

Morusin inhibits the growth of human colorectal cancer HCT116-derived sphere-forming cells via the inactivation of Akt pathway

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Abstract. The existence of colorectal cancer stem-like cells (CSC) is responsible for the failure of current treatments against colorectal cancer. Therefore, novel therapies need be developed to target CSCs. Some natural agents, including morusin have been proposed as possible candidates for this purpose. Morusin has been shown to exert antitumor effects. In the present study, it is demonstrated that morusin exerts antitumor effects on colorectal CSCs (CCSCs). The viability of human CCSCs was enhanced when the CCSCs formed spheroids in a serum-free and non-adhesive floating culture system. HCT116 sphere cells exhibited an increased proliferative capacity and a higher expression of stemness markers [octamer-binding transcription factor 4 (Oct4), Sox2 and Nanog]. Morusin inhibited the development of cancer spheroids and suppressed the growth of sphere cells via the induction of cell cycle arrest. Similarly, morusin decreased the expression levels of the stemness markers, Nanog and Oct4. The data partially revealed the molecular mechanisms involved: β -catenin signaling maintains the growth of CSCs and directly modulates the expression of Nanog and Oct4. Morusin suppressed the activity of β -catenin signaling via the inactivation of Akt; the executive β -catenin/TCF4 complex and the downstream targets, c-Myc, survivin and cyclin D1, were also downregulated. Moreover, the morusin-induced inactivation of Akt also increased the expression of p21Cip1/WAF1 and p27Kip, which can block the cell cycle by interacting with cyclin-dependent kinase (CDK) complexes. On the whole, the present study demonstrates that morusin inhibited the growth

of colorectal cancer sphere cells, which were enriched with CCSCs via the inactivation of the Akt pathway.

Introduction

Colorectal cancer represented almost 10% of the global cancer incidence burden in 2012, causing a mortality rate of 694,000 per year worldwide (1). Despite recent advances in medicine, approximately 50% of patients with colorectal cancer exhibit tumor recurrence (2), and the overall mortality rate associated with the disease is approximately 40% (3). Colorectal cancer stem-like cells (CCSCs) are considered to be one of the primary causes of tumor recurrence.

Cancer stem-like cells (CSCs) are defined as 'cells within a tumor that possess the capacity for self-renewal and that can cause the heterogeneous lineages of cancer cells that constitute the tumor' (4). At present, CSCs have been proven to exist in a variety of solid tumors, including colon cancer (5-7). It is considered that only CSCs can drive tumor initiation, proliferation and spreading (8). In addition, CSCs are not affected by chemotherapy or radiation (9). CSCs that survive chemoradiotherapy will initiate and maintain the regrowth of the tumor. A number of currently used protocols for cancer therapy are now understood to fail, as marked by the reappearance of disease, due to the inability to eradicate CSCs. Therefore, the requirement and the challenge in fighting cancer is to develop novel therapies targeting CSCs. To counteract the challenges, researchers have turned to natural products. Some natural agents, including morusin (10), have been proposed as candidates for the targeting of CSCs alone or in combination with chemoradiotherapy (11,12).

Morusin (2-(2,4-dihydroxyphenyl)-5-hydroxy-8,8-dimethyl-3-(3-methylbut-2-enyl)pyranol[2,3-h]chromen-4-one, C₂₅H₂₄O₆) (NCBI PubChem Compound database, CID:5281671) is a naturally occurring agent isolated from the traditional Chinese medicinal (TCM) herb, *Morus alba* L. (Sang Bai Pi), which is used as an antiphlogistic, an antipyretic, an expectorant, an antitussive, a diuretic and an antidiabetic medication (13,14). Researchers have demonstrated that morusin exerts antitumor effects in several types of neoplasms, including hepatocellular carcinoma (15,16),

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gastric cancer (16), cervical cancer (10) and neuroblastoma (17). To the best of our knowledge, the last and only study on the antitumor activity of morusin against colorectal cancer was published in 2008 (14), in which it was reported that morusin induced the nuclear factor (NF)- κ B-mediated apoptosis of HT-29 colorectal cancer cells. Based on previous research, it was hypothesized that morusin may exert antitumor effects on CCSCs. To confirm this hypothesis, in the present study, the effects of morusin treatment on CCSCs were examined and the potential underlying mechanisms were investigated.

Materials and methods

Drugs. Morusin (purity 99%) was purchased from the Shanghai Research Center of Traditional Chinese Medicine Standardization. Morusin was dissolved in dimethyl sulfoxide (DMSO) to a concentration of 4,800 μ M. The stock solution was stored at -20°C and diluted in the medium before the experiment. The final DMSO concentration did not exceed 1% throughout the study.

Reagents and antibodies. McCoy's 5A medium, Dulbecco's modified Eagle's medium (DMEM)/F12 medium, fetal bovine serum and B27 supplement (50X) were obtained from Gibco; Thermo Fisher Scientific, Inc. Penicillin G, streptomycin and Heparin Na salt were purchased from Sigma-Aldrich; Merck KGaA. Epidermal growth factor (EGF) and basic fibroblast growth factor (bFGF) were purchased from PeproTech, Inc. The primary antibodies [octamer-binding transcription factor 4 (Oct4) (cat. no. sc-5279), Sox2 (cat. no. sc-365823) and Nanog (cat. no. sc-293121)] used for immunofluorescence assay were purchased from Santa Cruz Biotechnology, Inc. The primary antibodies used for western blot analysis were purchased from the following companies: Cell Signaling Technology, Inc. [Oct4 (cat. no. 2750), Sox2 (cat. no. 3579), Nanog (cat. no. 4903), β -catenin (cat. no. 8480), transcription factor 4 (TCF4) (cat. no. 2569), c-Myc (cat. no. 18583) and survivin (cat. no. 2808)], Elabscience® [Akt (cat. no. E-AB-30471), phospho-Akt-473 (cat. no. E-AB-21135), glycogen synthase kinase (GSK)-3 β (cat. no. E-AB-20885)], American Research Products, Inc. [CDK2 (cat. no. A2439), cyclin D1 (cat. no. 10-M1033) and p21Cip1/WAF1 (cat. no. 24-1026-MSM3)] and ProteinTech Group, Inc. [(p27Kip1 (cat. no. 25614-1-AP), cyclin A2 (cat. no. 27242-1-AP), GAPDH (cat. no. 60004-1-Ig) and β -actin (cat. no. 20536-1-AP)].

Cells and cell culture. The HCT116 human colorectal cancer cell line was purchased from the Shanghai Type Culture Collection of the Chinese Academy of Sciences. HCT116 cells were cultured in McCoy's 5A medium supplemented with 10% fetal bovine serum, 100 U/ml penicillin G and 100 μ g/ml streptomycin. The cells were cultured in a humidified 5% CO_2 atmosphere at 37°C . The human fetal colon (FHC) cells (a gift from Dr Hu of Shanghai Funeng Biotechnology Co., Ltd.) were cultured in a 1:1 mixture of Ham's F12 and DMEM (Corning, Inc.) containing HEPES (25 mM), cholera toxin (10 ng/ml; Sigma-Aldrich; Merck KGaA), insulin (5 lg/ml), transferrin (5 lg/ml) and hydrocortisone

(100 ng/ml; Sigma-Aldrich; Merck KGaA), supplemented with 10% fetal bovine serum (FBS) (Invitrogen; Thermo Fisher Scientific, Inc.).

Spheroid formation. Spheroid formation was enriched using a serum-free and non-adhesive floating culture system. Prior to spheroid formation, viable cells were counted by trypan blue staining and seeded in 24-well ultra-low attachment plates (Corning Inc.) at a density of 2,000 cells per well in 500 μ l 1X serum-free medium (SFM) (DMEM/F12 medium supplemented with 20 ng/ml EGF, 20 ng/ml bFGF, 4 μ g/ml Heparin Na salt and 1X B27 supplement) for 8 days; subsequently, 50 μ l of 10X SFM medium per well was added on days 3, 5 and 7. Tumor spheroids were dissociated, digested and harvested on day 8, and the viable cells were stained with 0.4% trypan blue staining solution for 5 min at room temperature. Single-sphere cell suspensions were cultured in McCoy's 5A medium, then treated with the morusin (0, 9.1, 18.2 and 36.4 μ M). Spheroids were observed under an inverted fluorescent microscope (Olympus Corporation).

Cell proliferation assay/Cell Counting Kit (CCK)-8 assay. The proliferation of the HCT116 cells and FHC cells was measured using a CCK-8 assay (Dojindo Molecular Technologies, Inc.). There were 3 general groups: The blank group (McCoy's 5A medium without cells or drugs), the control group (cells were cultured in McCoy's 5A medium without the drug) and the experimental groups (cells were exposed to 2.5, 5, 10, 15, 20, 25, or 50 μ M morusin for 72 h, or to 9.1, 18.2 or 36.4 μ M morusin for 24, 48 or 72 h). Following treatment, the culture medium was discarded and 100 μ l of fresh medium including 10 μ l of CCK-8 was added to each well. Following 4 h of incubation in a humidified 5% CO_2 atmosphere at 37°C , the absorbance was measured at 450 nm using a Multiskan Spectrum Microplate Reader (Bio-Rad Laboratories, Inc.). The experiments were performed in triplicate. The percentage inhibitory rate (IR%) of the treated cells was calculated using the following formula: $\text{IR}\% = 1 - [(\text{EXP group OD}_{450 \text{ nm}} - \text{blank group OD}_{450 \text{ nm}}) / (\text{CTRL group OD}_{450 \text{ nm}} - \text{blank group OD}_{450 \text{ nm}})] \times 100\%$.

5-Ethynyl-2'-deoxyuridine (EdU) assay. Cells were seeded in Millicell EZ 8-well glass slides (EMD Millipore) at a density of 1×10^4 cells/well, and the cells were exposed to 9.1, 18.2 or 36.4 μ M Morusin for 24 h. Following treatment, the culture medium was discarded, and an EdU incorporation assay was performed using a Cell-Light™ EdU Apollo® 567 *In Vitro* Imaging kit (Ribobio Co.) according to the manufacturer's instructions. The stained cells were observed under a fluorescence microscope (Olympus Corporation), images were captured and analyzed using ImageJ software (<https://imagej.nih.gov/ij/download.html>).

Immunofluorescence assay. Cells were seeded in Millicell EZ 8-well glass slides (EMD Millipore) at a density of 1×10^4 cells/well, washed with phosphate-buffered saline (PBS), fixed and permeated using a BD Cytofix/Cytoperm™ kit (BD Biosciences). The cells were then incubated with Oct4, Sox2 and Nanog antibodies at a dilution of 1:50 in 1% bovine serum albumin (BSA)-PBS at 4°C overnight followed by

Table I. Primer sequences used for RT-PCR.

Gene	Primer sequences (5'-3')	Product length (bp)
Oct4	F: GGG TGG AGA GCA ACT CCG A R: GCT TGG CAA ATT GCT CGA G	150
Sox2	F: GTT CTA GTG GTA CGG TAG GAG CTT TG R: TTT GAT TGC CAT GTT TAT CTC GAT	150
Nanog	F: CCA GCT GTG TGT ACT CAA TGA ATT T R: TTC TGC CAC CTC TTA GAT TTC ATT C	150
18s rRNA	F: TCG GAG GTT CGA AGA CGA TC R: CAG CTT TGC AAC CAT ACT CCC	150

F, forward; R, reverse; Oct4, octamer-binding transcription factor 4.

incubation with the secondary antibody [goat anti-mouse IgG H&L (Alexa Fluor® 647) (Abcam, 1:200, cat. no. ab150115) and donkey anti-goat IgG H&L (Cy5®) (Abcam, 1:400, cat. no. ab97117)] in 1% BSA-PBS at room temperature in the dark for 1 h. The cells were then washed with PBS and stained with 10 µg/ml Hoechst 33342 (Abcam, cat. no. ab228551) at room temperature in the dark for 5 min. The stained cells were observed under a fluorescence microscope, and images were captured and analyzed using ImageJ software.

Detection of cell cycle distribution by flow cytometry. Following treatment with 9.1, 18.2 or 36.4 µM morusin for 24 h, the cells were harvested, washed with 4°C PBS and stored in cold 70% ethanol at -20°C overnight. The cells were centrifuged (200 x g) for 4 min at room temperature, then stained with propidium iodide (PI) treated with RNase A (100 U/ml) and detected by flow cytometry (FACScan; BD Biosciences) for cell cycle analysis.

Detection of apoptotic cells by flow cytometry. Cell apoptosis was detected using an Annexin V-Fluorescein Isothiocyanate (FITC) Apoptosis Detection kit I (BD Biosciences) according to the following protocol: The cells were seeded in 6-well plates at a density of 2x10⁵ cells/well and exposed to 9.1, 18.2 or 36.4 µM morusin for 24 h. The cells were digested, incubated with FITC-Annexin buffer at room temperature for 10 min, stained with PI and detected by flow cytometry (FACScan; BD Biosciences).

Reverse transcription-polymerase chain reaction (RT-PCR). Total RNA was extracted from the cells using the RNeasy Micro kit (Qiagen GmbH). Total RNA (0.5 µg) was used as a template to prepare cDNA (Reverse Transcription System, Promega Corporation; cat. no. A3500) according to the manufacturers' instructions. The amplification conditions were as follows: Stage 1 (holding 95°C for 10 min); stage 2 (40 cycles of denaturing at 95°C for 15 sec, annealing at 60°C for 45 sec and extending at 72°C for 30 sec); stage 3 (extension at 72°C for 7 min). The primer sequences are listed in Table I. The housekeeping gene GAPDH was used as an internal reference to normalize the results. All experiments were performed in triplicate. Finally, the 2^{-ΔΔC_q} method was performed to calculate the relative expression (18).

Semi-quantitative RT-PCR. Similarly, total RNA was extracted from the cells and the cDNAs were prepared as described above for RT-PCR. Subsequently, 1% of the cDNA samples were used for GAPDH, Oct4, Sox2, Nanog PCR. Sense primers were ³²P-radiolabeled with T4 polynucleotide kinase. The PCR reactions were performed with the following profile: 30 sec denaturation at 94°C, 15 sec annealing, 25 s extension at 72°C. An initial denaturation step at 94°C for 2.5 min and a final extension step at 72°C for 7 min were also performed. The PCR products were analyzed by 2% agarose gel electrophoresis and stained with ethidium bromide.

Western blot analysis. HCT116 sphere cells were seeded in 10-mm dishes at a density of 2x10⁶ cells per dish and were exposed to 9.1, 18.2, or 36.4 µM morusin for 24 h. Following treatment, the cells were washed with cold PBS at 4°C, lysed in RIPA lysis buffer including PhosSTOP (Roche Diagnostics) and 1 µM PMSF at 4°C for 30 min. Cell lysates were homogenized and centrifuged of 14,000 x g at 4°C for 10 min. The protein concentration was detected by Enhanced BCA Protein Assay kit (Beyotime Institute of Biotechnology). Equal amounts of total protein (40 µg) were boiled, loaded onto an 8% SDS-PAGE gel and electro-transferred to polyvinylidene fluoride (PVDF) membranes (EMD Millipore). After blocking with 5% non-fat milk in Tris-buffered saline (TBST) for 2 h at room temperature, the membranes were incubated with the primary antibodies at 4°C overnight and incubated with 1:4,000 dilutions of secondary antibodies labeled with horseradish peroxidase [rabbit anti-mouse IgG H&L (HRP), Abcam, cat. no. ab6728] at room temperature for 1 h. After being washed with TBST 3 times, the target proteins were detected using an Enhanced Chemi Luminescence (ECL) kit (Thermo Fisher Scientific, Inc.). Finally, images were captured by ChemiDoc™ XRS+ (Bio-Rad Laboratories, Inc.).

Statistical analysis. All data were analyzed using the Statistical Package Social Science (SPSS) 19.0 software and are expressed as the means ± SD. Statistical analysis was performed among multiple groups by one-way analysis of variance (ANOVA) followed by Tukey's post hoc test. Values of P<0.05 and P<0.01 were considered to indicate statistically significant and highly statistically significant differences, respectively.

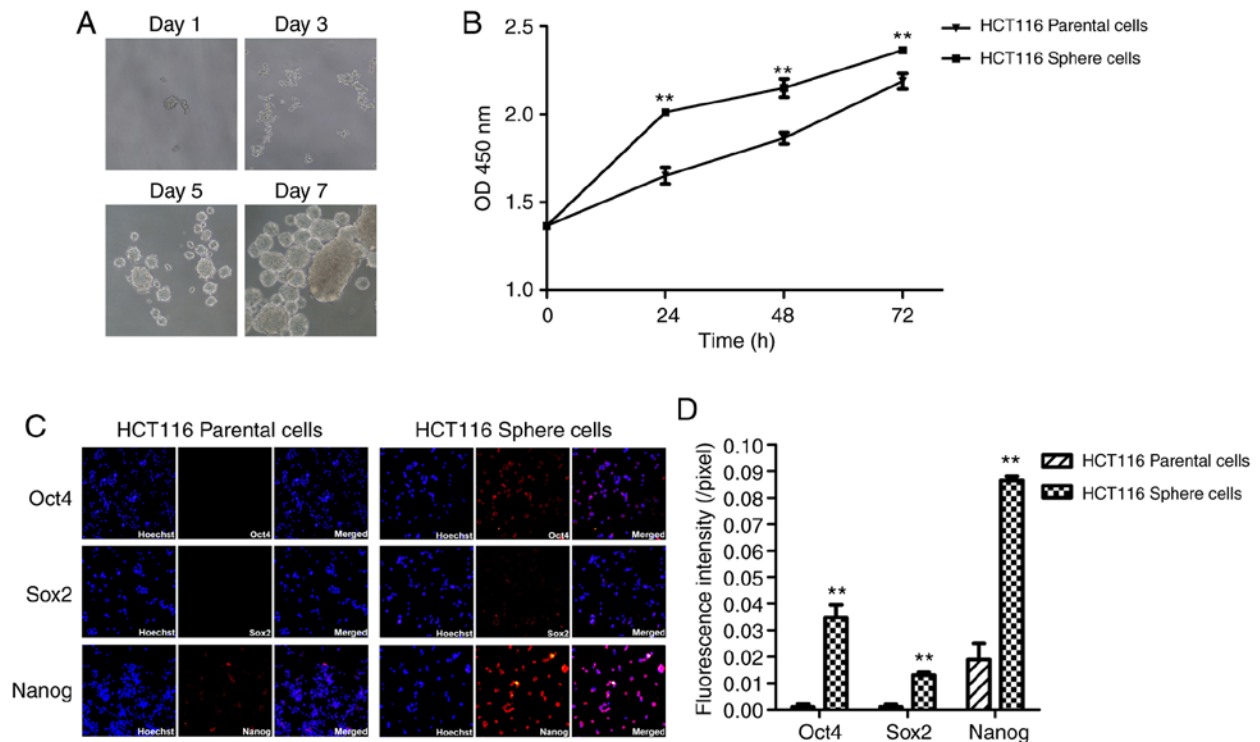


Figure 1. Enrichment and biological characterization of CCSCs. (A) Spheroid-formation state of HCT116 cells over a period of 1-7 days. (B) Cell growth inhibition assay revealed that HCT116 sphere cells had an increased proliferative capacity at 24, 48 and 72 h. (C) The expression of Oct4, Nanog, and Sox2 in HCT116 parental and sphere cells was assayed by immunofluorescent staining assay (magnification, x200). (D) Fluorescence intensity of Oct4, Nanog and Sox2 expression in HCT116 parental and sphere cells. All the data are expressed as the means \pm SD of data obtained in 3 parallel experiments; * $P < 0.01$ vs. HCT116 parental cells. CCSCs, colorectal cancer stem cells.

Results

Enrichment and biological characterization of CCSCs. Spheroid formation was induced in a serum-free and non-adhesive floating culture system (SFM) enriched with CCSCs (19), as it has been demonstrated that sphere cells are enriched with CSCs (20,21). Visible tumor spheroids were observed under an optical microscope on day 3 and were then gradually grown for the following 5 days (Fig. 1A). Following spheroid formation, the number of viable cells in each well increased from 2,000 to $65,500 \pm 18,358$. To evaluate the proliferative capacity of the HCT116 sphere cells, HCT116 parental and sphere cells were seeded in 96-well plates at a density of 5,000 cells per well in McCoy's 5A medium, and CCK-8 assay was performed at 24, 48 and 72 h. The sphere cells exhibited an increased proliferative capacity compared with the parental cells ($P < 0.01$; Fig. 1B). Subsequently, the expression levels of stemness markers (Oct4, Sox2 and Nanog) was detected by immunofluorescence staining. It was observed that Nanog mRNA and protein expression was detectable in the HCT116 parental cells, whereas Oct4 and Sox2 expression was undetectable. By contrast, the Oct4, Sox2 and Nanog expression levels were significantly increased in the HCT116 sphere cells, with a particular increase noted in Nanog mRNA and protein expression (Fig. 1C). As shown in Fig. 1D, the fluorescence intensity of Oct4, Sox2 and Nanog expression was significantly upregulated in the HCT116 sphere cells (each $P < 0.01$). Taken together, the results indicated that there was an upregulation in the expression of stemness-related transcription factors (Oct4, Sox2 and Nanog) in HCT116 sphere cells.

Inhibitory effects of morusin on the growth of HCT116 sphere cells (results of CCK-8 and EdU assays). The proliferation of HCT116 sphere cells was assessed by CCK-8 assay. The HCT116 sphere cells were treated with various concentrations of morusin (2.5, 5.0, 10.0, 15.0, 20.0, 25.0 and $50.0 \mu\text{M}$) for 72 h. When the concentration was $\geq 5.0 \mu\text{M}$, morusin significantly inhibited the proliferation of HCT116 sphere cells in a concentration-dependent manner ($P < 0.01$); an approximately 50% growth inhibition at 72 h was achieved at the concentration of $18.2 \mu\text{M}$ (IC_{50}) (Fig. 2A). Subsequently, the HCT116 sphere cells were treated with 9.1, 18.2, or $36.4 \mu\text{M}$ morusin for 24, 48 and 72 h. Morusin significantly decreased cell viability at each time point ($P < 0.01$; Fig. 2B and C). These results indicated that treatment with morusin led to a significant inhibition of cell growth in a concentration- and time-dependent manner.

To further evaluate the proliferative activity of HCT116 sphere cells following treatment with morusin, an EdU incorporation assay was also performed (Fig. 2D). Following treatment with 9.1, 18.2, or $36.4 \mu\text{M}$ morusin for 24 h, cell viability was decreased from 70.65 ± 5.33 to 29.54 ± 3.33 , 13.23 ± 1.19 and $4.12 \pm 1.13\%$, respectively. There was a significant decrease in EdU incorporation when the cells were incubated with morusin ($P < 0.01$; Fig. 2E), suggesting that morusin significantly inhibited the proliferative activity of the HCT116 sphere cells. In addition, the proliferation of normal FHC cells treated with morusin (9.1, 18.2 and $36.4 \mu\text{M}$) demonstrated that morusin did not affect normal colonic mucosa cell viability (Fig. 2F).

Morusin inhibits the initiation of colorectal cancer spheroid formation. To determine whether morusin inhibits the

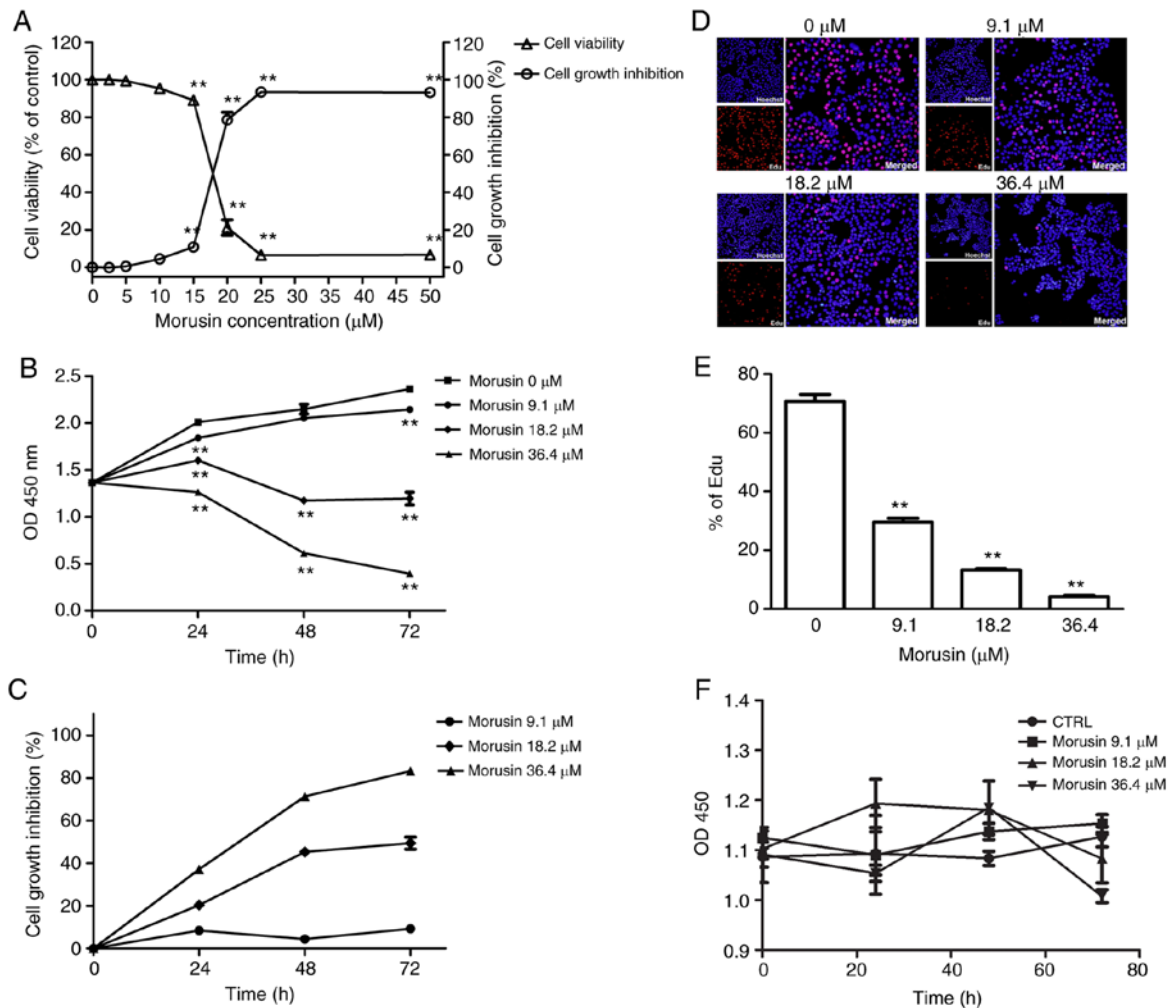


Figure 2. Inhibitory effects of morusin on the growth of HCT116 sphere cells. (A) Results of cell viability and cell growth inhibition assay in HCT116 sphere cells following treatment with increasing concentrations of morusin for 72 h. (B and C) Results of (B) OD450 nm and (C) cell growth inhibition assay in HCT116 sphere cells following treatment with morusin (9.1, 18.2 and 36.4 μM) for 24, 48 and 72 h. (D) Following treatment with morusin (4.6, 9.1, 18.2 and 36.4 μM) for 24 h, the proliferative ability of HCT116 sphere cells was measured by EdU incorporation assay (magnification, $\times 200$). (E) The number of positive cells in the EdU incorporation assay was counted. The EdU-positive rate is shown. (F) The proliferation of normal fetal colonic mucosa cells (FHC cells) treated with morusin (9.1, 18.2 and 36.4 μM) for 24, 48 and 72 h. All data are expressed as the means \pm SD of the data obtained in 3 parallel experiments; ** $P < 0.01$ vs. control.

initiation of cancer spheroids formation, HCT116 cells were pre-treated with 9.1, 18.2, or 36.4 μM morusin for 24 h. The culture medium was then removed and the cells were washed with PBS and collected; viable cells were seeded at a density of 2,000 viable cells per well. Following culture in CSC medium for 7 days, images of each cell group were obtained and the viable cells were counted by trypan blue staining. As shown in Fig. 3A, the HCT116 parental cells that were pre-treated with 9.1 μM morusin could only rarely form spheroids; in the higher concentration groups (18.2 and 36.4 μM), the HCT116 parental cells lost the ability to initiate tumor spheroid formation entirely. Compared with the control group, the number of viable cells in the experimental groups (morusin at 9.1, 18.2 and 36.4 μM) was significantly decreased ($P < 0.01$; Fig. 3B). These results indicated that the spheroid-forming capacity of the HCT116 cells was suppressed by morusin.

Effect of Morusin on the cell cycle and apoptosis of HCT116 cells. After being incubated with morusin for 24 h, the cell cycle status and apoptosis were detected by flow cytometry. Compared with the control group, morusin (9.1, 18.2 and

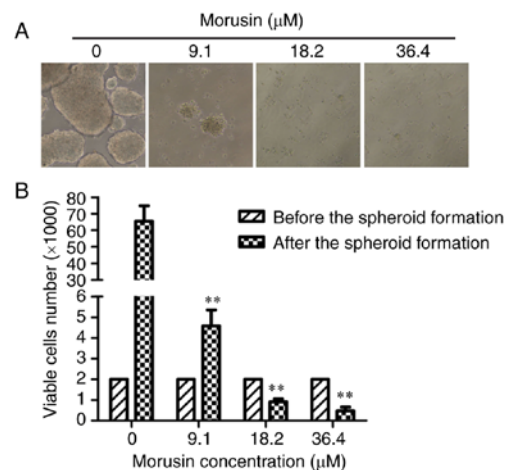


Figure 3. Morusin inhibits the initiation of cancer spheroid-formation. (A) Following pre-treatment with morusin (4.6, 9.1, 18.2 and 36.4 μM) for 24 h, the surviving viable cells were cultured in SFM at a density of 2,000 cells per well for 7 days; the spheroid-formation state is shown at day 8. (B) The number of viable cells was counted after harvesting the spheroids; data are expressed as the means \pm SD of the data obtained in 3 parallel experiments; ** $P < 0.01$ vs. control (no treatment).

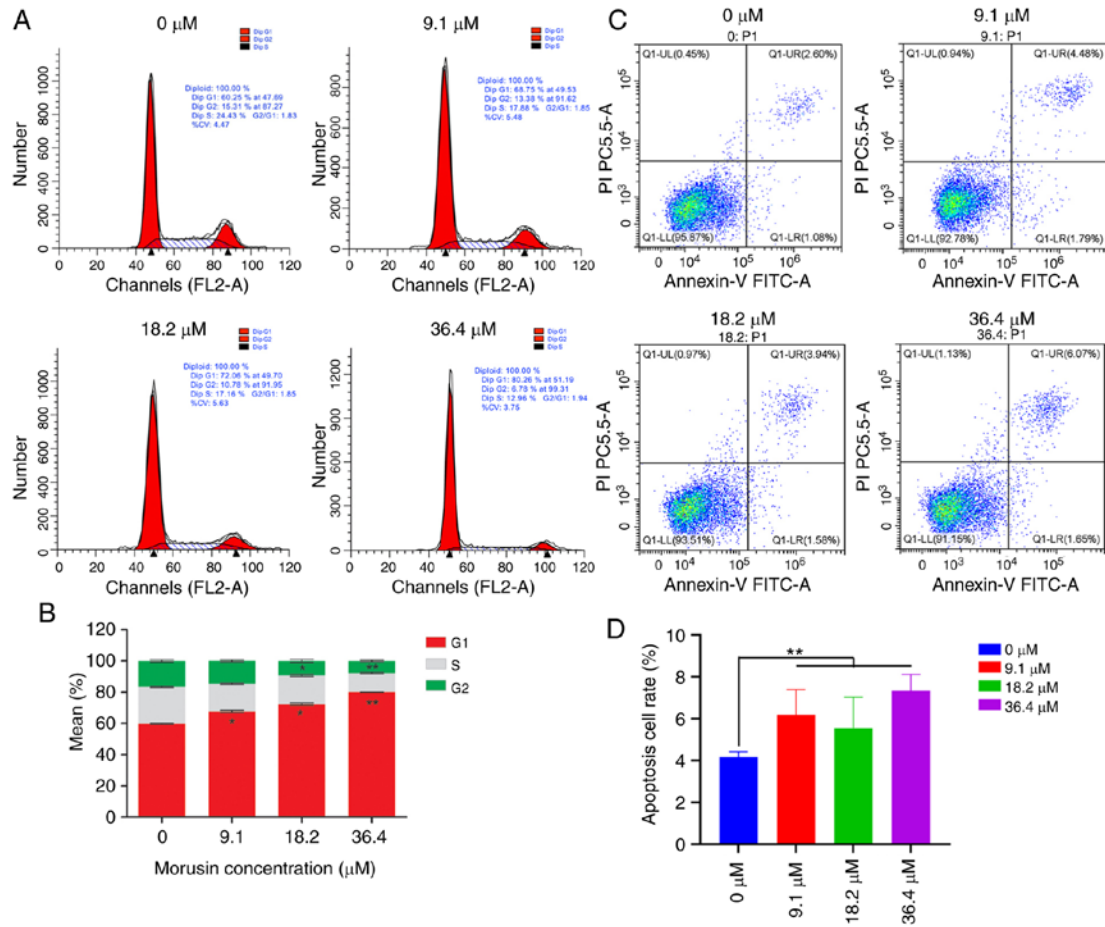


Figure 4. (A and B) Cell cycle status of HCT116 sphere cells treated with morusin (9.1, 18.2 and 36.4 μM) for 24 h. (C and D) Apoptosis of HCT116 sphere cells treated with morusin (9.1, 18.2 and 36.4 μM) for 24 h. All data are expressed as the means \pm SD of the data obtained in 3 parallel experiments; * $P < 0.05$, ** $P < 0.01$ vs. control.

36.4 μM) significantly increased the percentage of cells in the S phase ($P < 0.01$) and decreased the percentage of cells in the G0/G1 phase ($P < 0.01$); the higher concentrations of morusin (18.2 and 36.4 μM) decreased the percentage of cells in the G2/M phase ($P < 0.05$; Fig. 4A and B). These results indicated that morusin induced cell cycle arrest at the S phase in the HCT116 sphere cells. The percentages of apoptotic cells in each group are shown in Fig. 4C and D. It was found that morusin induced the apoptosis of HCT116 sphere cells in a dose depended manner. The apoptotic rate of the HCT116 sphere cells increased with the increased concentration of morusin.

Effect of Morusin on the expression of stemness markers. The expression levels of stemness markers (Oct4, Sox2 and Nanog) were detected by RT-PCR and western blot assay. There was a decreasing trend in Nanog mRNA expression following morusin treatment. In addition, the results of western blot analysis further confirmed that morusin inhibited the protein expression of Nanog in a concentration-dependent manner. Furthermore, the Oct4 protein expression level was decreased in the group treated with the higher concentration of morusin (36.4 μM); however, the Sox2 protein expression level was too low to be detectable (Fig. 5).

Morusin suppresses the activity of the Akt/ β -catenin pathway. Subsequently, the mechanisms involved in the morusin-induced

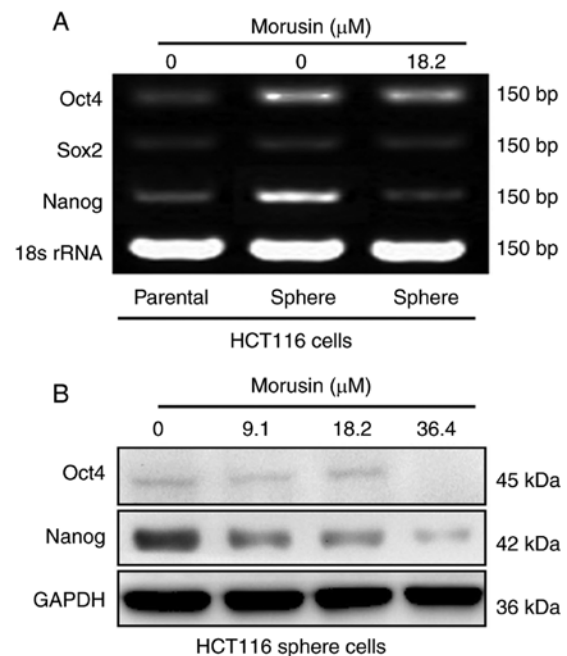


Figure 5. Effect of Morusin on the expression of stemness markers. (A) RT-PCR assays were performed for Oct4, Sox2 and Nanog mRNA expression in HCT116 parental cells, HCT116 sphere cells and morusin (18.2 μM)-treated HCT116 sphere cells. (B) Following treatment with morusin (9.1, 18.2 and 36.4 μM) for 24 h, western blot analysis was performed for Oct4 and Nanog protein expression in HCT116 sphere cells.

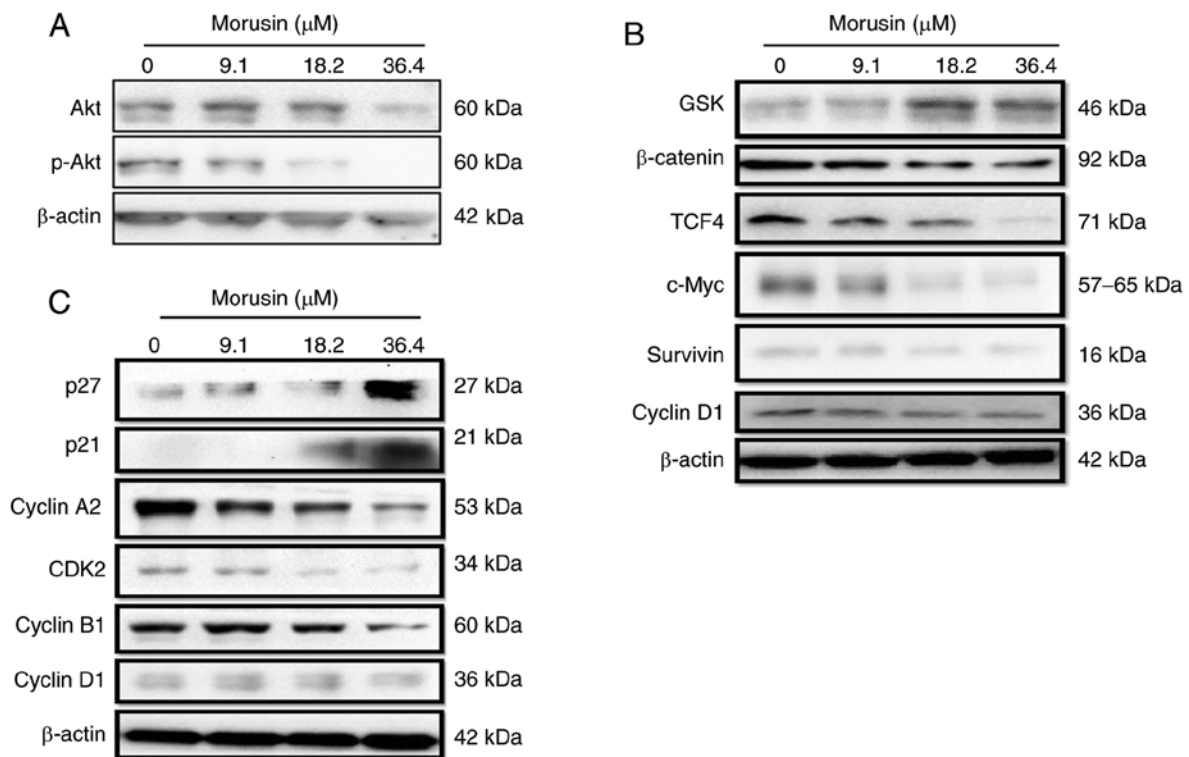


Figure 6. Effect of morusin on the expression of Akt and its target proteins in HCT116 sphere cells. (A) Following treatment with morusin (9.1, 18.2 and 36.4 μM) for 24 h, western blot analysis for Akt activity was performed. (B) Morusin-induced inactivation of β -catenin signaling and its downstream proteins. (C) Effect of morusin on the expression of p21Cip1/WAF1, p27Kip1, and other cell cycle-related proteins.

growth inhibition of HCT116 sphere cells were investigated. Following incubation with 9.1, 18.2 and 36.4 μM morusin for 24 h, the activity of Akt/protein kinase B and β -catenin signaling was detected by western blot analysis. The expression levels of total Akt and phosphorylated Akt at Ser473 are required for Akt activation (22). In the present study, morusin inhibited the expression of total Akt at the higher concentration (36.4 μM) and suppressed the phosphorylation of Akt in a concentration-dependent manner (Fig. 6A). As shown in Fig. 6B, the morusin-induced inactivation of Akt increased the expression of Gsk-3 β , which is a common target of the Akt and Wnt/ β -catenin pathway (23). Activated Gsk-3 β decreased the expression of β -catenin, followed by a reduction in TCF4 expression. Furthermore, morusin suppressed downstream targets of the Wnt/ β -catenin pathway (c-Myc and survivin) in a concentration-dependent manner, which decreased the expression of cyclin D1 at the higher concentration of morusin (36.4 μM).

Morusin modulates the expression of cell cycle-related proteins via the activation of p21Cip1/WAF1 and p27Kip1. Inactivated Akt can also increase the expression of p21Cip1/WAF1 and p27Kip1 (24). Morusin upregulated the expression of p21Cip1/WAF1 and p27Kip1 via the inactivation of Akt and then suppressed a number of cell cycle-related proteins, including cyclin A and CDK2 (Fig. 6C), that participate in the progression through the S and G2 phases cooperatively (25). The reduced cyclin A-CDK2 complex expression induced cell cycle arrest at the S phase, which was consistent with the results of the cell cycle distribution assay (Fig. 4).

Discussion

Colorectal cancer causes approximately 694,000 deaths per year worldwide (1). Despite recent advancements being made in therapies, the response rate to current systemic therapies is ~50%, resistance develops in nearly all patients (26). Although the mechanisms involved remain unclear, chemoresistant colorectal CSCs are an important cause. Conventional anti-tumor therapies that can eliminate the bulk of tumor cells, cannot target CSCs that may lead to a reduction of the tumor mass, but not the regression of the tumor (27). Natural products are proposed as candidates for targeting CSCs (28,29). The present study demonstrated that morusin exerts antitumor effects on CCSCs.

In the present study, spheroid formation was induced in a serum-free and non-adhesive floating culture system (SFM) enriched with CCSCs (19), as it has been demonstrated that sphere cells are enriched in CCSCs (20,21). Compared with parental HCT116 cells, sphere cells exhibited an increased proliferative capacity and a higher expression of pluripotent transcriptional factors (Oct4, Sox2 and Nanog), core components of the pluripotency regulation network that also maintain the stem-like properties of CSCs (20,30,31) and are considered to be colorectal CSCs markers (32). It was observed that morusin not only inhibited the growth of colorectal sphere cells, but also suppressed the initiation of colorectal cancer spheroid formation. Similarly, morusin decreased the expression of the stemness markers, Oct4 and Nanog. Subsequently, further experiments were performed to examine the mechanisms responsible for the effects of morusin on HCT116 sphere cells.

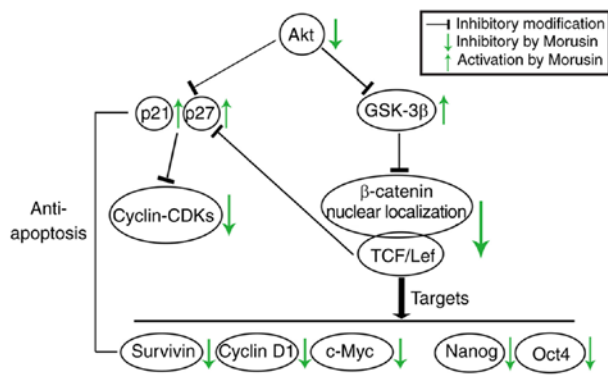


Figure 7. Model depicting the role of Morusin in Akt/ β -catenin and Akt/p21Cip1/WAF1 p27Kip1 pathway of CCSCs. CCSCs, colorectal cancer stem cells.

The potential molecular mechanisms of the effects of morusin on CCSCs were demonstrated in the present study. First, activated Akt downregulates the activity of GSK-3 β (33), which is also a signaling target of the canonical Wnt/ β -catenin pathway. It is recognized that the Wnt/ β -catenin pathway carries activating mutations in virtually all colon cancers (34) and plays an important role in CCSCs (2). The constitutive activation of the Wnt/ β -catenin pathway triggers the tumor-initiating potential (35,36) and maintains the growth of CSCs (tumorspheres *in vitro*) (20). The Akt-mediated inactivation of GSK-3 β results in the stabilization and accumulation of β -catenin in the cytoplasm, followed by the nuclear translocation of β -catenin, which leads to the activation of the canonical Wnt pathway. In the cell nucleus, β -catenin binds to members of the T-cell factor (TCF)/lymphoid enhancer factor (LEF) family to modulate the expression of target genes. In addition, Nanog and Oct4 are also downstream targets of β -catenin pathway (37,38), which may partly explain why Wnt/ β -catenin signaling plays a role in sustaining stemness.

In the present study, the expression levels of total Akt and phosphorylated Akt at Ser473, which is required for Akt activation (22), were assayed by western blot analysis. The morusin-induced inactivation of Akt significantly increased the expression of GSK and decreased the activity of β -catenin. In turn, the expression of TCF4 expression, which is a critical factor in response to β -catenin (39), was decreased, and the β -catenin/TCF4 complex is an important effector of Wnt/ β -catenin signaling. Morusin reduced the expression of the downstream proteins, c-Myc [a known promoter of cell proliferation and growth (40), survivin [a member of the inhibitor of apoptosis protein (IAP) family that inhibits caspases and blocks cell death (41), and cyclin D1 (which facilitates cell-cycle progression through the G1 phase (42)]. In particular, Nanog and Oct4 are directly regulated by the β -catenin/TCF complex (37,38). Nanog not only is a CSC marker, but also promotes cell proliferation (43,44); the results indicated that morusin downregulated Oct4 and Nanog expression via the suppression of the β -catenin/TCF4 complex. Taken together, the results demonstrated that Morusin inhibited the growth of HCT116 sphere cells and decreased stemness marker expression via the Akt/GSK/ β -catenin pathway.

According to the second pathway shown in Fig. 7, activated Akt directly or indirectly decreased the expression of p21Cip1/WAF1 and p27Kip1 (45,46), which belong to the CDK interacting protein/kinase inhibitory protein (CIP/KIP) family, a CDK inhibitor that can halt the cell cycle by interacting with a variety of cyclin-CDK complexes (47). The morusin-induced inactivation of Akt attenuated its suppression of the growth inhibitory activity of p21 and p27, which led to a reduction in cyclin A-CDK2 complex formation, which is required for cell cycle progression through the S and G2 phases (25). Furthermore, the morusin-induced reduction in TCF4 expression can also contribute to the increase in p27 expression cooperatively (48). As a result of the inactivity of cyclin A-CDK2, morusin induced cell cycle arrest at the S phase (Fig. 4), and the EdU incorporation assay further demonstrated that the proliferative ability was inhibited (Fig. 2C and D). According to the above, morusin blocked cell cycle progression in CCSCs via the Akt/p21Cip1/WAF1 p27Kip1 pathway to inhibit cell proliferation.

It has been reported that morusin induced cancer cell apoptosis (10,14). The results of the present study were consistent with those of these other studies. It was found that morusin promoted the apoptosis of HCT116 sphere cells in a concentration-dependent manner. In addition, it was found that morusin was able to increase the expression of p21Cip1/WAF1 and p27Kip1, resulting in inducing cell cycle arrest. Although the mechanism involved is unclear, it was hypothesized that it was associated with the upregulation of p21Cip1/WAF1 expression (49).

In conclusion, the present study demonstrated that Morusin inhibited the growth of colorectal cancer sphere cells, which were enriched with CCSCs. Morusin induced the inactivation of Akt followed by the suppression of β -catenin signaling, which resulted in a reduction in stemness marker expression and cell growth inhibition. Morusin also increased the expression of p21Cip1/WAF1 and p27Kip1 and induced cell cycle arrest. Morusin may thus be considered a novel antitumor agent targeting CSCs.

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Availability of data and materials

All data generated or analyzed during this study are included in this published article or are available from the corresponding author on reasonable request.

Authors' contributions

YZ performed the experiments and wrote the manuscript. XL assisted in performing the experiments and processed all the figures. MY designed of the study, provided funding support and revised the manuscript. All authors read and approved the final manuscript.

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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