

Water Extract of *Senecio scandens* Buch.-Ham Ameliorates Pruritus by Inhibiting MrgprB2 Receptor

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Background: *Senecio scandens* Buch.-Ham (*S. scandens*) belongs to the Compositae family. As a Traditional Chinese medicine, *S. scandens* has been used in China to treat conjunctivitis, mastitis and vaginitis, it also has the function of antibacterial and relieving itching.

Methods: Water extract of *S. scandens* (WSS) was prepared and its quality was controlled by HPLC. The antipruritic effects of WSS were evaluated by itch behavioral experiments. The oxazolone and compound 48/80 were induced to mice scratch behavior, scratch was recorded 30 min after sensitization. The relationship between the antipruritic mechanism and MrgprB2 on mast cell was studied by using mast cell-deficient Kit (W-sh) “Sash” mice and MrgprB2^{-/-} mice. The mast cells were observed by toluidine blue staining. In vitro, the effects of WSS on MrgprB2 were studied by calcium imaging; The whole-cell patch clamp method recorded the MrgprB2 mediate voltage-dependent currents in mast cells.

Results: The content of rutin (0.012%) and hyperin (0.014%) in the WSS were determined. WSS could ameliorate the pruritus induced by Oxazolone (inhibition was 41.19%, $p = 0.004$) and compound 48/80 (inhibition was 50.29%, $p = 0.001$). Meanwhile, WSS could reduce the number of mast cells in mice skin tissue with allergic contact dermatitis (ACD) ($p = 0.002$) or compound 48/80 ($p = 0.013$). In addition, WSS could inhibit the calcium influx (1 mg/mL: $p = 0.001$, 3 mg/mL: $p < 0.0001$) and the voltage-dependent currents induced by activation of MrgprB2 on mast cell. WSS also attenuated the calcium influx induced by compound 48/80 in HEK293 cells overexpressing MrgprB2/X2.

Conclusion: These results showed that WSS could ameliorate pruritus by inhibiting MrgprB2 receptor on mast cells.

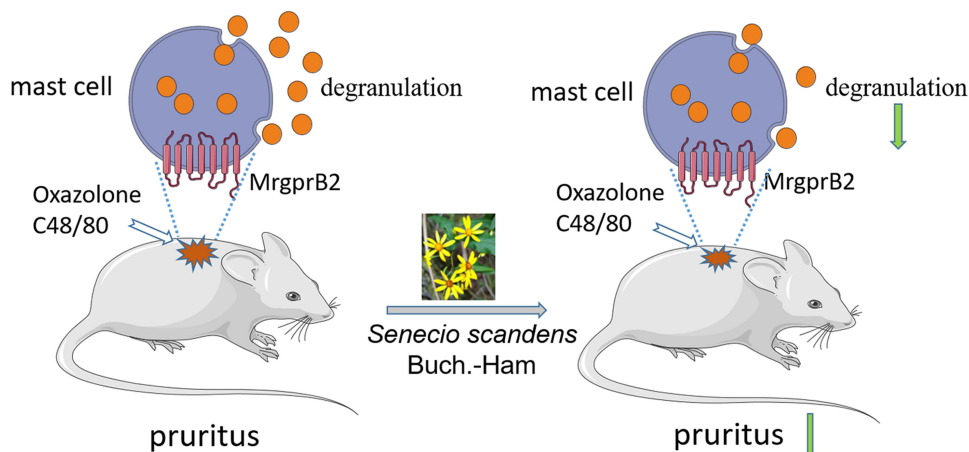
Keywords: *Senecio scandens* Buch.-Ham, pruritus, mast cell, MrgprB2 receptor

Introduction

The *S. scandens* is used in Chinese medicine as a treatment for antimicrobial, antioxidant and anti-inflammatory medicine.¹ Studies have shown that the *S. scandens* water extract could be commonly used to treat amniotic keratitis.² And could effectively alleviate the inflammation of keratoconjunctivitis in rats with allergic conjunctivitis via the NLRP3/Caspase-1/IL-1 β pathway.³ Petroleum ether extract of *S. scandens* participated in process of anti-inflammation response by inhibiting the expression of some pro-inflammatory mediators such as NO, IL-1 β and IL-6, and the activation of the NF- κ B and MAPK signaling pathways.⁴ *S. scandens* flavonoid could alleviate inflammation by inhibiting the production and release of inflammatory factor prostatin E2 (PGE2) and the infiltration of white blood cell.⁵ Three novel polysaccharides (SP2-1, SP2-2 and SP3-2) from *S. scandens* have recently displayed prominent antioxidant activity and potential immune response,⁶ however, the antipruritic mechanism of *S. scandens* needs to be further studied.

Pruritus is caused by a desire or reflex to scratch.⁷ About 25% of people experience itching during their lifetime,⁸ this common symptom has a serious impact on their way of life (such as working and sleeping).^{9,10} The skin immune diseases

Graphical Abstract



have pruritus symptoms, such as allergic dermatitis (AD), allergic contact dermatitis (ACD), and urticarial.¹¹ Since conventional anti-pruritus therapies, such as antihistamines, exert only limited effects, a drug of a treatment option for itch is urgently needed.^{12,13} Mast cell as sentinels, regulate innate and adaptive the immune responses and have a significant impact on disease.¹⁴ Currently, skin mast cells have been indicated to play a role of pruritus in autoimmune bullous disease.¹⁵ Mast cell activation by MrgprB2 mediates the neurogenic inflammation-evoked itch in mice, and promoted ACD-associated pruritus in mice.¹⁶ Activation of MrgprB2 receptors on mast cells, could lead to mast cell degranulation and release of histamines, tryptase, and serotonin, which are involved in pruritus. Therefore, in this study, we explored whether the antipruritic mechanism of *S. scandens* was related to the activity of MrgprB2.

Materials and Methods

Materials

Ethics Statement

The Experimental Animal Ethics Committee of Nanjing University of Chinese Medicine approved the development of this animal experiment. All animal experiments conformed to the Guidelines for the Care and Use of Laboratory Animals and were approved by the Animal Ethics Committee of Nanjing University of Chinese Medicine (Ethics licence ACU190601, 20,190,605).

Animals

C57BL/6J mice (WT), purchased from Charles River (Beijing, China), sash mice (C-kit mutant genetically mast cell-deficient Kit (W-sh) /Kit (W-sh)) and MrgprB2^{-/-} mice were donated by Johns Hopkins University. Male mice and 18–22 g for experimental animals were housed at constant humidity (40–60%) and temperature (22 ± 2°C).

Reagents

DMEM culture medium, Fetal bovine serum, Penicillin, Streptomycin and Trypsin (All purchased from Gibco, USA); Fibronectin (Sigma, USA), Stem cell growth factor (Sigma, USA), Percoll (Sigma, USA), Oxazolone (Sigma, USA), Compound 48/80 (Sigma, USA). Rutin and hyperin (Yuanye, Shanghai). H&E kit (Solarbio, Beijing), Toluidine Blue (Aladdin, Shanghai).

Methods

Water Extract of *S. scandens* (WSS)

S. scandens (Suzhou Tianling Chinese Herbal Medicine Co. Ltd. voucher no: 210722, dry and aboveground parts) were purchased from the Taizhou Hospital of Traditional Chinese Medicine, Jiangsu, and authenticated by Gongxi Chen. The preparation method of the water extract *S. scandens*: first, we weighed 30 g of *S. scandens* in ultrapure water with a volume of 600 mL; then we boiled them at 100 °C for 30 min to collect the extract. And the materials were boiled again in 600 mL of ultrapure water with 100 °C for 30 min, then we mixed the two extracts, boiled and evaporated them to 25 mL (equivalent to 1.2 g (*S. scandens*)/mL). We filtered the extract through a 0.22 µm filter and stored it at -20 °C.

Behavior Tests

Pruritus behavior test in ACD: Acetone and olive oil were mixed in a ratio of 4:1, and oxazolone was added to prepare solutions containing 3% and 1% oxazolone for presensitization and sensitization, respectively. C57BL/6J, Sash and MrgprB2^{-/-} mice were randomly divided into Vehicle group, ACD group, and ACD + WSS group, respectively. According to the human-mouse dose conversion formula, the dose of WSS was 3.9 g/kg/d (The optimal concentration of *S. scandens* was screened in [Supplementary Figure 1](#)), and the administration method was gavage. Four days before sensitization, presensitization was performed by adding 40 µL of 3% oxazolone solution to the hairless area of the abdomen of mice. The baseline of scratch behavior was recorded at day 0 for 30 min. Then, mice were sensitized with 1% oxazolone solution 40 µL in the middle of the hairless area of the back, and stimulated once every other day. Mice in the WSS group were given WSS 1 hour before sensitization every day. Scratch behavior was recorded 30 min after sensitization ([Supplementary Figure 2](#)).

Behavior test in compound 48/80-induced pruritus: The scratch behavior baseline of mice was recorded 1.5 h before modeling, and the WSS group was given *S. scandens* 1 h before modeling. We dissolved 50 µg compound 48/80 in 0.9% NaCl 25 µL and injected subcutaneously into the neck. Scratching behavior was observed for 30 min ([Supplementary Figure 3](#)).

Each group number of mice was not less than 6 in all experiments. The frequency of hind foot lift was used to evaluate scratching behavior. All experiments were conducted from August 2021 to the end of August 2022.

Histopathological Analysis

After the itching test, the skin in the model area (about 1 cm²) was removed and fixed with 4% paraformaldehyde (4°C, 12 h) and 30% sucrose solution (4°C, 12 h). After optimal cutting temperature compound (OCT) embedding, the tissue sections were prepared with a thickness of 15 microns. We used 1% toluidine blue staining to identify mast cells and hematoxylin-eosin (H&E) staining was used to evaluate skin inflammation.

Intracellular Calcium Measurement

Perfusion buffer configuration (1 L): (7.25 g NaCl, 222 mg KCl, 187 mg CaCl₂, 121 mg MgCl₂·6H₂O, 3.6032 g glucose, 2.383 g HEPES, 6.846 g sucrose, 112.8 mg NaHCO₃, PH was 7.4 by NaOH). Mast cells were treated with mixture (1 µL /mL fluo-4 AM and 0.02% Pluronic F-127 dissolve in buffer) for 30 min at 25 °C. The intracellular calcium ions were detected in anechoic chamber by 488-nm excitation.¹⁷ For HEK293 cells, the intracellular calcium ions were detected by 340 and 380 nm excitation.

Whole-Cell Currents Recording

The electrode resistance was 4–6 MΩ, and 5 µg/mL compound 48/80 as agonist. The currents signal was collected by Multiclamp 700 B amplifier (Molecular Devices, Inc, San Jose, USA), and the currents signal was digitized by Digidata 1440 A (Molecular Devices, Inc, San Jose, USA). Whole-cell currents were recorded and stored in computer. We used Clampex 10.3 for currents numerical analysis.

High Performance Liquid Chromatography (HPLC) Analysis

A mixed standard stock solution containing two reference substances of rutin and hyperin were prepared by dissolving them in methanol and their concentrations were as follows: 0.2 mg/mL rutin, 0.1 mg/mL hyperin. The mixed standard

stock solution was then diluted with methanol to a series of appropriate concentrations for construction of the calibration curves.

Analyses were performed by using a reversed-phase HPLC system with a C₁₈ column (250 mm × 4.6 mm, 5 μm). The mobile phase consisted of 0.1% formic acid (A) and acetonitrile (B) in a ratio specified by the following binary gradient with linear interpolation: 1–3 min, 5%B; 3–6 min, 5–15%B; 6–15 min, 15–20%B; 15–17 min, 20–70%B; 17–17.5 min, 70–5%B; 17.5–23 min, 5%B; The flow rate: 0.8 mL/min; temperature: 30°C; detection wave length: 360 nm; and sample volume: 1 μL.

Statistical Analysis

All results values are exported as mean ± standard error (SEM) from GraphPad Prism 8.0 software, two-tailed Student's *t*-test, or one-way ANOVA analysis for comparing different groups.

Results

Rutin and Hyperin Were Detected in WSS

To identify the quality and authenticity of *S. scandens*, the content of rutin and hyperin in the WSS were determined by HPLC.¹⁸ After calculation, the content of rutin was 0.012% and the content of hyperin was 0.014% (Figure 1A and B).

WSS Could Ameliorate Pruritus Mediated by MrgprB2 Receptor

Our experimental results showed that WSS is not relieve the skin damage (Figure 2A); it could significantly inhibit the scratching behavior of ACD model in C57BL/6J mice (Figure 2B, the inhibition was 41.19%, *P* = 0.004). In addition, results also showed that compared with WT mice, the inhibition ratio of WSS on the scratching behavior in sash (Figure 2C, the inhibition was 27.96%, *P* = 0.033) or MrgprB2^{-/-} mice (Figure 2D, the inhibition was 12.83%, *P* = 0.046) was reduced.

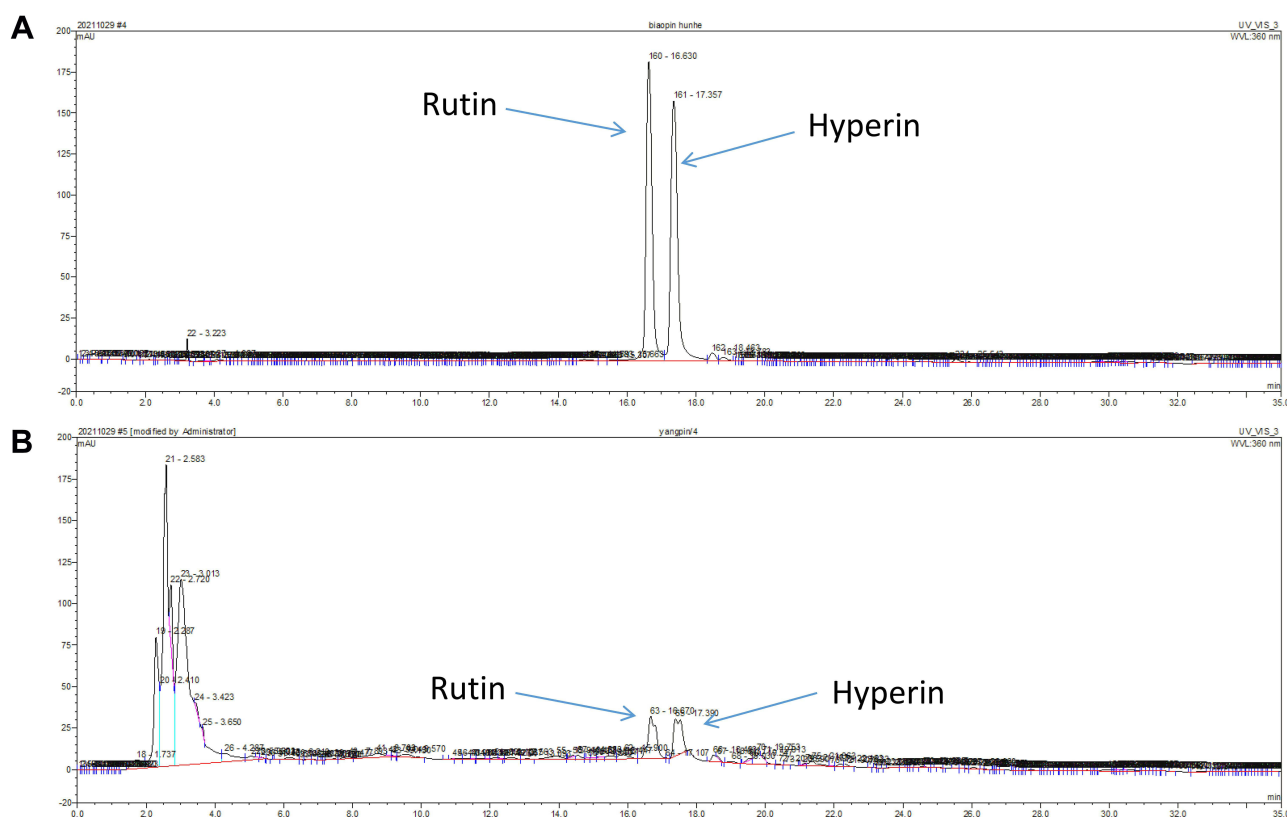


Figure 1 High-performance liquid chromatography of WSS. **(A)** High-performance liquid chromatography rutin and hyperin (Reference substances). **(B)** High-performance liquid chromatography of WSS.

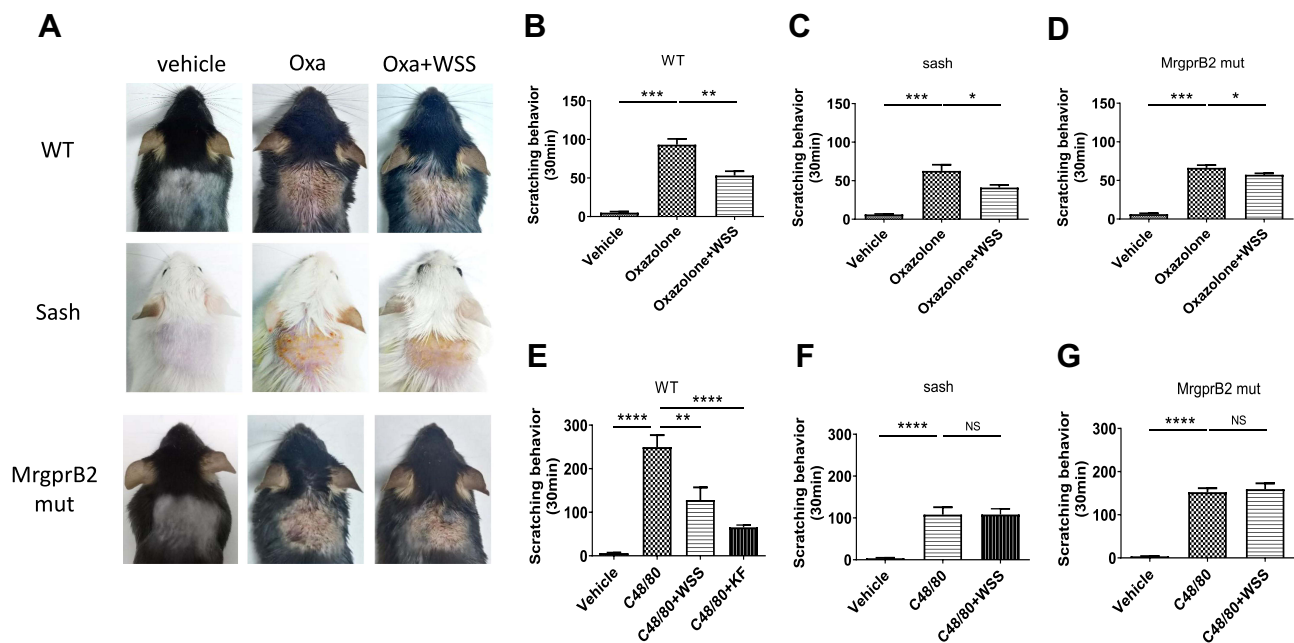


Figure 2 WSS ameliorated pruritus by inhibiting MrgprB2 receptor on mast cell. **(A)** The skin lesions in mice induced by oxazolone. **(B)** The WSS could significantly reduce the scratching behavior of ACD model in C57BL/6 mice. **(C)** The WSS could slightly reduce scratching behavior in ACD model of Sash mice. **(D)** The WSS could slightly reduce scratching behavior in ACD model of MrgprB2^{-/-} mice (**p* < 0.05, ***p* < 0.01, ****p* < 0.001, one-way ANOVA analysis followed by Dunnett's multiple comparisons test). **(E)** WSS and ketotifen (10 mg/kg, ig) could reduce the scratching behavior of C57BL/6 mice induced by compound 48/80 (***p* < 0.01, *****p* < 0.0001, one-way ANOVA analysis followed by Dunnett's multiple comparisons test). **(F and G)** WSS had no synergistic effect on the scratching behavior in Sash or MrgprB2^{-/-} mice induced by compound 48/80.

Abbreviation: NS, no significant.

These results suggest that the mechanism of WSS alleviating pruritus induced by allergic contact dermatitis may be related to the MrgprB2 receptor on mast cells.

To verify the mechanism of WSS anti-pruritus was related to mast cell and MrgprB2 receptor, we examined the effect of WSS on scratching behavior induced by compound 48/80. Our results showed that mast cell stabilizer (ketotifen) could reduce the scratching bouts (Figure 2E, *P* < 0.0001), and WSS could reduce the scratching bouts induced by compound 48/80 (Figure 2E, inhibition was 50.29%, *P* = 0.001). In addition, WSS could not reduce the scratching bouts induced by compound 48/80 in sash mice (Figure 2F, *P* = 0.991) or in MrgprB2^{-/-} mice (Figure 2G, *P* = 0.672). Therefore, we believed that the anti-pruritus mechanism of WSS may be related to MrgprB2 receptor.

The Infiltration of Mast Cell Was Inhibited by WSS

Toluidine blue staining results showed that WSS could reduce the number of mast cells in mice skin tissue with ACD (Figure 3A and B, *P* = 0.002). However, WSS had no significant inhibitory effect in MrgprB2^{-/-} mice (Figure 3C, *P* = 0.613). Meanwhile, WSS could reduce the number of mast cells in C57BL/6J mice induced by compound 48/80 (Figure 3D and E, *P* = 0.013). However, WSS did not show significant inhibitory effect in MrgprB2^{-/-} mice (Figure 3F, *P* = 0.975). In addition, HE staining showed that WSS did not improve inflammatory cell infiltration in C57BL/6J mice (Supplementary Figure 4).

WSS Inhibited MrgprB2-Mediated [Ca²⁺]_i Elevation

Next, we verified the effect of WSS on MrgprB2-mediated [Ca²⁺]_i influx (Figure 4A). The results showed that WSS inhibited [Ca²⁺]_i influx of mast cells, and depended on concentration (Figure 4B–C, 300 ng/mL: *P* = 0.13, 1 mg/mL: *P* = 0.001, 3 mg/mL: *P* < 0.0001). Meanwhile, WSS had no effect on the [Ca²⁺]_i influx in mast cells from MrgprB2^{-/-} mice (Figure 4D–F, 300 ng/mL: *P* = 0.382, 1 mg/mL: *P* = 0.961, 3 mg/mL: *P* = 0.572). In addition, WSS inhibited [Ca²⁺]_i influx of mast cells, which had nothing to do with the cytotoxicity (Supplementary Figure 5).

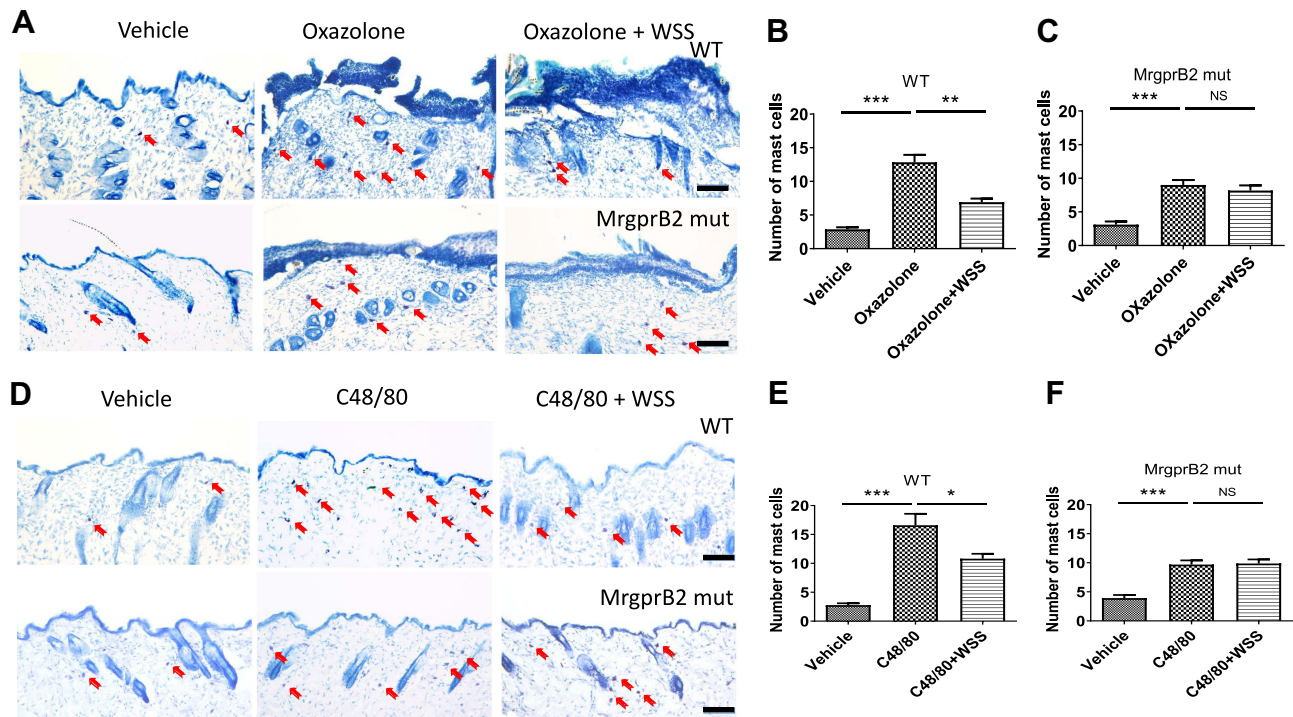


Figure 3 The infiltration of mast cell was inhibited by WSS. **(A)** Toluidine blue staining of different groups of mice skin tissue of ACD. **(B)** The number of mast cell was significantly decreased in Oxazolone + WSS group. **(C)** WSS did not affect the number of mast cells in *MrgprB2*^{-/-} mice. **(D)** Toluidine blue staining of different groups of mice skin tissue induced by compound 48/80. **(E)** The number of mast cell was significantly decreased in C48/80 + WSS group, **(F)** WSS did not affect the number of mast cells in *MrgprB2*^{-/-} mice. (**p* < 0.05, ***p* < 0.01, ****p* < 0.001, one-way ANOVA analysis followed by Dunnett's multiple comparisons test), scale bar: 100 μ m.

Abbreviation: NS, no significant.

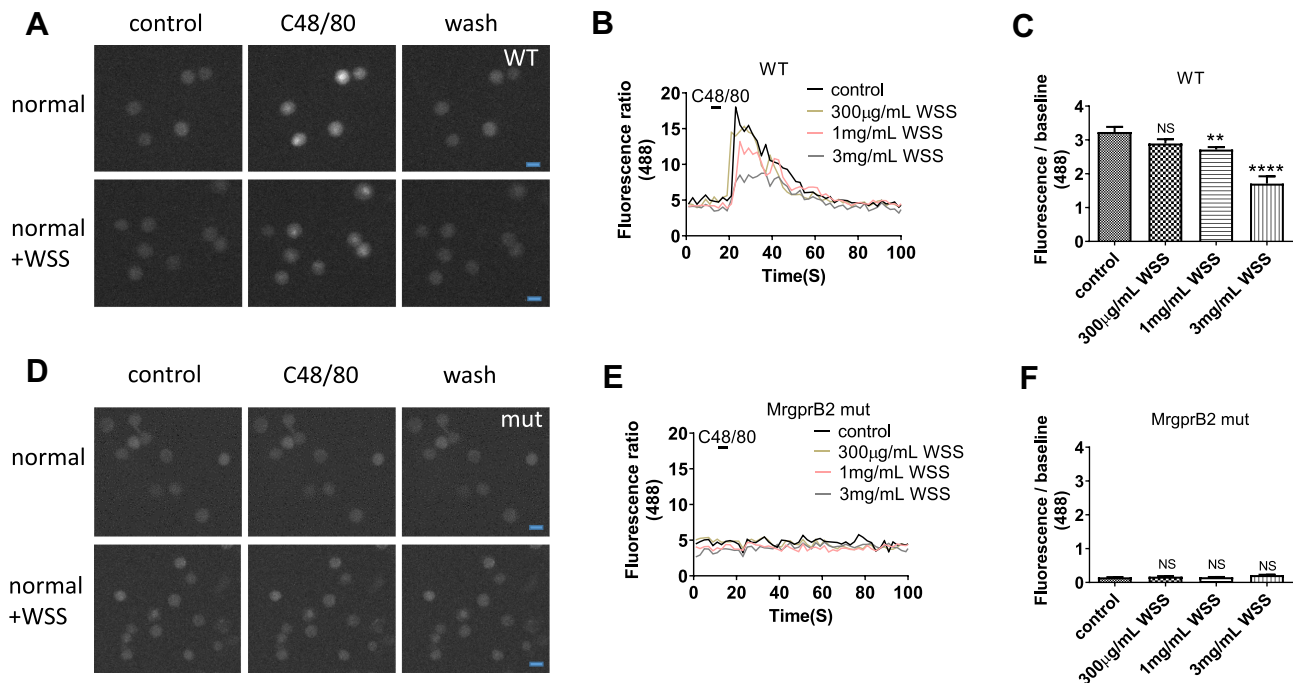


Figure 4 Effect of WSS on the calcium influx caused by compound 48/80 in mouse peritoneal mast cells isolated from C57BL/6j or *MrgprB2*^{-/-} mice. **(A and D)** Representative trace for changes in $[Ca^{2+}]_i$ induced by compound 48/80 (5 μ g/mL), as assayed by Fluo-4 imaging in different extracellular solution. **(B and E)** The representative traces showing changes in $[Ca^{2+}]_i$ as measured by Fluo-4 imaging for C57BL/6j or *MrgprB2*^{-/-} mice peritoneal mast cells. **(C)** Compared with the normal external solution, calcium influx induced by compound 48/80 in the solution contain WSS was significantly reduced. (***p* < 0.01, 1 mg/mL WSS vs normal, *****p* < 0.0001, 3 mg/mL WSS vs normal, one-way ANOVA analysis followed by Dunnett's multiple comparisons test). **(F)** WSS had no effect on calcium influx induced by compound 48/80 in mast cells isolated from *MrgprB2*^{-/-} mice. Scale bar = 10 μ m.

Abbreviation: NS, no significant.

WSS Inhibited the Generation of Currents Mediated by MrgprB2 Receptor

The MrgprB2 receptor mediated whole-cell currents were evoked by compound 48/80 at different voltage commands. The whole-cell currents (baseline) are shown in **Figure 5A**; compound 48/80 (5 $\mu\text{g/mL}$) has increased voltage dependent currents (**Figure 5B**). Currents increased from -41.66 ± 4.11 pA to -209.0 ± 53.89 pA at -130 mV; Currents increased from 67.29 ± 7.71 pA to 1053 ± 176.4 pA at $+130$ mV (**Figure 5C**, $n=16$, $P = 0.0007$). 1 mg/mL WSS has inhibited the currents increased by compound 48/80 (**Figure 5D and E**). Currents increased from 118.9 ± 13.24 pA to 306.2 ± 77.17 pA at $+130$ mV (**Figure 5F**, $n=10$, $P = 0.024$); WSS has decreased current amplitude of variation (**Figure 5G**, $P = 0.004$). Furthermore, compound 48/80 has no effect on currents in mast cells come from MrgprB2^{-/-} mice (**Figure 5H**, $n=6$, $P = 0.893$) (**Figure 5I**, $n=6$, $P = 0.552$).

WSS Inhibited the Activation of MrgprB2/X2 Receptor

In addition, we also found that WSS could reduce the calcium influx in HEK-293 cells over-expressing MrgprB2 receptor (**Figure 6A–D**), or MrgprX2 receptor (**Figure 6E–H**), induced by compound 48/80 in a concentration-dependent manner. These results indicate that WSS could inhibit the function of MrgprB2/X2 receptor. MrgprB2: At the 300 $\mu\text{g/mL}$, 1 mg/mL, 3 mg/mL WSS, the inhibition was 53.47%, 68.92%, 77.42%; MrgprX2: At the 300 $\mu\text{g/mL}$, 1 mg/mL, 3 mg/mL WSS, the inhibition was 62.05%, 69.64%, 85.11%.

Discussion

Our experiment results show that WSS could ameliorate MrgprB2-mediated pruritus in vivo. Meanwhile, WSS also inhibited MrgprB2 receptor activity in vitro by intracellular calcium measurement and whole-cell currents recording experiments.

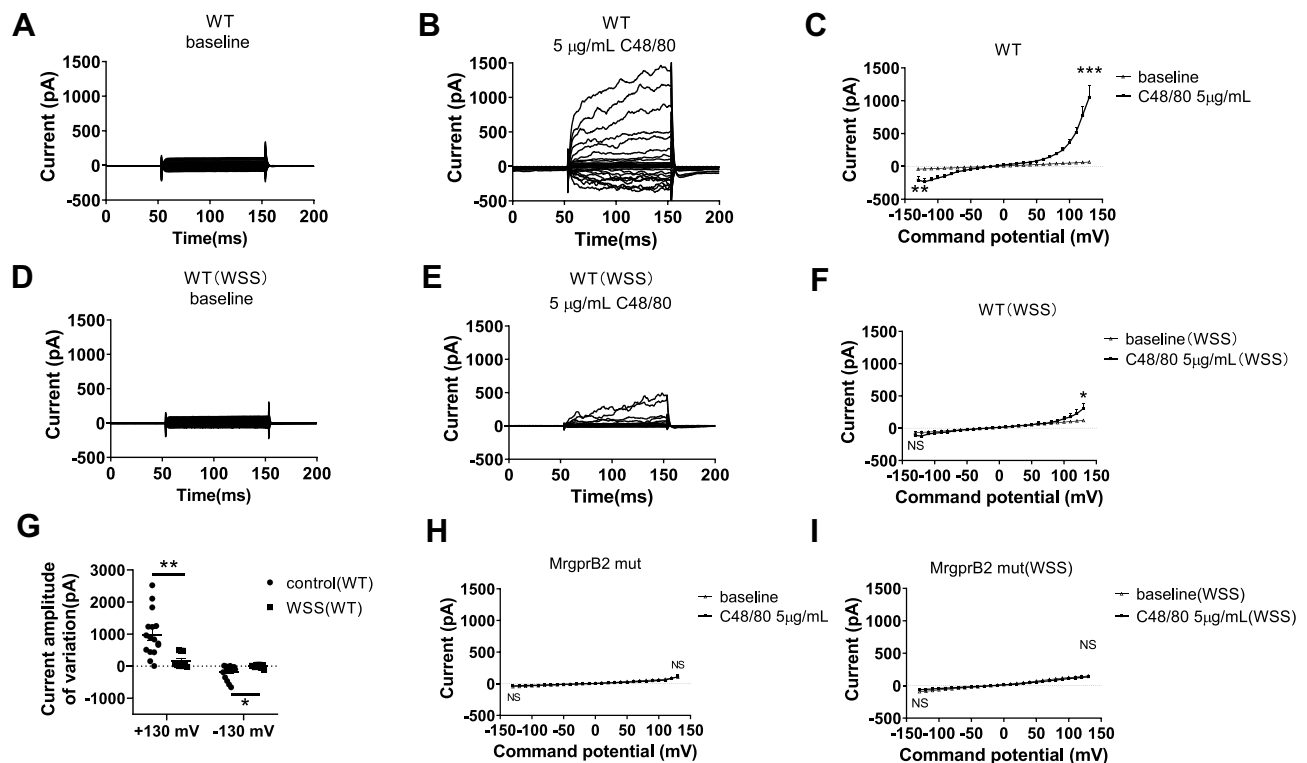


Figure 5 The effect of WSS on the voltage dependent currents induced by compound 48/80. (**A** and **B**) Under the normal extracellular solution and the extracellular solution containing WSS (**D** and **E**), typical whole-cell membrane current recorded from a mast cell, which clamped at a holding potential of -20 mV, and 100 ms voltage commands were applied from -130 to $+130$ mV in 10 mV step. (**C**) The voltage dependent currents induced by compound 48/80 for mouse peritoneal mast cells in the standard extracellular solution (** $p < 0.01$, *** $p < 0.001$, paired Student's *t*-test). (**F**) The voltage dependent currents induced by compound 48/80 for mouse peritoneal mast cells in the extracellular solution contain WSS (* $p < 0.05$, paired Student's *t*-test). (**G**) The current amplitude in different external solutions (* $p < 0.05$, ** $p < 0.01$, unpaired Student's *t*-test). (**H** and **I**) The voltage dependent currents for MrgprB2^{-/-} mast cells in different external solutions (paired Student's *t*-test, NS: no significant).

Abbreviation: NS, no significant.

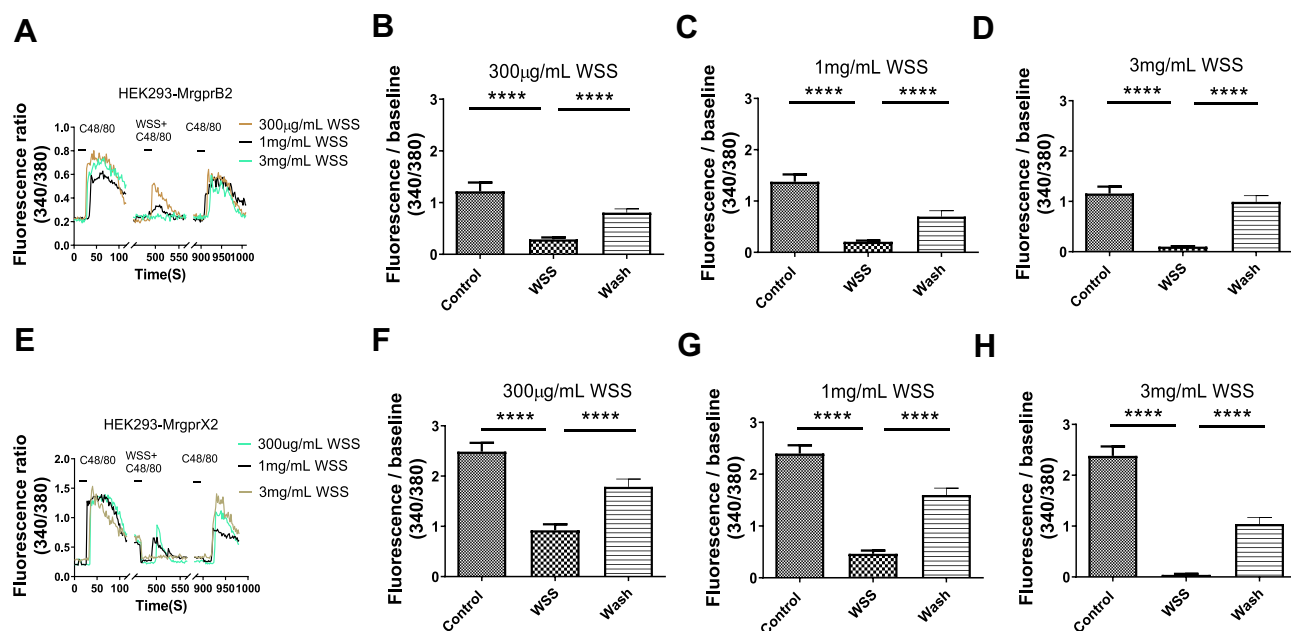


Figure 6 WSS could inhibit the calcium flux induced by compound 48/80 in MrgprB2/X2 over expressing HEK293 cells. **(A)** The representative traces showing changes in $[Ca^{2+}]_i$ in MrgprB2 over-expressing HEK293 cells. **(B–D)** 300 μ g/mL WSS, 1 mg/mL WSS, 3 mg/mL WSS inhibited the calcium flux in MrgprB2 over-expressing HEK293 cells induced by compound 48/80. **(E)** The representative traces showing changes in $[Ca^{2+}]_i$ in MrgprX2 over-expressing HEK293 cells. **(F–H)** 300 μ g/mL WSS, 1 mg/mL WSS, 3 mg/mL WSS inhibited the calcium flux in MrgprB2 over-expressing HEK293 cells induced by compound 48/80. (**** $p < 0.0001$, paired Student's t-test).

Pruritus are major characteristic of skin pathologies mainly including allergic contact dermatitis, but the underlying molecular mechanisms are not clear.¹⁹ ACDs are common skin disorders that are characterized by crusting and pruritus.²⁰ Allergic contact dermatitis affects up to 20% of adults and children in America.²¹ Although itch is the major symptom of dermatitis, which often leads to excessive scratching, tissue damage, the mechanisms that drive strong pruritus in ACD remain largely unknown.²² Mast cells are believed to play an important role in ACD pathogenesis.²³ A large number of studies have shown that Traditional Chinese Medicine can relieve itching in ACD by inhibiting mast cell degranulation. For example, ethanol extract of *Sanguisorbae Radix* inhibits mast cell degranulation.²⁴ Root bark of *Dictamnus dasycarpus* ACD via decreasing degranulation of mast cells by inhibiting p38 MAPK pathway.²⁵ *Astragalus membranaceus* and *Nepeta Tenuifolia* could inhibit skin hyperplasia and mast cell infiltration and ACD.²⁶ In our experiments, we found that WSS alleviated the pruritus in the ACD model and the pruritus induced by compound 48/80, a mast cell degranulation agent.²⁷ In addition, our results show that compared with C57BL/6J mice, pruritus induced by ACD is significantly reduced in mast cell-deficient Kit (W-sh) /Kit (W-sh) sash mice (sash) ([Supplementary Figure 6](#), $P < 0.05$). The antipruritic effect of WSS was decreased in Sash mice. These results indicated that the anti-pruritus mechanism of WSS is related to mast cells.

MrgprB2 receptor is mainly expressed on connective tissue-type mast cells in the skin and lungs, and can be activated by a broad range of cationic ligands, including basic secretagogues such as compound 48/80, substance P (SP), numerous FDA-approved drugs, and endogenous protein fragments.²⁸ MrgprB2 has been implicated in allergic and chronic inflammatory diseases, such as allergic contact dermatitis,¹⁶ imiquimod-related dermatitis.²⁹ In addition, our results show that compared with C57BL/6J mice, pruritus induced by ACD is significantly reduced in MrgprB2^{-/-} mice ([Supplementary Figure 6](#), $P < 0.05$). The number of mast cells was used as the index of pruritus in mice skin tissue with ACD.¹⁶ Compared with C57BL/6J mice, the number of mast cells in MrgprB2^{-/-} mice induced by compound 48/80 were significantly reduced ([Supplementary Figure 6](#), $P < 0.01$); The antipruritic effect of WSS was decreased in MrgprB2^{-/-} mice. These results indicated that the anti-pruritus mechanism of WSS is related to MrgprB2 receptor.

Furthermore, we also found that the calcium flux induced by compound 48/80 in mouse peritoneal mast cells or MrgprB2/X2 over expressing HEK293 cells could be inhibited by WSS. These results suggest that the antipruritic mechanism of WSS is by inhibiting the activity of MrgprB2. Therefore, we believe that the antipruritic effect of WSS

depends on the MrgprB2 receptor. *S. scandens* contained numerous valuable compounds, such as flavonoids, alkaloids and phenolic acids. Flavonoids mainly include quercetin, kaempferol, luteolin, hyperin, rutin, isorhamnetin and emodin, etc.¹ Quercetin, Kaempferol and luteolin could inhibit Mrgprx2-induced Ca²⁺ fluctuations.^{30–32} We speculated that the material basis of WSS inhibition of the MrgprB2 receptor was related to quercetin, kamanol and luteolin; however, we need further investigate which active component in WSS can inhibit MrgprB2 receptor.

Conclusion

In conclusion, our study revealed that the water extract of *S. scandens* (WSS) could ameliorate pruritus, and that the antipruritic mechanism of WSS is by inhibiting the activity of MrgprB2 receptor in mast cells.

Abbreviations

ACD, Allergic contact dermatitis; HEK293, Human Embryonic Kidney 293; H&E, Hematoxylin-eosin staining; HPLC, High performance liquid chromatography; MrgprB2, Mas-related G-protein-coupled receptor B2; *Senecio scandens* Buch.-Ham, *S. scandens*; WSS, Water extract of *S. scandens*.

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Author Contributions

All authors made a significant contribution to the work reported, whether that is in the conception, study design, execution, acquisition of data, analysis and interpretation, or in all these areas; took part in drafting, revising or critically reviewing the article; gave final approval of the version to be published; have agreed on the journal to which the article has been submitted; and agree to be accountable for all aspects of the work.

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Disclosure

The authors declare that there are no conflicts of interest in this work.

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