Salvianolic acid B reverses multidrug resistance in nude mice bearing human colon cancer stem cells

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Abstract. Salvianolic acid B (SalB) is a water-soluble phenolic compound, extractable from Salvia miltiorrhiza, and has previously been demonstrated to reverse tumor multidrug resistance (MDR) in colon cancer cells. Cancer stem cells (CSCs) are closely associated with drug resistance. Therefore, establishing a nude mouse model bearing human colon CSCs is important for the study of the mechanisms underlying colon cancer drug resistance as well as the reversal of drug resistance. The present study aimed to establish a nude mouse model bearing human colon CSCs and to investigate the effects of SalB on the drug resistance exhibited by the nude mouse model as well as determine its underlying mechanism. Cells from two colon cancer cell lines (LoVo and HCT-116) were cultured in serum-free medium to obtain CSCs-enriched spheroid cells. Following this, nude mice were transplanted with LoVo and HCT-116 colon CSCs to establish the CSC nude mouse model, which was subsequently demonstrated to exhibit MDR. The results of the present study revealed that following treatment with SalB, the chemotherapeutic drug resistance of xenografts was reversed to a certain extent. Western blot analysis was performed to investigate the expression levels of cluster of differentiation (CD)44, CD133, transcription factor sox-2 (SOX2) and ATP-binding cassette sub-family G member 2 (ABCG2) proteins, and the results demonstrated that treatment with SalB downregulated the expression of CD44, SOX2 and ABCG2 proteins in both LoVo and HCT-116 colon CSCs xenografts. In conclusion, the results of the present study revealed that a serum-free suspension method can be performed to successfully isolate colon CSCs. In addition, a nude mice bearing colon CSCs animal model was successfully established, and associated tumors were confirmed to exhibit MDR. Furthermore, SalB was demonstrated to successfully reverse MDR in nude mice bearing LoVo and HCT-116 colon CSCs, as well as suppress the expression of CD44, SOX2 and ABCG2 proteins.

Introduction

Cancer multidrug resistance (MDR) is a major factor affecting chemotherapy efficacy, which may ultimately lead to the failure of chemotherapy (1). Cancer stem cells (CSCs) represent a cancer cell population with stem cell-like properties. Following the development of the cancer stem cell hypothesis as well as increasing research of stem cells in the oncology field, numerous studies have demonstrated that drug resistance exhibited by CSCs is an important factor affecting cancer MDR (2,3). Therapy targeting CSCs provides a novel therapeutic approach for clinical cancer therapy (4). Therefore, establishing an adaptable and feasible animal xenograft model bearing human colon CSCs is essential for the study of the underlying mechanisms of CSCs in order to develop novel and effective anticancer drugs.

At present, there is no agreement regarding the best method to acquire colon CSCs for the establishment of an animal model. There are three main methods routinely performed for the isolation and identification of colon CSCs. Firstly, specific expression of surface markers for CSCs can be determined, predominantly by flow cytometry (FCM) (5) and immunomagnetic beads sorting (6). In this method, isolation and identification are performed using expressed cell surface markers present on colon CSCs, including cluster of differentiation (CD)44, CD166, CD133, Nanog and transcription factor sox-2 (SOX2). This method has been widely used to identify and define CSCs. Secondly, Hoechst
Salvianolic acid B (SalB) is a water-soluble phenolic compound and is extractable from *Salvia miltiorrhiza* (11). SalB has previously been demonstrated to reduce the toxicity and enhance the efficacy of radiochemotherapy for the treatment of colon cancer, and its role in reversing tumor MDR has generated increasing attention (12,13). Stem cell factors, including SOX2, CD24, organic cation/carnitine transporter 4 (OCT4), CD29, CD44 and ABCG2; serve an important role in maintaining the morphology and function of colon CSCs, and are closely associated with the proliferation, drug resistance, invasion and migration of colon cancer cells (14-17). Investigation into the regulatory effects of SalB on CSC marker expression will further the understanding of the mechanisms underlying the effect of SalB on MDR reversal.

Two colon cancer cell lines (LoVo and HCT-116) were used in the present study to investigate the effect and underlying mechanism of SalB. The present study aimed to isolate and identify colon CSCs, establish a nude xenograft mouse model bearing colon CSCs, perform H&E staining, investigate the MDR of the xenografts and determine the efficacy of chemotherapeutic drugs on the mouse model. Furthermore, the present study aimed to investigate the effect of SalB on drug resistance exhibited by xenografts, determine the expression levels of CD44, CD133, SOX2 and ABCG2 following treatment with SalB, and investigate the underlying mechanism of this effect.

**Materials and methods**

**Cell lines.** Human colon cancer cell lines LoVo and HCT-116 were purchased from the Cell Resource Center, Shanghai Institutes for Biological Science (Shanghai, China).

**Animals.** A total of 100 specific pathogen free (SPF) BALB/c male nude mice were purchased from Shanghai SLAC Laboratory Animal Co., Ltd. [Shanghai, China; license no. SCXK (Hu) 2007-005]. Mice were aged 4 weeks and had a body weight of 20±2 g. All animal experiments were performed according to the guidelines of the Chinese Experimental Animals Administration Legislation and were approved by the Ethics Committee for Animal Experiments of Shanghai University of Traditional Chinese Medicine (Shanghai, China; reference no. SYXK (Shanghai) 201504023). Mice were fed in separate cages under specific pathogen-free conditions in a laminar flow chamber in the Lab Animal Center, Shanghai University of Traditional Chinese Medicine (Shanghai, China). Standard water in drinking bottles and pelleted food were freely available to the mice. The temperature was maintained at 18-25°C at a relative humidity of 40-60% and a 12/12 h light/dark cycle. The cages, padding, feed and water were autoclaved at 121°C for 30 min. The padding was replaced at least twice a week. Animals were acclimatized for 1 week prior to the initiation of the experiments.

**Drugs and reagents.** SalB was purchased from Shanghai Winher Medical Technology Co., Ltd. (Shanghai, China). A total of 100 mg of oxaliplatin (L-OHP) was purchased from Jiangsu Hengrui Medicine Co., Ltd. (Lianyungang, China; cat. no. H20040817). 5-Fluorouracil (5-FU) was purchased from Shanghai Xudong Haipu Pharmaceutical Co., Ltd. (Shanghai, China; 25 g/l; cat. no. 20070802). PRMI-1640 and DMEM/F12 were purchased from Hyclone; Thermo Fisher Scientific, Inc. (Waltham, MA, USA). Penicillin (5,000 IU/ml) and streptomycin (5,000 µg/ml) were purchased from Invitrogen; Thermo Fisher Scientific, Inc. (Waltham, MA, USA). Fetal bovine serum (FBS), L-glutamine, β-mercaptoethanol and 2-hydroxyethyl methacrylate were purchased from Sigma-Aldrich; Merck KGaA (Darmstadt, Germany). Recombinant human basic fibroblast growth factor, recombinant human epidermal growth factor, KnockOut Serum Replacement and 1% Non Essential Amino Acid (NEAA) were purchased from Gibco; Thermo Fisher Scientific, Inc. (Waltham, MA, USA). FcR Blocking Reagent was purchased from Miltenyi Biotec GmbH (Bergisch Gladbach, Germany). Anti-human CD133-PE and Mouse IgG2b kappa isotype control were purchased from eBioscience; Thermo Fisher Scientific, Inc., A biocincheminic acid assay (BCA) kit, Hematoxylin and Eosin Staining kit and an immunochemistry kit were purchased from Shanghai Beyotime Biological Science & Technology Co., Ltd. (Shanghai, China). Mouse monoclonal antibody ABCG2, mouse monoclonal antibody CD24, rabbit monoclonal antibody CD133, rabbit monoclonal antibody OCT-4, rabbit monoclonal antibody CD44, goat anti-mouse immunoglobulin G (IgG) and goat anti-rabbit IgG were all purchased from Abcam (Cambridge, MA, USA). Rabbit monoclonal antibody SOX2 was purchased from Cell Signaling Technology, Inc. (Danvers, MA, USA). RNA reverse kit and RNA amplification kit were purchased from Takara Bio, Inc. (Otsu, Japan).

**Cell culture and spheroid formation.** LoVo and HCT-116 cells were separately cultured in F-12K and McCoy’s5A medium, respectively. Both cultures contained 10% FBS, 100 µg/ml penicillin and 100 µg/ml streptomycin at 37°C, in a humidified 5% CO₂ incubator. Tumor cell spheroids were isolated using the serum-free suspension culture method: Logarithmic growth phase LoVo and HCT-116 cells were trypsinized from plates and re-suspended with serum-free medium: Dulbecco’s Modified Eagle Medium (DMEM)/F12 + KnockOut Serum Replacement (10-20%) + 1% NEAA + 1% L-glutamine + 0.1 mM β-mercaptoethanol + 20 µg/l recombinant human basic fibroblast growth factor + 20 µg/l recombinant human epidermal growth factor] and the cell suspension was then added to a petridish coated with 2-hydroxyethyl methacrylate in a
Table I. DNA primer sequences.

| Primer name | Forward | Reverse |
|-------------|---------|---------|
| CD24        | CCAGGGCAATGATGAAATGAGA | GGGAGGCTGAGCACGAGAAT |
| CD44        | GGAATGGACAGTTTGTGCGACGG | GGTACACCCCAATCTCTCATGTC |
| CD90        | GACTGCCGCTAGAATAACACC | CGGCCGCCAGACTTGAA |
| CD133       | GATTCTACTGTTGGCTCGGATG | GCAAGTGGAAGTGCGCTAAGT |
| SOX2        | TCTGATCTCAACTTTCAT | ACATCGATTCTCGGCACAG |
| ABCG2       | GGTTTCCAAGCCTTATCAAA | TAGCCCCAAATATAAGGCACCTA |
| OCT4        | CGAAGAGAAAAGCGAACCAGTATC | AGAACCACACTGAGCCACATC |
| Nanog       | GCACAAAGGAGAACAGGGTCC | CCTTCTGCGTACACCATTG |
| GAPDH       | GGTGGTCTCCTCCTGACTTCAACA | CCAATTTCGTTGTCAATCCGAAATG |

CD, cluster of differentiation; SOX2, transcription factor Sox2; ABCG2, ATP binding cassette subfamily G member 2; OCT4, organic cation/carnitine transporter 4.

Experiments were performed in triplicate. Primer sequences for the targeting of the selected genes are presented in Table I.

Western blot analysis. Western blot analysis was performed to determine the expression levels of OCT4, CD24, CD44, CD133, SOX2 and ABCG2 proteins. Cytoplasmic and nuclear proteins were extracted using the protein extraction kit (cat. no. P0013B; Shanghai Beyotime Biological Science & Technology Co., Ltd.), and protein levels were then determined using a BCA protein assay kit (cat. no. P0010; Shanghai Beyotime Biological Science & Technology Co., Ltd.). Protein samples (30 µg) were run on a 10% SDS-PAGE gel, and then transferred to polyvinylidene fluoride membrane. Membranes were then blocked with 5% bovine serum albumin (cat. no. W029; Shanghai Bogu Biotechnology Co., Ltd., Shanghai, China) blocking solution for 2 h at room temperature, and then incubated overnight at 4°C with the following primary antibodies: Rabbit monoclonal antibody OCT4 (1:1,000; cat. no. ab200834), rabbit monoclonal antibody CD44 (1:1,000; cat. no. ab51037), mouse monoclonal antibody CD24 (1:1,000; cat. no. ab76514), rabbit monoclonal antibody CD133 (1:1,000; cat. no. ab216323), rabbit monoclonal antibody SOX2 (1:1,000; cat. no. ab3579) and mouse monoclonal antibody ABCG2 (1:1,000; cat. no. ab13024). Following overnight incubation, the membranes were washed three times with Tris-buffered saline with 0.1% Tween-20 (TBST), and goat anti-mouse IgG (1:1,000; cat. no. ab205719) and goat anti-rabbit IgG (1:1,000; cat. no. ab205718) were incubated at 37°C for 2 h. The membrane was then washed three times with TBST, and then visualized using enhanced chemiluminescent reagents (cat. no. P0018; Shanghai Beyotime Biological Science & Technology Co., Ltd.) according to the previously published protocol (19). ImageJ software (version 1.48; National Institutes of Health, Bethesda, MD, USA) was used for densitometric analysis of target protein bands. Experiments were performed in triplicate.

Establishment of the xenograft nude mouse model. Spheroid cells generated from LoVo and HCT-116 cells were labeled as LoVocsc and HCT-116csc, respectively. LoVo, LoVocsc,

Identifying the colon CSCs. FCM was performed to determine CD133-positive expression. A single cell suspension (10⁶ cells/ml) containing 10⁶ cells/ml was prepared and treated with mouse IgG2b kappa FcR blocking reagent in an ice bath for 10 min and then incubated in the dark at 4°C for 30 min with phycoerythrin tagged CD133 (1:100; cat. no. 12-1394-41; eBioscience; Thermo Fisher Scientific, Inc.) and mouse IgG2b kappa isotype control (1:100; cat. no. 12-4732-81; eBioscience; Thermo Fisher Scientific, Inc.). Subsequently, the cells were washed twice with PBS and centrifuged at 300 x g for 10 min at 4°C. The supernatant was aspirated and 500 µl PBS (cat. no. C0221A; Shanghai Beyotime Biological Science & Technology Co., Ltd.) was added for analysis of the cells using a FACScalibur (BD Biosciences, Franklin Lakes, NJ, USA).

Reverse transcription-quantitative polymerase chain reaction (RT-qPCR). RT-qPCR was performed to investigate the expression levels of CD90, CD24, CD44, CD133, SOX2, ABCG2, OCT4 and Nanog. Following the culture of adherent LoVo and HCT-116 cells in serum-free suspension media for ~5-6 days, suspended regular stem cell spheroids were visible, and spheroid cell suspension and adherent cells were simultaneously collected. Following this, total RNA was extracted using the RNAeasy extraction kit (cat. no. 9108Q; Takara Biotechnology Co., Ltd., Dalian, China). Total RNA (1 µg) was subjected to RT at 42°C for 15 min followed by 95°C for 3 min using a FastQuant RT Kit (cat. no. KR106; Tienang, Biotec Co., Ltd., Beijing, China). qPCR was performed using SYBR Green PCR Master Mix (cat. no. FP302; Tienang, Biotec Co., Ltd.) in a ABI 7000 PCR instrument (Eppendorf, Hamburg, Germany) under the following thermocycling conditions: Initial denaturation at 95°C for 2 min; followed by 40 cycles of 95°C at 10 sec and 60°C at 30 sec. The threshold cycle (Cq) values of each sample were calculated using the 2−ΔΔCq method (18). GAPDH was used as an internal control for the normalization of mRNA expression. However, the threshold cycle (Cq) values of each sample were calculated using the 2−ΔΔCq method (18). GAPDH was used as an internal control for the normalization of mRNA expression. Instead, the threshold cycle (Cq) values of each sample were calculated using the 2−ΔΔCq method (18). GAPDH was used as an internal control for the normalization of mRNA expression.
HCT-116 and HCT-116csc cells were harvested, rinsed twice with ice-cold PBS and then re-suspended with normal saline in order to obtain cell suspensions with a concentration of 2x10^6 cells/ml, which were then placed on ice for subsequent injections. A total of 0.2 ml of cell suspension of LoVo, LoVoCsc, HCT-116 and HCT-116csc groups were injected subcutaneously into the right fossa axillaris of 12 randomly selected nude mice (each cell suspension group injected into 3 nude mice). When the xenografts had grown to ~0.8-1.5 cm, the mice were sacrificed and the tumors were isolated. The tumor tissue was then dissected into 1 mm^3 blocks and inoculated subcutaneously into the right fossa axillaris of nude mice under sterile conditions. Tumors formed from LoVo and HCT-116 cells were subsequently inoculated into 24 nude mice (12 mice per tumor type). Tumors formed from LoVoCsc and HCT-116csc cells were inoculated into 64 nude mice (32 mice per tumor type). Every 2 days, major and minor axes of the tumors were measured using a vernier caliper, denoted as a and b, respectively; to determine the tumor volumes (V) using the following formula:

$$V (\text{mm}^3) = 1 / 2a^2b^2 (20)$$

When xenografts had grown to a size of 100 mm^3, 8 nude mice bearing LoVo, HCT-116, LoVoCsc and HCT-116csc xenografts were randomly selected (2 mice per tumor type). These mice were sacrificed via cervical dislocation, and the tumor tissues were obtained for subsequent experimentation.

H&E staining. Tissues were collected, fixed with 4% paraformaldehyde for 10 min at room temperature, paraffin-embedded and sectioned (5 µm thick sections). Following de-waxing with xylene and rehydrated using a descending ethanol series (100, 90, 80 and 70%), sections were stained using a H&E staining kit (cat. no. C0105; Shanghai Beyotime Biological Science & Technology Co., Ltd.), in accordance with the manufacturer’s protocol, and subsequently dehydrated with gradient ethanol for 10 sec per concentration gradient (70, 80, 90 and 100%). Sections were then mounted and histopathological changes were observed under a light microscope (magnification, x100 and x400) (21).

Grouping and SalB dosing. The remaining mice bearing LoVoCsc xenografts were randomly assigned into 6 groups (5 mice per group): LoVoCsc group, LoVoCsc + L-OHP group, LoVoCsc + SalB-low dose (L) group, LoVoCsc + SalB-medium dose (M) group, LoVoCsc + SalB-high dose (H) group and LoVoCsc + L-OHP + SalB-L group. The remaining mice bearing LoVo xenografts were randomly assigned into 2 groups (5 mice per group): LoVo group and LoVo + L-OHP group. The mice bearing HCT-116csc and HCT-116 xenografts were grouped using the aforementioned protocol. Mice in the LoVoCsc + L-OHP and LoVo + L-OHP groups were intraperitoneally injected with L-OHP (0.5 mg); Mice in the LoVoCsc + SalB-L, LoVoCsc + SalB-M and LoVoCsc + SalB-H groups were intragastrically administered low, medium and high doses of SalB solution (0.36, 0.72 and 1.44 g, respectively), at a dosage of 0.4 ml; Mice in the LoVoCsc + L-OHP + SalB-L group were intraperitoneally injected with L-OHP (0.5 mg) combined with intragastric administration of low dose of SalB (0.36 g). Mice in the LoVo, LoVoCsc, HCT-116 and HCT-116csc groups were intragastrically administered 0.4 ml of normal saline. L-OHP and 5-FU were administered to respective groups once every two days, whereas SalB was administered every day. Treatment lasted for a total duration of 28 days. The activity and skin color of nude mice were recorded every day for 28 days and were then euthanized by cervical dislocation when they reached the pre-determined measureable end points: Weight loss exceeding 15%.

Determination of tumor volume, growth curves and the rate of tumor inhibition. At 3 day time intervals, major and minor axes of the tumors were determined using a vernier caliper to calculate the tumor volume. Using these data, growth curves of the tumors were determined. Furthermore, the inhibitory rate was determined using the following formula: Inhibitory rate=(mean tumor volume of the control group-mean tumor volume of the test group)/mean tumor volume of the control group *100%. To investigate the interaction between SalB with L-OHP and 5-FU chemotherapeutic agents, the method described by Jin et al (22) was used. This method provides a ‘Q’ value, according to which the interaction between two drugs can be classified as exhibiting an antagonistic effect (Q<0.85), additive effect (0.85≤Q<1.15) or synergistic effect (Q≥1.15). The formula used to calculate the Q value is as follows: Q=E_{ab}/(E_{a}+E_{b}-E_{a}E_{b}), where E_{ab} represents the mean effect of combination treatment, and E_{a} and E_{b} represent the effect of drug A and drug B alone, respectively.

Tumor CD44, CD133, SOX2 and ABCG2 expression levels determined by western blot analysis. Tumor tissues were lysed in radioimmunoprecipitation assay buffer (1 mg tumor tissue for 10 µl lysis buffer; cat. no. P0013B; Shanghai Beyotime Biological Science & Technology Co., Ltd.), and liquid nitrogen was quickly added to grind the tissues to extract protein. This was followed by centrifugation at 10,000 x g for 5 min at 4°C, and the supernatant was collected. Procedures for western blot were performed according to the aforementioned protocol. β-actin was used as an internal control.

Statistical analysis. Statistical analyses were performed using SPSS software (v. 18.0; SPSS, Inc., Chicago, IL, USA). Data are presented as mean ± standard deviation for at least three separate experiments. Comparisons between two groups were performed using a Student’s t-test, and multiple comparisons between groups was performed using one-way analysis of variance followed by the Newman-Keuls post hoc test. P<0.05 was considered to indicate a statistically significant difference.

Results

Growth of spheroid CSCs. Human colon cancer cell lines LoVo and HCT-116 were cultured for 2 days (Fig. 1A and B) and then cultured in stem cell culture medium ES. Following
suspension culture for 2-3 days, only a small portion of cells survived. At the day 4 time interval, a small number of floating cell spheroids were visible, with a number of cells closely clustered. These cell spheroids were oval shaped and exhibited a clustered appearance, with a clear intercellular space (data not shown). The spheroids increased in size in a time-dependent manner and at days 5-6 time interval, suspended regular stem cell spheroids were visible and an increased number of cells, blurred intercellular space and strong refractivity was observed using a light microscope (Fig. 1C and D). Necrosis was observed in the center of the spheroids at the >10 day time intervals (data not shown). Spheroids were collected on days 5 and 6 for further experimentation.

Verifying CSC formation. FCM was performed to determine the expression of cell surface marker CD133 in populations of adherent and spheroid primary colon cancer cell populations. CD133 has been previously used as a marker for stem cells and to isolate colon CSCs from spheroids (23,24). The results demonstrated that there were significant differences in CD133 expression levels between the adherent LoVo (1.48±0.30%) and LoVocsc spheroid cells (87.44±4.32%; P<0.01; Fig. 2A). The levels of CD133+ cells in the adherent HCT-116 cells (1.25±0.24%) was also significantly decreased compared with HCT-116scs spheroid cells (92.53±3.56%; P<0.01; Fig. 2A).

RT-qPCR and western blot analyses were performed to determine the expression levels of the following stem cell markers in the parental and CSCs: CD24, CD44, CD90, SOX2, ABCG2, OCT4 and Nanog. As revealed in Fig. 2B, mRNA expression levels of CD24, CD44, CD133, SOX2 and ABCG2 were significantly increased in LoVocscs cells compared with LoVo cells (P<0.01; Fig. 2B). In addition, the protein expression levels of CD24, CD133, SOX2 and ABCG2 were significantly increased in LoVocscs cells compared with LoVo cells (P<0.01; Fig. 2B); whereas the expression of CD24 did not exhibit a significant difference (Fig. 2B). Furthermore, mRNA expression levels of OCT4, CD44, CD133, SOX2 and ABCG2 were significantly increased in HCT-116scs cells compared with HCT-116 cells (P<0.01; Fig. 2C), and their protein expression levels were also significantly higher in HCT-116scs cells compared with HCT-116 cells, with the exception of OCT4 (P<0.01; Fig. 2C).

Histopathology of xenografts. 10-15 days following the subcutaneous injection of LoVo and HCT-116 cells, grain-like bulges in the right axillary appeared. By contrast, LoVocsc and HCT-116csc cells exhibited decreased oncogenic time, grain-like bulges appeared ~6-8 days post-injection (data not shown). When the xenografts grew to ~0.8-1.5 cm (measured at the major axis of the tumor), whole tumors were harvested, cut into sections and transplanted into the right armpit of the remaining nude mice. Soft, grain-like solid bulges were visible 4-5 days post-transplant, and the xenografts grew to a size of 100 mm³ 2 weeks post-injection (data not shown). No nude mice died during the modeling.

The results of HE staining revealed glandular cavities in the xenografts of the LoVo and HCT-116 group, whereas irregular growth was observed in the LoVocsc and HCT-116scs groups. Compared with the LoVo and HCT-116 groups, xenografts of the corresponding LoVocsc and HCT-116csc groups exhibited an increased level of irregular structures, a decreased level of differentiation, a marked increase in nuclear atypia and an increased level of mitosis (Fig. 3), all of which indicated a higher malignancy in the tumors in the LoVocsc and HCT-116csc groups.

Treatment with L-OHP and 5-FU inhibits rate of tumor growth. No mice died during drug administration. Nude mice in the LoVocsc + L-OHP, HCT-116scs and HCT-116scs+5-FU groups gradually lost weight, with the HCT-116scs+5-FU group exhibiting the greatest weight loss. Furthermore, these groups exhibited lower activity as well as skin color deterioration; however, these conditions were not observed in the other groups (data not shown). Some of the mice suffered from ulcerations on the tumor surface when reaching the end of drug intervention. Such mice did not exhibit clinical evidence of pain or distress. In addition, the depth and size of the ulcer is very limited and did not exhibit any manifestation of infection
or hemorrhage, and so no clinical intervention regarding ulceration was performed. Mice belonging to the LoVosc and HCT-116csc groups demonstrated a reduced oncogenic time (data not shown), and the growth rates of the tumors were markedly increased compared with their corresponding parental cells. Treatment with L-OHP was revealed to induce an inhibitory effect on tumor volumes in the LoVo and LoVosc groups, where its inhibitory rate in the LoVo group was 44.98%, which was markedly higher compared with that exhibited by the LoVosc group (33.92%; Fig. 4A and Table II). Treatment with 5-FU was also demonstrated to induce an inhibitory effect on tumor volume in HCT-116 and HCT-116csc cells, which demonstrated inhibitory rates of 43.92 and 22.18%, respectively (Fig. 4B and Table III). These results suggested that xenografts of the LoVosc and HCT-116csc groups exhibited drug resistance, thus suggesting that the model was successfully established.

SalB attenuates drug resistance exhibited by colon CSCs xenografts. Prior to drug administration, tumor size was not significantly different compared with the different groups (P>0.05; Fig. 5 and Table IV). Following L-OHP and SalB administration, tumor volumes exhibited by the LoVosc + L-OHP and LoVosc + L-OHP + SalB-L groups were significantly reduced and demonstrated statistically significant differences compared with the LoVosc group (P<0.01; Fig. 5A and Table IV). However, the LoVosc + SalB-L group and the LoVosc group did not demonstrate a significant difference in tumor volume post-drug administration.

Figure 2. Identification of colon CSCs. (A) Flow cytometry analysis and quantification of CD133 expression in spheroid cells and corresponding parental cells. Cells in Q2-2 correspond to CD133-positive cells. Results are representative images of three independent experiments. **P<0.01 vs. parental cells. (B) Relative expression of stem cell markers at mRNA and protein levels in LoVo and LoVosc cells. **P<0.01 vs. LoVo cells. (C) Relative expression of stem cell markers at mRNA and protein levels in HCT-116 and HCT-116csc cells. **P<0.01 vs. HCT-116 cells. Data are presented as mean ± standard deviation. CSC, cancer stem cell; CD, cluster of differentiation; SOX2, transcription factor sox-2; ABCG2, ATP-binding cassette sub-family G member 2.
value was revealed to be 1.68 (data not shown), thus suggesting that SalB and L-OHP may exhibit a synergistic effect on the suppression of tumor volume, and that treatment with SalB may significantly reverse the drug resistance exhibited by LoVocsc xenografts in nude mice (Fig. 5A and Table IV). Furthermore, the results demonstrated that the tumor volumes of the LoVocsc + SalB-M and LoVocsc + SalB-H groups were 1,638.22±130.41 and 1,270.15±108.58 mm³, respectively; which were significantly smaller than that of the LoVocsc group (2,456.69±216.35 mm³) and LoVocsc + SalB-L group (2,293.45±203.64) (P<0.01, data not shown). These results suggested that medium and high doses of SalB suppressed tumor volume in LoVocsc xenografts in a dose-dependent manner, whereas low doses of SalB did not exhibit marked antitumor activity.

Following 5-FU and SalB administration, tumor volumes exhibited by the HCT-116csc + SalB-L, HCT-116csc + 5-FU and HCT-116csc + 5-FU + SalB-L groups demonstrated statistically significant differences compared with that exhibited by the HCT-116csc group (P<0.05 and P<0.01; Table V). In addition, the tumor volume of the HCT-116csc + 5-FU + SalB-L group was revealed to be suppressed compared with that exhibited by the HCT-116csc + 5-FU group (P<0.01; Fig. 5B and Table V). A Q value of 1.75 suggested that SalB and 5-FU may exhibit a synergistic effect on the suppression of tumor volume, and that treatment with SalB may significantly reverse the drug resistance exhibited by HCT-116csc xenografts in nude mice (Fig. 5B and Table V). Tumor volumes exhibited by the HCT-116csc, HCT-116csc + SalB-L, HCT-116csc + SalB-M and HCT-116csc + SalB-H groups were 3,756.20±416.92, 3,289.15±383.58, 2,857.24±255.62 and 2,564.73±241.08 mm³ respectively, indicating that SalB suppressed tumor volume in HCT-116csc xenografts in a dose-dependent manner (P<0.01, data not shown).

SalB was revealed to attenuate drug resistance exhibited by colon CSCs xenografts in nude mice, thereby increasing tumor sensitivity to chemotherapeutic agents. In addition, SalB was revealed to inhibit tumor growth in a dose-dependent manner.

CD44, CD133, SOX2 and ABCG2 protein expression is regulated by SalB. Western blot analyses were used to determine the protein expression of stem cell markers CD44, CD133 and SOX2, as well as the drug resistance marker ABCG2, in all experimental groups. The results revealed that protein expression levels of CD44, CD133, SOX2 and ABCG2 exhibited by the LoVocsc and HCT-116csc groups were significantly increased compared with those exhibited by the LoVo and HCT-116 groups (Fig. 6A and B; P<0.01). Expression levels of these four markers exhibited by the LoVocsc + SalB-L group were significantly decreased compared with the LoVocsc group (P<0.01; Fig. 6A); whereas the expression levels of only three of these markers (CD44, SOX2 and ABCG2) were revealed to be significantly decreased in the HCT-116csc + SalB-L group compared with those exhibited by the HCT-116csc group (P<0.01; Fig. 6B). These results demonstrate that low doses of SalB may significantly suppress CD44, SOX2 and ABCG2 protein expression in colon CSCs xenografts, which may contribute to the attenuation or reversal of drug resistance.

In addition, treatment with SalB was revealed to have an inhibitory effect on the protein expression levels of
CD44, SOX2 and ABCG2 in the LoVoAxc xenografts in a dose-dependent manner (Fig. 6C; P<0.05 and P<0.01). As presented in Fig. 6C, protein expression levels of these three markers in the LoVoAxc + SalB-L, LoVoAxc + SalB-M and LoVoAxc + SalB-H groups were significantly decreased in a dose-dependent manner and exhibited a statistically significant difference compared with associated expression levels exhibited by the LoVoAxc group (Fig. 6C; P<0.05 and P<0.01). CD133 expression levels did not demonstrate a significant difference between the LoVoAxc + SalB-M and LoVoAxc + SalB-H groups (Fig. 6C; P>0.05); however, it was significantly decreased compared with the expression levels exhibited by the LoVoAxc and LoVoAxc + SalB-L groups (Fig. 6C; P<0.01). Furthermore, there were no significant differences in the expression levels of CD44, CD133, SOX2 and ABCG2 proteins exhibited by the HCT-116csc + SalB-L, HCT-116csc + SalB-M and HCT-116csc + SalB-H groups; however, the expression levels of CD44, SOX2 and ABCG2 were significantly suppressed compared with the HCT-116csc group (Fig. 6D; P<0.05).

**Discussion**

Cancer MDR refers to the phenomenon that tumor cells are resistant to the effects of numerous antineoplastic drugs, regardless of their chemical structure or target. MDR in tumors is a complex process and involves numerous mechanisms (25). 

CSCs are specialized cell populations of cancer cells with unlimited potential to proliferate, self-renew and differentiate into numerous cell types (24). CSCs promote rapid proliferation of tumors, and can induce tumor differentiation into malignancies that are of greater maturity, both in function and phenotype (26). CSCs exhibit the primary characteristics of drug resistance, as they express high levels of ABC transporter families on their cell membrane, which tends to enhance the efflux of chemotherapeutic drugs, thus...
resulting in greater resistance of cancer cells to cytotoxicity and apoptotic induction by a variety of chemotherapeutic agents (27,28). ABCG2 is an important member of the ABC transporter family that is directly involved in the induction of drug resistance; however, it also maintains the stem cell characteristics of cancer cells. Thus, ABCG2 may represent a marker for cancer stem cells in numerous cancer types (29,30).

Deng et al (31) demonstrated that CSCs present in malignant fibrous fibrohistiocytoma expressing high levels of ABCG2 were able to self-renew and exhibited a strong resistance to doxorubicin and cisplatin. SOX is a transcription factor protein family characterized by a homologous sequence called the high mobility group-box. SOX2 is a member of the SOX protein family, and is an important stem cell marker for inducing stem cell formation, maintaining the characteristics of stem cells as well as inhibiting the differentiation of stem cells (32). The association between SOX2 and CSC resistance has been extensively demonstrated (33,34). In addition, CD44 is a transmembrane glycoprotein belonging to the adhesion molecule family and serves a role in numerous physiological and pathological processes. CD44 has been demonstrated to serve a significant role in tumor invasion, metastasis and drug resistance (35). Yan et al (36) and Bourguignon et al (37) have demonstrated that CD44 is closely associated with drug resistance exhibited by colon CSCs.

With the increasing interest and research regarding CSCs, colon CSCs have been successfully isolated from colon cancer solid tumors, colon cancer ascites and colon cancer cell lines (38). Furthermore, a number of studies have revealed that colon CSCs are closely associated with primary and secondary multidrug resistance of colon cancer (39,40). Despite the existence of CSCs having been demonstrated in the 1970s, the gold standard for isolating CSCs, as well as the development of animal xenograft models, have not been fully established. LoVo and HCT-116 cells were selected for investigation in the present study for the two reasons: Firstly, cell lines are readily available as a pure population compared with cells from colon cancer metastases; Second, the cells exhibit stable biological characteristics; The results of the present study suggested that colon CSCs LoVoCsc and HCT-116Csc derived from LoVo and HCT-116 cells stably express stem cell-like characteristics and xenografts generated by subcutaneous inoculation of colon CSCs LoVoCsc and HCT-116Csc can be serially passaged in mice. In addition, the results of the present study demonstrated that xenograft tumors exhibited drug resistance, rapid proliferation and other malignant biological characteristics. During modeling and drug administration, none of the mice died, however, some did exhibit weight loss. This suggested that the technique used to isolate CSCs and the method used to establish the nude mouse model was effective.

The results of the present study revealed that the serum-free suspension culture method may be used to isolate colon CSCs from LoVo and HCT-116 adherent cells. Furthermore, it was revealed that the xenografts obtained by subcutaneous

Table IV. Inhibitory rate of tumor growth in the L-OHP, low doses of SalB, and L-OHP combined with low doses of SalB in xenografts obtained from the LoVoCsc group (n=5).

| Group                  | n | Pre-administration | Post-administration | Tumor inhibition rate (%) |
|------------------------|---|--------------------|---------------------|---------------------------|
| LoVoCsc               | 5 | 112.69±13.89       | 2,456.82±216.35     | /                         |
| LoVoCsc + L-OHP       | 5 | 106.07±17.10       | 1,623.58±147.86     | 33.92                     |
| LoVoCsc + SalB-L      | 5 | 95.14±7.58         | 2,293.45±203.64     | 6.65                      |
| LoVoCsc + L-OHP + SalB-L | 5 | 110.29±8.43        | 875.30±106.21       | 64.37                     |

*P<0.01 vs. LoVoCsc group; aP<0.01 vs. LoVoCsc + L-OHP group. L-OHP, oxaliplatin; csc, cancer stem cells; n, number of nude mice; SalB-L, low dose of salvianolic acid B.

Table V. Inhibitory rate of tumor growth following treatment with 5-FU, low doses of SalB, and 5-FU combined with low doses of SalB in xenografts obtained from the HCT-116Csc group (n=5).

| Group                  | n | Pre-administration | Post-administration | Tumor inhibition rate (%) |
|------------------------|---|--------------------|---------------------|---------------------------|
| HCT-116Csc             | 5 | 108.54±10.62       | 3,756.20±416.92     | /                         |
| HCT-116Csc + 5-FU      | 5 | 110.69±9.06        | 2,923.17±347.33     | 22.18                     |
| HCT-116Csc + SalB-L    | 5 | 107.33±10.14       | 3,289.15±383.58     | 12.43                     |
| HCT-116Csc + 5-FU + SalB-L | 5 | 110.86±9.15        | 1,667.37±106.81     | 55.61                     |

*P<0.05 and bP<0.01 vs. HCT-116Csc group; cP<0.01 vs. HCT-116Csc + 5-FU group. csc, cancer stem cells; n, number of nude mice; 5-FU, fluorouracil; SalB-L, low dose of salvianolic acid B.
inoculation of LoVocsc and HCT-116csc cells could be serially passaged in mice. The growth rate of the tumors in the LoVocsc and HCT-116csc groups was demonstrated to be increased compared with rates exhibited by the LoVo and HCT-116 groups. Furthermore, the xenografts exhibited a high malignancy as determined by H&E staining and were resistant to chemotherapy drugs, such as L-OHP and 5-FU. In addition, the derived CSCs from the LoVocsc and HCT-116csc groups exhibited significantly increased expression levels of CD44, CD133, SOX2 and ABCG2 proteins compared with corresponding LoVo and HCT-116 groups. In vivo experiments revealed that treatment with L-OHP or 5-FU combined with SalB had an inhibitory effect on tumor growth and demonstrated a greater efficacy compared with treatment with L-OHP or 5-FU alone. The determined Q values were >1.15, which, based on the method by Jin et al (22), suggested that SalB, when used in combination with L-OHP or 5-FU, exhibited a synergistic inhibition of tumor growth. This suggested that SalB could significantly attenuate the drug resistance of xenografts, and thus improve the efficacy of chemotherapeutic agents. Furthermore, the results of the present study demonstrated that SalB suppressed tumor growth in
a dose-dependent manner. In addition, western blot analysis revealed that treatment with SalB significantly decreased the expression levels of CD44, SOX2 and ABCG2 proteins in LoVoCsc and HCT-116Csc xenografts, and this was exhibited in a dose-dependent manner in LoVoCsc xenografts.

In conclusion, the results of the present study revealed that serum-free suspension cultures may be used to effectively isolate colon CSCs from LoVo and HCT-116 cells. Nude mice models bearing LoVoCsc and HCT-116Csc cells were successfully established, and treatment with SalB was demonstrated to effectively attenuate MDR exhibited by of colon CSCs xenografts via the suppression of CD44, SOX2 and ABCG2 protein expression levels.

A limitation of the present study was the absence of non-cancerous cell line to be used as a control. Future studies should focus on the observation of dynamic alterations of xenograft growth in the orthotopic transplant tumor model of colon CSCs in mice using in vivo optical imaging, as well as tumor invasion and metastasis of liver, lung, brain and other organs. Such investigations would serve to deepen the understanding of in vivo SalB treatment as a novel therapeutic strategy for the treatment of MDR in colon CSCs. In addition, in future studies we will aim to further investigate the anti-MDR effect of SalB using in vitro models as well as determine the effect of SalB on morphological changes of colon CSCs, including the cell refractive index, the cell cycle, drug resistance, proliferation and apoptosis. Furthermore, we will aim to uncover the molecular mechanism underlying the anti-MDR effect of SalB.

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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
PG, YL, JW and WG contributed to the study design. PG, YL, JW, WG, XL, SW, LX and BW performed the experiments. XL, SW and BW analyzed data and contributed to the writing of the manuscript. All authors reviewed the manuscript.

Ethics approval and consent to participate
All animal experiments were performed according to the guidelines of the Chinese Experimental Animals Administration Legislation and were approved by the Ethics Committee for Animal Experiments of Shanghai University of Traditional Chinese Medicine (Shanghai, China; reference no. SZY 201504023).

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.

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