Isoliquiritigenin inhibits the proliferation and induces the differentiation of human glioma stem cells

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Abstract. Glioma stem cells (GSCs) have been proven to be resistant to various therapeutic strategies, such as temozolomide chemotherapy and radiotherapy, leading to glioma recurrence. Isoliquiritigenin (ISL), a member of the flavonoids isolated from liquorice has been found to be a potent stimulator of cell differentiation and has potential application for treating various types of cancer including human brain glioma. However, the antitumor activity of ISL on GSCs and the signaling pathway underlying its therapeutic effects are poorly understood. In the present study, GSCs were isolated from SHG44 human glioma cells by serum-free culture and treated with ISL or DAPT (a Notch/γ-secretase inhibitor). It was found that ISL dose-dependently inhibited GSC growth after 72 h of treatment and decreased the formation of tumor spheres. Meanwhile, GSCs differentiated into astrocytes and neurons. Furthermore, these therapeutic effects were accompanied by downregulation of Notch1 and Hes1 at the protein and mRNA levels. Taken together, our results demonstrated that ISL exhibits antitumor effects on GSCs by inhibiting proliferation and inducing differentiation. The therapeutic effect may be related to downregulation of the Notch1 signaling pathway. Application of ISL presents potential benefits for the treatment of human brain glioma.

Introduction

Eighty percent of malignant tumors that develop in the central nervous system (CNS) are malignant gliomas, which display histological similarities to glial cells, including astrocytes and oligodendrocytes (1). Glioblastoma multiforme (GBM) is the most common primary malignant adult brain tumor. GBM patients have a median survival of 13-16 months even following therapy (2). The current standard of care consists of maximal safe surgical resection with a combination radiotherapy and adjuvant temozolomide chemotherapy. However, the patient 5-year survival rate is <10% (3). Malignant gliomas are aggressive brain tumors with limited therapeutic options, possibly because of the highly tumorigenic subpopulations of glioma stem cells (GSCs) (4). GSCs have been proven to be resistant to various chemotherapeutic agents, such as temozolomide, the first-line treatment for glioblastoma, allowing these cells to survive therapy, leading to radioresistance and tumor relapse (5,6). As the current treatment modalities of GBM are not specifically designed to kill GSCs and the effectiveness is limited, new agents targeting GSCs for potential use in GBM clinical therapy have attracted attention.

Isoliquiritigenin (ISL), extracted from Glycyrrhiza spp., is a promising candidate as a natural, alternative method of cancer therapy (7). It is a flavonoid compound with efficient antitumor activity (8). In particular, many studies both in vitro and in vivo, have demonstrated the antitumor efficacy of ISL against various types of cancer (9-12). It has been suggested that ISL can inhibit proliferation and induce apoptotic and necrotic cell death of glioma cells (13). In the present study, the antitumor activity of ISL on human GSCs was examined.

Notch signaling has been reported to be involved in fetal development and tissue homeostasis by directing many cellular functions, including cell growth and differentiation, cell fate determination and regulation of stem cell maintenance (14). Notch signaling pathway is an attractive target for cancer therapy as targeting Notch signaling could overcome multi-drug resistance (MDR) (15). It was found that a Notch/γ-secretase inhibitor (GSI) in combination with radiotherapy and temozolomide attenuated proliferation, decreased glioma stem cell markers and resulted in a marked reduction in clonogenic survival in primary and established glioma cell lines (16). Here, we hypothesized that the anticancer effects of ISL are related to the Notch1 signaling pathway. The present study proposes the possibility of the therapeutic application and an intracellular antitumor mechanism of ISL in GSC-targeted treatment for human brain glioma.

Materials and methods

Isolation of GSCs. SHG44 human glioma cells from the American Type Culture Collection (ATCC, Manassas, VA, USA) were cultured in Dulbecco’s modified Eagle’s medium...
(DMEM) supplemented with 10% fetal bovine serum (FBS), 100 U/ml penicillin and 100 µg/ml streptomycin at 37°C in a humidified incubator with 5% CO₂. Tumor cells grown to 80% confluence were rinsed with sterile phosphate-buffered saline (PBS) (0.01 M, pH 7.4) and digested at 37°C with 0.25% trypsin for 1 min, then resuspended in a serum-free neural stem cell medium consisting of D-MEM/F-12, 2% B27 (Gibco BRL, Gaithersburg, MD, USA), 20 ng/ml EGF (PeproTech, Inc., Rocky Hill, NJ, USA), 20 ng/ml bFGF (PeproTech, Inc.), and then seeded into 6-well suspension cell culture plates (Beaver, China). After 7 days of culture at 37°C in 5% humidified CO₂ atmosphere, the medium was removed by centrifugation and GSC-forming spheres were digested with 0.25% trypsin into a single-cell suspension. GSCs were subcultured and enriched three times before being used for assays.

Identification of surface phenotypes of GSCs. GSCs grown on polylysine-coated coverslips were fixed using 4% paraformaldehyde for 15 min and permeabilized with 0.5% Triton X-100 (PBST) for 10 min. The coverslips were blocked with 5% bovine serum albumin (BSA) in PBS (0.01 M, pH 7.4) for 30 min at room temperature, followed by incubation with primary antibodies targeted against rabbit anti-CD133 (#ab121663; 1:200; Abnova), mouse anti-nestin (#ab18102; 1:200; Abcam) and rabbit anti-Bcl-2 (#ab32124; 1:100; Abcam) overnight at 4°C. After three washes with PBS, the coverslips were incubated with PE-labeled rabbit anti-mouse IgG (#ab7000; 1:100; Abcam) and FITC-labeled goat anti-rabbit IgG (#ab6717; 1:1000; Abcam) for 120 min at 37°C. Nuclear DNA was labeled in blue with DAPI. The staining results were imaged on a Nikon C-1 laser-scanning confocal microscope (Nikon, Tokyo, Japan).

Quantitative real-time PCR (qPCR). The cultured cells were harvested and lysed at 48 h after ISL and DAPT treatment. Total cellular RNA was extracted using TRIzol reagent (Takara Biotechnology Co., Ltd., Dalian, China). After removing any DNA contamination, RNA samples were treated with Recombinant DNase I (Takara Biotechnology). Single-stranded cDNA was synthesized from total RNA with the PrimeScript™ RT Master Mix (Takara Biotechnology). qPCR was performed for HES1 and internal control GAPDH using the SYBR Premix Ex Taq™ II (Takara, Shiga, Japan) on a 7300 real-time PCR system (Applied Biosystems, Singapore, Singapore). All reactions were carried out in triplicate and expression data (after being calibrated with GAPDH levels) were analyzed using the 2^(-ΔΔCt) method. Optimal oligonucleotide primers used in the above PCR assays were designed and synthesized by Takara Biotechnology based on published human cDNA sequences. The primer sequences were as follows: HES1 sense 5'-GTGT CAACACGACCGGATAAC-3' and antisense 5'-CAGA GTGTCGGCTCTTCCAGC-3'; GAPDH sense 5'-GCACCGT CAAGGCTGAGAAC-3' and antisense 5'-TGGTGAAAGAC GCCAGTGGA-3'. All reactions were run in triplicate.

Western blot analysis. The protein expression levels of differentiation markers (GFAP and β III tubulin) and Notch1 signaling pathway markers (Notch1 and Hes1) were examined by western blotting. β-actin levels were evaluated as a loading control. Briefly, cells were lysed and homogenized in RIPA lysis buffer (Solarbio, Beijing, China). The supernatants were obtained by centrifugation at 4°C for 10 min at 12,000 rpm and the total protein contents were quantified using a BCA kit (Solarbio). For electrophoresis, equal amounts of protein (15 µg/lane) were separated by sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) analysis on 8, 10 or 12% polyacrylamide gel depending on the target protein and transferred onto PVDF membranes (Millipore, Billerica, MA, USA) using a wet electroblotting system (Mini Trans-Blot; Bio-Rad). For immunoblotting, the membranes were blocked in 5% non-fat dry milk for 2 h at room temperature and incubated overnight at 4°C with one of the following primary antibodies: GFAP (#ab4648; 1:400; Abcam), β III tubulin (#ab78078; 1:800; Abcam), Notch1 (#ab128076; 1:800; Abcam), Hes1 (#ab19776; 1:800; Abcam) and β-actin (#ab0060; 1:3,000; Bioworld Technology Inc.). After three washes in Tris-buffered saline.
saline containing 0.5% Tween-20 (TBST), the samples were treated with HRP-conjugated goat anti-rabbit (#E030120-01; 1:10,000; EarthOx Life Science, Millbrae, CA, USA) or goat anti-mouse secondary antibodies (#ab6728; 1:10,000; Abcam) for 1 h at room temperature. The antibody labeling was visualized using enhanced chemiluminescence reagent (Millipore) and exposed on X-ray film (Kodak, China). For the immunoblot analyses, densitometry was performed with Image-Pro Plus 6.0 software. The optical densities (ODs) of the protein bands were calculated relative to the ODs of the reference protein β-actin.

Statistical analysis. Data are expressed as the mean ± SD. All statistical analyses were performed using the SPSS software program (version 21.0; IBM Corp., Armonk, NY, USA). The differences between the experimental groups were analyzed by one-way analysis of variance (ANOVA). P<0.05 was considered statistically significant.

Results

Glioma cells and GSC-forming spheres. Within 48-72 h of serum-free culture SHG44 human glioma cells yielded a minority fraction of cells that possessed extensive proliferative and self-renewal potential, and generated free-floating neurosphere-like brain tumor spheres. These spheres gradually enlarged as the time of the culture increased (Fig. 1B and C). In comparison, glioma cells cultured in DMEM with 10% FBS grew adherently (Fig. 1A).

Characterization of surface phenotypes of GSCs. Expression of GSC surface markers were examined in human GSC-forming spheres. As shown in Fig. 1D, the cells expressed characteristic neural stem cell (NSC) markers CD133 and Nestin. SHG44 human glioma cells and tumor neurospheres were positive for the proto-oncogene Bcl-2 which distinguished them from normal neurons and NSCs (Fig. 1E and F). The immunofluorescence staining data indicated that the tumor spheres expressed typical surface markers of GSCs and thus were used for experiments as described below.

ISL inhibits cell proliferation in a dose-dependent manner. To investigate the influence of ISL on GSC proliferation, we firstly examined the activities of cells treated with various concentrations of ISL for different times. As shown in Fig. 2A, within 24 and 48 h of treatment the proliferative activity of the cells was significantly higher in the 10, 20 and 40 µM ISL groups than that noted in the 0 µM ISL group. There was no significant difference between the 0 and 80 µM ISL group within 24 h, but the activity of cells began to decline after 48 h. After 72 h, the cell proliferation of the ISL treatment groups were obviously inhibited in a dose-dependent manner, with a half inhibitory concentration IC50 of 102.744 µmol/l.
Formation of neurospheres is a morphological feature of GSCs. Then we examined the influence of ISL on GSC-forming spheres on the 9th day of treatment. It was found that the 0 µM ISL group had the largest diameter and highest number of tumor spheres formed from the GSCs (Fig. 2B-D). The diameters and the numbers were both decreased by ISL in a dose-dependent manner, and 160 µM ISL completely prevented neurosphere formation, displaying a similar tendency with cell activity at 72 h.

**Differentiation-promoting action of ISL on GSCs.** To determine whether ISL promotes the differentiation of GSCs, intracellular differentiation markers GFAP and β III tubulin were examined after 48 h of ISL treatment by immunofluorescence staining. As shown in Fig. 3, the numbers of differentiation marker-positive cells were higher in the ISL group and DAPT group than that in the control group after 48 h. As a comparison, FBS effectively induced the differentiation of GSCs.

To determine whether ISL inhibits Hes1 (Notch1 target gene) in the differentiated cells, GFAP and Hes1 were assessed together after the cells were treated for 48 h. The relative intensity of Hes1 was decreased in the GFAP+ cells induced by ISL (Fig. 4).

The protein expression levels of differentiation markers were also examined after 48 h of ISL treatment. As expected, the protein expression levels of GFAP and β III tubulin in the DAPT group and ISL groups were higher than those of the control, and DAPT induced higher GFAP and β III tubulin protein expression levels compared to the ISL groups. However, there was no obvious difference between the low dose (20 µM) and high dose (80 µM) ISL groups (Fig. 5A and B).

**Ability of ISL to downregulate the Notch signaling pathway.** To examine whether ISL inhibits the proliferation and induces the differentiation of human GSCs through Notch1 signaling, Notch1 and Hes1 were analyzed 48 h after GSCs were respectively treated with ISL and DAPT. The predicted molecular
Figure 3. Differentiation-inducing effect of ISL (20 µM) on GSCs. The intracellular expression of differentiation markers at 48 h was examined by immunofluorescence staining. GFAP (A) and β III tubulin (B) were assessed by staining with PE-labeled rabbit anti-mouse IgG (red), and the nuclei were stained with DAPI (blue); scale bar, 50 µm. (C and D) The number of GFAP+ and β III tubulin+ cells per visual field were determined from 5 random fields of view by Image-Pro Plus 6.0. Data are summarized as mean ± SD. ***P<0.001 vs the control (CON). GSCs, glioma stem cells. ISL, isoliquiritigenin.

Figure 4. A decrease in Hes1 is noted in differentiated cells induced by ISL (20 µM). (A) The intracellular expression of Hes1 and GFAP was examined by immunofluorescence staining. Hes1 was assessed by staining with FITC-labeled goat anti-rabbit IgG (green), GFAP was assessed by staining with PE-labeled rabbit anti-mouse IgG (red), and the nuclei were stained with DAPI (blue); scale bar, 50 µm. The relative intensity of Hes1 (B) and GFAP (C) in the cells was measured in 5 random fields of view by Image-Pro Plus 6.0. Data are expressed as mean ± SD. ***P<0.001 vs. the control (CON). ISL, isoliquiritigenin.
weight of Notch1 is 272 kDa. In the present study, the band observed at 120 kDa could potentially be a truncated form of Notch1, following cleavage by furin-like protease. The protein expression levels of Notch1 in the DAPT group and ISL groups were lower than those of the control (Fig. 5C). The mRNA and protein expression levels of Hes1 were also significantly decreased compared to the control (Fig. 5D and E). DAPT induced lower Notch1 and Hes1 expression levels compared to the ISL groups.

**Discussion**

It has been reported that supplementation with ISL may be an effective approach to inhibit the growth of tumors and counteract the adverse effects of existing drugs for cancer patients (17). Despite the studies demonstrating the efficacy of ISL in tumor therapy, its anticancer activities against GSCs remain unclear. In the present study, we identified that ISL inhibited proliferation and induced differentiation of human GSCs, and inhibited the activation of the Notch1 pathway.

Serum-free culture is a simple and convenient method for neural stem cell (NSC) enrichment and has been applied for cancer stem cell (CSC) isolation from brain tumors and many other types of cancers (18). Using serum-free medium with supplemental factors B27, EGF and bFGF, the present study isolated tumor spheres from SHG44 human glioma cells on suspension cell culture plates. CD133 and Nestin are important cell surface markers present on NSCs. A high level of CD133 expression may be an independent risk factor for glioma patient prognosis and high expression of Nestin tends to correlate with a worse outcome for glioma patients (19). As previously described, GSCs develop high chemoresistance and radioresistance. In this respect, a high level of CD133 or/and Nestin expression in glioma cells may imply a high proportion of GSCs which contribute to adverse prognosis. In this study, CD133 and Nestin were used together to examine the stem

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**Figure 5.** Effect of ISL treatment on differentiation markers and Notch1 signaling pathway in GSCs. The protein expression levels of GFAP (A) and β III tubulin (B) were examined after the cells were treated for 48 h. The protein expression level of Notch1 (C) and the mRNA and protein expression level of Hes1 (D and E) were examined at 48 h after ISL treatment. The band observed at 120 kDa could potentially be a truncated form of Notch1, following cleavage by furin-like protease. For the immunoblotting analyses, densitometry was performed with Image-Pro Plus 6.0 software. The optical densities (ODs) of the protein bands were calculated relative to the ODs of the reference protein β-actin. Data are summarized as mean ± SD "P<0.01, "**P<0.001 vs. the control (CON). GSCs, glioma stem cells. ISL, isoliquiritigenin.
cell property of tumor spheres and the expression of Bcl-2 provided further evidence that these neurospheres were GCSs derived from the SHG44 cell line.

It has been reported that the inhibition rate of U87 human glioma cells can be ~80% after a 72-h treatment with 40-60 µM ISL (13,20). Our data indicated that ISL can dose-dependently inhibit the proliferation of human SHG44 GSCs 72 h after treatment. GSC proliferation was obviously inhibited after a 72-h incubation with 10 µM ISL. However, the IC50 of 102.744 µmol/l seemed much higher than that of other studies mentioned above. This might be because GSCs are proven to be more resistant to various chemotherapeutic agents. Some recent studies have suggested the use of ISL as a natural candidate to reduce CSC-like populations and colony formation, accompanied by enhanced CSC chemosensitivity (21,22). Here, we investigated the inhibitory effect of ISL on the formation of glioma neurospheres for the first time. In terms of long-term efficacy, the diameters and the numbers of GSC-forming spheres on the 9th day of 20 µM ISL treatment were significantly decreased. Thus, we believe that 20 µM ISL can be sufficient to inhibit GSCs. Unfortunately, there are few toxicological evaluations reported for ISL treatment. Our findings will provide the theoretical and experimental bases for ISL clinical use in the treatment of glioma. Further studies are still required to investigate the clinical side-effects in vivo.

Here, we also showed the stimulating effect of ISL on GSC differentiation into downstream neural lineages. It has been reported that ISL facilitated the differentiation of human promyelocytic leukemia and mouse melanoma cells (23,24). The present study first showed that ISL induced GSCs to differentiate into astrocytes (GFAP-positive) (25) and neurons (β III tubulin-positive) (26). As a result, after differentiation induced by ISL, the majority of GSCs were able to be killed by ISL, leading to decreased cell numbers. Neurons are highly differentiated cells and rich in tubulin isotypes (27). Another study found that β-tubulin is required for cell viability and neurite outgrowth, and the suppression of β I isotype resulted in significant loss of cell viability during cell differentiation (28). In the present study, it is perhaps not coincidental that ISL increased the viability of GSCs at 24 and 48 h when a proliferation inhibitory effect was not dominant. Therefore, GSC proliferation was activated following 20 µM ISL at 48 h when GSC viability was mainly increased in the differentiating cells. On the contrary, GSC proliferation was inhibited by 80 µM ISL at 48 h as a high dose of ISL exhibited a strong growth inhibitory effect which abrogated the increase in viability of the differentiated cells. GSCs and NSCs express similar signatures, such as infinite self-renewal ability, neurosphere formation, and multipotential differentiation capacity (6,29). GSCs in the control group treated without ISL still generated a small population of differentiated cells, but were much less than those in the ISL groups. Further studies are required to investigate the precise mechanism of how ISL induces GSCs to differentiate into each phenotype.

Recently, an accumulating body of evidence has emerged to suggest that the Notch1 signaling pathway plays an important role in cell fate decisions, and functions as a key regulator of cell growth, differentiation and proliferation (30,31). Fan et al. (32) observed that inhibition of the Notch1 signaling pathway by a γ-secretase inhibitor resulted in the apoptosis and differentiation of medulloblastoma cell lines, and reduced tumor progression. In the present study, the Notch1 pathway was both inhibited by DAPT and ISL. The mRNA and protein expression levels of Notch1 were altered in a similar trend with cell proliferation and in a reverse trend with GFAP and β III tubulin, indicating that Notch1 signaling was blocked directly or indirectly by ISL and therefore changed the proliferation and differentiation of GSCs. However, there was no obvious difference in Notch1 pathway expression between low dose (20 µM) and high dose (80 µM) ISL groups. This is possible as other pathways may be involved. In terms of differentiation, cells treated with low dose and high dose ISL had the same protein level of differentiation markers. The possible reason may be that cell differentiation-induced action reached a plateau. DAPT induced lower Notch1 and Hes1 expression levels compared to the ISL groups. It is proposed that DAPT and ISL might have different target sites of downregulation of Notch1. Further studies are still required to investigate the crosstalk between Notch and other signaling pathways, including the NF-κB, Wnt and PI3K/Akt/mTOR pathways in GSCs.

ISL is a flavonoid compound with efficient antitumor activity. However, the drug clearance has not been measured in recent studies. Yang et al. (33) firstly reported the transport of ISL and other cardio-cerebral vascular protection flavonoids using the CaCo-2 and blood-brain barrier (BBB) cell models in vitro. An increasing number of studies have shown that flavonoids are capable of crossing the blood-brain barrier via different BBB models such as RBE-4 cells, hCMEC/D3 cells, and ECV304/C6 coculture (34-36). In situ (rat) models have demonstrated that flavonoids are able to traverse the BBB in vivo. In recent years, new strategies such as the use of carrier-mediated transport systems and nanotechnology-based approaches enable drug transport into the brain (37). These will hopefully reduce the excretion of ISL by the body and lead toward the successful treatment of glioma patients. Nevertheless, further studies are still required to reveal the underlying mechanisms and confirm the benefit of ISL treatment in vivo.

In conclusion, the present study confirmed our hypothesis that ISL can effectively inhibit the proliferation and induce the differentiation of human GSCs. The therapeutic effect may be related to downregulation of the Notch1 signaling pathway. However, further studies are needed to identify the target/receptor molecules of ISL on the cell surface, and to confirm the therapeutic effect in vivo. These studies will help to screen for the optimal parameters and elucidate the mechanism of ISL as a promising agent for the treatment of human glioma.

References


