ASMq protects against early burn wound progression in rats by alleviating oxidative stress and secondary mitochondria-associated apoptosis via the Erk/p90RSK/Bad pathway

HANLEI ZHOU1*, QUAN FANG2*, NAN LI2, MEIRONG YU3, HONGMING CHEN4 and SONGXUE GUO2

Departments of 1Vascular Surgery and 2Plastic Surgery, The Second Affiliated Hospital Zhejiang University School of Medicine; 3Clinical Research Center, The Second Affiliated Hospital Zhejiang University School of Medicine; 4Department of Plastic Surgery, International Medical Center of The Second Affiliated Hospital Zhejiang University School of Medicine, Hangzhou, Zhejiang 310009, P.R. China

Received August 7, 2020; Accepted February 23, 2021

DOI: 10.3892/mmr.2021.12029

Abstract. Burn wounds present an evolutionary progression, in which the initial wound tissue deepens and expands following thermal injury. Progressive tissue damage in the zone of stasis may worsen burn injury, which is associated with oxidative stress and secondary apoptosis, and worsen the prognosis of patients with burn wounds. The mitochondrial apoptotic pathway is involved in receiving oxidative signals and regulating tissue apoptosis. Previously, Abnormal Savda Munziq (ASMq), a natural compound of traditional Uyghur Medicine, which includes ten types of herb, has been reported to exhibit a number of effects, including anti-inflammatory, antioxidative and anti-apoptotic activities. The present study demonstrated that ASMq protected against early burn wound progression following thermal injury in rats; this effect may be mediated by its ability to attenuate oxidative stress-induced mitochondria-associated apoptosis. The present study may provide a novel therapeutic method to prevent early burn wound progression following burn injury.

Introduction

Burn wound progression, which was first introduced by DM Jackson in 1953, refers to the deterioration of tissue damage in the zone of stasis, especially during the first few days following thermal injury (1). According to Jackson's model of thermal wound injury, the burn wound is divided into three concentric zones: A central zone of coagulation, an intermediate zone of stasis and an outer zone of hyperemia. The central zone is damaged by thermal injury directly and suffers irreversible coagulation necrosis with thrombosed vessels; the zone of hyperemia is characterized by spontaneous recovery, while the zone of stasis is considered salvageable via optimal intervention (2). The recovery of the zone of stasis is vital in preventing irreversible damage and determining the final severity of the wound. Burn wound progression contributes to the development of wound contraction, hypertrophic scarring, wound infection and sepsis. Therefore, appropriate and timely interventions to prevent burn wound progression are of clinical significance for treatment of the wound and prognosis of patients with burns (3), as well as for the early prevention of hypertrophic scar formation due to burn injury (4).

Multiple mechanisms are involved in burn wound progression, including oxygen free radical-induced damage, hypoperfusion, micro-thrombosis and vasoconstriction, which can lead to early inflammatory cascades and cell death (apoptosis or necrosis) (3,5,6). Mitochondria are important organelles in eukaryotic cells that are susceptible to oxidative stress by regulating energy metabolism and apoptosis (7). Our previous studies demonstrated the effectiveness of protective strategies against oxidative damage and tissue apoptosis in preventing burn wound conversion (8,9).

Abnormal Savda Munziq (ASMq), a traditional herbal formula in Uyghur medicine, primarily consists of ten herbs, including Lavandula angustifolia, Foeniculum vulgare, Anchusa italica, Euphorbia humifusa, Melissa officinalis, Adiantum capillus-veneris, Glycyrrhiza uralensis, Cordia dichotoma, Ziziphus jujuba and Alhagi pseudoalhagi (10). Previous reports demonstrated that ASMq exerts antioxidative, anti-inflammatory and anti-apoptotic properties to inhibit the formation of hypertrophic scars (4) and treat diabetes, cardiovascular disease, pulmonary disorders and various types of cancer, such as cervical cancer and hepatic carcinoma (10-17); to the best of our knowledge, however, few studies have investigated its potential effects on burn injury. In consideration of the critical role of reactive oxygen species (ROS)-induced...
oxidative stress and subsequent mitochondria-associated apoptosis in the pathophysiological development of early burn wounds, it was hypothesized that ASMq may prevent early burn wound progression in rats by suppressing oxidative stress and apoptosis. The present study aimed to evaluate the possible mechanisms of action and regulation.

The present study aimed to demonstrate that ASMq may be an effective drug to ameliorate early burn wound progression, thus avoiding or alleviating adverse outcomes associated with burn wounds.

Materials and methods

Experimental animals. Adult male Sprague-Dawley (SD) rats weighing 200-220 g were used according to an experimental protocol approved by The Second Affiliated Hospital, Zhejiang University, School of Medicine, Institutional Animal Care and Use Committee (approval no. 2018-095) in accordance with the National Institutes of Health Guidelines for the Care and Use of Laboratory Animals (18). The animals were housed in a 12-h light/dark cycle in a filtered-air unit with consistent temperature (20-26˚C) and humidity (50-65%) and ad libitum access to food and water. All animals were purchased from Shanghai SLAC Laboratory Animal Co., Ltd.

‘Comb’ burn model establishment, animal grouping and drug administration. To create a row of four bands of full-thickness thermal burn injuries (20x10 mm) with three interspaces of uninjured skin (20x5 mm), a customized rectangular brass comb was boiled in 100°C water for 5 min, and then applied to the shaved dorsum skin of rats for 10 sec following anesthesia with intraperitoneal injection of 1% pentobarbital sodium at a dose of 30 mg/kg, as previously described (19-22). The breathing and heart rates of experimental rats were monitored during anesthesia. The rats received immediate postoperative analgesia with buprenorphine (0.03 mg/kg, subcutaneous), followed by continuous analgesia with buprenorphine (0.03 mg/kg, subcutaneous) every 12 h. The interstices of undamaged skin represented the zone of stasis. Animals in the sham group were treated with exposure to 25˚C water and food intake ad libitum before the operation. All animals were sacrificed at 24 h by overdose of sodium pentobarbital (200 mg/kg, intraperitoneal), according to a previous study (25).

Tissue collection and preparation. Skin tissue specimens from wounds were harvested 48 h after operation, according to our previous study (8,9). One band of the interspace skin with 2 mm burned tissue on each side (10x9 mm) was collected from each burn model and stored in 10% formalin at 4°C for subsequent immunohistochemistry analysis, while the remaining two bands of interspace skin (without burn tissue; 10x5 mm) from each burned rat were stored at -80°C for molecular biological assessments. A total of three corresponding areas of unburnt skin from each rat of the sham group were used as controls.

Histological assessment. Rat skin tissue was prepared by fixation in 4% formaldehyde solution for 12 h at 4°C. The tissue samples were then dehydrated in a graded series of ethanol solutions, embedded in paraffin and cut into 3-μm sections. The hematoxylin and eosin (H&E) staining was performed according to routine procedure. Histological changes were evaluated under a light microscope (DM2500; Leica Microsystems GmbH). A total of five fields (magnification, x200) were selected randomly for observation from each slice.

Oxidative stress evaluation. The frozen tissue samples from rats were crushed, ground and homogenized for 8 min using a tissue homogenizer with homogenization buffer (4˚C), which included 1 mmol/l phenylmethylsulfonyl fluoride, 1 mg/ml pepstatin A, 1 mg/ml aprotinin and 1 mg/ml leupeptin in phosphate-buffered saline (PBS; pH, 7.2) containing 0.5% sodium deoxycholate and 1% Triton X-100, at a ratio of 100 mg tissue/ml. Following incubation at 4°C for 1 h, the final homogenate was centrifuged at 10,000 x g for 15 min at 4°C and the supernatants were used for detection kit assays. The total protein concentration was determined using a BCA assay kit (Nanjing KeyGen Biotech Co., Ltd.).

In order to evaluate lipid peroxidation of skin tissue, malondialdehyde (MDA) levels were measured. Skin tissue homogenates from the burn wounds were incubated with a thiobarbituric acid reactive species assay kit reagent (cat. no. KGT003-1; Nanjing KeyGen Biotech Co., Ltd.) to determine MDA levels, which were expressed in nmol/mg protein. Levels of tissue superoxide dismutase (SOD) and glutathione peroxidase (GPx) were evaluated to determine the oxidative stress status in the skin tissues of burn wounds and were detected using commercial assay kits from Nanjing KeyGen Biotech Co., Ltd. (cat. nos. KGT00150 and KGT-014, respectively) according to the manufacturer's protocols. The results
were expressed in U/mg for SOD and nmol/min/mg for GPx. Absorbance values were measured using a microplate reader (MDA, 532 nm; SOD, 550 nm; GPx, 412 nm) (Model 680 Microplate Reader; Bio-Rad Laboratories, Inc.).

**Immunofluorescence staining.** In order to determine the location and quantity of the specific antigens in the interspace skin following burn injury, the prepared slices were washed in PBS for 10 min, followed by boiling at 95˚C in 0.01 mmol citrate buffer (pH, 6.0) for 10 min for antigen retrieval. Then, the slices were incubated with hydrogen peroxide for 10 min at room temperature, and 5% bovine serum albumin ( Gibco; Thermo Fischer Scientific, Inc.) was applied later as the blocking solution for 20 min at room temperature. Without being washed, the sections were incubated with anti-phosphorylated (p)-Erk (1:200; cat. no. 4370; Cell Signaling Technology, Inc.) or anti-p-Bad (1:200; cat. no. sc-12969-R; Santa Cruz Biotechnology, Inc.) antibodies overnight at 4˚C. After being rinsed with PBS, the sections were incubated with FITC-(1:50; cat. no. BA1105; Wuhan Boster Biological Technology, Ltd.) or Cy3-(1:50; cat. no. BA1032; Wuhan Boster Biological Technology, Ltd.) labeled goat anti-rabbit secondary antibodies for 2 h at 37˚C in the dark. The sections were rinsed and stained with DAPI (100 ng/ml; Wuhan Boster Biological Technology, Ltd.) for 8 min at room temperature, and then mounted with VECTASHIELD® mounting medium (Wuhan Boster Biological Technology, Ltd.). All slices were observed and photographed under a fluorescence microscope (magnification, x200) (DM5500B; Leica Microsystems GmbH).

Terminal deoxynucleotidyl-transferase-mediated dUTP nick end labeling (TUNEL) staining. TUNEL staining was performed using a commercial cell death detection kit (Roche Diagnostics) according to the manufacturer’s protocols. Briefly, after deparaffinization and rehydration, the slices were pretreated by proteinase K (PCR grade; Roche Diagnostics) for 15 min at room temperature. Then, the permeabilization was conducted with fresh prepared 0.1% Triton-X-100 and 0.1% sodium citrate for 8 min at room temperature. Then, incubation with trypsin (0.01 N HCL) was applied for 30 min at 37˚C. Subsequently, the slices were transferred into a container with 0.1 M citrate buffer (pH, 6) and received microwave irradiation (350 W) for 5 min. Rinsed slices were treated with TUNEL mixture for 60 min at 37˚C in a humidified atmosphere in the dark. Finally, the signal conversion was handled with the help of converter-POD and DAB substrate. The stained slices were observed by a light microscope (magnification, x400; DM2500; Leica Microsystems GmbH) and images were captured. The index of apoptosis was calculated as the percentage of apoptotic cells among all cells counted. Cells with dark brown-stained nuclei were counted by two independent investigators who were blinded to the group assignments. At least three visual fields per slide and five slides per group were evaluated by the investigators.

**Western blot analysis.** Frozen skin samples were harvested from rats for western blotting. Briefly, samples were cut into pieces (~1x1 mm) and lysed with RIPA lysis buffer (cat. no. AR0105; Wuhan Boster Biological Technology, Ltd.) for 1 h on ice, then the lysates were centrifuged at 14,000 x g for 10 min at 4˚C. The protein samples (50 µg) were mixed with loading buffer and subjected to SDS-PAGE on 6-12% gels, and subsequently transferred onto nitrocellulose membranes by electrophoresis, while aliquots of samples were used to determine the protein concentration of each sample using a bicinchoninic acid kit (cat. no. KGPBCA; Nanjing KeyGen Biotech Co., Ltd.). Subsequently, membranes were incubated in blocking buffer (5% skimmed milk) for 2 h at room temperature, and incubated overnight at 4˚C with the following primary antibodies: Anti-XO (1:5,000; cat. no. ab109235; Abcam), anti-Nox4 (1:2,000; cat. no. ab109225; Abcam), anti-Bad (1:1,000; cat. no. sc-8044; Santa Cruz Biotechnology, Inc.), anti-p-Bad (1:1,000; cat. no. sc-12969-R; Santa Cruz Biotechnology, Inc.), anti-cytochrome c (Cyto C; 1:1,000; cat. no. sc-7159; Santa Cruz Biotechnology, Inc.), anti-cleaved caspase (CC):3 (1:1,000; cat. no. 9664; Cell Signaling Technology, Inc.), anti-CC9 (1:1,000; cat. no. 9509; Cell Signaling Technology, Inc.), anti-Erk1/2 (1:1,000; cat. no. 9102; Cell Signaling Technology, Inc.), anti-p-Erk1/2 (1:1,000; cat. no. 9101; Cell Signaling Technology, Inc.) and anti-p90 ribosomal S6 kinase (p90RSK; 1:1,000; cat. no. ab32114; Abcam). β-actin (1:2,000; cat. no. SC-47778; Santa Cruz Biotechnology, Inc.) was used as a control. Following incubation with an HRP-conjugated secondary antibody (1:5,000; cat. no. 31160; Pierce; Thermo Fisher Scientific, Inc.) and anti-phosphorylated (p)‑Erk (1:1,000; cat. no. 9101; Cell Signaling Technology, Inc.) and anti-p90 ribosomal S6 kinase (p90RSK; 1:1,000; cat. no. sc-7159; Santa Cruz Biotechnology, Inc.) antibodies overnight at 4˚C (Model 680 Microplate Reader; Bio-Rad Laboratories, Inc.).

**Statistical analysis.** GraphPad Prism version 7 software (GraphPad Software, Inc.) was used for statistical analysis. Data are presented as the mean ± SD. Multiple comparisons among the sham and burn groups were analyzed with one-way ANOVA followed by Bonferroni’s post hoc test. P<0.05 was considered to indicate a statistically significant difference.

**Results**

**ASMq alleviates tissue damage of skin in the zone of stasis of rat burn wounds.** H&E staining demonstrated thinning of the epidermis layer, swelling of the dermis layer, alteration of collagen structure, infiltration of inflammatory cells and tissue destruction in the interspaces between the two wounds following thermal injury. ASMq alleviated these changes in a dose-dependent manner (Fig. 1).

**Effect of ASMq on early elevated oxidative stress in the zone of stasis of rat burn wounds.** MDA, an indicator of lipid peroxidation, indicates the severity of oxidative damage (26). There was no significant difference in the levels of MDA between the sham + high dosage ASMq (1,600 mg/kg) and sham groups. Although low dosage ASMq (400 mg/kg) did not significantly alleviate the increase of MDA levels, high and medium dosage ASMq significantly decreased the levels of MDA in a dose-dependent manner (P<0.01; Fig. 2A).

Low dosage ASMq (400 mg/kg) did not influence the activity levels of endogenous antioxidant enzymes (GPx and SOD) in...
ZHOU et al: ASMq PREVENTS BURN-WOUND PROGRESSION BY ALLEVIATING OXIDATIVE STRESS AND APOPTOSIS

Figure 1. Histological assessment of burn skin samples in the early stages following severe burn and ASMq application. Representative hematoxylin and eosin staining images show the typical histological changes in the different groups (n=8/group). The progression boundaries are indicated by dotted lines, the stasis zones are marked with # and the initial burn regions are marked with *, presenting notable hyalinization. Scale bar, 200 µm. ASMq, Abnormal Savda Munziq.

Figure 2. Assessment of oxidative stress within the zone of stasis and the effects of ASMq. (A-C) ASMq downregulates MDA levels and upregulates the activity of endogenous antioxidant enzymes (GPx, SOD) in wounds and (D) downregulates the expression of XO and Nox4 in a dose-dependent manner following burn injury. Data are presented as the mean ± SD (n=8/group). **P<0.01 vs. sham; #P<0.05, ##P<0.01. ASMq, Abnormal Savda Munziq; XO, xanthine oxidase; Nox, NADPH oxidase; ns, not significant; MDA, malondialdehyde; SOD, superoxide dismutase; GPx, glutathione peroxidase.

the healthy skin of rats. High dosage AMSq (1,600 mg/kg) upregulated the activity of endogenous antioxidant enzymes (GPx and SOD) in the zone of stasis of rat burn wounds, while no significant differences were detected in medium and lower dose groups (Fig. 2B and C).

XO, which is associated with the generation of ROS (27), was semi-quantified by western blotting (Fig. 2D). The results demonstrated that XO expression levels in the zone of stasis of rat burn wounds were significantly downregulated by ASMq. There was no significant difference in the expression levels
of XO between medium and low dosage or medium and high dosage ASMq (P>0.05). XO levels in the high dosage ASMq group were lower than those in the low dosage ASMq group (P<0.05). The effect of decreasing XO levels induced by burn injury was more significant in the high dosage ASMq group (P<0.01). Nox4 is an important member of the Nox family that can be used to indicate mitochondrial status (28). The effect of ASMq on Nox4 expression was similar to that of XO: Specifically, both high and medium dosage ASMq significantly decreased expression levels of Nox4 in the zone of stasis (Fig. 2D).

ASMq dose-dependently decreases cell apoptosis in burn wound tissue of rats. TUNEL staining demonstrated that the number of apoptotic cells with brown-stained nuclei in burn wounds increased. According to the random multi-field cell count, high dosage ASMq significantly decreased the number of apoptotic cells in the zone of stasis of skin tissue compared with the number observed in the burn group, while statistically insignificant differences were observed in the medium and low dosage groups (Fig. 3A).

Western blot analysis was performed to determine the protein expression levels of CC3 and CC9, which are key signaling molecules in the execution phase of mitochondria-associated cell apoptosis (29). Burn injury significantly increased CC3 and CC9 expression levels. Increasing doses of ASMq resulted in decreased CC3 expression levels, with a peak observed in the group treated with a dose of 1,600 mg/kg. The expression levels of CC9 were significantly decreased by high and medium dosage ASMq (Fig. 3B).

Pretreatment with PD98059 partially reversed the mitigative effect of high dosage ASMq on cell apoptosis and expression levels of CC3 and CC9 in the zone of stasis of skin tissue following burn injury. PD98059 reversed the decrease in pro-apoptotic protein (Cyto C) in the burn wound induced by high dosage ASMq (Fig. 3A). These results suggested a potential involvement of Erk signaling in regulating tissue apoptosis in the zone of stasis.

ASMq dose-dependently increases the ratio of p-Bad/Bad and release of mitochondrial Cyto C. The aforementioned results indicated that the anti-apoptotic effect of ASMq may be associated with the regulation of mitochondria-associated apoptosis pathways involved in Bad. Western blotting results showed that ASMq altered the Bad/Cyto C signaling molecules cascade. The ratio of p-Bad to Bad increased significantly in skin tissue following burn injury. All three dosages of ASMq increased the ratio of p-Bad/Bad in a dose-dependent manner. Secondly, a significant increase in expression levels of Cyto C was observed following burn injury, which was alleviated by medium and high dosage ASMq in a dose-dependent manner. PD98059 antagonized the effect of high-dosage ASMq (Fig. 4).
Erk/p90 ribosomal S6 kinase (p90RSK) pathway acts upstream in the regulation of ASMq on mitochondria-associated apoptosis in the zone of stasis. As indicated by immunofluorescence staining, the distribution of cells positive for p-Erk was demonstrated by an elevation in green-stained cells in the interspace skin following burn injury (Fig. 5). All three dosages of ASMq notably enhanced the phosphorylation of Erk, and the highest dose (1,600 mg/kg) exerted the strongest effect (Fig. 5).

The results of western blotting were consistent with immunofluorescence staining: ASMq dose-dependently increased protein expression levels of p-Erk and p90RSK. PD98059 reversed the elevated expression of p-Erk and p90RSK induced by ASMq (Fig. 6).

Discussion

Uygur medicine, one of the most important branches of Chinese traditional medicine, provides ways to manage emergency situations and disease (16,30-32). In the systemic theory of Uygur medicine, the liver produces bodily fluids with different effects that can be divided into four types: Sapra, kan hiliti, phlegm and sawda (15,16). There is a complementary balance between these fluids, which serves a role in the maintenance of normal physiology and the occurrence and development of disease. When the internal environment changes, these four body fluids become AS, which can be harmful to various organs (10). With advancements of science and technology, research into Uygur traditional medicine has become more in-depth, with significant progress in drug extraction and component analysis (33-35). ASMq, a natural compound of traditional Uygur Medicine, has been reported to exhibit anti-inflammatory, antioxidant and anti-apoptotic effects, and decreases oxidative damage by free radicals (10-17). The production of ASMq is subject to a patent application (no. ZL02130082.8) in China (10).

Deep burn injury tends to induce the formation of scars, especially hypertrophic scars, which are related to damage of the dermis (3,6). Early intervention before progressive changes of initial deep burn wounds can prevent further deepening and extension, decrease damage of the dermis and subcutaneous tissue and prevent the generation of secondary hypertrophic scars (36). Based on previously reported anti-inflammatory, anti-apoptotic and antioxidant effects of ASMq in the treatment of various diseases (4,15,23,37) and the underlying mechanism of early burn wound progression, it was hypothesized that ASMq may have a potential therapeutic effect on the progressive deepening of early burn wounds. Oxygen free radicals are considered to be important mediators of tissue damage progression following thermal injury (9,38). In addition to thermal injury, XO and Nox participate in the regulation of oxygen free radicals (9). ROS induce cell death in the zone of stasis via lipid peroxidation and protein degradation (8). Moreover, ROS have been reported to disrupt the mitochondrial outer membrane, leading to the release of Cyto C and the activation of apoptosis (39,40). The current focus of research is the administration of antioxidants to alleviate oxidative stress of burn wounds, and thus prevent the progress of burn wound progression.
of wound deterioration (34,41). Guo et al (9) demonstrated that hydrogen-rich saline, a specific free radical scavenger, decreases lipid peroxidation and enhances the activity of endogenous antioxidant enzymes, thus alleviating early wound progression following deep burn injury in vivo. Previous reports demonstrated that ASMq attenuates mitochondrial damage caused by free radicals (12,13). Certain ingredients (such as tannins, vitamins and flavones) in ASMq prevent free radical chain reaction, scavenge free radicals and decrease the production of -OH (11). Ischemia/reperfusion injury elevates levels of oxidative stress in tissue and induces cell damage, such as lipid peroxidation (37,42). Abudunaibi et al (13) reported that ASMq effectively ameliorates lipid peroxidation and oxidative stress in rats with myocardial ischemia/reperfusion injury. ASMq also shows significant dose-dependent antioxidative effects (37). In the present study, it was observed that both medium and high dosage ASMq attenuated lipid peroxidation of skin tissue in the zone of stasis, and the high dosage group showed more significant effects. Although low and medium dosage ASMq showed limited effects on increasing burn-induced decreases in endogenous antioxidative enzyme activity, high dosage ASMq significantly enhanced the activity of endogenous antioxidant enzymes (GPx and SOD) in the zone of stasis of rat burn wounds. Additionally, by investigating potential in vivo effects of ASMq on the generation pathway of oxidative free radicals XO and Nox, it was found that ASMq had a notable effect on decreasing oxidative free radicals by dose-dependently down-regulating the expression levels of XO and Nox.

Apoptosis serves a key role in early burn wound progression secondary to burn-induced oxidative stress (43). Before irreversible cell inactivation occurs, appropriate drug intervention can alleviate cell apoptosis in the zone of stasis (9,41). Studies have attempted to ameliorate the progressive deepening of burn wounds by inhibiting apoptosis (6,41). Our previous study (9) illustrated that hydrogen-rich saline exerted a potent effect on ameliorating apoptosis in the zone of stasis and prevented wound deepening and extension. In the present study, TUNEL staining was performed to investigate the distribution of apoptotic cells in the zone of stasis directly and evaluate the effect of ASMq on the number of apoptotic cells in the burn wound. The results indicated that ASMq dose-dependently decreased cell apoptosis in the burn wound tissue of rats. The caspase family are important signaling molecules of the cell apoptotic pathway (29). Immunofluorescence staining demonstrated that medium and high dosage ASMq decreased the distribution of CC3 significantly. In agreement with the aforementioned results, western blotting showed that ASMq dose-dependently reversed the increased expression of CC3 and CC9 in burn wounds. These results supported the protective effect of ASMq on tissue apoptosis in the zone of stasis. In addition, the release of Cyto C was also detected in the burn wound. As a proapoptotic protein, Cyto C is released by mitochondria under cellular stress and combines with apoptotic peptidase activating factor 1 and caspase-9 to form an activated complex, which cleaves caspase-3 into CC3 (an activated state) and induces apoptosis (29). Here, ASMq exhibited the ability to lower cytosolic levels of Cyto C at all dosages.

As a member of the proapoptotic BH3-only subfamily, Bad is involved in the induction of apoptotic cascades (44,45). Dephosphorylated Bad binds to anti-apoptotic Bcl-2 and Bcl-xL proteins, then inactivates these proteins and induces pro-apoptotic Bax protein translocation from the cytosol to mitochondrion. The release of cytochrome C from the mitochondrion can activate the caspase cascade, leading to apoptosis. Here, ASMq exhibited a significant effect on decreasing the expression of Bad in the zone of stasis. Apoptosis is a key mechanism in the progression of burn wounds, and inhibiting apoptosis can effectively alleviate burn wound progression. Therefore, ASMq may be a promising drug for the treatment of burn wounds.
the mitochondria, which results in the release of Cyto C from mitochondria into the cytoplasm, and subsequently activates the caspase cascade (42,46). p-Bad can inhibit the combination of Bad with anti-apoptotic Bcl-2 and Bcl-xL proteins at the mitochondrial outer membrane, thereby blocking the Bax-induced Cyto C/caspase-mediated apoptotic pathway (46,47). In the current study, ASMq increased the ratio of p-Bad/Bad, which indicated that Bad dissociated from Bcl-2 and Bcl-xL via phosphorylation of the Serine-136 residue of Bad (48,49), thus downregulating the expression levels of Cyto C. In all, the mitochondria-associated apoptotic pathway seemed to be activated in the process of burn wound conversion. MAPK, a type of serine/threonine protein kinase in eukaryotes, serves a vital role in transducing extracellular stimuli into intracellular signals, which cause various biochemical reactions, such as proliferation, differentiation and apoptosis in cells (45,50). Erk is an important member of the MAPK signaling pathway (45). p90RSK, a downstream target of MAPK that is phosphorylated and activated by p-Erk1/2, has been reported to be involved in the phosphorylation of Bad and subsequent inactivation of the pro-apoptotic function of Bad (42,46,47). The present study found that the phosphorylation of Erk and expression of p90RSK was increased in the zone of stasis following burn injury, which indicated that the Erk/p90RSK signaling pathway may be an endogenous defense mechanism against apoptosis caused by thermal injury. Furthermore, the Erk inhibitor PD98059 notably reversed the beneficial effect of ASMq on cell apoptosis and expression levels of CC3 and CC9 in the zone of stasis following burn injury. Meanwhile, the effect of PD98059 was associated with its inhibition of Erk phosphorylation and p90RSK expression levels and promotion of Bad-induced Cyto C release. These results suggested that the Erk/p90RSK/Bad signaling pathway may play a crucial role in the underlying mechanism of ASMq in the regulation of mitochondria-associated apoptosis. A schematic diagram of the potential mechanisms uncovered by the present study is presented in Fig. 7.

However, there are a number of limitations of the present study: Quantitative analysis of histological changes were not performed, ultrastructural changes of mitochondria and the metabolites of ASMq in vivo were not detected, expression levels of ROS, nitric oxide (NO) and inducible NO synthase were not detected directly and changes in other potential signaling molecules were not assessed. Further investigation should be considered to obtain in-depth details of the underlying mechanism. Overall, ASMq administration contributed to the attenuation of burn wound progression following thermal injury in a dose-dependent manner. The protective mechanism was shown to be associated with the antioxidative and anti-apoptotic effect of ASMq and the regulation of the mitochondria-associated apoptotic pathway by ASMq, with the involvement of upstream Erk/p90RSK activation.

Acknowledgements

Not applicable.

Funding

The present study was supported by the National Natural Science Foundation of China (grant nos. 81671909 and 8190195), Zhejiang Provincial Natural Science Foundation of China (grant nos. LY18H150004, LY19H150004 and LY20H150010) and Key Research and Development Project of Sichuan Science and Technology Department (grant no. 2018SZ0380).

Availability of data and materials

The analyzed datasets generated during the study are available from the corresponding author on reasonable request.

Authors' contributions

HZ, QF and SG conceptualized and designed the present study and provided administrative support. HZ, NL, HC and MY made contributions to the acquisition of data. HZ and SG analyzed and interpreted the data. HZ, QF and NL were involved in drafting the manuscript. HC, MY and SG revised the manuscript critically for important intellectual content. QF and SG confirm the authenticity of all the raw data. All authors read and approved the final manuscript.

Ethics approval and consent to participate

The present study was protocol approved by The Second Affiliated Hospital, Zhejiang University, School of Medicine, Institutional Animal Care and Use Committee (approval no. 2018-095) in accordance with the National Institutes of Health Guidelines for the Care and Use of Laboratory Animals.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

References


