Abstract
Itch is a major indicator of psoriasis, but the underlying mechanisms behind this symptom are largely unknown. To investigate the neuronal mechanisms of psoriatic itch, we tested whether mice subjected to the imiquimod-induced psoriasis model exhibit itch-associated behaviors. Mice received daily topical applications of imiquimod to the rostral back skin for 7 days. Imiquimod-treated mice exhibited a significant increase in spontaneous scratching behavior directed to the treated area as well as touch-evoked scratching (alloknesis). To characterize this model, we measured the mRNA expression levels of pruritogens and itch-relevant receptors/channels using real-time reverse transcription PCR. The mRNA expression of MrgprA3, MrgprC11, and MrgprD decreased gradually over time in the dorsal root ganglion (DRG) cells. There was no significant change in the mRNA expression of TRPV1 or TRPA1 in DRG cells. TRPV4 mRNA expression was transiently increased in the DRG cells, whereas TRPM8 mRNA was significantly decreased. The mRNA expression levels of histidine decarboxylase and tryptophan hydroxylase 1, as well as the intensity of histamine and serotonin immunoreactivity, were transiently increased in the skin on day 2, returning to baseline by day 7. Histamine H1-receptor antagonists, chlorpheniramine and olopatadine, significantly inhibited spontaneous scratching on day 2, but not day 7. Neither chlorpheniramine nor olopatadine affected alloknesis on day 2 or day 7. These results may reflect the limited antipruritic effects of histamine H1-receptor antagonists on human psoriasis. The imiquimod-induced psoriasis model seems to be useful for the investigation of itch and its sensitization in psoriasis.

Keywords: Psoriasis, Chronic itch, Histamine, Alloknesis, Scratching

1. Introduction
Psoriasis is a chronic inflammatory skin disease affecting approximately 2% to 3% of the world’s population. Psoriasis is characterized by the presence of scaly skin plaques that display histological features including a thickened stratum corneum (hyperkeratosis), retention of nuclei within corneocytes (parakeratosis), and infiltration of inflammatory cells. The term psoriasis is derived from the Greek word “psora,” meaning “itching condition.” Pruritus has been reported in 60% to 90% of psoriasis patients. One of the aggravating factors of psoriatic itch is contact with clothes.

The pathogenesis of psoriasis has been investigated through the study of human and animal models. Repeated topical application of the drug imiquimod on the skin induces psoriasis-like inflammation in mice as well as humans and is considered a valid model for psoriasis. The pathogenesis of itch in psoriasis is unclear. A recent study found that there is no correlation between the severity of the disease and the intensity of pruritus in patients with psoriasis.

Recent studies have revealed a variety of molecules which are relevant to itch signal transmission. Inflammatory mediators including histamine and serotonin are known to act as pruritogens. However, histamine is less likely to be involved in psoriatic itch, as histamine H1-receptor (H1R) antagonists are reported to have limited or weak effects on psoriatic itch. Sensory neurons expressing Mas-related G-protein-coupled receptors (Mrgprs) mediate nonhistaminergic itch. Sensory neurons expressing Mas-related G-protein-coupled receptors (Mrgprs) mediate nonhistaminergic itch. MrgprD-expressing neurons exhibited enhanced excitability in a mouse model of allergic contact dermatitis. In addition, thermosensitive TRP channels including TRPV1, TRPA1, TRPV4, and TRPM8 are activated downstream of GPCRs to mediate or modulate itch.

In this study, we investigated whether an imiquimod-induced psoriasis mouse model exhibited spontaneous scratching and itch in response to light touch. We also characterized this model by measuring the gene expression levels of itch-relevant molecules. Further, we pharmacologically evaluated this model by testing the effects of H1R antagonists on spontaneous and touch-evoked itches.

2. Materials and methods
2.1. Imiquimod application
Experiments were performed using adult male C57BL/6J mice (21-28 g) under a protocol approved by the Temple University Animal Care and Use Committee. Fur on the rostral back was
trimmed with electric clippers and then removed with an electric shaver. The mice received a daily topical application of 62.5 mg Aldara cream (5% imiquimod) on the shaved back skin (2.5 × 2 cm) for 7 consecutive days. Control mice were treated similarly with a control vehicle cream (Vaseline Lanette cream, Fargon).

2.2. Behavioral tests

Mice were habituated twice to a Plexiglas recording arena for 60 minutes before testing. Twenty to 22 hours after each topical application, mice were videotaped from above for 60 minutes. The number of videotaped scratch bouts was counted by a trained observer blinded to the treatment condition. A scratch bout was defined as 1 or more rapid back-and-forth hind paw motions directed toward and contacting the treated area, ending with licking or biting of the toes or placement of the hind paw on the floor. Hind paw movements directed away from the treated area (e.g., ear scratching) and grooming movements were not counted. Following the 60-minute recording period, allodynia was assessed as follows: the mouse received 5 separate innocuous mechanical stimuli delivered using a von Frey filament (bending force: 0.7 mN) to 5 randomly selected sites along the border of the cream application area. The 0.7 mN von Frey filament was selected because it does not elicit scratch bouts in naive mice, and it was the minimum strength to elicit scratch bouts when delivered to skin surrounding the site of either histamine injection or dry skin treatment. The presence or absence of a positive response (a hindlimb scratch bout directed to the site of mechanical stimulation) was noted for each stimulus. The allodynia score was the total number of positive responses elicited by the 5 stimuli (0-5). In H1R antagonist experiments, each animal received a subcutaneous injection of either vehicle (isotonic saline), chlorpheniramine (10 mg/kg; Alfa Aesar, Ward Hill, MA), or olopatadine (10 mg/kg; Sigma-Aldrich, St. Louis, MO), 30 minutes before the behavior recording on either day 2 or day 7.

2.3. Real-time quantitative reverse transcription PCR

Cervical dorsal root ganglia (DRG) and skin samples were isolated from mice, submerged in RNAlater (Qiagen, Valencia, CA), and stored at – 80°C. Total RNA was extracted using Direct-zol RNA Mini Prep (Zymo Research, Irvine, CA). Reverse transcription of 0.5 μg total RNA was performed using the ProtoScript II First Strand cDNA Synthesis Kit (New England Biolabs, Ipswich, MA). Amplification of GAPDH cDNA was used for normalization. Real-time reverse transcription PCR was performed using the Fast Plus EvaGreen qPCR Master Mix (Biotium, Hayward, CA) on a 7500 Real-Time PCR System (Applied Biosystems, Grand Island, NY). Forty cycles of amplification were performed involving sequential denaturation at 95°C for 5 seconds, annealing at 65°C to 60°C for 5 seconds, and extension at 72°C for 33 seconds. Assays were validated using serial dilutions and confirmation of equal amplification efficiencies of the cDNA of interest and the GAPDH cDNA. Fold differences in expression were calculated using the comparative cycle threshold method by standardizing against GAPDH expression. All primer pairs are listed in Table 1. Gene expression levels of enzymes that are responsible for the synthesis of pruritogenic inflammatory mediators were measured in skin, whereas gene expression levels of receptors/channels involved in itch transduction (e.g., Mrgprs and transient receptor potential channels) were measured in DRG.

2.4. Immunohistochemistry

Animals were euthanized under sodium pentobarbital anesthesia, and the skin was acutely dissected. Skin was fixed in 4% paraformaldehyde followed by 30% sucrose, frozen in an optimal cutting temperature compound (Tissue-Tek; Sakura Finetek, Torrance, CA), and cut into 10-μm sections on a cryostat. The sections were incubated with 5% donkey serum and then immunostained with a rabbit histamine antibody (1:500; ImmunoStar Inc, Hudson, WI) and a rabbit serotonin antibody (1:1000; ImmunoStar Inc) at 4°C overnight, followed by incubation with the corresponding secondary antibody conjugated with Alexa Fluor 488 (1:300; Life Technologies Inc, Grand Island, NY) for 2 hours. Specificity of the primary antibodies was confirmed by preabsorption of the antibodies with either histamine (Sigma-Aldrich) or serotonin (Alfa Aesar). For triple staining, sections were additionally stained with rhodamine-labeled avidin (1:200; Vector Laboratories, Burlingame, CA), a marker for mast cells, chlorpheniramine (10 mg/kg; Alfa Aesar, Ward Hill, MA), or olopatadine (10 mg/kg; Sigma-Aldrich, St. Louis, MO), 30 minutes before the behavior recording on either day 2 or day 7.

Table 1

<table>
<thead>
<tr>
<th>Gene name</th>
<th>Primer sequence (5’ → 3’)</th>
</tr>
</thead>
<tbody>
<tr>
<td>GAPDH</td>
<td>FP: TCACCTGCGGTCTCTCAC</td>
</tr>
<tr>
<td></td>
<td>RP: GCAGAGATGTGACCCCTTT</td>
</tr>
<tr>
<td>Histidine decarboxylase (HDC)</td>
<td>FP: CSTGATATCTACGCCATGAGG</td>
</tr>
<tr>
<td></td>
<td>RP: ACTGCTGCAATGCDCAAGG</td>
</tr>
<tr>
<td>Tryptophan hydroxylase 1 (TPH1)</td>
<td>FP: ACAAAGACCATTCCTCDGAAAG</td>
</tr>
<tr>
<td></td>
<td>RP: TGACAGACGCTACATGATCTC</td>
</tr>
<tr>
<td>MrgprA3</td>
<td>FP: CTCAAGTTTCCCCCTCCAAAAGG</td>
</tr>
<tr>
<td></td>
<td>RP: CCAGAAGAATAACCATCCAGAA</td>
</tr>
<tr>
<td>MrgprC11</td>
<td>FP: ACTCTCTGCTACGGATCATTGA</td>
</tr>
<tr>
<td></td>
<td>RP: TGATTGCTGATTGGCTCTAGATA</td>
</tr>
<tr>
<td>MrgprD</td>
<td>FP: TTTTICAGSTACCTCTGCGC</td>
</tr>
<tr>
<td></td>
<td>RP: GCACATAGACAGGAGGGAGA</td>
</tr>
<tr>
<td>TRPV1</td>
<td>FP: CCAGAGCGAGAGATAGGTGCA</td>
</tr>
<tr>
<td></td>
<td>RP: TCAATTGCAAATGGTATGGTCG</td>
</tr>
<tr>
<td>TRPV4</td>
<td>FP: CGTGATACGGAGACAGAGAT</td>
</tr>
<tr>
<td></td>
<td>RP: GGATGCGGCCGTTAGGACAGAT</td>
</tr>
<tr>
<td>TRPM8</td>
<td>FP: AGACGTGCTCTACAGCTGAC</td>
</tr>
<tr>
<td></td>
<td>RP: GCTCTGGGCCATAACGACCTT</td>
</tr>
</tbody>
</table>

2.5. Histology

Skin was fixed in 4% paraformaldehyde, embedded in paraffin, and cut into 10-μm sections on a microtome. The sections were stained with hematoxylin and eosin (H&E) or toluidine blue using standard procedures. The number of mast cells per skin sample (total area of each image = 0.845 mm²) was quantified.
2.6. Data analysis

Between-group comparisons were made by 1-way or 2-way analysis of variance followed by the Tukey post-test. In all cases \( p < 0.05 \) was considered to be significant.

3. Results

Imiquimod or the control cream was applied on the rostral back skin for 7 consecutive days (Fig. 1A). Imiquimod treatment elicited erythema, scaliness, thickening, and infiltration of inflammatory cells (Fig. 1B–E), consistent with previous studies.\(^{37,38}\) Toluidine blue staining showed that there was no change in the number of mast cells in the control (8.8 ± 1.0 cells/mm\(^2\), \( n = 3; \) Fig. 1F arrows) vs the imiquimod-treated group (8.2 ± 2.1 cells/mm\(^2\), \( n = 3; \) Fig. 1G arrows).

With imiquimod treatment, counts of spontaneous scratch bouts were consistently higher than those in control mice and increased gradually over time (\( P < 0.001, F_{(7,94)} = 6.837; \) Fig. 2A). Vehicle-treated mice exhibited little spontaneous scratching, which increased only slightly over the tested time period. In imiquimod-treated mice, the alloknesis score increased significantly by day 2 and remained at a plateau through day 7 (\( P < 0.001, F_{(7,80)} = 23.111; \) Fig. 2B). Vehicle-treated mice exhibited an alloknesis score of 0 on all 7 test days.

To investigate whether histamine is upregulated in the skin of imiquimod-treated mice, mRNA expression of histidine decarboxylase, the catabolic enzyme of histamine synthesis, was determined in the skin. A transient increase in histidine decarboxylase mRNA expression was observed, with maximal expression on day 2 (Fig. 3H). Histamine immunoreactivity was detected in mast cells and keratinocytes on day 0 (Fig. 3A). Figure 3G shows a time course for fluorescence intensity of histamine in the epidermis of imiquimod-treated mice. The fluorescence intensity increased transiently on day 2 (Fig. 3B) and returned to the basal level by day 7 (Fig. 3C). Histamine fluorescence intensity did not change over time in the skin of vehicle-treated mice. Histamine immunoreactivity was completely abolished by preabsorption of primary antibody with histamine (data not shown). Seventy-nine percent (\( n = 4 \)) of histamine-immunoreactive cells were colocalized with avidin, a marker for mast cells.

Another pruritogenic inflammatory mediator, serotonin, is synthesized by tryptophan hydroxylase 1. A transient increase in tryptophan hydroxylase 1 mRNA expression was observed, with maximal expression on day 2 (Fig. 4H). Serotonin immunoreactivity was detected in mast cells as well as keratinocytes on day 0 (Fig. 4A). Figure 4G shows a time course for fluorescence intensity of serotonin in the epidermis of imiquimod-treated mice. The fluorescence intensity increased significantly on day 2 (Fig. 4B) and day 5 but returned to the basal level by day 7 (Fig. 4C). Serotonin fluorescence intensity did not change over time in the skin of vehicle-treated mice. Serotonin immunoreactivity was completely abolished by preabsorption of primary antibody with serotonin (data not shown).
Ninety-three percent (n = 4) of serotonin-immunoreactive cells were colocalized with avidin, a marker for mast cells. There was no significant change in the number of histamine- or serotonin-immunoreactive mast cells over time in imiquimod-treated mice (histamine, 4.79-8.89 cells/mm²; serotonin, 3.34-6.19 cells/mm²).

MicroRNA expression levels of MrgprA3 (Fig. 5A), MrgprC11 (Fig. 5B), and MrgprD (Fig. 5C) all decreased gradually over time in the DRG cells of imiquimod-treated mice. MrgprC11 mRNA showed a significant reduction on days 4, 5, and 7, whereas MrgprD mRNA was significantly reduced on day 5 alone.

There was no significant change in the mRNA expression of TRPV1 (Fig. 6A) or TRPA1 (Fig. 6B) over time in imiquimod-treated mice. A transient increase in TRPV4 mRNA expression was observed in the DRG cells, with maximal expression on day 2 (Fig. 6C). Interestingly, TRPV4 mRNA decreased significantly over time in the skin of imiquimod-treated mice (data not shown). TRPM8 mRNA in the DRG cells began to decrease on day 4 with maximal reduction on day 5 (Fig. 6D).

Finally, we investigated the effects of H1R antagonists on itch in this model. Mice were treated with either chlorpheniramine or olopatadine on day 2 of imiquimod treatment, when the histamine expression level had peaked, or on day 7 when the histamine expression level had returned to baseline. The number of spontaneous scratch bouts was significantly lower in H1R antagonist-treated mice compared with vehicle-treated mice on day 2 (Fig. 7A). In contrast, there was no significant difference in the number of spontaneous scratch bouts on day 7 (Fig. 7A).

Neither chlorpheniramine nor olopatadine inhibited allodynia on either day 2 or day 7 (Fig. 7B).

Figure 2. Time-dependent changes in scratch bouts and allodynia score in imiquimod-treated mice. (A) Spontaneous scratching was measured on pretreatment day 0 and imiquimod treatment days 1 to 7. Black dots (•) and white squares (○) show, respectively, imiquimod- and vehicle- (Vaseline Lanette cream) treated groups. Error bars are SEM (n = 6-8). *P < 0.05, significant difference from day 0, 2-way analysis of variance followed by Bonferroni test. (B) As in A for allodynia score.

Figure 3. Time-dependent changes in histamine in imiquimod-treated mice. (A-E) Typical examples of histamine (green) and DAPI (blue) expression in the skin treated with imiquimod on days 0 (A), 2 (B), and 7 (C), and vehicle on days 2 (D), and 7 (E). Scale bar indicates 500 μm. (F) Triple fluorescence immunohistochemistry for expressions of histamine (green), avidin (red), and DAPI (blue). Scale bar indicates 250 μm. (G) Immunofluorescence intensity of histamine in the epidermis was measured using Image J in imiquimod-treated mice. Error bars are SEM (n = 4). *P < 0.05, significant difference from day 0, one-way analysis of variance followed by Tukey test. (H) Levels of histidine decarboxylase mRNA in the skin were measured using quantitative reverse transcription PCR. Error bars are SEM (n = 6). *P < 0.05, significant difference from day 0, one-way analysis of variance followed by Tukey test.
Chronic itch occurs in the patients with a variety of pathological conditions, including inflammatory skin, systemic, and neurological diseases. Recent studies have revealed the neurotransmitters, receptors, and signal pathways involved in acute itch transduction. However, it is crucial for these findings to be tested in clinically relevant models that mimic specific pathological conditions. Although many animal models for itch in atopic dermatitis have been reported, the availability of animal models of itch related to other diseases is limited.

Previously, we reported that mouse models for itch associated with other skin diseases, such as atopic dermatitis and dry skin, exhibited spontaneous scratching and alloknesis, common manifestations of chronic itch. However, this study represents the first time in which itch has been characterized in a model of psoriasis. We found that the imiquimod-induced psoriasis model exhibited spontaneous scratching and alloknesis, and we characterized the expression of several principal itch-related mediators across the timeline of the model, providing a framework for future work to examine the neuronal mechanisms underlying psoriatic itch and its sensitization.

The number of mast cells did not change over time in the imiquimod-induced psoriasis model. Previous studies reported that there was no significant change in the number of mast cells in lesional vs nonlesional skin from psoriasis patients, though the number of mast cells was increased after elicitation of the Koebner phenomenon (the development of isomorphic...
pathological lesions in the traumatized, uninvolved skin of patients with psoriasis. Previous work has also found a pronounced infiltration of lymphocytes into lesional skin compared with nonlesional skin of patients with psoriasis. In this study, we similarly observed an increase in lymphocyte infiltration into imiquimod-induced psoriatic skin.

The interstitial histamine concentration in psoriatic skin has previously been analyzed using microdialysis. Two studies showed that dialysate histamine levels in psoriatic skin were higher than those in the skin of healthy subjects. However, a later study reported that similar levels of histamine were observed in the skin from healthy subjects, lesional psoriatic skin, and perilesional psoriatic skin using a novel rapid and sensitive chromatographic method. In this study, we found that the expression level of histamine peaked in the early phase and returned to the basal level in the late phase. Moreover, histamine H1-receptor antagonists inhibited spontaneous scratching in the early phase but not the late phase, suggesting that histamine might play a role in spontaneous itch only in the initial stage of psoriasis.

**Figure 6.** Time-dependent changes in TRP channels in imiquimod-treated mice. (A-D) Levels of TRPV1 (A), TRPA1 (B), TRPV4 (C), and TRPM8 (D) in the dorsal root ganglion cells were measured using quantitative reverse transcription PCR. Error bars are SEM (n = 6). *P < 0.05, significant difference from day 0, one-way analysis of variance followed by Tukey test.

**Figure 7.** Effects of histamine H1-receptor antagonists on ongoing (spontaneous) scratching and alloknesis score (touch-evoked scratching). (A) A histamine H1-receptor antagonist (chlorpheniramine or olopatadine) or vehicle (isotonic saline) was subcutaneously injected on day 2 or day 7 in imiquimod-treated mice. Mice were videotaped from above for 60 minutes. White, gray, and black columns show, respectively, vehicle, chlorpheniramine, and olopatadine groups. Error bars are SEM (n = 6-8). *P < 0.05, significant difference from vehicle-treated group, one-way analysis of variance followed by Tukey test. (B) As in (A), for alloknesis score.
Seronin is involved in itch associated with dry skin through the 5-HT2 and possibly 5-HT7 receptors. In the imiquimod-induced psoriasis model, we observed that serotonin was expressed by mast cells and keratinocytes. The expression levels of serotonin peaked in the early phase and returned to the basal level in the late phase. A similar chronological change in serotonin expression was observed in patients with psoriasis. 

Serotonin-positive expression was stronger in the stratum granulosum of the lesions in the progressive stage of psoriasis than in the static stage. Overall, serotonin does not seem to be a major itch mediator in the static phase of psoriasis.

It has recently been reported that Mrgrps present in subsets of small-diameter sensory neurons play an important role in itch. Unexpectedly, gene expression levels of MrgrpA3, MrgrpD, and MrgrpC11 were decreased in the imiquimod-induced psoriatic itch model. It would be of interest to test whether Mrgrp-expressing neurons exhibit hyperexcitability associated with an increase in sodium current in the imiquimod-induced psoriasis model, as observed in a contact dermatitis model.

There are very few studies regarding the role of TRP channels in psoriatic itch. The RNA-seq database indicates a down-regulation of TRPV4 mRNA in psoriatic skin. This study confirms that TRPV4 mRNA is downregulated in the skin of our imiquimod-induced psoriasis model. In contrast, TRPV4 gene expression was transiently increased in the DRG cells. This alteration paralleled the time course of serotonin expression in the skin. Previously, we showed that TRPV4 seems to function as a downstream transducer of the serotonin-induced itch signal. Increased gene expression levels of TRPV4 in DRG cells may be related to increased expression of serotonin. In contrast, the cold-sensitive ion channel TRPM8 is believed to be involved in itch relief and is negatively regulated following MrgrpA3 agonist stimulation. Interestingly, TRPM8 gene expression was decreased in the DRG cells of the imiquimod-induced psoriasis model. Altered TRPM8 expression may be involved in the enhancement of itch in the primary sensory neurons through disinhibition of itch signaling. TRPV1 and TRPA1 are required for histaminergic and nonhistaminergic itch, respectively. The proportion of TRPA1-expressing DRG neurons was increased in an interleukin-13–induced inflammatory skin model that exhibited spontaneous scratching. Neither TRPV1 nor TRPA1 gene expression levels significantly changed in the DRG cells of our psoriasis model.

This study reveals that imiquimod-treated mice represent a useful model to examine the neuronal mechanisms underlying itch and its sensitization in psoriasis for the following reasons: (1) Imiquimod-treated mice exhibit psoriasis-like skin conditions such as erythema, scales, epidermal thickness, and inflammatory cell infiltration, consistent with previous studies. (2) Imiquimod-treated mice exhibit an increase in spontaneous scratching, suggesting that chronic itch occurs in this model. In addition, imiquimod-treated mice exhibit alloknesis, which is a sign of itch sensitization and is observed in patients with psoriasis. (3) Histamine H1-receptor antagonists inhibited spontaneous scratching only on day 2 and not on day 7 and failed to inhibit alloknesis on either day 2 or day 7. This presumably reflects the limited antipruritic effects of H1R antagonists on psoriatic itch.

The imiquimod-induced psoriatic itch model should prove useful in future studies to examine the mechanisms underlying histaminergic (as seen in the early phase) and nonhistaminergic itch (as seen in the late phase) as well as the sensitization of itch signaling pathways in psoriasis.

Conflict of interest statement
Y.G. has served as a consultant for Pfizer. The remaining authors have no conflicts of interest to declare.

Acknowledgements
The authors thank Dr. Inami for preparation of CDNA samples. The work was supported by grants from the National Institutes of Health (AR063228) and Pfizer.

Article history:
Received 6 March 2016
Received in revised form 9 June 2016
Accepted 13 July 2016
Available online 19 July 2016

References


