Abstract. Progressive macrophage dysfunction and apoptosis are some of the major events that occur during atherogenesis. To further investigate the intrinsic association between atherosclerosis (AS) and macrophage apoptosis and autophagy, cholesterol crystals (CHCs) were used to stimulate RAW264.7 macrophages to establish a macrophage model of advanced AS. Cells in the CHC group were treated with salvianolic acid B (Sal B) to evaluate its protective effects and reveal its underlying molecular mechanism. The results demonstrated that treatments with Sal B significantly improved autophagy dysfunction and reduced the apoptotic rate of CHC-induced macrophages. Furthermore, Sal B significantly attenuated CHC-induced release of proinflammatory factors (TNF-α and IL-6) by macrophages. Treatment of macrophages with a specific inhibitor of autophagy (3-methyladenine) significantly reversed Sal B-mediated effects on autophagy, suggesting that Sal B-induced autophagy may display a protective effect in CHC-induced macrophages. Furthermore, pretreatment of CHC-induced macrophages with insulin significantly decreased Sal B-induced autophagy, indicating that the Akt/mTOR signaling pathway may serve as a critical mediator in regulating Sal B-mediated cell death. Taken together, the present study demonstrated that Sal B improved autophagic dysfunction and reduced the apoptosis of CHC-induced macrophages via inhibiting the Akt/mTOR signaling pathway.

Introduction

According to the World Health Organization, among the 56.9 million deaths recorded worldwide in 2016, 15.2 million were caused by ischemic heart disease and stroke. Therefore, cardiovascular diseases have consistently ranked first in the list of global causes of death during the last 15 years (1). Atherosclerosis (AS) is an important foundation for these two diseases. The accumulation of foam cells and apoptotic residues, which are produced by foam cells, at the lesions are landmark events of both heart disease and stroke (2,3). In the early stage of AS, macrophages reduce the lipids and apoptotic residues to limit plaque growth via endocytosis. During disease progression, the function of macrophages is decreased until it is lost, which is accompanied by increased cell apoptosis, eventually leading to necrosis, inflammation and plaque instability (4). The aforementioned findings suggested that reducing macrophage apoptosis and maintaining their normal function could be considered as a promising approach for the prevention and treatment of AS.

Autophagy is a cellular self-defense mechanism that has been widely studied in several organisms and is characterized by the intracellular degradation of organelles, proteins and other cytoplasmic components by lysosomes (5). When cells are exposed to external stimuli, such as oxidative stress and starvation, the lipid structure of the bilayer membrane encapsulates cytoplasmic substances, such as damaged organelles and misfolded proteins, to form autophagosomes. Subsequently, the autophagosome fuses with the lysosome to form an autophagosomal, which in turn catalyzes cytoplasmic materials, including those internalized by endocytosis, through its acidic environment and proteases, thereby maintaining normal cell activity, morphology and structure, and delaying the aging process (6). Autophagy is also closely associated with the pathogenesis and prevention of malignant tumors, as well as cardiovascular, neurodegenerative and immune system diseases (7). For instance, it has been reported that macrophage autophagy exhibits an antiatherosclerotic effect via inhibiting inflammation and promoting cholesterol efflux (8,9).
Recently, an increasing number of researchers have turned their attention to the regulation of autophagy in apoptosis (10,11). However, the intrinsic mechanisms underlying the association between autophagy and apoptosis have not yet been elucidated.

As a traditional Chinese medicine, *Salvia miltiorrhiza* is widely used in Asia for the prevention and treatment of cardiovascular diseases via promoting blood circulation (12). Salvianolic acid B (Sal B) is one of the main water-soluble components of *Salvia miltiorrhiza* (13). It has been reported that Sal B exerts anti-inflammatory, antioxidant and anti-tumor effects (12). In cardiovascular diseases, Sal B serves a significant role in protecting against myocardial infarction, improving myocardial ischemia-reperfusion injury and inhibiting the formation of atherosclerotic plaques (13-15). However, whether the antiatherosclerotic effects of Sal B are associated with macrophage apoptosis is not completely understood.

In the present study, RAW264.7 macrophages were treated with cholesterol crystals (CHCs) to investigate the intrinsic association between macrophage autophagy and apoptosis in AS. Furthermore, the mechanism underlying the protective effects of Sal B on macrophages in AS was explored.

### Materials and methods

**Cell culture.** The RAW264.7 cell line was purchased from the American Type Culture Collection and cultured in high glucose DMEM (Gibco; Thermo Fisher Scientific, Inc.) supplemented with 10% (v/v) FBS (Biological Industries) at 37°C in a 5% CO₂ incubator.

**Experimental design.** In order to confirm whether CHCs can induce macrophage apoptosis and autophagy dysfunction, different concentrations of CHCs (0, 400, 800, 1,200 µg/ml) were used to culture cells, followed by verification via Hoechst staining and western blotting. Next, different concentrations of Sal B (0, 40, 80, 120, 200, 240, 280, 320, 360 µM) were used to culture cells and toxicity was evaluated via an MTT assay. A total of three different concentrations (100, 150, 200 µM) were selected for subsequent experiments. Afterwards, cells were divided into five groups (untreated control; 800 µg/ml CHC; 800 µg/ml CHC + 100 µM Sal B; 800 µg/ml CHC + 150 µM Sal B; and 800 µg/ml CHC + 200 µM Sal B) and then it was evaluated whether Sal B could prevent CHC-induced apoptosis, autophagy and inflammation via Hoechst, flow cytometry, western blotting and ELISA. In addition, the effect of Sal B on the autophagic flow of CHC-treated cells was detected via an Ad-GFP-LC3B adenovirus infection experiment [experimental groups were as follows: Untreated control; 800 µg/ml CHC; 800 µg/ml CHC + 200 µM Sal B; and 800 µg/ml CHC + 200 µM Sal B + 5 mM 3-methyladenine (3-MA; Selleck Chemicals)] and the relationship between the reduction in apoptosis and the improvement of autophagy mediated by Sal B was explored via western blotting and flow cytometry. Ultimately, cells were divided into four groups.

**Preparation of CHCs.** Cholesterol powder (Beyotime Institute of Biotechnology) was diluted in 95% ethanol to a final concentration of 12.5 mg/ml and then heated to 65°C. Once sufficiently dissolved, the cholesterol solution was filtered through filter paper to remove undissolved impurities. Subsequently, the solution was cooled in an ice bath to precipitate crystals and then filtered through filter paper to obtain crystals. The aforementioned steps were repeated five times. Subsequently, all crystals were collected, grounded using a mortar, dispersed in PBS, adjusted to a final concentration of 5 mg/ml and stored at -20°C following autoclaving.

**Cell viability assay.** RAW264.7 cells (5x10⁴/well) were seeded into 96-well plates and treated with different concentrations of Sal B for 24 h. Then, the activity of cells was evaluated using an MTT assay. Subsequently, each well was supplemented with 5 µl MTT solution (5 mg/ml; Beyotime Biotechnology). Following incubation for 3 h, the medium was discarded and 150 µl DMSO (Sangon Biotech Co., Ltd.) was added to each well to dissolve the formed crystals. Finally, the optical density of each well was measured using a microplate reader (Molecular Devices, LLC) at a wavelength of 490 nm.

**Cell apoptosis analysis.** Following treatment with the corresponding treatments for 24 h, RAW264.7 cell apoptosis was evaluated by performing Hoechst staining. Briefly, cells (4x10⁴ cells/well) were seeded into 6-well plates and cultured for 24 h, then the medium was discarded. After washing with PBS, cells were fixed with 4% paraformaldehyde (500 µl/well; Boster Biological Technology) for 15 min at 4°C. Cells were washed again with PBS and then stained with 500 µl Hoechst 333258 (Beyotime Institute of Biotechnology) for 10 min at room temperature in the dark. Finally, after washing with PBS, morphological changes were observed under a fluorescence microscope (Thermo Fisher Scientific, Inc.). In addition, RAW264.7 cells (5x10⁴ cells/well) were seeded into 6-well plates and treated with CHCs for 24 h. Then, the apoptosis rate was detected by conducting Annexin V-FITC/PI double staining. Briefly, the medium was aspirated, and cells were washed with PBS, digested with trypsin (Beyotime Institute of Biotechnology), collected into a centrifuge tube and centrifuged at 400 x g for 4 min at 4°C. Cells (1x10⁴ cells/ml) were resuspended in 1X Binding Buffer (BD Biosciences). Subsequently, 100 µl cell suspension was supplemented with 5 µl Annexin V-FITC and 5 µl PI (both from BD Biosciences). Following staining for 15 min at room temperature in the dark, 400 µl 1X Binding Buffer was added and the apoptotic rate was measured within 30 min via flow cytometric analysis (BD FACSymphony™ S6; Becton, Dickinson and Company). FlowJo software version 10.5.3 (FlowJo LLC) was used for analysis.
Detection of the inflammatory factors TNF-α and IL-6. Following treatment with the corresponding intervention drugs for 24 h, the levels of the inflammatory factors TNF-α and IL-6 were measured by performing ELISAs. Cells were centrifuged at 400 x g for 20 min at 4°C. Subsequently, the levels of TNF-α (cat. no. JM-02415M2) and IL-6 (cat. no. JM-02446M1) in the supernatant were measured using the appropriate ELISA kits (Jingmei Biotechnology) according to the manufacturer’s instructions.

Cell transfection with Ad-GFP-LC3B adenovirus. To assess autophagy, RAW264.7 cells were infected with Ad-GFP-LC3B adenovirus (cat. no. C3006; Beyotime Institute of Biotechnology). The cells were inoculated into 12-well plates, then the culture medium was changed and an appropriate volume of virus solution was added 24 h later (MOI 20). After culturing at 37°C for 24 h, cells were treated with the corresponding intervention drugs for 24 h. Cell proliferation and the expression of fluorescent protein were observed under a fluorescence microscope.

Western blotting. Following treatment of RAW264.7 cells with the corresponding intervention drugs, the protein expression levels of cleaved-caspase 3 (1:500; cat. no. ab2302; Abcam), pro-caspase 3 (1:1,000; cat. no. ab32150; Abcam), Bax (1:2,000; cat. no. ab182733; Abcam), Bel-2 (1:2,000; cat. no. ab182858; Abcam), LC3-II (1:2,000; cat. no. ab192890; Abcam), beclin-1, p62 (1:10,000; cat. no. ab109012; Abcam), p-mTOR (1:1,000; cat. no. 5536; Cell Signaling Technology, Inc.), mTOR (1:1,000; cat. no. 2972; Cell Signaling Technology, Inc.), p-Akt (1:1,000; cat. no. 13038; Cell Signaling Technology, Inc.) and Akt (1:1,000; cat. no. 9272; Cell Signaling Technology, Inc.) were determined by performing western blotting. Briefly, the cells were seeded into 6-well plates at 40x10^4 cells/well. After 24 h, intervention drugs were added and cells were incubated for another 24 h. Afterwards, total protein was extracted from cells using lysis buffer consisting of RIPA lysate (Beyotime Institute of Biotechnology), 5X protease inhibitor reagent (Roche Diagnostics) and 1% phenylmethylsulfonyl fluoride (Beyotime Institute of Biotechnology) for 30 min on ice. Subsequently, cells were centrifuged (400 x g, 4°C, 20 min) to obtain the supernatant. The protein concentration was measured using a BCA protein quantification kit (Beyotime Institute of Biotechnology) and then the proteins were denatured. Proteins (10 µg) were separated via SDS-PAGE (15% gel for LC3 and 12% gel for other proteins) and transferred onto a PVDF membrane (EMD Millipore). Following blocking with 5% skimmed milk powder at room temperature, the membrane was incubated with the aforementioned primary antibodies for 24 h at 4°C. Subsequently, the membrane was incubated with corresponding secondary antibodies (Goat anti-Rabbit; 1:10,000; cat. no. bs-0295Gs; BIOSS) for 1 h at room temperature. Following washing with TBST (0.1% Tween-20; Beijing Solarbio Science & Technology Co., Ltd.), protein bands were visualized using an ECL developing solution (EMD Millipore) and evaluated using a Bio-Rad gel imaging system (Bio-Rad Laboratories, Inc.). β-actin (1:5,000; cat. no. bs-0061R; BIOSS) was used as the loading control. ImageJ (v1.52; National Institutes of Health) was used for semi-quantification.

Statistical analysis. Experiments were repeated in triplicate. Statistical analyses were performed using SPSS 17.0 (SPSS, Inc.), GraphPad Prism 8.0.2 (GraphPad Software, Inc.) software. Data are presented as the mean ± SD. Comparisons among multiple groups were analyzed using one-way ANOVA followed by Tukey’s post hoc test. P<0.05 was considered to indicate a statistically significant difference.

Results

Treatment with CHCs induces apoptosis and autophagic dysfunction in RAW264.7 macrophages. To investigate whether CHCs could induce macrophage apoptosis, Hoechst 33258 staining assays were conducted to observe the morphological alterations of cell nuclei. Following treatment of RAW264.7 cells with different concentrations of CHCs for 24 h, apoptosis-related morphological changes were markedly increased in the cell nuclei, including chromatin condensation and nuclear fragmentation, compared with the control group (Fig. 1A). Subsequently, western blotting was performed to evaluate the effect of CHCs on the expression of key apoptotic factors. The results revealed that, compared with the untreated control group, the ratios of cleaved-caspase 3/pro-caspase 3 and Bax/Bcl-2 were significantly increased in CHC-treated (800 and 1,200 µg/ml) RAW264.7 macrophages (Fig. 1B).

Furthermore, to determine whether CHCs could induce autophagic dysfunction in RAW264.7 macrophages, their effect on the expression of autophagy-related proteins was evaluated. The expression levels of LC3-II and beclin-1 were significantly downregulated, whereas that of p62 was significantly upregulated in CHC-induced RAW264.7 cells at doses of 800 and 1,200 µg/ml compared with those in the untreated control group (Fig. 1C). The aforementioned findings suggested that CHCs (800 µg/ml) promoted apoptosis and autophagic dysfunction in RAW264.7 macrophages.

Sal B reduces CHC-induced apoptosis and the expression of proinflammatory cytokines in RAW264.7 macrophages. To evaluate the toxicity of Sal B, an MTT viability assay was performed. Following treatment of RAW264.7 macrophages with different concentrations of Sal B for 24 h, the MTT assay results revealed that Sal B (>240 µM) significantly attenuated the viability of macrophages compared with the untreated control group (Fig. 2A). Therefore, the concentrations of 100, 150 and 200 µM Sal B were selected for subsequent experiments. The western blotting results demonstrated that the expression levels of autophagy-related proteins were significantly altered by Sal B treatment at the aforementioned concentrations compared with those in the untreated control group (Fig. 2B).

In addition, the effect of Sal B on CHC-induced apoptosis in RAW264.7 macrophages was evaluated using Hoechst 33258 staining. Treatment of RAW264.7 macrophages with Sal B notably reduced CHC-induced nuclear apoptosis-related morphological changes in cell nuclei (Fig. 3A). Additionally, the western blotting results demonstrated that, compared with the CHC group, the cleaved-caspase 3/pro-caspase 3 and Bax/Bcl-2 ratios were significantly decreased in CHC-induced cells treated with Sal B for 24 h (Fig. 3B).

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To acquire quantification of cell apoptosis, an Annexin V-FITC/PI dual staining assay was performed. As shown in Fig. 3C, the apoptotic rate in the untreated control group was 7.76±0.38%. Compared with the untreated control group, the apoptotic rate in CHC-induced cells was significantly increased (26.42±0.59%). Additionally, treatment of CHC-induced RAW264.7 macrophages with 100, 150 and 200 µM Sal B significantly decreased the apoptotic rate to 23.20±0.71, 18.95±1.13 and 13.33±0.51%, respectively, compared with cells treated CHC alone. Furthermore, the secretion levels of proinflammatory cytokines (TNF-α and IL-6) were significantly reduced in Sal B-treated, CHC-induced RAW264.7 macrophages compared with those in the CHC + LPS group (Fig. 3D). These results indicated that Sal B attenuated CHC-induced RAW264.7 macrophage apoptosis and the production of proinflammatory cytokines.

Sal B improves CHC-induced autophagic dysfunction in RAW264.7 macrophages. To assess whether Sal B could ameliorate CHC-induced autophagy dysfunction in RAW264.7 macrophages, western blotting was performed to detect the expression levels of apoptosis- and (C) autophagy-related proteins in CHC-induced RAW264.7 macrophages (n=3).

To acquire quantification of cell apoptosis, an Annexin V-FITC/PI dual staining assay was performed. As shown in Fig. 3C, the apoptotic rate in the untreated control group was 7.76±0.38%. Compared with the untreated control group, the apoptotic rate in CHC-induced cells was significantly increased (26.42±0.59%). Additionally, treatment of CHC-induced RAW264.7 macrophages with 100, 150 and 200 µM Sal B significantly decreased the apoptotic rate to 23.20±0.71, 18.95±1.13 and 13.33±0.51%, respectively, compared with cells treated CHC alone. Furthermore, the secretion levels of proinflammatory cytokines (TNF-α and IL-6) were significantly reduced in Sal B-treated, CHC-induced RAW264.7 macrophages compared with those in the CHC + LPS group (Fig. 3D). These results indicated that Sal B attenuated CHC-induced RAW264.7 macrophage apoptosis and the production of proinflammatory cytokines.

Sal B improves CHC-induced autophagic dysfunction in RAW264.7 macrophages. To assess whether Sal B could ameliorate CHC-induced autophagy dysfunction in RAW264.7 macrophages, western blotting was performed to detect the expression levels of autophagy-related proteins. Compared with the CHC group, the results demonstrated that the expression levels of LC3-II and beclin-1 were significantly increased, whereas those of p62 were significantly decreased in RAW264.7 macrophages co-treated with Sal B and CHCs for 24 h (Fig. 4A).

In addition, to further verify the Sal B-mediated autophagic flux, RAW264.7 macrophages were infected with Ad-GFP-LC3B for 24 h. Following cell treatment with BafA1, an autophagosome-lysosome degradation inhibitor, the specific green fluorescent point distribution of LC3B was observed under a microscope. As shown in Fig. 4B, a
markedly enhanced accumulation of green fluorescent puncta was observed in CHC-induced and Sal B-treated RAW264.7 macrophages treated with 20 nM BafA1 for 4 h compared with the CHC + Sal B group. These findings suggested that Sal B improved CHC-induced autophagic dysfunction in RAW264.7 macrophages.

Sal B attenuates CHC-induced RAW264.7 macrophage apoptosis partially via improving autophagic dysfunction. The inhibitor 3-MA is widely used to selectively inhibit the activity of type III PI3K, thereby blocking the formation of autophagosomes (16). To further explore the association between Sal B-mediated decreases in apoptosis and improvements in autophagy, RAW264.7 macrophages were co-treated with Sal B and 3-MA for 24 h. Subsequently, western blotting was performed to detect alterations in the expression levels of autophagy- and apoptosis-related proteins. As shown in Fig. 5A, compared with the Sal B + CHC group, LC3-II and beclin-1 expression levels were significantly downregulated, but p62 was significantly upregulated in CHC-induced RAW264.7 macrophages co-treated with Sal B and 3-MA.

More importantly, 3-MA-mediated autophagy inhibition significantly increased the cleaved caspase 3/pro-caspase 3 and Bax/Bcl-2 ratios in Sal B-treated CHC-induced RAW264.7 macrophages (Fig. 5B). Consistent with the western blotting results, the Annexin V-FITC/PI dual staining assay results revealed that the apoptotic rate of RAW264.7 macrophages treated with Sal B and 3-MA was also significantly enhanced compared with that in the Sal B + CHC group (Fig. 5C). Overall, the aforementioned results indicated that Sal B attenuated CHC-induced apoptosis in RAW264.7 macrophages partially via improving autophagic dysfunction.

Sal B improves CHC-induced autophagic dysfunction in RAW264.7 macrophages partially via inhibiting the Akt/mTOR signaling pathway. The Akt/mTOR signaling pathway serves an important role in the regulation of autophagy (17). Therefore, the present study aimed to investigate whether the Akt/mTOR signaling pathway was involved in Sal B-induced autophagy. The western blotting results demonstrated that, compared with the untreated control group, the p-mTOR/mTOR and p-Akt/Akt ratios were significantly elevated in the CHC group (Fig. 6A). However, the p-mTOR/mTOR and p-Akt/Akt ratios were significantly decreased in the Sal B groups compared with the CHC group. These findings revealed that Sal B exerted an effective inhibitory effect on the Akt/mTOR signaling pathway, thus improving CHC-induced autophagy dysfunction in RAW264.7 macrophages.

Subsequently, RAW264.7 macrophages were co-treated with Sal B and insulin, an activator of the PI3K/Akt/mTOR
Figure 3. Sal B attenuates CHC-induced apoptosis and the secretion of proinflammatory cytokines in RAW264.7 macrophages. The following groups are presented: CONT, untreated control; CHC, 800 µg/ml CHC; CHC + Sal B 100, 800 µg/ml CHC + 100 µM Sal B; CHC + Sal B 150, 800 µg/ml CHC + 150 µM Sal B; and CHC + Sal B 200, 800 µg/ml CHC + 200 µM Sal B. (A) Morphological features of CHC-induced and Sal B-treated RAW264.7 cells were evaluated by performing Hoechst 33258 staining. White arrows indicate apoptotic cells (scale bar, 100 µm). (B) Western blotting was performed to determine the expression levels of apoptosis-related proteins in CHC-induced RAW264.7 macrophages following treatment with Sal B (n=3). *P<0.05 vs. untreated control group; #P<0.05 vs. CHC group. (C) Annexin V-FITC/PI double staining was performed to assess the effect of Sal B on the apoptotic rate of CHC-induced RAW264.7 macrophages (n=3). *P<0.05 vs. untreated control; #P<0.05 vs. CHC. Sal B, salvianolic acid B; CHC, cholesterol crystal.

signaling pathway, for 24 h. As shown in Fig. 6B, compared with the Sal B + CHC group, the p-mTOR/mTOR and p-Akt/Akt ratios were significantly increased in Sal B + insulin-treated RAW264.7 macrophages, whereas the expression levels of LC3-II and beclin-1 were significantly downregulated. In conclusion, these findings revealed that insulin partially restored Sal B-induced autophagy via activating the Akt/mTOR signaling pathway, which suggested that Sal B partially inhibited the Akt/mTOR signaling pathway to improve CHC-induced autophagy dysfunction in RAW264.7 macrophages.

Discussion

The role of apoptosis in the pathological process of AS has received increasing attention. Several factors further promote macrophage inflammatory responses and apoptosis, secondary inflammation and necrosis of other cells at the site
of the plaque, thereby leading to increased plaque instability, and eventually thrombosis (18-20). Therefore, reducing apoptosis and maintaining normal endocytic capacity in macrophages could be considered as a potential therapeutic approach for advanced AS. CHCs are commonly found in the necrotic core and atherosclerotic subendothelium (21). Emerging evidence has suggested that the massive accumulation of CHCs at the lesions induces inflammatory responses and increases plaque instability (22-25). Sergin et al (26) demonstrated that CHCs could increase the expression of p62 and ubiquitinated proteins at the site of the lesion. However, whether CHCs can promote macrophage apoptosis in atherosclerotic lesions and their specific underlying mechanism remains to be elucidated. The present study revealed that CHCs induced macrophage apoptosis and autophagy dysfunction.

As a highly conserved and widespread physiological and defense mechanism in organisms (27), the effects of autophagy on the development of several diseases have been the focus of research. Liao et al (8) suggested that macrophage-mediated autophagy displayed an inhibitory effect on the progression of AS. In addition, the role of traditional Chinese medicine in provoking and suppressing autophagy for treating different diseases has also received increasing attention. In traditional Chinese medicine, Salvia miltiorrhiza has been widely used in the treatment of cardiovascular diseases by relieving pain and promoting blood circulation (28).

Sal B, one of the primary water-soluble components in the Salvia miltiorrhiza extracts, also exerts protective effects against various cardiovascular diseases (29-31). However, no study has yet investigated the possible benefits of Sal B in CHC-induced apoptosis. The present study showed that Sal B offered an important protective effect to alleviate macrophage autophagy dysfunction and apoptosis in advanced AS. The protective effects of Sal B were determined in the current study via the Hoechst 33258 staining assay, as demonstrated by the reduction in the apoptosis of CHC-insulted macrophages and the decreased Annexin V-FITC/PI staining. The western blotting results indicated that Sal B significantly increased autophagy in CHC-induced macrophages, which was consistent with the results obtained by Jing et al (32). Surprisingly, the ELISA results also demonstrated that Sal B significantly attenuated the release of proinflammatory cytokines (TNF-α and IL-6) by CHC-induced macrophages, and we hypothesized that the aforementioned mechanism was closely associated with Sal B-mediated improvements in macrophage autophagy. However, the specific mechanism requires further investigation.

It has been suggested that autophagy is an upstream response to apoptosis, indicating that apoptosis is mediated by the activation of autophagy (33-35). However, it has also been suggested that enhancing autophagy may attenuate apoptosis (36). For example, Wang et al (37) demonstrated that apigenin could simultaneously induce autophagy and...
apoptosis in macrophages, whereas suppressing autophagy could increase macrophage apoptosis, thus negatively regulating AS. The role of autophagy in apoptosis remains controversial. The present results indicated that Sal B significantly decreased macrophage apoptosis by improving autophagy. Macrophages were co-treated with Sal B and the autophagy inhibitor, 3-MA. The western blotting results suggested that 3-MA successfully attenuated the effect of Sal B on autophagy, partially compensating for the protective effect of Sal B on apoptosis. This finding indicated that the role of Sal B in reducing the apoptotic rate of RAW264.7 macrophages was partially due to its effect on relieving CHC-induced autophagy dysfunction.

Autophagy is regulated by a variety of signal transduction pathways. The adenylylate-activated protein kinase/mTOR and PI3K/Akt/mTOR signaling pathways are two common cellular regulatory pathways of autophagy (38). The Akt/mTOR signal transduction pathway is closely associated with the occurrence of autophagy and regulation of apoptosis, and is considered as the only inhibitory signal transduction pathway regulating autophagy (39). Zhai et al (40) demonstrated that specifically inhibiting the Akt/mTOR signaling pathway notably
enhanced macrophage autophagy, eventually attenuating the infiltration of macrophages and enhancing the stability of the atherosclerotic plaque. To investigate the molecular mechanism underlying the effect of Sal B on improving autophagy function, macrophages were treated with Sal B and insulin, an agonist of the Akt/mTOR signaling pathway. Insulin inhibited Sal B-mediated inactivation of Akt/mTOR signaling in CHC-induced macrophages, which partially restored Sal B-induced autophagy. The results of the present study suggested that CHCs induced autophagy dysfunction in macrophages via activating the Akt/mTOR signaling pathway, whereas Sal B partially improved autophagy dysfunction in RAW264.7 macrophages via blocking the Akt/mTOR signaling pathway.

In summary, the results of the present study revealed that Sal B partially improved RAW264.7 macrophage autophagy dysfunction via inhibiting the Akt/mTOR signaling pathway to attenuate macrophage apoptosis. In addition, treatment of macrophages with Sal B inhibited CHC-induced release of proinflammatory cytokines (TNF-α and IL-6). Overall, the present study may provide novel insight into the molecular mechanism underlying the beneficial effects of Sal B on AS.

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Availability of data and materials
The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.
Authors' contributions

MS, SW and YH contributed to designing the study. MS, WY and ZY performed the experiments. YY analyzed the data. MS, YY and YH drafted the manuscript. All authors read and approved the final manuscript. MS and SW confirm the authenticity of all the raw data.

Ethics approval and consent to participate

Not applicable.

Patient consent for participation

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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