INTRODUCTION

Psoriasis is a common chronic T-cell mediated autoimmune dermal disease, which affects approximately 2%–3% of the global population [1]. Reports have shown that psoriasis mostly affects the skin at the head (scalp), arms, elbow, knees, and shoulder [2]. Hence most of the psoriatic patients experience social discrimination and stigma (mental stress) due to their appearance and thus pose a huge negative impact on the quality of life [3]. The major dermal pathological events in psoriasis include hyperproliferation of epidermis (acanthosis) abnormal or hyper-differentiation of keratinocytes (parakeratosis) or plasmacytoid dendritic cells, excess infiltration of leukocytes (neutrophils) along with enhanced pro-inflammatory cytokines production and angiogenesis [4,5]. In particular, Th17 (T helper cells) is directly involved in the production of various inflammatory cytokines (interleukin [IL]-17 and 23) and has a direct influence on psoriasis [6]. At present only a few therapeutic strategies are used for treating or managing psoriasis patients.
rhiza dried root has several biologically active phytocomponents, 
and chronic pain [11]. Radix Salvia miltiorrhiza (Danshen/red sage; dried root) is a 
popular Chinese herb, which is recommended for treating dermal 
diseases, cardiovascular and cerebrovascular diseases as well as 
rheumatoid arthritis and chronic pain [11]. Radix Salvia miltiorrhiza 
dried root has several biologically active phytocomponents, 
out of which salvianolic acid B (SAB) is one of the abundant active 
phytocomponent with many beneficial properties like anti-inflammatory, 
antioxidant, anti-tumor/cancer, anti-diabetic [12-14]. Also, SAB is beneficial against various auto-immune dis- 
orders including rheumatoid arthritis and alopecia owing to its 
potent immunomodulatory and anti-inflammatory activities [15]. 
In addition, salvianolic acid B was reported to suppress melanin 
proliferation (production) and thus exhibit its derma protective 
property [16]. Based on the above reports, we hypothesize that 
salvianolic acid (potent immunomodulatory, anti-inflammatory 
and derma protective properties) would abolish psoriatic changes 
by lowering inflammatory markers in imiquimod (IMQ)-induced psoriasis.

METHODS

Chemicals and drugs

SAB (98% pure), sodium dodecyl sulfate (SDS), paraformaldehyde, 
glycerol, and bromophenol blue were bought from Sigma Aldrich (St. Louis, MO, USA). Topical IMQ (5%) and methotrexate (MTX) were purchased from Sichuan Mingxin Pharmaceuticals Co., Ltd. (Sichuan, China). All the other chemicals and reagents used in this study are of either analytical or HPLC grade.

Experimental animal and ethical approval

Healthy BALB/c mice of male gender aged between 7 to 8 weeks, weighing 23 ± 5 g were procured from Suzhou Fengshi Laboratory Animal Equipment and Merchant Co., Ltd. (Suzhou, China). All the mice were maintained at optimal laboratory conditions (22°C ± 2°C; 55%–60% humidity) in the polycarbonate cage. Mice have full access to mice fed and water (ad libitum) under 12 h light/dark cycle every day. All the mice were shaved at the back (3 × 4 cm) and allow for a week as an assimilation period. All the animal procedures and protocols were approved by the institutional animal ethical board of Wuhan Hospital of Traditional Chinese Medicine (No. WH/12a-34-2018). Mice were handled with extreme care based on guidelines put forth by National Institutes of Health for Care and Use of laboratory animals.

Induction of psoriasis

Psoriasis in BALB/c mice was induced by topical application of IMQ (5%; 62.5 mg) on the shaved back of all mice (except control and SAB alone group mice) for 7 days as previously mentioned by El Malki and his co-workers [17].

Experimental animals grouping

Totally 50 healthy male BALB/c mice were divided into 5 groups with 10 in each group. Control group mice (n = 10) received the only saline for 7 days, drug control group mice received only SAB (40 mg/kg via i.p. mixing with saline) for all for 7 days. Whereas, IMQ-induced psoriasis model mice were topically applied with 5% IMQ as indicated in the above section (induction of psoriasis) for 7 continuous days. The treatment group was pre-treated with either SAB (40 mg/kg) or MTX (1 mg/kg) for 7 days and followed by 7 days of IMQ induction as well as co- treatment with SAB or MTX. Overall, 7 days of pre and 7 days of co-treatment with SAB or MTX. But SAB alone and control mice received only 7 days of either saline or SAB.

Sample collection and processing

After 7 days of IMQ induction, all mice were fasted (overnight) and on 8th day morning all mice were weighed and euthanized under strong sodium pentobarbital at a dose of 55 mg/kg and the shaved area (back) were removed (epidermis) softly and gently as well as spleen were collected immediately and dry weight were weighed (to check spleen index: spleen weight/body weight). The dermal sample was segregated into two portions, one for biochemical/molecular analysis and other portion for histopathological analysis (fixed in 10% paraformaldehyde). The dermal tissue was homogenized with phosphate saline buffer solution (pH 7.4) and centrifuged (15,000 g at 4°C for 10 min) and the supernatant was used for biochemical and molecular analysis. The psoriasis area severity index (PASI) score (0 to 4) including the skin erythema and scaling were measured (lesioned area) to monitor the grade of psoriasis severity on the 2nd, 4th, 6th, and 8th day based on scoring range from 0 to 4. Where 0 represents none, 1 represents slight, 2 represents moderate, 3 represents marked/high and 4 represents very marked/very high as mentioned previously by Zhang et al. [18]. Moreover, the skin (dermal) thickness was measured on the 8th day (before sacrifice) using thickness gauge/caliper bought from Guanglu Digital Caliper Manufacturer Co.,
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Lipid peroxidation products and antioxidants

The skin lipid peroxidation product (malondialdehyde, MDA) and antioxidant activities (catalase, CAT; superoxide dismutase, SOD) were determined using commercial assay kit bought from Nanjing Bio-engineering Institute, Co. (Nanjing, China) based on suppliers’ protocol.

Inflammatory markers/mediators

The concentration of various inflammatory markers like IL-23/22/17A/1β and TNF-α in dermal tissue homogenate were measured using commercial ELISA kit purchased from R & D Systems (Minneapolis, MN, USA) in accordance with manufacturers instruction.

Western blot

The dermal tissue homogenate was incubated with RIPA lytic buffered solution containing various proteases and centrifuged and the resultant supernatant was used to estimate the protein level using Pierce BCA protein assay kit (Abcam, Cambridge, UK). An equal amount of protein sample (each group) was resolved and separated by 8% (keratin) and 10% (PI3k and Akt) SDS-PAGE apparatus and electro-transferred onto polyvinylidene difluoride (PVDF) membrane. The PVDF membrane was washed with PBS and followed by treatment with non-fat milk (5%), Tween 20, PBS. Then the membrane was probed with primary antibodies (bought from Cell Signaling Technology, Danvers, MA, USA) such as rabbit monoclonal anti-K16 (1:1,000), anti-K17 (1:1,200), rabbit polyclonal anti-pPI3K110 and PI3K (1:1,000), rabbit monoclonal anti-pAkt Ser473 and Akt (1:1,000) as well as housekeeping protein mouse polyclonal GADPH (1:1,500) for overnight at 4°C. Then the membrane was washed twice with PBS to remove unbound antibodies. Finally, the PVDF membrane was again incubated with secondary antibody-anti-horseradish peroxidase (HRP) anti-donkey IgG antibody (1:10,000) for 1 h at 37°C. The protein intensity (band) in the PVDF membrane was developed using Enhanced Chemiluminescence (ECL) detection system and the picture of the protein bands were quantified using Image Lab software; (V6.0) from Bio Rad Laboratories, Inc. (Hercules, CA, USA).

Histopathological analysis

As mentioned in the sample section, a portion of dermal tissue was fixed in 10% paraformaldehyde and treated with graded series of ethyl alcohol (rehydrate) and embedded with liquid paraffin wax to make a solid dermal tissue block. Tissue block was sectioned into a thin slice (4 μm) using ultra-microtome and mixed with xylene and mounted on a microscopic slide and stained with hematoxylin and eosin (H&E) stain for 12 h at room temperature. Then, the stained tissue slides were visualized using a light microscope at a magnification of 400× and pictured (BX43; Olympus Co., Tokyo, Japan) to examine any skin morphological changes with the help of expert pathologist who is blinded from grouping or experiment.

Data analysis

For the present study, the data analysis was conducted using Statistical Package for the Social Software-SPSS (ver 21; IBM, Armonk, NY, USA). Values are expressed as the mean ± standard deviation. The statistical difference (p-value) between the experimental groups was assessed by one-way ANOVA followed by Duncan’s multi-comparison test. p < 0.05 is deemed statistically significant and were expressed with different symbols.

RESULTS

Impact of SAB/MTX on spleen index

The spleen index was calculated by dividing spleen weight (dry weight)/body weight to assess the dynamic body changes. Both control (0.005) and drug control (0.005) mice showed a lesser spleen index. Whereas, IMQ exposed mice (0.012) displayed higher spleen index (p < 0.01) than control mice. Nevertheless, IMQ-induced mice pre-treated and co-treated with SAB (0.008) and MTX (0.008) significantly reduced (p < 0.01) the spleen index than IMQ-exposed BALB/c mice.

Impact of SAB/MTX on PASI, erythema, scaling and skin thickness

Fig. 1 showcase the impact of SAB on cumulative PASI (scaling/erythema), dermal scaling and erythema as well as skin thickness (dorsal skin) in control and IMQ-induced mice. Topical application of IMQ (5%) resulted in elevated psoriatic plaque, erythema, scaling and inflamed dermal lesion, which are reflected in a significant increase (p < 0.01) in the levels of PASI (Fig. 1A), scaling (Fig. 1B), erythema (Fig. 1C) scores and skin thickness (Fig. 1D) than control mice. Whereas, 7 days of pre and co-treatment with SAB or MTX showed a significant decrease (p < 0.01) in the levels of PASI, scaling, erythema scores and skin thickness as compared to IMQ-induced mice. Both the control and SAB alone (drug control) mice did not show any psoriatic changes (symptoms) with normal skin texture.

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Table 1. Exemplifies the skin lipid peroxidation products and antioxidant activities in control and IMQ-induced mice

| Parameters      | MDA (nmol/mg protein) | CAT (U/mg protein) | SOD (U/mg protein) |
|-----------------|------------------------|--------------------|--------------------|
| Control         | 10.64 ± 1.80           | 21.50 ± 2.40       | 39.22 ± 4.50       |
| SAB             | 11.02 ± 1.75           | 21.00 ± 2.25       | 38.85 ± 4.09       |
| IMQ             | 28.95 ± 3.70           | 09.88 ± 1.10       | 20.40 ± 2.20       |
| IMQ + SAB       | 18.61 ± 2.35           | 17.10 ± 3.00       | 33.68 ± 3.80       |
| IMQ + MTX       | 20.05 ± 2.03           | 14.75 ± 2.05       | 29.79 ± 3.15       |

Values are expressed as the mean ± standard deviation. IMQ, imiquimod; MDA, malondialdehyde; CAT, catalase; SOD, superoxide dismutase; SAB, salvianolic acid B; MTX, methotrexate. p-value (\*p < 0.05, \*\*p < 0.01): where ‘a’ denotes the comparison between IMQ vs. Control, ‘b’ denotes the comparison between IMQ + SAB vs. IMQ, ‘c’ denotes the comparison between IMQ + MTX vs. IMQ, ‘d’ denotes the comparison between IMQ + SAB vs. IMQ + MTX.

Table 2. Exemplifies the skin inflammatory markers in control and IMQ-induced mice

| Parameters      | IL-23 (pg/mg protein) | IL-22 (pg/mg protein) | IL-17A (pg/mg protein) | IL-1β (ng/mg protein) | TNF-α (ng/mg protein) |
|-----------------|-----------------------|-----------------------|------------------------|-----------------------|-----------------------|
| Control         | 15.80 ± 1.75          | 1.10 ± 0.15           | 7.55 ± 0.75            | 7.42 ± 0.71           | 5.10 ± 0.50           |
| SAB             | 16.00 ± 2.10          | 1.05 ± 0.10           | 7.40 ± 0.58            | 7.45 ± 0.80           | 4.97 ± 0.52           |
| IMQ             | 41.25 ± 4.60          | 3.87 ± 0.40           | 13.98 ± 1.80           | 25.10 ± 2.45          | 15.04 ± 1.85          |
| IMQ + SAB       | 23.50 ± 3.00          | 2.02 ± 0.20           | 9.75 ± 1.10            | 11.21 ± 1.35          | 7.14 ± 0.88           |
| IMQ + MTX       | 25.18 ± 2.10          | 2.22 ± 0.22           | 9.50 ± 1.12            | 13.76 ± 1.50          | 8.95 ± 0.88           |

Values are expressed as the mean ± standard deviation. IMQ, imiquimod; IL-23/22/17A/17B, interleukins 23/22/17A/17B, tumour necrosis factor alpha; SAB, salvianolic acid B; MTX, methotrexate; NS, non-significant. p-value (\*p < 0.05, \*\*p < 0.01): where ‘a’ denotes the comparison between IMQ vs. Control, ‘b’ denotes the comparison between IMQ + SAB vs. IMQ, ‘c’ denotes the comparison between IMQ + MTX vs. IMQ, ‘d’ denotes the comparison between IMQ + SAB vs. IMQ + MTX.
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Impact of SAB/MTX on lipid peroxidation products and antioxidants

Table 1 exemplifies the skin lipid peroxidation products and antioxidant activities in control and IMQ-induced mice. An exponential increase in the levels of lipid peroxidation and substantial decrement in the activity of dermal antioxidants like CAT and SOD were observed in IMQ-induced mice (p < 0.01). However, the mice injected with SAB or MTX (before and after IMQ-induction) showed a notable decrease in the levels of lipid peroxidation as well as a considerable increment (p < 0.01) in the antioxidant’s activity of CAT and SOD on equivalent with IMQ-induced psoriatic mice. In the case of antioxidant status, the SAB exhibits potent antioxidant activity than MTX.

Impact of SAB/MTX on inflammatory markers/mediators

As shown in Table 2, the concentration of various dermal inflammatory markers/mediators like IL-23, 22, 17A, 1β and TNF-α were exponentially elevated in IMQ-exposed mice (p < 0.01) in parallel to control mice. Nevertheless, those elevated skin inflammatory markers/ mediators like IL-23, 22, 17A, 1β, and TNF-α were significantly (p < 0.01) abolished after treatment with SAB or MTX (before and after IMQ induction). Also, on a comparison between both treatment groups, the standard MTX showed slightly better anti-inflammatory activity than SAB.

Impact of SAB/MTX on the protein expression of keratin markers and PI3K/Akt signaling molecules

The protein expression of keratin markers (K16 and K17; Fig. 2) and PI3K/Akt signaling molecules (Fig. 3) were quantified using the western blot technique [19]. The mice topically applied with 5% IMQ (p < 0.01) showed greater upregulation of keratin markers (K16, K17) and PI3K/Akt signaling molecules (pAkt/Akt and pPI3K/PI3K ratio). Whereas, mice supplemented with SAB (p < 0.01) or MTX (p < 0.01), significantly downregulated the protein expression of keratin markers (K16, K17) and PI3K/Akt signaling molecules (pAkt/Akt and pPI3K/PI3K ratio) as compared to IMQ-exposed mice and thus exhibiting potent anti-psoriasis activity. Nonetheless, MTX showed superior anti-psoriasis activity than SAB due to its strong immunomodulatory property.

Impact of SAB/MTX on histopathological changes in dermal tissue

Fig. 4 illustrates the histopathological changes in dermal tissue

![Fig. 2. Illustrate the protein expression of skin keratin markers (K16 and K17) in control and IMQ-induced mice. Values are expressed as the mean ± standard deviation. IMQ, imiquimod; SAB, salvianolic acid B; MTX, methotrexate; K16 and K17, keratin 16/17. p-value (p < 0.05, *p < 0.01): where 'a' denotes the comparison between IMQ vs. Control, 'b' denotes the comparison between IMQ + SAB vs. IMQ, 'c' denotes the comparison between IMQ + MTX vs. IMQ, 'd' denotes the comparison between IMQ + SAB vs. IMQ + MTX.](Image 1)

![Fig. 3. Illustrate the protein expression of skin PI3K/Akt signaling molecules in control and IMQ-induced mice. Values are expressed as the mean ± standard deviation. IMQ, imiquimod; SAB, salvianolic acid B; MTX, methotrexate; PI3K, phosphatidylinositol-3-kinase; Akt: protein kinase B. p-value (p < 0.05, *p < 0.01): where 'a' denotes the comparison between IMQ vs. Control, 'b' denotes the comparison between IMQ + SAB vs. IMQ, 'c' denotes the comparison between IMQ + MTX vs. IMQ, 'd' denotes the comparison between IMQ + SAB vs. IMQ + MTX.](Image 2)

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upon staining with H&E in control and IMQ-induced mice. Control (Fig. 4A) and SAB (Fig. 4B) mice dermal tissue slide display normal smoother epidermis without any inflammation or lesion. The IMQ induced mice dermal section (Fig. 4C) showed thickened epidermis (hyperplasia and acanthosis-circle) with elevated immune cell infiltration (arrow mark). Nevertheless, the mice dermal section treated with SAB (D) and MTX (E) portrait lesser thick epidermis (circle) with decreased immune cell infiltration (arrow mark) than IMQ-induced mice section. Scale bar:100 μm. IMQ, imiquimod; SAB, salvianolic acid B.

**DISCUSSION**

Ample amount of studies indicated that topical application of IMQ (5%) could trigger immune system via toll-like receptor 7/8 (TLR 7/8) which results in hyperstimulation of dendritic cells and hyperproliferation of keratinocytes (epidermis) and eventually upregulate various inflammatory cytokines (modulating IL-23/17 axis) and resembles human-plaque like psoriasis [1,18]. Therefore, the IMQ-exposed psoriasis-like mice model is a well-accepted psoriasis model (similar pathophysiology) to examining the anti-psoriasis activity of any drug [6]. Hence, the IMQ-induced psoriasis mouse model (BALB/c mice- highly susceptible to immune changes) was employed for this study to assess the anti-psoriatic activity of SAB and compared with a standard drug-like MTX (potent immunomodulatory/suppressive property). Before this study, we also conducted dose-dependent preliminary studies with SAB in IMQ-model to choose an effective dose, the result indicated that 40 mg/kg of SAB can effectively lower PASI...
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and histological changes (data not shown). Therefore, the 40 mg/kg of SAB was only used and compared with MTX. The outcome of this animal study clearly indicated that BALB/c mice treated with SAB or MTX significantly lowered the levels of PASI, skin thickness, lipid peroxidation product formation (free radicals), inflammatory markers as well as modulating the protein expressions of keratin markers and PI3K/Akt signaling pathway.

The spleen index was determined to check the dynamic body changes as it is expressed based on spleen and body weight ratio. Mice topically applied with IMQ showed splenomegaly (enlarged spleen) through systemic effect and thus the spleen index was notably increased. Nevertheless, BALB/c mice SAB or MTX treatment significantly reduces the spleen index by reversing splenomegaly through decreasing cellularity (T cells) of periarteriolar lymphoid sheaths (PALS) and by lowering immune cell infiltration in the spleen. Previously, Elmore [20], also indicated the involvement of increased cellularity of PALS for splenomegaly in IMQ-induced psoriasis model. Aforementioned, IMQ induces hyperproliferation of dendritic cells and enhances keratinocyte proliferation which subsequently leads to scaling, redness/erythema as well as thickening of the epidermis (acanthosis) and increased inflammatory cell infiltration [18]. Likewise, during this study also we also encountered similar results with elevated psoriatic plaque, erythema, scaling with increases PASI, scaling, erythema scores along with increased skin thickening along with considerable psoriatic morphological changes (hyperplasia and acanthosis with elevated neutrophil infiltration) in IMQ-induced mice. While, 7 days of supplementation with SAB or MTX significantly suppressed those elevated PASI, scaling, erythema scores, skin thickness and psoriatic morphological changes (lesser hyperplasia with reduced neutrophil infiltration) due to its anti-inflammatory, immunomodulatory and dermaprotective activities [12,15].

A growing body of evidence indicates that excessive free radical production (reactive oxygen species) with abolished antioxidants (oxidative stress) is one of the major pathological events which contributes to psoriasis [21,22]. Hence, the antioxidant activity including CAT and SOD in dermal tissue as well as lipid peroxidation products (MDA) was determined for this study. The levels of lipid peroxidation products like MDA were elevated along with decreased antioxidants like SOD and CAT in IMQ applied mice due to excessive free radical generation. Nevertheless, the mice treated with SAB or MTX significantly decreased the production of free radicals and thus improve antioxidant activity with lowered lipid peroxidation product (MDA) formation due to potent antioxidant and free radical scavenging as well as anti-lipid peroxidative activities [14,23]. Moreover, Zhao and others [24], hinted that salvianolic acid B display potent free radical scavenging activity owing to its free hydroxyl group. For the present study, we plan to examine various inflammatory markers like IL-23, 22, 17A, 1β and TNF-α. Since those pro-inflammatory cytokines (markers) plays a pivotal role in the pathogenesis of psoriasis along with oxidative stress (as mentioned before). A pronounced increase in the concentration of various dermal inflammatory markers/mediators (IL-23, 22, 17A, 1β and TNF-α) was observed in IMQ-induced mice. But, those skin inflammatory markers/mediators (IL-23, 22, 17A, 1β, and TNF-α) were significantly lowered upon supplementation with SAB or MTX in BALB/c mice. Previously, Zhang and Wang [25], also demonstrated that salvianolic acid B could significantly inhibit various inflammatory markers like IL-1β, 6, and TNF-α in human aortic smooth muscle cells because of its anti-inflammatory and antioxidant properties.

Many scientists have demonstrated that hyperproliferation of keratinocytes (epidermis) via upregulating various keratins especially K16 and K17 is recognized as a hallmark of psoriasis [26,27]. However, the expression of these keratins (K16 and K17) in normal keratinocytes were zero or minimal and are tightly regulated by various signaling pathway [28]. K16 is a type 1 cytokeratin, which strongly pairs with K6 and plays a major role in intermediate filament formation especially during the psoriatic condition. Whereas, K17 is also a type 1 cytokeratin (a cytoskeletal protein) and display a similar role as K16. Both K16 and 17 were found in the abnormal sebaceous gland, nail beds, and hair follicles and directly contribute to psoriasis as they both act as immunogenic antigens and trigger an immune response [29,30]. Hence, for the present study, the protein expression of both keratin K16 and K17 were examined using the western blot technique. The protein expression of both keratins (K16 and K17) were significantly upregulated in IMQ applied BALB/c mice due to immunogenic stimuli (IMQ) via TLR 7/8 by upregulating pro-inflammatory cytokines like IL-17 and 23 which results in upregulation of K16 and K17 and subsequently end up in hyperproliferation of keratinocytes. The above results are in corroboration with the results of Zhang and his co-workers [31], also confirmed that induction of IMQ would significantly upregulate K16 and K17. But mice treated with SAB or MTX showed considerable downregulation of these keratins (K16 and K17). The reason might be a potent anti-inflammatory and immune-suppressive activity of both SAB and MTX [15,32].

Based on the above results it’s clear that SAB could downregulate keratin expression and thus favor anti-hyperproliferation of keratinocytes, but the mechanism is still unknown. Moreover, numerous research groups hinted that the PI3K/Akt pathway inhibitors might prevent the psoriasis development in various psoriasis model particularly in the IMQ-induced mice model [33,34]. Therefore, the author hypothesizes that SAB would exert anti-hyperproliferation of keratinocytes (anti-psoriasis) by regulating PI3K/Akt signaling pathway. Topically application of 5% IMQ on the back of mice showed greater upregulation of PI3K/Akt signaling molecules (pAkt/Akt and pPI3K/PI3K ratio). Likewise, Chamcheu et al. [35], showed that mice induced with IMQ showed enhanced protein expression of PI3K/Akt signaling molecules in the IMQ-induced mice model. Nonetheless, mice administered with SAB or MTX, significantly suppressed the activation of PI3K/
Akt signaling molecules by lowering phosphorylation of PI3K 110 and Akt Ser^{12} (serine residue 473) which reflects in lowered pAkt/Akt and pPI3K/PI3K ratio. Salvianolic acid B was reported to inhibit oral squamous cell carcinoma by downregulating PI3K/ Akt signaling molecules (inactivate PI3K/Akt signaling pathway) and thus protect cells from hyperproliferation or hyperplasia [13]. Overall, SAB display potent anti-psoriasis activity by downregulating Akt signaling molecules (inactivate PI3K/Akt signaling pathway) to inhibit oral squamous cell carcinoma by downregulating PI3K/ Akt signaling molecules. A few limitations of this study are the avoidance of measurement of the dermal concentration of SAB or MTX, as well as the expression of various immune cells and its proliferative property.

This animal study clearly demonstrated that BALB/c mice treated with SAB or MTX significantly lowered the levels of PASI, skin thickness, lipid peroxidation product production (free radicals), inflammatory markers and keratin markers through modulating PI3K/Akt signaling pathway and thus exhibit its strong anti-psoriatic property. However, clinical trials are needed to confirm the anti-psoriatic property of SAB before recommending to psoriasis patients.

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CONFLICTS OF INTEREST

The authors declare no conflicts of interest.

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