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β -Escin inhibits the proliferation of osteosarcoma cells via blocking the PI3K/Akt pathway

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β -Escin exhibits anticancer effects on a panel of established cancer cells. However, the effects of β -escin on human osteosarcoma (OS) are still unknown. The aim of the present study was to investigate whether β -escin was effective against OS both *in vivo* and *in vitro*. Our results showed that β -escin induced dose- and time-dependent effects against MG-63, OS732, U-2OS, HOS and SAOS-2 cell proliferation. β -Escin also exhibited excellent anti-proliferative and pro-apoptotic effects in an established OS xenograft model. β -Escin and cytotoxic drugs, including cisplatin, methotrexate (MTX), doxorubicin (Dox) and ifosfamide (Ifos), synergistically inhibited proliferation of MG-63 and OS732 cells *in vitro*. Moreover, β -escin induced apoptotic death, activated caspase-3, caspase-8 and caspase-9, and regulated expression of Bax and Bcl-2 in MG-63 cells. In addition, our results showed that β -escin treatment reduced expression of p-PI3K, p-Akt and p-mTOR both in MG-63 cells and in an MG-63 xenograft OS model. Interestingly, SC79, which is an Akt activator, inhibited the anti-proliferative effects of β -escin on MG-63 cells. Taken together, our data support the conclusion that β -escin effectively inhibits OS proliferation both *in vivo* and *in vitro*. The inhibitory effect of β -escin, at least in part, is due to the inactivation of the PI3K/Akt signalling pathway.

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Introduction

Osteosarcoma (OS) is a common primary malignant bone tumour that is most prevalent in children and young adults.¹ Currently, OS is the eighth most-common type of cancer among children, accounting for 2.4% of all malignancies in paediatric patients and 35% of all bone cancers.¹ During the past decades, a combination of surgical resection and chemotherapy has made significant progress in preventing OS development, and the 5 year survival rate of localized OS is approximately 65–70%. However, OS patients with metastatic or recurrent disease at the time of diagnosis have poor prognoses, with a 5 year survival rate of only 20%.^{2,3} More strikingly, radiotherapy and chemotherapy cause serious systemic toxicity, for which OS patients have low tolerance.^{1,4} Therefore, a new effective drug for OS treatment is certainly needed.

In recent years, traditional Chinese medicines have received wide attention for their multisided, multi-target pharmacological actions and fewer side effects in the treatment of tumours. β -Escin (Fig. 1) is a pentacyclic triterpenoid extracted from horse chestnut (*Aesculus hippocastanum*) seeds and is currently used to treat encephalitis, trauma,

surgery-induced turgidity, and disordered venous reflux.^{5–7} In previous investigations, β -escin has been reported to possess an anti-arthritic effect, and anti-oedematous and anti-inflammatory properties in various disease models.^{8–11} More strikingly, it is widely reported that β -escin effectively inhibits proliferation of various human cancer cell lines (cholangiocarcinoma, prostate cancer cells, gastric adenocarcinoma, colon adenocarcinoma, lung cancer cells, hepatocellular carcinoma, leukaemia) *in vivo* and *in vitro* through multiple signalling pathways.^{12–19} However, whether β -escin has inhibitory effects on OS is poorly understood. Therefore, to explore the possibility of developing β -escin as a therapeutic agent, in the present study we investigated its effect on human OS cells.

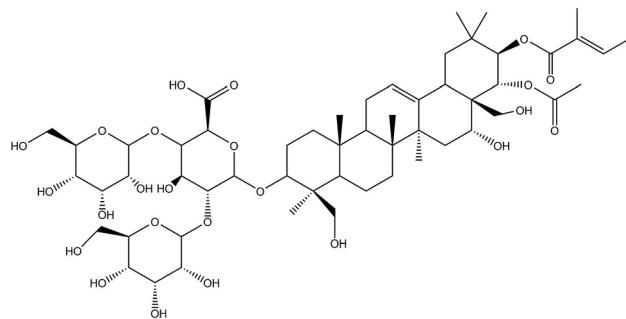


Fig. 1 Chemical structure of β -escin. Molecular weight: 1131.





Materials and methods

Cell culture

Human osteosarcoma cell lines MG-63, OS732, U-2OS, HOS and SAOS-2 were purchased from the Cell Bank of Shanghai Institute of Biochemistry and Cell Biology, Chinese Academy of Sciences (Shanghai, China). The cells were grown in Dulbecco's modified Eagle's medium (DMEM) (Gibco, Invitrogen, Carlsbad, CA, USA) supplemented with 10% heat-inactivated FBS, 100 U mL⁻¹ penicillin and 100 µg mL⁻¹ streptomycin. All cells were cultivated in a humidified atmosphere containing 5% CO₂ at 37 °C. The *in vitro* study was approved by the Ethics Committee on scientific research of the First Affiliated Hospital of Wenzhou Medical University.

MTT assay

All cell lines proliferated up to 70–80% confluence and then seeded in 96-well plates at a density of 1 × 10⁴ cells per well. The cells were treated with β-escin (Aladdin Chemistry Co, Shanghai, China, 95% purity) at various concentrations (0, 5, 10, 20, 50 or 100 µM) for indicated times (24–72 h). Four hours before the end of each incubation period, MTT solution (5 mg L⁻¹, Sigma-Aldrich, St. Louis, MO, USA) was added to each well and incubated for 4 h at 37 °C. Then, the cultured medium was replaced with DMSO. The optical density (OD) was recorded at a wavelength of 570 nm by an enzyme immunoassay analyser (Bio-Rad, Hercules, Calif., USA). To investigate whether the combinations of β-escin and anti-OS cytotoxic drugs (methotrexate: MTX; doxorubicin: Dox; ifosfamide: Ifos; cisplatin; Sigma-Aldrich, St. Louis, MO, USA) exhibit stronger inhibitory effects on OS proliferation, MG-63 and OS732 cells were cultured in 96-well plates, and then treated with different concentrations of cytotoxic drugs in the presence of β-escin (5 µM or 10 µM), or with different concentration of β-escin in the presence of cytotoxic drugs for 48 h. In addition, to study the effects of protein kinase B (Akt) activator SC79 (purity > 97%, Sigma) on the anti-proliferative action of β-escin, MG-63 cells were treated with β-escin in the presence of 20 µM SC79 for 48 h. The rate of inhibition of cellular proliferation was calculated as [1-A570 (test)/A570 (control)] × 100%. IC₅₀ was calculated based on nonlinear regression analysis of survival curves (SPSS software, version 13.0, Chicago, IL, USA). All data were collected from three independent experiments.

Apoptosis assay

MG-63 cells were maintained in the medium containing β-escin (2.5–10 µM) for 48 h. Then, the cells were harvested and washed twice with PBS and re-suspended in 500 µL binding buffer. The cells were stained with Annexin V-fluorescein isothiocyanate (20 µg mL⁻¹, FITC, 5 µL, Life Technologies, Waltham, MA, USA) and propidium iodide (PI) (50 µg mL⁻¹, 5 µL, Life Technologies, Waltham, MA, USA) for 20 min. The apoptotic rate of MG-63 cells was analysed by flow cytometry using a BD fluorescence activated cell sorting (FACS) flow cytometer (BD Biosciences, USA).

The apoptotic cells in the tumour tissue were detected using terminal deoxynucleotidyl transferase dUTP nick end labelling (TUNEL) assay according to the manufacturer's instructions (Promega, Madison, WI, USA). For nuclear staining, 4',6-diamidino-2-phenylindole (DAPI) was used. TUNEL-positive and DAPI-positive staining was visualized in a confocal laser scanning microscope (SP8, Leica Microsystems, Germany). The apoptotic index was calculated as a ratio of an average percentage of positive nuclei, including apoptotic bodies, to a total number of nuclei from five random observation fields at 400× magnification.

Caspase activity assay

Caspase-3, caspase-8 and caspase-9 activities were detected by a commercial caspase assay kit according to the manufacturer's instruction (R&D systems, Minneapolis, MN). To investigate the effect of β-escin on caspase activation, MG-63 cells (1 × 10⁶ cells per mL) received β-escin treatment (2.5, 5.0, 10.0 µM) for 48 h. Then, the supernatant of lysed cells was added to the reaction mixture containing dithiothreitol and caspase substrates (for caspase-8, caspase-9 and caspase-3), and the mixture was incubated for 1 h at 37 °C. Finally, the chromophore absorbance was measured and recorded at 405 nm. The data are shown as the fold-changes relative to control.

Western blot analysis

MG-63 cells were harvested and homogenized in an ice-cold cell lysis buffer (Bio-Rad, Hercules, CA, USA). Cellular lysates were centrifuged at 12 000 × g for 30 min to separate the supernatant. The protein extraction from the tumour tissues was performed in the same way. A bicinchoninic acid protein assay kit (Pierce, Rockford, IL, USA) was used to determine the total protein concentration. Then, equal amounts of total proteins were separated on 12% sodium dodecyl sulfate-polyacrylamide gels and transferred to polyvinylidene difluoride membranes (Millipore Corporation, Billerica, MA, USA). The membranes were blocked with 5% skimmed milk for 2 h and then probed with specific primary antibodies against Bax, Bcl-2, phosphatidyl inositol 3-kinase (PI3K), p-PI3K, Akt, p-Akt, mammalian target of rapamycin (mTOR) and p-mTOR (Abcam, Cambridge, UK) at 4 °C overnight. After washing three times, the membranes were further incubated with secondary antibodies coupled to horseradish peroxidase. The blots were visualized using electrochemiluminescence reagent (Millipore, Bedford, MA, USA). β-actin was used for normalization.

In vivo tumour model

The animal study was carried out according to the U.K. Animals (Scientific Procedures) Act, 1986 and associated guidelines and approved by the Ethics Commission of Wenzhou Medical University. Male Balb/c nude mice aged 6 weeks (Model Animal Research Center of Nanjing University, Nanjing, China) were subcutaneously injected into left side with MG-63 cells (5 × 10⁶ cells). When the tumours reached a size of approximately 150 mm³ (20 days after transplantation), mice were treated with β-escin (5 mg kg⁻¹, 10 mg kg⁻¹) by daily oral administration for 3

weeks. The tumour volume during treatment was determined at an interval of two days by caliper measurement [$V = 1/2 \times (L \times W^2)$]. Additionally, the tumour tissues were weighted at the end of study.

Immunohistochemistry

The tumour tissues were removed at the end of study, and 5 μm -thick sections of paraffin-embedded, formalin-fixed tumour tissues were stained. In brief, the sections were dewaxed with xylene and hydrated in decreasing concentrations of ethanol. Endogenous peroxidase activity was quenched using 3% H_2O_2 (MSDS88597, Sigma) in absolute methanol for 20 min at room temperature. Sections were then blocked for 1 h using a HistostainTM-SP Kit (SP9002, ZSGB Biotechnology Co. LTD, Beijing, China; <https://www.zsbio.com/product/SPN-9002>). The p-PI3K and p-Akt staining was performed according to the manufacturer's instruction (ab182651, ab81283, Abcam, Cambridge, UK). After washing with TBS solution (Tris 10 mM, NaCl 0.15 M, pH 7.5), localization of the antigen-antibody complexes was performed by adding diaminobenzidine (D3939, Sigma).

Statistical analysis

All experiments were repeated a minimum of three times. Data were presented as the mean (M) \pm standard deviation (SD). For analysis of difference between two groups, Student's *t* test (SPSS software, version 13.0, Chicago, IL, USA) was performed. A *P* value less than 0.05 was considered statistically significant.

Results

β -Escin inhibits the proliferation of OS cells *in vitro*

Cytotoxic effects of β -escin on MG-63, OS732, U-2OS, HOS and SAOS-2 cells were investigated by MTT. β -Escin showed the dose- and time-dependent proliferation inhibition effects on all OS cells (Fig. 2A–E, *P* < 0.01). The OS732 and MG-63 cells were more sensitive to β -escin treatment, whereas HOS & SAOS-2

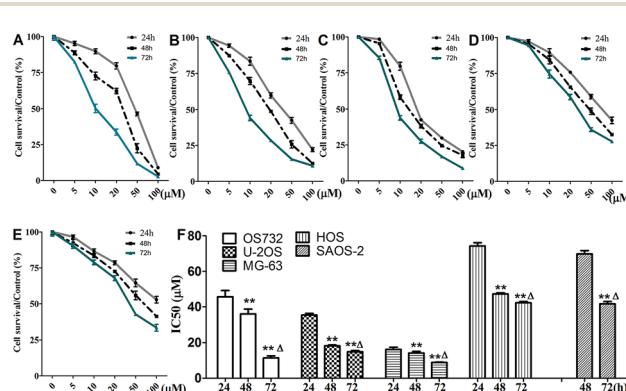


Fig. 2 Viability of OS cells following treatment with β -escin. OS732 (A), U-2OS (B), MG-63 (C), HOS (D) and SAOS-2 (E) cells were treated with β -escin (0, 5, 10, 20, 50 or 100 μM) for the indicated time. The cell viability was evaluated by MTT, and IC₅₀ values (F) were calculated based on nonlinear regression analysis of survival curves. Data were collected from three independent experiments. ***P* < 0.01 vs. the 24 h group; Δ *P* < 0.01 vs. the 48 h group.

exhibited less response (Fig. 2F). These data indicate that β -escin effectively reduces the proliferation of OS cells.

β -Escin and cytotoxic drugs synergistically inhibit OS cell proliferation *in vitro*

Based on the results showed in Fig. 2, we found β -escin exhibited stronger inhibitory effects on proliferation of MG-63 and OS732 cells than the other cell lines. Therefore, MG-63 and OS732 cells were preferentially used to evaluate anti-proliferative effects of β -escin with anti-OS cytotoxic drugs together. Cisplatin, MTX, Dox and Ifos are commonly used anti-OS drugs. As seen in Fig. 3A–D, β -escin (5 or 10 μM) significantly reduced IC₅₀ values of cisplatin, MTX, Dox and Ifos (*P* < 0.01). For example, β -escin (5 μM) decreased IC₅₀ value of cisplatin from 31.7 μM to 18.9 μM , decreased IC₅₀ value of MTX from 3.13

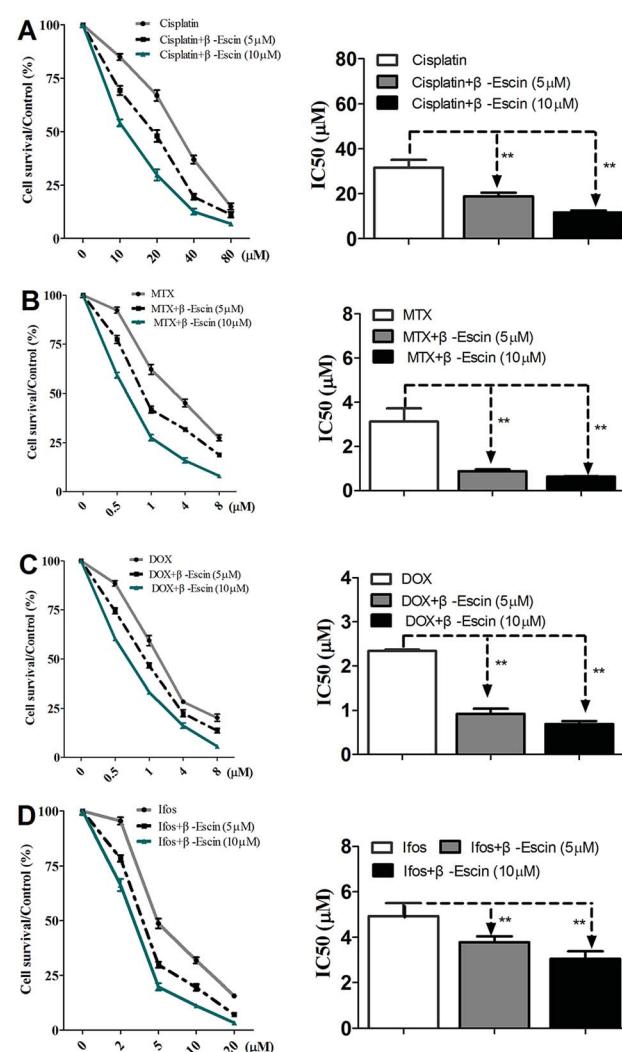


Fig. 3 Effects of β -escin on cytotoxic drug-induced inhibition of proliferation of MG-63 cells *in vitro*. MG-63 cells were treated with cisplatin (A), MTX (B), Dox (C) and Ifos (D) in the presence of β -escin (5 μM or 10 μM) for the indicated time, and then the cells viability was measured by MTT. Data are presented as the mean (M) \pm standard deviation (SD), and the results were collected from three independent experiments. ***P* < 0.01 vs. the chemotherapeutic drug only group.

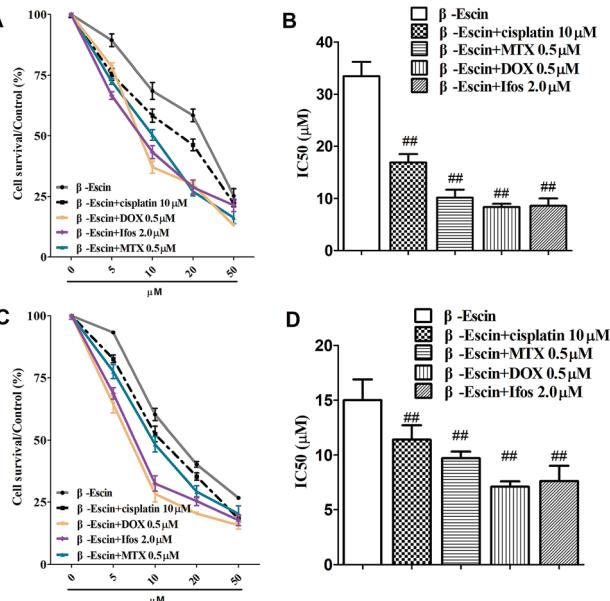


Fig. 4 Effects of cytotoxic drugs on β -escin-induced inhibition of proliferation of OS732 and MG-63 cells *in vitro*. OS732 (or MG-63) cells were treated with β -escin in the presence of cytotoxic drugs (cisplatin, MTX, Dox or Ifos) for the indicated time, and the cell viability was measured by MTT. The OS732 cell survival curves are shown in (A), and the IC₅₀ values of β -escin in the presence of cytotoxic drugs are shown in (B). The curves for MG-63 cells are shown in (C), and the IC₅₀ values of β -escin in the presence of cytotoxic drugs are shown in (D). Data are presented as the mean (M) \pm standard deviation (SD), and the results were collected from three independent experiments. $^{##}P < 0.01$ vs. the β -escin only group.

μ M to 0.88 μ M, and reduced IC₅₀ value of Ifos from 4.92 μ M to 3.78 μ M. Moreover, it should be noted that a combined use of reduced concentration of cytotoxic drugs and β -escin could maintain their therapeutic effects. For instance, a combined therapy with cisplatin (10 μ M) and β -escin (5 μ M) showed the same inhibitory effect as with cisplatin (20 μ M) only. OS cells received a combined therapy with MTX (0.5 μ M) and β -escin (10 μ M) showed the same survival rate (59.2% vs. 62.1%, $P > 0.05$) as cells received MTX (1 μ M). Moreover, as seen in Fig. 4A–D, the cytotoxic drugs studied here significantly reduced IC₅₀ value of β -escin when treating MG-63 and OS732 cells. These data suggest that β -escin and anti-OS cytotoxic drugs synergistically inhibit proliferation of human OS cells.

β -Escin induces apoptosis of MG-63 cells

As shown in Fig. 5, following treatment with β -escin (2.5 μ M, 5 μ M or 10.0 μ M), the earlier and late apoptosis rates of MG-63 cells were both increased ($P < 0.01$). Our results suggest that β -escin induces apoptotic death in the MG-63 cells.

β -Escin induces activation of caspase-3, caspase-8 and caspase-9 in MG-63 cells

As shown in Fig. 6, caspase-3 in β -escin pre-treated MG-63 cells was remarkably activated ($P < 0.01$). Moreover, caspase-8 and caspase-9 were both activated by the treatment with β -escin ($P < 0.01$). These results indicate that the inhibitory effect of β -escin

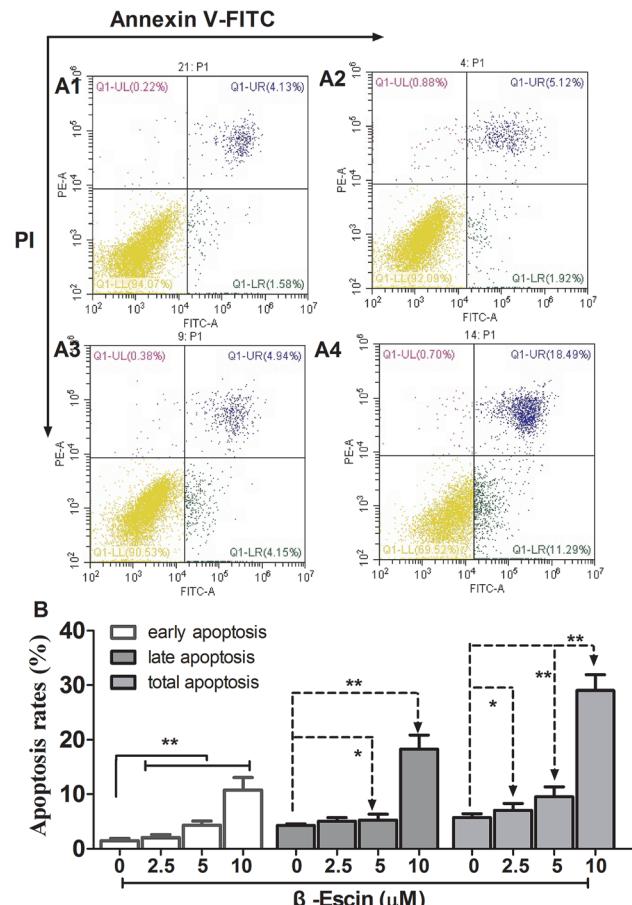


Fig. 5 Apoptosis of MG-63 cells following treatment with β -escin. MG-63 cells were treated with β -escin (2.5, 5 or 10 μ M) for 48 h, and then double-stained by Annexin V/FITC, and analysed by flow cytometry. Representative images are shown in A1 (0 μ M), A2 (2.5 μ M), A3 (5 μ M) and A4 (10 μ M). The apoptotic rates of MG-63 cells treated with β -escin are shown in (B). Data were collected from three independent experiments. $^{**}P < 0.01$ vs. the A1 group.

on cell proliferation was mediated through activation of caspase activity. It is proposed that β -escin triggers both the extrinsic and intrinsic apoptotic pathways in MG-63 cells.

β -Escin down-regulates expression of apoptosis-related proteins

As shown in Fig. 7A, after treatment with β -escin, the ratios of Bax/Bcl-2 were up-regulated in MG-63 cells. Specifically, 5 and 10 μ M β -escin caused the 168% and 440% increases in the Bax/Bcl-2 ratio, respectively, compared with that of the control group ($P < 0.01$).

β -Escin inhibits tumour growth in MG-63-xenografted mice

To determine whether β -escin could inhibit tumour growth *in vivo*, we further investigated the effects of β -escin on tumour growth in MG-63 xenografted mice. The results showed that β -escin treatment significantly reduced the tumour volume and weight in MG-63 xenografted nude mice (Fig. 9A and B, $P < 0.01$). Furthermore, β -escin treatment significantly increased the apoptotic rates of tumour cells in the mice (Fig. 10B, $P < 0.01$).



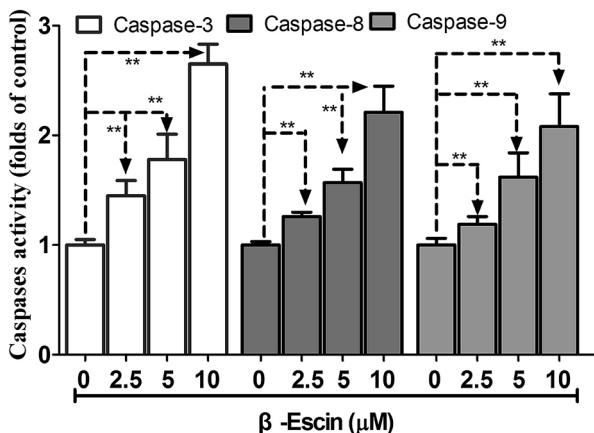


Fig. 6 β -Escin induces activation of caspase-3, caspase-8 and caspase-9 in MG-63 cells. MG-63 cells were treated with β -escin for 48 h, and then were lysed. Supernatants of the lysed cells were reacted with caspase substrates, and the levels of caspase-3, caspase-8 and caspase-9 activity were expressed as the chromophore absorbance at 405 nm, which was determined by a microplate reader. Data were collected from three independent experiments. ** $P < 0.01$ vs. the group without β -escin treatment.

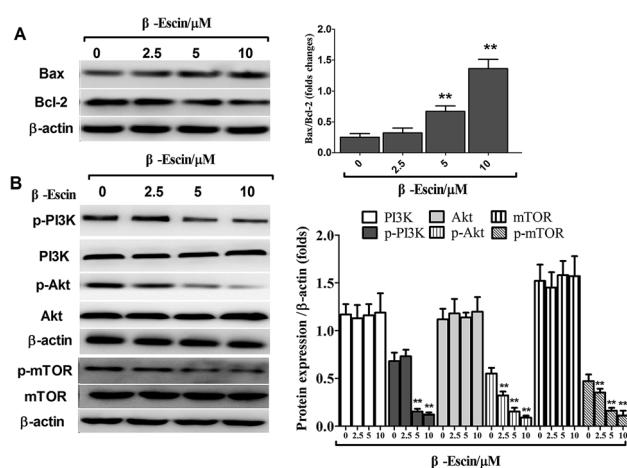


Fig. 7 β -Escin regulated Bax and Bcl-2 expression and blocked the PI3K/Akt pathway in MG-63 cells. (A) MG-63 cells were treated with β -escin for 48 h, and the levels of Bax and Bcl-2 were measured by western blotting. (B): MG-63 cells were treated with β -escin for 48 h, and intracellular levels of PI3K, p-PI3K, Akt, p-Akt, mTOR and p-mTOR were analysed by western blotting. Data were collected from three independent experiments. ** $P < 0.01$ vs. the group without β -escin treatment.

β -Escin blocks the PI3K/Akt pathway in MG-63 cells and MG-63-transplanted nude mice

To elucidate the underlying anti-proliferative mechanism of β -escin, we evaluated the β -escin effects on the PI3K/Akt pathway in MG-63 cells. As seen in Fig. 7B, no obvious difference was observed in the expression of PI3K and Akt between the β -escin group and the control group ($P < 0.01$). Interestingly, we noticed that the levels of p-Akt, p-PI3K and p-mTOR in MG-63 cells were significantly decreased compared with the control ($P < 0.01$). To

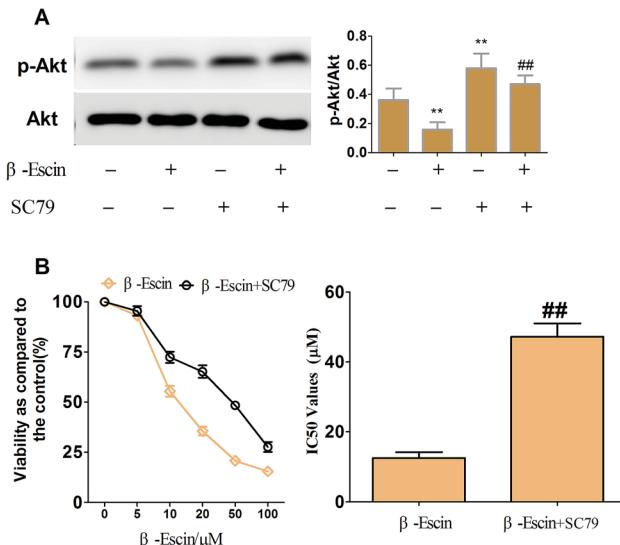


Fig. 8 SC79 reduced the β -escin-induced inhibitory effects on MG-63 cell proliferation. MG-63 cells were treated with β -escin in the presence (+) or the absence (−) of 20 μ M SC79 for 48 h, and then the cell viability was assayed by MTT. The expression of Akt and p-Akt was detected by western blotting. (A) Typical blot bands of target proteins were quantified. (B) SC79 increased the IC₅₀ value of β -escin. Data were collected from three independent experiments. ** $P < 0.01$ vs. the group without β -escin and SC79 treatment. ## $P < 0.01$ vs. the group of β -escin in the absence of SC79 treatment.

further characterize the role of the PI3K/Akt pathway in the anti-proliferative effect exerted by β -escin, we used Akt activator SC79 to interfere with the action of β -escin. As seen in Fig. 8A and B, SC79 significantly increased p-Akt levels in MG-63 cells treated with β -escin and exerted an antagonistic effect to the anti-OS action of β -escin ($P < 0.01$). Moreover, we have also observed that oral administration of β -escin significantly reduced the expression of p-Akt, p-PI3K and p-mTOR in the tumour tissues (Fig. 9C, Fig. 10A, $P < 0.01$). Therefore, our data suggest that β -escin exerts its anti-OS activity by affecting the PI3K/Akt pathway.

Discussion

β -Escin is a well-known pentacyclic triterpenoid extracted from horse chestnut (*Aesculus hippocastanum*) seeds. In the present study, we showed that β -escin reduced the proliferation of OS cells in a dose- and time dependent manner. Moreover, we have also observed that β -escin exhibited anti-proliferative effects in an established OS xenografted mice model. The outcomes reported here are consistent with previous results. For instance, β -escin showed significant inhibitory effects on proliferation of hepatocellular carcinoma cells (IC₅₀: 65.7 μ g mL⁻¹)¹², ovarian cancer cells (IC₅₀: 10.0 μ g mL⁻¹)¹³ and cholangiocarcinoma cells (IC₅₀: 33–63 μ M).¹⁴ Our results showed that IC₅₀ values of β -escin ranged from 14.2 μ M to 69.8 μ M in the tested OS cells. These data indicate that OS cells are highly sensitive to β -escin treatment and that β -escin is a promising anti-OS drug. Further experiments were performed to investigate a combined use of β -

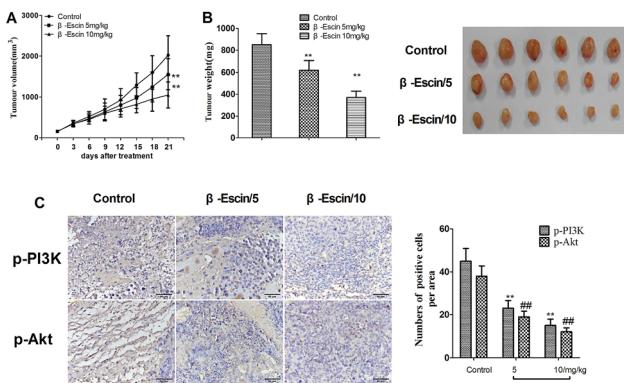


Fig. 9 β -Escin effectively inhibited tumour growth in MG-63-transplanted nude mice. MG-63-transplanted nude mice were established, and orally treated with β -escin for 3 weeks. The expression of p-PI3K and p-Akt in the tumour tissue was detected by immunohistochemical staining. (A) Tumour volume; (B) tumour weight; (C) the p-PI3K and p-Akt-positive cells in tumour tissue. Data are presented as the mean (M) \pm standard deviation (SD), $n = 6$, ** $P < 0.01$ or **# $P < 0.01$ vs. the control group (the group without β -escin treatment).

with cisplatin and ADR.⁴ β -Escin is a low-toxic compound.²² Previous *in vitro* studies reported^{16,23} that the survival of normal prostate epithelial cell line was minimally affected by escin at a cytotoxic concentration (24 μ M) and that the survival of normal rat skeletal muscle cells was hardly influenced by escin at a high concentration (50 μ M). These reports imply that cytotoxic effects of β -escin have high selectivity between normal and cancer cells. In the present study, we first showed that the combined application of β -escin and anti-OS cytotoxic drugs (cisplatin, MTX, Dox or Ifos) synergistically inhibited proliferation of MG-63 or OS732 cells *in vitro*. We speculate that the main reasons are as following: first β -escin and chemotherapy possess beneficial effects in inhibiting OS cell proliferation based on their different molecular mechanisms; second, it is well known that P-glycoprotein is highly expressed in tumour cells, and pumps many anti-tumour drugs out of the cells to induce tumour resistance.²⁴ In a previous study, β -escin (5 μ M and 10 μ M) has been reported to reduce the expression of P-glycoprotein in cholangiocarcinoma.²⁵ Therefore, it is suggested that β -escin may have beneficial effects on absorption of the anti-OS cytotoxic drugs in the tumour cells, which results in enhanced anti-tumour effects. However, further research is required to test this possibility.

Apoptosis is one of the major mechanisms of cell death. It is well known that caspase signalling cascades control the process of apoptosis. Extrinsic and intrinsic cascades are triggered by caspase-8 and caspase-9, respectively. Caspase-8 activates downstream caspases and leads to apoptosis through the death receptor pathway, whereas activation of caspase-9 further activates the executioner caspase-3 and induces apoptosis through the mitochondrial pathway.^{26,27} The members of Bcl-2 family, such as Bcl-2 and Bcl-xL, regulate the mitochondrial apoptosis pathways by changing mitochondrial permeability.²⁸ β -Escin-mediated apoptosis has been previously described in other cells.^{12,19} Consistent with the previous results, our data revealed that β -escin promoted the apoptosis process both at the early and late stages as determined by FACS of Annexin V/PI-labelled cells. Moreover, we observed that β -escin significantly increased activities of caspase-3, caspase-8 and caspase-9 and simultaneously regulated the expression of Bax and Bcl-2. The current findings are consistent with the previously published results for colon cancer cells, hepatocellular carcinoma cells and human cholangiocarcinoma cells.^{12,14,29} However, it should be noted that some cases of chemotherapy-mediated apoptosis differ from those found by us. For instance, the cisplatin-induced apoptosis in Hep-2 cells is *via* modulation of caspase-9 activity instead of caspase-8,³⁰ and 5-fluorouracil enhances activation of caspase-8 rather than of caspase-9 in SMMC-7721 cells.¹² Therefore, our findings may explain why β -escin and chemotherapies showed synergistic inhibitory effects on the proliferation of OS cells.

The PI3K/Akt signalling pathway is well known for its key role in the cell proliferation and survival. Normally, PI3K generates PIP3 at plasma membrane, and PIP3 phosphorylates Akt to form p-Akt. Subsequently, p-Akt regulates expression of many molecules to affect cell proliferation and apoptosis.³¹ In OS, the PI3K/Akt pathway is deregulated in most cases of

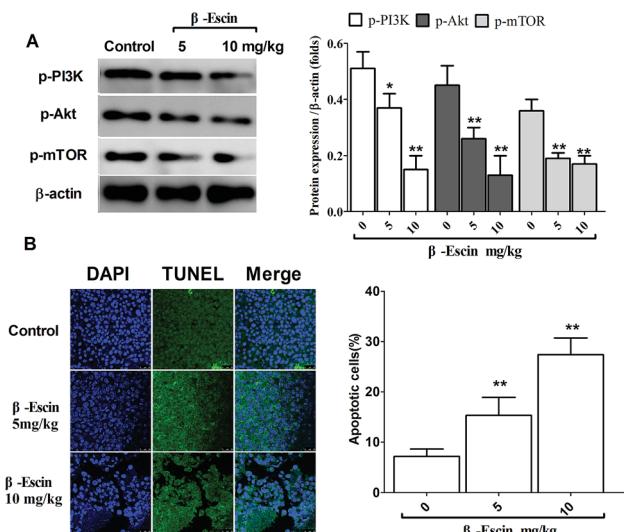


Fig. 10 β -Escin promoted apoptotic death and blocked the PI3K/Akt pathway in the MG-63-transplanted nude mice. The levels of p-PI3K and p-Akt in the tumour tissue were detected by western blotting, and the apoptotic death was evaluated by the DAPI/TUNEL staining, which was visualized in a laser confocal microscope. (A) Typical blot bands of p-PI3K, p-Akt and p-mTOR in the tumour tissues were quantified. (B) Apoptotic death in the tumour tissue was analysed by DAPI and TUNEL double-staining. Data are presented as the mean (M) \pm standard deviation (SD), $n = 6$, * $P < 0.05$, ** $P < 0.01$ vs. the control group (the group without β -escin treatment).

escin and commonly used anti-OS cytotoxic drugs. In the 1970s, chemotherapy started to be used for treating OS.^{20,21} Since then, combination chemotherapies have been introduced to enhance curative effects but caused significant side-effects, such as nausea, emesis, alopecia and myelosuppression. For example, it has been previously shown that most patients with high-grade OS experienced grade III-IV nausea, vomiting and haematologic toxicities after pre-operative combination chemotherapy



localized disease and in 100% cases of advanced-stage disease,³² which implies that alterations in this pathway may be a pre-requisite for OS progression. Three reports^{33–35} have shown that p-Akt markedly increased c-Myc expression alongside an increase in the OS cell proliferation. Moreover, silencing miR-223 increases protein levels of PI3k and p-Akt and promotes MG-63 cell proliferation, whereas knockdown of CXCR7 inhibits proliferative activity of OS cells, partially through the decreased expression of PI3K and Akt. It has been previously confirmed that in the apoptotic process in OS cells increasing HSP70 expression leads to activation of PI3K/Akt, which partially prevents human OS cells from undergoing apoptosis.³⁶ Furthermore, knockdown of Akt2 increases the apoptotic sensitivity of SAOS-2 cells in response to cisplatin treatment.³⁷ Therefore, the PI3K/Akt pathway is a promising anti-OS therapeutic target. In previous studies, it has been concluded that β -escin chemosensitizes pancreatic cancer cells *via* deactivation of the nuclear factor-kappa B pathway³⁸ and induces apoptosis of hepatocellular carcinoma cells *via* down-regulation of the STAT3/JAK2 signalling pathway.³⁹ In the present study, we found that β -escin effectively reduced expression of p-PI3K, p-Akt and p-mTOR both *in vitro* and *in vivo*. Induction of Akt activation by SC79 exerted an antagonistic effect to the anti-OS effects of β -escin. Therefore, our results indicate that β -escin inhibits OS cell proliferation, at least in part, by inactivation of the PI3K/Akt pathway.

Conclusions

Taken together, in this study we found that β -escin inhibits proliferation of human OS cells *in vitro* and *in vivo*. Further work showed that β -escin blocked the PI3K/Akt pathway in MG-63 cells and MG-63 xenograft OS model. Our findings suggest that β -escin and cytotoxic drugs may be used together to treat OS patients.

Conflicts of interest

The authors report no conflicts of interest.

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