established glioblastoma cell line of unknown origin U87-MG (ATCC HTB-14) (26), primary glioblastoma cell line SMC448 (kindly provided and STR-profiled by Dr Do-Hyun Nam, Samsung Medical Center) (27), and proneural patient-derived xenograft GBM line D456 (kindly provided and STR-profiled by Dr G. Yancey Gillespie, Department of Neurosurgery, University of Alabama at Birmingham) (28). A mouse neural stem cell line NE-4C (ATCC CRL-2925) was also used. The cells were grown in either serum-containing or serum-free culture media. Serum-free culture cells were grown in neurobasal-A media supplemented with 1 mM glutamine (Life Technologies; Thermo Fisher Scientific, Inc.), 8 µg/ml heparin (J.T. Baker Chemical Co.), 0.5X N-2 and 0.5X B-27 (both from Gibco; Thermo Fisher Scientific, Inc.), 1% penicillin/streptomycin (Corning, Inc.), 20 ng/ml epidermal growth factor and 10 ng/ml fibroblast growth factor (both from Shenandoah Biotechnology, Inc.) (NBE media). Serum culture cells were grown in Eagle's minimal essential medium supplemented with 10% fetal bovine serum and 1% penicillin/streptomycin (Corning, Inc.) (EMEM media).

Light microscopy. VistaVision Microscope (VWR #82026-630) at x4 magnification was used to observe the morphology of cells in serum and serum-free culture. The microscope at a magnification of x10 was also used to observe sphere formation ability.

Salinomycin treatment. For all experiments, the same salinomycin treatment procedure was followed. Cells were seeded in appropriate media (NBE for serum-free culture, EMEM for serum culture) and incubated for 24 h at 37°C. After 24 h, salinomycin (MP Biomedicals, LLC) reconstituted in dimethyl

sulphoxide (DMSO) or mock control (DMSO alone) was added to a final concentration of 0.1% v/v DMSO. Cells were incubated for an additional 48 h at 37°C and then collected for analysis.

Viability assay. For the viability assay, 1×10^4 cells in 100 µl of media per well were seeded in 96-well plates. Cell Counting Kit-8 (CCK-8), a water-soluble tetrazolium salt-based colorimetric assay, was used to determine the cell viability as per the manufacturer's protocol (Sigma Aldrich-Merck KGaA). Briefly, 10 µl of CCK-8 solution per well was added to salinomycin-treated and mock-treated cells, incubated in a humidified incubator at 37°C for 4 h, and the absorbance was measured at $\lambda=450$ nm.

Flow cytometry. For flow cytometry, 3×10^5 cells in 3 ml of media per well were seeded in 6-well plates. Flow cytometry was used to assess the expression of SOX2. NE-4C, U87-MG, D456, and SMC448 were cultured in NBE (serum-free culture) or EMEM (serum culture) for 48 h with 1 µM salinomycin or mock control (DMSO). Cells were then pelleted, resuspended in 4% paraformaldehyde, then in a permeabilization buffer (BD Cytotfix/Cytoperm™; cat. no. 554722; BD Biosciences) on ice for 15 min, and stained with Alexa Fluor 647 anti-SOX2 antibody (cat. no. 562139; dilution 1:10 dilution; BD Biosciences) on ice for 30 min. Flow cytometry was then performed analyzing 20,000 events per sample using BD Accuri C6 flow cytometer (BD Biosciences). Results were analyzed using Accuri C6 software (v1.0; BD Biosciences).

Flow cytometry was also used to assess apoptosis. U87-MG, D456, and SMC448 were cultured in NBE (serum-free culture) or EMEM (serum culture) for 48 h with 1 µM salinomycin or mock control. Cells were then pelleted, dissolved in Annexin V binding buffer, and double-stained with Annexin V (Enzo Life Sciences) and 7-AAD (EMD Millipore). Flow cytometry was performed analyzing 20,000 events per sample using BD Accuri C6 flow cytometer. Specific apoptosis was calculated as established previously using the following equation: (Late apoptotic population with 1 µM Sal-late apoptotic population with 0 µM Sal)/(100-late apoptotic population with 0 µM Sal) x100.

Finally, flow cytometry was used to assess caspase-3 cleavage. Caspase-3 cleavage was determined using NucView® 488 Caspase-3 assay kit (Biotium; cat. no. 30029). The assay was conducted according to manufacturer's protocol. Flow cytometry was performed analyzing 20,000 events per sample using BD Accuri C6 flow cytometer.

Real-time quantitative PCR (RT-qPCR). For RT-qPCR, 3×10^5 cells in 3 ml of media per well were seeded in 6-well plates. Primers were designed by retrieving nucleotide sequences from the NCBI gene database for SOX2 (NM_003106), Nestin (NM_006617), Prominin1/CD133 (NM_001145847), and ACTB (NM_001101; see Table I). ACTB was used as a housekeeping gene control. Primers were synthesized by Eurofins Genomics. RNA isolation was performed using PureLink RNA Mini kit (Invitrogen, Thermo Fisher Scientific, Inc.) according to the manufacturer's protocol for mammalian cultured cells. RNA quantification was performed using NanoDrop (Thermo Fisher Scientific, Inc.). Complementary

Table I. Primer sequences.

Human	
<i>ACTB</i>	
F	CTCGTCGCCCACATAGGAA
R	AGGGCTTCTTGTCTTTCTCTC
<i>CD133</i>	
F	GGTCCCTTCTGTGAACCAAC
R	CAGATAAGTCAGCCAGGGAGC
<i>Nestin</i>	
F	GGTCCCTTCTGTGAACCAAC
R	CAGATAAGTCAGCCAGGGAGC
<i>SOX2</i>	
F	AAGCCCTGAAAGCGCAAGTCCTCAA
R	GGCAGTGGTAGTGGTGGCATTAGCAG
Mouse	
<i>ACTB</i>	
F	AAGAAGGCTATAGTCACCTCGG
R	TGGTAATAATGCGGCCGGT
<i>Nestin</i>	
F	CTGGAAGAAGTTCCCAGGCTT
R	GAAGATGTGGAAGGAGAGCGT
<i>SOX2</i>	
F	ACAGCATGTCCTACTCGCAG
R	CCTCGGACTTGACCACAGAG

F, forward; R, reverse; SOX2, SRY (sex determining region Y)-box transcription factor 2.

DNA (cDNA) was synthesized using qScript cDNA SuperMix (Quanta Biosciences) and Mastercycler nexus gradient (Eppendorf) according to manufacturers' protocols. Quantitative real-time PCR was performed using PerfeCTa SYBR Green FastMix (Quanta Biosciences) using Eco Real-Time PCR system (Illumina) according to manufacturers' protocols. The thermocycling conditions were as follows: Forty cycles of 2-step cycling were performed with denaturing at 95°C for 15 sec and annealing at 60°C for 1 min. Relative quantification was determined using the $2^{-\Delta\Delta C_q}$ method (29).

Sphere formation assay. A sphere formation assay was used to qualitatively assess stemness. Dissociated single cells were seeded at either 10 or 100 cells per well in a 96-well plate with or without the addition of 1 μ M salinomycin (n=20). Cells were incubated at 37°C for 7 days and then were evaluated for sphere formation using light microscopy at a magnification of x10.

Limiting dilution assay. A limiting dilution assay was used as a functional assay for confirming stemness (2). Dissociated single cells were plated into 96-well plates with various

seeding densities (1-10 cells/well with 20 wells per condition) in NBE media with and without the addition of 1 μ M salinomycin (n=8). Sphere formation was visually confirmed within each well. A diameter >50 μ m was used as the criterion for counting spheres.

Statistical analysis. All experiments were performed with at least triplicates for each condition. Data were analyzed by a 2-tailed t-test with equal or unequal variance and ANOVA with Dunnett's multiple comparisons test. An f-test was used to determine variance prior to the t-test or ANOVA. Data with $P < 0.05$ were considered to indicate a statistically significant difference.

Results

As demonstrated in Fig. 1A, cells grown with serum adhered to the bottom of the flask and assumed an extended morphology. Alternatively, cells grown in serum-free culture remained suspended in the media and adhered to each other, forming spheres. Such a growth environment enriches the culture for GSCs. To validate this model, GSC markers *SOX2*, *Nestin*, and *CD133* were analyzed by RT-qPCR. As revealed in Fig. 1B-D cells grown in serum-free culture had higher expression of these markers. For the U87 cell line, the difference in GSC marker expression was statistically significant for *SOX2* and *Nestin*, but not for *CD133*. SMC448 exhibited statistical significance for all three GSC markers with $P < 0.01$ for both *SOX2* and *Nestin*. D456 exhibited differences in GSC expression for all markers, albeit not at statistically significant levels.

Having established that serum-free culture enriched GSC-like cells, this culture method was used to assess the effectiveness of salinomycin against GSCs. Increasing doses of salinomycin were added to cells grown in both serum and serum-free culture. All three GBM cell lines exhibited a decreasing viability with increasing doses of salinomycin (Fig. 2). In both the U87 and SMC448 lines, the cells grown in serum-free culture had lower viabilities, suggesting the GSCs were more affected than the differentiated cancer cells in the serum culture. However, lower viabilities were observed in serum culture than in serum-free culture for the D456 cell line at all concentrations of salinomycin except at the highest 5 μ M. This result was unexpected and may indicate the importance of the GBM subtype on the effectiveness of salinomycin. It could also be explained by the lack of a statistically significant difference in GSC markers between D456 cells grown in serum vs. serum-free culture. The non-cancerous neural stem cell line NE-4C was also investigated as a negative control (Fig. 2A). However, NE-4C cells exhibited significant toxicity to salinomycin in both serum and serum-free culture. This finding highlights the importance of understanding the mechanism of action of salinomycin and its effect on the GSC population.

To more thoroughly assess the effect of salinomycin on the GSC population, the mRNA expression of the GSC markers *SOX2*, *Nestin*, and *CD133* was evaluated. Cells grown in sphere culture were treated for two days in triplicate with either 1 μ M salinomycin or 0.1% DMSO. Salinomycin treatment significantly decreased the *SOX2* expression in U87 and SMC448 cells and resulted in a non-statistically

Figure 1. Serum-free culture enriches GSCs. GBM cells were grown in both serum and serum-free media and their expression of GSC markers were evaluated. (A) Representative bright-field images of NE-4C, U87, SMC448, and D456 cells after four days of growth in serum-free and serum-containing media exhibiting marked differences in cell morphology. Scale bar, 100 μ m. (B-D) RT-qPCR analysis of the gene expression of (B) *SOX2*, (C) *Nestin*, and (D) *CD133* for NE-4C, U87, SMC448, and D456 GBM cells. Error bars represent 95% confidence intervals (n=3). * $P < 0.05$, ** $P < 0.01$ for each gene compared to the serum culture counterpart. These results indicated increased stemness characteristics when GBM cells were grown in serum-free media. GSCs, glioblastoma stem cells; GBM, glioblastoma; *SOX2*, SRY (sex determining region Y)-box transcription factor 2.

significant decrease in *SOX2* expression in D456 cells (Fig. 3A). However, the effect of salinomycin on *Nestin* and *CD133* was inconsistent (Fig. 3B and C). Salinomycin increased the expression of *Nestin* in U87 cells, decreased its expression in SMC448 cells, and had no observable effect on its expression in D456 cells. None of these results were statistically significant. For *CD133*, salinomycin caused a statistically significant increase in U87 cells but minimal changes in SMC448 and D456. The variance in marker expression brings into question whether these markers are in fact assessing the same types of cells.

To further investigate the effect of salinomycin on GSCs, the protein expression and GSC functionality were assessed. Since salinomycin caused a decrease in *SOX2* mRNA expression for all three GBM cell lines, *SOX2* protein expression was analyzed using flow cytometry (Fig. 4). A statistically significant decrease in *SOX2*-positive population in D456 expression, with a similar decrease in U87 and SMC448 cells (albeit not at statistically significant levels) was revealed. Placed in context with the transcriptional results,

these data indicated that while most of the cells continue to express *SOX2*, salinomycin is reducing the total amount of *SOX2*-positive population. The functional impact of salinomycin on GSC clonogenicity was assessed via sphere formation assay. Qualitatively, 1 μ M of salinomycin was sufficient to prevent sphere formation at both 10 and 100 cells/well seeding densities, even though these densities were otherwise sufficient to form spheres without salinomycin (Fig. 5A). Quantitatively using the extreme limiting dilution assay (30), a significant reduction of clonogenic population was also observed. Clonogenic potential of U87 cells was altered from 1 of 1.04 to 1 of 8.58 cells after 1 week of 1 μ M of salinomycin treatment (Fig. 5B). Similarly, the clonogenic potential of SMC448 was altered from 1 of 1.76 to 1 of 6.84 and D456 changed from 1 of 1.04 to 1 of 7.01. Notably, the clonogenic potential of NE-4C mouse neural stem cells was also altered but to a lower extent from 1 of 1.15 to 1 of 4.63 (comparison of slopes of lines in Fig. 5B). Collectively, these data indicated that salinomycin significantly and selectively impacted GSC clonogenicity.

Figure 2. Salinomycin depletes populations of GBM cells. Cell viabilities of (A) NE-4C, (B) U87, (C) SMC448, and (D) D456 treated with varying concentrations of salinomycin were measured after 48 h (n=8, mean $\pm$ 95% confidence intervals). *P<0.05, **P<0.01 between serum and serum-free conditions. GBM, glioblastoma.

Figure 3. Effect of salinomycin on GSC marker mRNA expression. The ability of salinomycin (1 μ M) or mock control (0 μ M) to decrease the expression of the GSC markers (A) *SOX2*, (B) *Nestin*, and (C) *CD133* at the mRNA level after 48 h of treatment was determined using RT-qPCR. *ACTB* was used as the housekeeping gene. Expression levels were normalized to those of D456 control (n=3, mean $\pm$ 95% confidence intervals). *P<0.05. GSC, glioblastoma stem cell; *SOX2*, SRY (sex determining region Y)-box transcription factor 2.

To investigate the mechanism of action of salinomycin, an Annexin V/7-AAD assay was performed (Fig. 6A). This assay assesses the percentage of cells that are healthy (lower left quadrant), early apoptotic (lower right quadrant), late apoptotic (upper right quadrant), and necrotic (upper left quadrant). For all cell lines, salinomycin treatment led to an increased percentage of cells in the early apoptotic and late apoptotic quadrants, but no increase in the necrotic quadrant. This finding indicated that salinomycin induced cell death via

apoptosis rather than necrosis. The cells grown in serum-free culture, which are GSC enriched, led to higher rates of specific apoptosis than those grown with serum, reinforcing the prior observation that salinomycin can to a certain extent preferentially deplete GSCs. For U87, the difference in the apoptotic rates between the serum-free culture and serum culture was statistically significant. The GBM lines U87, SMC448, and D456 all exhibited higher rates of specific apoptosis than the non-cancerous NE4C cell line, providing evidence that

Figure 4. Effect of salinomycin on SOX2 protein expression. Percentage of NE-4C, U87, SMC448, and D456 serum-free cultured cells positive for SOX2 was analyzed via flow cytometry after 48 h of treatment with 1 μ M salinomycin (gray) or mock control (black). (A) Representative histograms for each cell line. Events to the right of the vertical line indicate SOX2-positive cells and (B) their quantification ($n=3$, mean $\pm$ 95% confidence intervals). * $P<0.05$. SOX2, SRY (sex determining region Y)-box transcription factor 2.

salinomycin has greater toxicity for cancer cells than their non-cancerous counterparts. To further investigate this potential mechanism, flow cytometry was used to identify caspase-3 cleavage activity in serum-free and serum culture cells with or without the addition of salinomycin (Fig. 6B). Salinomycin treatment led to statistically significant increases in caspase-3 cleavage for SMC448 cells in serum culture as well as for both SMC448 and D456 cells in serum-free culture. The cells in serum and serum-free culture exhibited similar trends in caspase-3 cleavage, indicating that the increased apoptosis in U87 serum-free vs. serum culture may be caused by a caspase-3-independent pathway (31,32).

Discussion

We have previously postulated the potential of salinomycin to selectively target GSCs (20). While salinomycin is yet to be the subject of any full-scale clinical trials in cancer patients, its emergence as a key anticancer agent has propelled it into the focus of pre-clinical models using patient-derived cancer

cells (33). The aim of the present study was to investigate the impact of salinomycin on GSC-like cells in GBM to assess the efficacy of the drug as a potential chemotherapy for GBM patients. GSCs are a distinct subpopulation of cells within glioblastomas which have the ability to initiate tumor growth and self-renew (6,8,11). Conventional therapies do not adequately target these cell populations in glioblastoma, which may lead to inefficient tumor elimination and cancer reemergence (11,34). The antibiotic salinomycin had been first identified to target CSCs nearly a decade ago, and subsequent studies on salinomycin have been centered on its potential to treat similar stem cell-like populations in various cancers (14). Additionally, drug delivery systems have been implemented *in vitro* with salinomycin to demonstrate the ability of the drug to deplete glioblastoma populations while also crossing the blood-brain barrier (35).

In the present study, the potential of salinomycin as a chemotherapeutic agent in the treatment of SOX2-positive GSCs was demonstrated. GSC populations within the tumor microenvironment must be targeted to adequately eliminate the tumor. One established human glioblastoma cell line (U87-MG) and two patient-derived glioblastoma cells lines (SMC448 and D456) were used in the present study. U87-MG is a well-established and widely used glioblastoma cell line of unknown origin that has been in culture since the 1960s (26). SMC448 is a radioresistant xenograft cell line derived from a high-grade glioma of unknown subtype (27). D456 is a glioblastoma of the proneural subtype that was derived from a human pediatric fronto-parietal GBM directly implanted into the flank of immunocompromised mice (28). Additionally, the mouse neural stem cell line NE-4C was used as a non-cancerous control. The present results revealed that salinomycin depleted the SOX2-positive GSC-enriched cell populations. Notably, the GSC markers *Nestin* and *CD133* did not demonstrate the same consistent trend. This finding could indicate that *SOX2*, *Nestin*, and *CD133* are not measuring the same GSC population. This conclusion could be a result of a heterogeneous GSC population or one or more of these markers not correctly identifying GSCs. An alternative conclusion could be that protein expression or conformation but not mRNA expression is indicative of the GSC population. Such a conclusion has been previously suggested for the detection of CD133 in colon cancer CSCs (36). Salinomycin was also demonstrated to decrease neurosphere formation and decrease the clonogenicity of GSC-like cells. Finally, it was demonstrated that salinomycin induced apoptotic cell death in GBM cells.

Salinomycin has previously been linked to apoptotic activity in leukemia, prostate, breast, and ovarian cancer (16,17,37,38). There is great potential for targeting CSCs, including manipulating pro- and anti-apoptotic pathways in CSC populations (39). Salinomycin has additionally been demonstrated to overcome anti-apoptotic cellular mechanisms in human cancer cells (20). In the case of ovarian cancer specifically, it is known that the salinomycin-induced apoptotic pathway relies on the activation of caspase-8 and death receptor 5 (40). Salinomycin has also been demonstrated to enhance the apoptotic activity of TRAIL in TRAIL-resistant GBM cells by modulating caspase-3 expression (41). The apoptotic effects of salinomycin have been shown to be retained even in biodegradable drug-delivery systems, an indication that this

Figure 5. Salinomycin reduces clonogenicity. (A) Representative bright-field images of U87 cells seeded with either 10 or 100 cells/well and treated with either a mock control (0 μ M) or with 1 μ M salinomycin qualitatively revealing the loss of sphere formation after 1 week (scale bar, 50 μ m). (B) Limiting dilution assay of NE-4C, U87, SMC448, and D456 was performed after 1 week of culture with 1 μ M salinomycin (Sal) and without (Ctrl). The steepness of the slope represents the frequency of clonogenic stem cells (dotted lines represent 95% confidence intervals). There was a significant reduction of clonogenic stem cells with salinomycin treatment ($P < 0.05$).

Figure 6. Salinomycin induces apoptosis in GBM. (A) Annexin V/7-AAD apoptosis assay was performed in NE-4C, U87, SMC448, and D456 serum and serum-free cultured cells. Representative flow cytometric plots of U87 cells with (1 μ M) and without (0 μ M) 48 h treatment of salinomycin (left) and quantified specific apoptosis (right). The cells in serum-free culture (GSC-enriched) exhibited higher specific apoptosis than those grown in serum culture (differentiated cells) ($n=3$, mean $\pm$ 95% confidence intervals). $^{**}P < 0.01$. (B) Caspase-3 cleavage was determined for all cell lines cultured in serum (left) and serum-free (right) conditions ($n=3$, mean $\pm$ 95% confidence intervals). $^{*}P < 0.05$, $^{****}P < 0.0001$. GBM, glioblastoma; GSC, glioblastoma stem cell.

pathway could serve as a point of interest in more advanced pre-clinical trials (42). Future study on salinomycin as a GBM treatment option should regard the apoptotic pathway as a factor that could help differentiate GSC death from normal neural cell death. Future studies should assess the effect of salinomycin on GBM *in vivo* to better understand its potential therapeutic benefits before translation to clinical testing.

In the development of new drugs to treat glioblastomas, it is important to consider the impact of these novel treatments on the function of normal brain tissue. Cases of neural toxicity induced by accidental salinomycin overdoses have been documented in humans and animals previously (43-45). Although our findings indicated that salinomycin decreased the sphere formation potential for the non-cancerous stem cell line NE-4C, they also revealed that salinomycin toxicity took effect at a higher concentration for this neural stem cell line than the GBM cell lines assessed. Additionally, the shifts in apoptotic activity induced by salinomycin were not significant for NE-4C where large shifts were observed in the glioblastoma lines. It has also been observed that salinomycin triggers calcium-induced apoptosis in Schwann cells and dorsal root ganglia, but this effect can be inhibited by $\text{Na}^+/\text{Ca}^{2+}$ exchanger inhibitors (46). A potential avenue that may prove useful for subsequent salinomycin studies is the implementation of synthetic structural analogs of the drug that exhibit lower levels of toxicity towards healthy cells (47). Salinomycin has also been demonstrated to have synergistic DNA damage effects when combined with the common chemotherapy drug temozolomide, highlighting the potential of salinomycin in combination therapies (48). As the drug is researched more thoroughly within glioblastoma studies and other cancers, our findings reaffirm that salinomycin toxicity towards healthy cells must be a contributing factor towards the implementation of the ionophore as a chemotherapeutic agent.

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Availability of data and materials

All data generated or analyzed during this study are included in this published article.

Authors' contributions

JWM and YK conceived and designed the study. JWM and WRR conducted the experiments and acquired the data.

JWM, WRR and YK were involved in the experimental design, data analysis, interpretation of data, and drafted and revised the manuscript. All authors read the final manuscript and agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.

Ethics approval and consent to participate

Not applicable.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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