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The Experience of Itch in Autistic Adults: An Online Survey

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Abstract

Background

The self-reported experience of itch for autistic people has not been studied. Anecdote, theory, and emerging research suggest that itch experience is different in autistic people, with negative consequences.

Methods

An online survey was completed by participants who self-categorised as diagnosed autistic (n=158), self-identified autistic (n=37) or not autistic (n=85). We asked about three categories of itch: spontaneous, provoked, and itch caused by a medical condition, with questions focussed on severity, timing and duration, and impact on daily activities.

Results

Across all categories of itch there was a pattern of greater severity, duration, and impact in diagnosed autistic people compared to non-autistic people. Those with self-identified autism largely fell between the diagnosed autistic and not autistic groups. Itch severity was associated with self-reported autistic traits. As expected, we found more dermatitis-related itch in autistic respondents, but this did not drive differences in spontaneous and provoked itch.

Conclusion

Findings suggest that the self-reported itch experience of autistic people is more severe, longer lasting and of greater daily relevance, differences that cannot be attributed to the higher prevalence of skin dermatitis found in autism. Future studies are required to understand itch experience within the broader picture of sensory differences, and to develop management strategies for what can be a highly morbid experience.

Community Brief

What was the purpose of this study?

Differences in sensory experiences in autistic people are well documented. However, the experience of itch for autistic people has been largely unresearched. Our study is the first to formally ask "Is itch experience different for autistic people?".

Why is this an important issue?

It is known that sensory differences can have a notable impact on autistic peoples' quality of life. It is also known that long-term itch can be detrimental to physical and mental health. What isn't known is whether autistic people suffer itch, with its negative impacts, more than non-autistic people.

What did the researchers do?

We distributed an international online survey that asked autistic and non-autistic adults about their experience of itch and the impact of itch on daily life. We asked about three types of itch: 1) spontaneous itch, 2) itch provoked by either contact with others, objects, hot and cold items, or itch experienced instead of pain, and 3) itch caused by medical conditions.

What were the results of the study?

Autistic adults reported significantly greater itch severity and itch-related impact on their daily life for all itch types. Heightened spontaneous and provoked itch experience in autistic people was not explained by higher rates of eczema-related itch.

What do these findings add to what was already known?

As the first of its kind, our study highlights that itch experience is different in autistic versus non-autistic adults.

What are potential weaknesses in the study?

Our study population was predominantly females over 30 years old, all of whom could access and complete an online survey. It will therefore be important to validate the findings in other autistic populations.

How will these findings help autistic adults now or in the future?

These new findings highlight the need to better understand the experience of itch for autistic people and are the foundation for beginning to translate this into support for autistic people who are being negatively impacted by itch.

Background [6,000]

Itch, "... a sensation in the skin that provokes the desire to scratch", can vary from a mild and transitory annoyance to a distressing and chronic experience.¹ When itch is experienced as distressing it has been associated with poorer quality of life, including depressed mood.² Despite differences in sensory experience between autistic and non-autistic people being commonly described,³ the experience of itch for autistic people is little explored. Itch can be described as an interoceptive sensation, which refers to sensations that convey information about one's "physiological condition".⁴ Interoceptive sensations are distinct from exteroceptive sensory modalities that inform people about the external world (such as discriminative touch). Differences between autistic and non-autistic people have been reported for the interoceptive somatosensory modalities of pain, temperature and affective touch,⁵⁻⁸ however, the interoceptive experience of itch for autistic people remains largely unexplored.

Somatosensory interoceptive modalities have a common neurobiology: peripheral, unmyelinated c-fibres synapsing in the superficial layers of spinal cord dorsal horn. They are also known to interact at multiple levels of the nervous system.⁹⁻¹⁸ Many theories of autistic perception predict the existence of interoceptive differences,^{19,20} and in light of the multimodal interoceptive differences found in autistic people,¹⁹ it would be surprising if itch experience were unaffected. It is important to note that exteroceptive sensory systems are also affected in autism, and a global mechanism might be driving interoceptive and exteroceptive changes in sensory sensitivity processing in autism (e.g. see Monday et al 2023²¹). Further, a recent study has suggested a genetic link between itch and autism in a knock-out mouse model of *CNTNAP2*.²²

Aside from neurobiological and mechanistic predictions, there is also limited but compelling evidence that itch experience is different for autistic people. Anecdotal, autobiographical reports from autistic adults make mention of itch, including Temple Grandin's accounts (Grandin, 1996, 2008).^{23,24} However, these allusions to a different, and often notably unpleasant, itch experience for autistic people

have not been followed-up with research studies. This is perhaps due to a passive assimilation of 'itch' under other terminology, such as 'tactile defensiveness',²⁵ 'self-injurious behaviour',²⁶ 'hypersensitivity' (archetypically to clothes labels)²⁷ and 'repetitive behaviours'²⁸. Indeed, the importance of efforts to formulate a standardised taxonomy to aid future autism sensory research have been highlighted elsewhere.²⁹ Kyriacou et al. (2021) presented qualitative data on the autistic experience of itch in relation to textiles, and highlighted some of the negative consequences of itch, such as self-injurious scratching.²⁷ However, textile-provoked *itch* hypersensitivity was not explored as a separate autistic phenomenon, and was nested under 'tactile defensiveness' and 'hyper-responsiveness' terminology.²⁷ The only study to explicitly quantify itch in autism is by Helt et al. (2021), who explored the phenomenon of 'itch contagion', which refers to itch that is elicited when watching others scratch.³⁰ Their baseline data confirmed that autistic children scratch more often than age-matched controls, and that scratching increased more for autistic children when exposed to images of others scratching.

The current study

Despite theoretical, physiological, and anecdotal accounts converging to suggest that the experience of itch may be different for autistic people, there has been very little exploration of self-reported experiences of itch. The aim of the current study was to address this need by conducting an international survey of itch experience in autistic and non-autistic people.

We used a bespoke online survey that drew on items in the 5-D Itch Scale³¹ to explore the experiences of three types of itch: provoked, spontaneous and medically induced. Provoked itch refers to an experience of itch that is provoked by an external stimulus. We included itch provoked by textiles, reflecting reports from autistic adults of problematic itch induced by clothing.²⁷ Based on existing understanding of the neurobiology of interoceptive somatosensory modalities (Craig, 2003; Marshall et al., 2019),^{4,32} we also asked about itch provoked by activation of other c-fibre modalities, such as contact

with hot or cold objects, social touch, or pain. For completeness, we also asked about itch provoked by contact with *any* object.

We additionally asked about itch occurring without provocation (i.e. spontaneous itch). There are suggestions that autistic people have difficulty attenuating interoceptive information,²⁰ which could increase vulnerability to spontaneous itch.

Our final category focussed on medical causes of itch, reflecting that itch can be caused by a wide range of medical conditions, including psoriasis, dermatitis (including eczema), liver disease and diabetes.³³ Autistic people have higher rates of poor physical health compared to non-autistic people,³⁴ and autistic people have repeatedly been shown to be of atopic dermatitis compared to non-autistic people.³⁵

To gain a breadth of insight, our survey asked respondents to reflect on the severity, timing and duration, and impact on daily life of each type of itch. Based on previous theoretical, physiological and anecdotal accounts, we predicted that the experience of each type of itch would be significantly more pronounced in autistic adults compared to non-autistic adults. We additionally predicted that people with dermatitis would be more susceptible to other forms of spontaneous and provoked itch (i.e., induced hypersensitivity) compared to those without dermatitis and that dermatitis would be over-represented in autistic adults.³⁶ This follows the predictions by Helt et al. (2021) who note that contagious itch is heightened in those with atopic dermatitis,^{30,37} and the predictions of Mishra et al. (2022) who speculate a common genetic cause for dermatitis and autism.²² We also collected self-report of autistic traits and general sensory differences, to explore their association with itch phenomena. We predicted that itch features would positively correlate with autistic traits and sensory features. Finally, in light of known associations between itch and anxiety and between anxiety and autism, an exploratory analysis was undertaken to look at the relationship between reported itch and anxiety diagnosis (Hollocks et al 2019; Sanders & Akiyama, 2018).^{38,39}

Methods

Participants

Our online survey was launched out of two autism research centres: Wales Autism Research Centre (WARC, UK) and Olga Tennison Autism Research Centre (OTARC, Australia). It was disseminated via means of research databases (academic and charitable, including Autistica: <https://www.autistica.org.uk/>), and via social media posts (TwitterTM and FacebookTM). Participants were invited to take part if they were over 18 years old and autistic or non-autistic. People who took part were invited to take part in a prize draw to win one of several cash prizes (GBP£50 or AUS\$90).

We received 280 fully completed responses. Based on participants' self-report we categorised them into three groups: Clinically diagnosed as autistic (AUT-D; $N = 158$); Self-identified autistic (AUT-S; $N = 37$); not autistic (N-AUT; $N = 85$). Age, sex and educational attainment did not differ across groups; white ethnicities were over-represented in AUT-D. There was a lower proportion of AUT-D to N-AUT in Australian participants compared to UK. Details of demographics are shown in Table 1, S5a & S5b. Our study was ethically approved by the Cardiff University School of Psychology Research Ethics Committee (EC.21.10.12.6416R), with reciprocal ethical approval from La Trobe University.

Materials

Demographic questions

We collected data on age, gender, sex, country or residence, ethnicity, and education level. Additional questions asked whether participants had a clinical diagnosis of autism or self-identified as autistic, as well as whether they have ever been diagnosed with a range of medical, neurodevelopmental and psychiatric conditions (See table S5c). Each participant was able to select multiple conditions. AUT-D had higher proportions of diagnosed anxiety (AUT-D: 74%; AUT-S: 46%; N-

AUT: 33%; χ^2 $p < 0.01$), depression (AUT-D: 66%; AUT-S: 35%; N-AUT: 24%; χ^2 $p < 0.01$) and asthma (AUT-D: 20%; AUT-S: 16%; N-AUT: 5%; χ^2 $p < 0.01$).

Itch Survey

Questions about itch experience were adapted from a previously validated measure used to evaluate medical causes of itch, the 5-D Itch Scale.³¹

Domains of Itch Measured. Our itch survey adapted four domains from the 5-D Itch Scale: Duration, Degree, Disability, and Distribution. Data on the distribution of itch (i.e., where on the body itch was experienced) was captured but will be reported elsewhere. We reformulated the remaining three domains as: severity (Degree), timing and duration (Duration), and impact (Disability), asking about a person's lifetime experience ("...at any point now or in the past...") rather than 'during the past 2 weeks' as in the 5-D Itch Scale. Most questions recorded responses on a Likert scale, with scales adjusted from the original where necessary to reflect our specific aims.

Severity of itch was measured by asking about the 'greatest intensity' of itch experienced, which ranged from 1 ('mild') to 4 ('very severe').

Timing and duration of itch was measured using three questions: the duration of the longest period of continuous itch ('longest episode'; ranging from 1 = 'less than six hours a day' to 5 = 'all day (or more)'); the longest duration of episodes ('longest cluster'; ranging from 1 = 'up to 1 week' to 6 = 'more than 5 years'); and how long ago the most recent episode was experienced ('most recent episode'; ranging from 1 = 'within the last week' to 6 = 'longer than 5 years ago').

The impact of current or prior itch on four different activities was measured. These were: sleep, leisure or sports activities, housework or errands, work or school activities. For the latter three activities the options ranged from 0 = 'never affects this activity' to 4 = 'always affects this activity', with a 'N/A'

response also available. For sleep, the choices ranged from 0 = 'never affected sleep' to 4 = 'delayed falling asleep and frequently wakes me up at night'.

Categories of Itch Measured. The survey questions were presented in a fixed order for three separate categories of itch: Spontaneous itch; Provoked itch; Medical itch. The itch categories were presented in a random order. For spontaneous itch, participants were asked: "Have you ever felt itchy (or wanted to scratch) for no obvious reason?", to which they could respond 'yes' or 'no'. If 'yes', they were then presented with the questions probing severity, timing and duration, and impact.

For provoked itch, participants were asked: "Have you ever felt itchy (or wanted to scratch) when something or someone touches (or injures) your skin?". If "yes", participants were asked to select the subtypes of provoked itch that reflected their experiences (Table 2). Multiple subtypes of provoked itch could be selected. They were then asked about each subtype in turn using severity, timing and duration, and impact questions. For most analyses, provoked itch types were coalesced and analysed as a single, parent category ('provoked itch'), alongside analyses of subtypes.

For itch of medical cause, participants were asked: "Have you ever felt itchy (or wanted to scratch) because of a medical condition? E.g. eczema". As for provoked itch, those identifying an itch of medical cause were asked to further specify its cause (Table 2). Multiple subtypes of medical itch could be selected. Subsequently, they were then asked about each subtype in turn using the severity, timing and duration, and impact questions. For the purposes of analysis, medical itch categories were coalesced into a single, parent category ('medical itch'), as well as into broad sub-categories (Table 2).

Throughout the survey, free-text questions were also asked about each itch (e.g. "Can you described this itch in more detail?"). These data will be analysed separately.

Other Measures

Additional validated measures were presented after completion of the itch questions. The 10-item version of the Autism Spectrum Quotient (AQ-10)⁴⁰ was administered to measure autistic traits. Scores can range from 1 to 10 with higher score indicating more autism traits. Clinical guidelines recommend score >5 should prompt referral for autism assessment. We opted for this over the full 50-question Autism Quotient (AQ)⁴¹ to reduce drop-out from questionnaire fatigue. The Glasgow Sensory Questionnaire (GSQ)⁴² was administered to explore general sensory features related to autism. This is a 42-item questionnaire covering 12 sensory domains. Within each domain there are two questions, one probing hyposensitivity and the other hypersensitivity. Scores can range from 0 to 168 with higher scores indicating more autism-related sensory features. Floor and ceiling effects for AQ-10 were present: highest (ceiling; AQ-10=10) values made up 15.1% of scores, whilst lowest (floor; AQ-10=0) values made up 3% of scores. GSQ did not have floor or ceiling effects.

Analysis and Statistics

Participant Characteristics

Age and performance on the AQ-10 and GSQ were compared across groups using one-way ANOVA, with post-hoc Tukey testing. Sex differences were evaluated with chi-square. Normal distribution of data was assessed using the Shapiro-Wilk test.

Itch Counts

Alongside the measures of severity, timing and duration, and impact, we wanted to explore the frequency with which itch was reported across groups. We calculated the number of participants

reporting each itch type (total participant count), *and* the number of itch types per participant (itch count per participant, inclusive of subtypes).

Total participant counts were compared across groups and evaluated with chi-square. Post-hoc paired chi-squares were performed where whole-group chi-squares were significant and Bonferroni corrected for multiple comparisons.

Itch counts per participant were non-normally distributed and therefore compared across groups using Kruskal-Wallis chi-square and post-hoc pairwise Wilcoxon signed rank sum test, Bonferroni corrected for multiple comparisons.

Comparisons of Itch Metrics Across Groups

Severity, timing and duration and impact were compared across the three groups, collapsed across all categories of itch and for separate parent itch categories and itch subtypes. The normality of itch severity, timing and duration, and impact scores could not be assumed.⁴³ Furthermore, Likert-scale type data are traditionally assessed with non-parametric tests. Therefore, Likert scores for the itch categories were compared using the omnibus Kruskal-Wallis chi-square. Post-hoc testing was performed using pairwise Wilcoxon signed rank sum test and Bonferroni corrected for multiple comparisons. Means were reported as we felt these better represent numerically small differences between groups for low-resolution Likert scales. Where itch types were coalesced into a parent category, within-participant maxima were used as the summary statistic. Those stating an absence of an itch type, at any point now or in the past, were given an 'anytime' itch severity of zero. Where an itch type was identified but a Likert scale metric was not completed, or an available 'N/A' option was selected, these were excluded from statistical evaluation.

Comparisons of Itch Metrics Across Itch Categories

Comparisons of severity, timing and duration, and impact measures across the three itch categories were performed using chi-square comparison of proportions and Kruskal-Wallis chi-squares. Detailed results of these comparisons can be found in the Supplemental Material.

Influence of Dermatitis

To ascertain the effect of dermatitis, known to be more prevalent in autistic people, a medical sub-category 'dermatitis' was formed. This combined the medical itch subtypes of eczema and contact dermatitis.

We compared dermatitis participant counts across groups and evaluated with chi-squares. Then, having excluded participants reporting dermatitis from our dataset, we re-ran count and severity, timing/duration and impact analyses for spontaneous and provoked forms of itch, and compared metrics with and without dermatitis sufferers with Mann-Whitney U test.

Influence of Anxiety and Other Diagnosed Conditions

We compared participant counts of diagnosed conditions across groups, evaluated with chi-square. Severity metrics, with and without diagnosed conditions, were compared with Kruskal-Wallis.

Relation Between Itch Metrics and Autistic and Sensory Traits

Correlations between the AQ-10 and GSQ with itch severity timing/duration and impact measures were evaluated with Spearman's rho (ρ).

Following this, manual hierarchical linear regression models were used to explore the relationships between itch characteristics and the AQ-10 and GSQ. The process was not automated, and its end-goal was *not* to produce the most-predictive model but rather to facilitate understanding of the associations within this explorative and highly-nested dataset. To encourage unbiased inference, every iteration of the hierarchical regression model is provided in Supplemental tables S11 and S12. The

variance inflation factor (VIF) was calculated for every independent variable within each model; none exceeded our cut-off of 5 and therefore no independent variables were excluded on the grounds of collinearity.

Two sets of manual hierarchical linear regression models were built. The first set of regression models used the AQ-10 or GSQ as the outcome variable. These were built iteratively, initially including all three itch categories as predictor variables and taking each type of itch characteristic in turn (i.e. severity, timing/duration, impact). At each iteration, predictor variables with p values that were below statistical significance (defined as $p < 0.05$) were removed whilst retaining and exploding into subtypes (where possible, e.g. provoked subtypes) other variables. At each stage, the explanatory power of the model was reassessed (adjusted R^2), with the aim of optimising this value. Our purpose in optimising R^2 was to provide a window into meaningful associations within our explorative dataset rather than to build the most-predictive model (e.g. to test on a future dataset). This is important to note as artificial inflation of R^2 is an inevitable consequence of using an iterative approach.⁴⁴ Again, because of the explorative nature of these models, model p -values were not corrected for multiple comparisons (all raw p -values can be found in Supplemental tables S11 and S12).

Combined models, taking the most predictive itch categories and characteristics from subordinate models were also evaluated. The effect of co-regressing group, AQ-10 (for GSQ models) and GSQ (for AQ-10 models) was also assessed and compared with models built with these regressors alone.

For all regression model independent variables, an absent value (i.e. because of omission or irrelevance) was substituted with a zero. Impact categories were summed to create a 'total impact' score for the purposes of regression.

Effect Sizes and Significance

Effect sizes were reported using Cramer's V for Chi-square, rank Eta-squared for Kruskal-Wallis, and rank biserial for Mann-Whitney's U. Significance was defined as $p < 0.05$. For post-hoc tests, adjustment of p-values for multiple comparisons were performed using the Bonferroni method.

Community Involvement

The idea and impetus for this study came was based on feedback from autistic participants we had previously worked with within other sensory studies. The charity Autistic UK aided in recruiting two autistic adults who advised on question relevance, appropriateness, and wording.

Results

Itch Counts

Comparison of Total Participant Counts Across Groups

Looking at the total participant counts across groups, itch was reported by more AUT-D participants compared to AUT-S and N-AUT ($\chi^2(2,265) = 81.00, p < 0.001$). All three itch categories had higher participant counts for AUT-D compared to N-AUT and omnibus evaluations were all significant. Post-hoc pairwise significant differences were demonstrated for spontaneous and provoked itch, but not for medical itch (See Table S1a). Results are summarised in Figure 1, with the number of counts additionally tabulated in Table S1b. Further analysis probed the participant counts for the subtypes of both provoked and medical itch. A similar pattern of participant counts across group was found for the provoked itch subtypes of material/clothing, objects, other's touch, and instead of pain. These provoked itch subtypes had significantly higher counts for the AUT-D group compared to the N-AUT group (Tables S6a and S7a). For medical itch subtypes, when coalesced into broad sub-categories the dermatological sub-category participant counts were different across groups with significantly higher counts in AUT-D

versus N-AUT post-hoc (Table S6b and S7b). The other broad medical itch sub-categories (Table 2) had no significant participant count differences across groups.

Provoked and Medical Itch Subtypes

The most commonly reported provoked itch subtype across all three groups was material/clothing (n=169), followed by other's touch (n=103) and objects (n=88), with fewer reporting itch instead of pain (n=62), contact with warm or cold objects (n=42) or other provoked causes (n=24). Of those with itch in response to contact with warm or cold objects, the majority were in response to either hot *or* cold objects (n=23), with fewer reporting itch in response to specifically warm (n=12) or cold objects (n=5) (Table 2).

Participants reported a total of 16 types of itch with medical cause (including 'other'), the commonest was eczema (n=83), followed by dry skin (n=68), sunburn (n=47), dandruff (n=43) and contact dermatitis (n=39) (Table 2).

Comparison of Itch Counts per Participant across Groups

The number of itch types reported per participant across all three categories were significantly different across AUT-D, AUT-S and N-AUT groups, with significantly more itch types per participant for AUT-D versus N-AUT (AUT-D, Md=5; AUT-S, Md=4, N-AUT, Md=2; $H(2) = 29.797$, $p < 0.001$; Post-hoc: AUT-D vs. N-AUT, $p < 0.001$).

Itch Severity

Comparison of Severity Across Groups

Omnibus evaluation of severity revealed significant differences in severity between groups for all itch categories (spontaneous: $H(2) = 44.39$, $\eta^2 = 0.16$, $p < 0.001$; provoked: $H(2) = 42.56$, $\eta^2 = 0.15$; p

<0.001; medical: $H(2) = 10.25$, $\eta^2 = 0.03$; $p < 0.006$), with significantly greater severity reported for AUT-D compared to N-AUT. Results are summarised in Figure 1 and Table S2.

For provoked itch subtypes, similar patterns of significance were found for the material/clothing, objects, other's touch and instead of pain subtypes (Table S8a). Of the medical itch subcategories, only dermatological itch showed significant differences across groups, with greater severity in AUT-D versus N-AUT (Table S8b).

Timing and Duration of Itch

Comparison of Timing and Duration Across Groups

Omnibus evaluation of timings and durations showed significant differences across groups. For spontaneous itch, the longest episode of itch was significantly different between groups ($H(2) = 13.07$, $\eta^2 = 0.05$, $p < 0.001$). For provoked itch, the longest episode of itch, the longest cluster of itch, and the most recent episode of itch were all significantly different (longest episode: $H(2) = 10.72$, $\eta^2 = 0.04$, $p = 0.005$; longest cluster: $H(2) = 10.24$, $\eta^2 = 0.04$, $p = 0.006$; most recent: $H(2) = 7.76$, $\eta^2 = 0.03$, $p = 0.021$), with significant differences in object and material subtypes for cluster length and most recent episode (Table S3 and S9a). In general, these reflected longer durations for AUT-D compared to N-AUT, although episodes were more recent for AUT-S. Results are summarised in Figure 2 and Table S3. Although omnibus evaluation did not reveal significant differences for the medical itch, the dermatological sub-category demonstrated significantly longer episodes for AUT-D compared to AUT-S (Table S9b).

Prevalence estimates of current itch (defined as 'within the last week') and chronic itch (defined as current itch of >4 weeks duration) are included in the Supplement (Table S13).

Impact of Itch on Everyday Life

Comparison of Impact of Itch Across Groups

There were significant differences for all types of impact and within all itch categories across groups, with the exception of the impact of medical itch on sleep. The pattern of significance was driven by itch having a significantly greater impact on those in the AUT-D group compared to the N-AUT group. Here, post-hoc comparisons were significant for all types of itch with the exception of the impact of medical itch on sleep and on work/school. The AUT-D group reported a significantly greater impact of itch compared to the AUT-S group on a more limited range of itch categories and impact types, most notably for housework where all itch categories had a significantly greater impact. There were only a few significant differences between the AUT-S and AUT-N group. Results are summarised in Figure 3 and Table S4.

Provoked itch subtypes largely replicated that of the provoked itch parent category (Table S10a). Notably, the dermatological medical itch subcategory showed significantly greater impacts for those in the AUT-D group, whilst no significant differences were found for other medical itch subcategories (Table S10b).

Exploring the effect of dermatitis on non-medical itch experience

There were 122 participants with dermatitis (formed by combining eczema with contact dermatitis). Counts were significantly different across groups (AUT-D 68/158, AUT-S 12/37, N-AUT 20/85; $\chi^2(2,280) = 9.36$, Cramer's $V=0.16$, $p=0.009$), with significantly more participants in the AUT-D group compared to the N-AUT group (post-hoc; 43% versus 23%; $p=0.012$).

When excluding dermatitis sufferers from analysis of spontaneous and provoked itch counts, significant differences across groups (i.e. greater numbers in AUT-D) remained for provoked itch but not

for spontaneous itch with a reduced effect size (Cramer's V, 0.16 with dermatitis excluded, previously 0.24).

When excluding dermatitis sufferers from analysis of spontaneous and provoked itch severity, significant differences remained for both ($H(2) = 26.27$, $\eta^2 = 0.14$, $p < 0.001$ and $H(2) = 28.16$, $\eta^2 = 0.15$, $p < 0.001$, respectively), with post-hoc tests showing greater severity in AUT-D.

Additionally, no significant differences were found between median severity scores for spontaneous or provoked itch. This was true for the whole sample, and for separate groups.

Exploring the Association of Anxiety and Depression on Itch Experience

Anxiety and depression were significantly associated with the presence of spontaneous itch (χ^2 , both $p < 0.05$); anxiety, depression and asthma were significantly associated with the presence of provoked itch (χ^2 , all $p < 0.01$); depression and asthma were significantly associated with the presence of medical itch (χ^2 , both $p < 0.01$). Severity was also significantly higher for those with an anxiety diagnosis for spontaneous and provoked itch (Kruskal-Wallis, $p < 0.001$), for those with depression for all itch types (Kruskal-Wallis, $p < 0.001$), and for asthma for provoked and medical itch types (Kruskal-Wallis, $p < 0.01$).

Exploring the Relation Between the Experience of Itch and Autistic Traits and Sensory Differences

The AQ-10 and GSQ differed significantly across group, with higher self-report of autistic traits and sensory differences in AUT-D versus AUT-S, and AUT-S versus N-AUT (Table 1). Severity, duration and impact variables for parent Itch categories positively correlated with AQ-10 and GSQ scores, with small but negative correlations for most recent episode between spontaneous itch and AQ-10 and for both spontaneous and provoked itch and GSQ. AQ-10 scores for AUT-S were normally distributed (Shapiro test, $p > 0.05$), whilst AUT-D and N-AUT were non-normal, with skews consistent with ceiling and floor effects (respectively). See Supplement, Tables S11 and S12 for detailed iterative model outputs.

The Relation Between Autistic Traits and Itch Features

Across all models, medical itch did not associate with autistic traits. Greater self-reported spontaneous and provoked itch severity were significantly associated with higher AQ-10 scores, producing a significant regression model that explained 27% of AQ-10 variance (Adj. $R^2(2,270) = 0.27$, $p < 0.001$).

Longer provoked itch episode durations were associated with higher scores on the AQ-10. The resultant significant model explained 10% of AQ-10 variance (Adj. $R^2(2,276) = 0.10$, $p < 0.001$).

Similarly, greater total impact from spontaneous itch and provoked itch caused by material/clothing were associated with greater AQ-10 scores and produced a significant model explaining 17% of AQ-10 variance (Adj. $R^2(2,276) = 0.17$, $p < 0.001$).

A combined model, including relevant severity, duration and impact variables from earlier iterations, isolated greater severity scores (spontaneous and provoked itch) as the only variable significantly associated with higher AQ-10 scores (Adj. $R^2(6,266) = 0.26$, $p < 0.001$). Their effects remained significant after controlling for the effect of group, and subsequent removal of spontaneous and itch severity variables led to a lower adjusted R^2 (i.e., itch severity added explanatory power beyond the group effect). Adding GSQ scores to the model removed the significant effect of spontaneous itch, however provoked itch remained significantly associated.

The Relation Between Sensory Differences and Itch Features

Across all models, medical itch did not associate with sensory differences. Greater severity of spontaneous and provoked itch (material/clothing, other's touch, objects or instead of pain) were significantly associated with higher total GSQ score, explaining nearly 50% of GSQ variance (Adj. $R^2(5,258) = 0.43$, $p < 0.001$).

Longer duration of provoked itch episodes and longer duration of provoked and spontaneous clusters of itch episodes were all significantly associated with GSQ score, together explaining 23% of GSQ variance (Adj. $R^2(3,273) = 0.23$, $p < 0.001$).

Total impact from spontaneous itch and itch provoked by both material/clothing and other's touch were significantly associated with GSQ, together explaining 40% of GSQ variance (Adj. $R^2(3,273) = 0.40$, $p < 0.001$).

A combined model, including relevant severity, duration and impact variables from earlier iterations, explained greater variance. Within this model, greater severity of spontaneous itch and itch provoked by objects and instead of pain, and greater total impact of spontaneous itch associated with higher levels of sensory differences (Adj. $R^2(4,261) = 0.43$, $p < 0.001$). The effect of these itch metrics remained significant after regressing out groups or AQ-10, and in both instances, the adjusted R-squared values were lower with itch measures excluded (i.e., itch severity and impact measures added predictive power beyond group).

Discussion

This is the first study to extensively and quantitatively explore the self-reported experience of itch in autistic people. We asked participants about their lifetime experience of itch across three categories: spontaneous, provoked and medical itch. Our questions targeted the feeling of itchiness rather than the behavioural act of scratching an itch. Compared to a non-autistic comparison group, autistic people experienced more severe and long-lasting itch, with a greater negative impact on daily life. Importantly, there was no evidence that itch caused by dermatitis, an itchy skin condition more prevalent in autistic populations, contributed to their heightened experience of non-medical itch.

Autistic participants had a heightened experience of provoked itch. The neurobiology of mechanically-induced itch (i.e., itch induced by a physical skin stimulus), is not fully understood but the role of c-fibres is increasingly recognised.^{45,46} There is evidence for mechanical itch-specific c-fibre

pathways as well as evidence for c-fibre pathways shared with other modalities, including pain and affective touch.^{14,47} In our study, itch caused by clothing and material was more common and severe in those with an autism diagnosis, as well as itch caused by another's touch or typically painful stimuli. This raises the possibility that itch-specific *and* non-specific pathways are sensitised in autism and may implicate a central location causing these differences.^{48,49} However, the limit to biology inferences an explorative questionnaire can make is quickly reached and future efforts exploring CNS structures, including the spinal cord (accessible using spinal cord fMRI), will be required to draw conclusions.

Similar to provoked itch, spontaneous itch also affected significantly more AUT-D participants at significantly greater severity, duration and impact compared to the non-autistic group. This finding is compatible with hypotheses proposing differences in attenuating interoceptive information in autism²⁰ but needs further work to confirm if this is the case.

As predicted, itch caused by medical conditions was experienced across all participant groups as more severe than spontaneous and provoked itch. Comparing the groups, it was more severe for those with a clinical diagnosis of autism and also had a greater impact on daily life, with the exception of sleep. No differences for timing and duration were found. Although the total number of people endorsing the different types of medical itch (participant counts) were significantly different by group, post-hoc pairwise tests were not significant. Itch caused by the sub-category dermatitis *did* affect significantly more AUT-D participants, in line with the literature and our hypothesis.³⁵ However, contrary to predictions,^{22,30} the severity of dermatitis itch did not influence severity scores of other co-existent non-medical itch types. Further, removing participants with a history of dermatitis did not affect significant group differences in itch severity for spontaneous or provoked itch, but did reduce the effect size for spontaneous itch group differences in participant counts. Finally, within our regression models, medical itch features were not associated with the degree of autistic traits reported by participants (AQ-10), nor their overall sensory differences (GSQ). As such, our prediction that autistic propensity to dermatitis

would exacerbate spontaneous and provoked itch experience was not reflected in the data. Instead, we suggest that medical itch features are largely independent of autistic traits and that provoked and spontaneous itch are distinct forms of itch that are experienced differently in autistic people. The data thus highlight that *both* medical itch and other forms of itch are important components of the autistic experience. Our results suggest that more autistic people will present with medical itch, with greater severity and impact compared to non-autistic people, and that this will negatively contribute to existing disadvantages faced by autistic people in healthcare.⁵⁰ It is therefore essential that heightened, medically-caused itch is researched *alongside* other itch-types in autism, and that relevant findings are made accessible to patients and healthcare professionals to guide their management.

Impacts on daily life were similarly and significantly overrepresented in our autistic participants with a clinical diagnosis. For example, over 75% of AUT-D with provoked or spontaneous itch reported affected sleep, compared with less than 50% of non-autistic respondents. Differences found in both the impact on sleep and other daily activities provide support that itch experience is not simply better attended to by autistic people but a highly relevant component of their wellbeing. It is known that poor sleep directly impacts physical and mental health and one's ability to work,⁵¹⁻⁵⁴ suggesting that unpleasant itch may have both direct and indirect effects on quality of life.

Our study allowed respondents to self-identify as autistic (AUT-S), that is they thought they were autistic but had not received a formal diagnosis. The scores for this group lay between the AUT-D and N-AUT groups for almost all measures, including AQ-10 and GSQ. We have limited information on the AUT-S participants and the reasons why they do not have (or have not sought) an autism diagnosis. It is likely many may be on a waiting list for diagnostic assessment, and some may have had a diagnostic assessment but not been diagnosed. Despite generally presenting with an intermediate itch presentation, a couple of subtle exceptions to this pattern were found. For spontaneous itch, participant counts and severity were the same (statistically) for those who self-identified they had autism compared

to those who were diagnosed, whilst for provoked itch they were significantly lower. The presence of severe provoked itch is therefore associated with autism diagnosis, whilst severe spontaneous itch is a more general marker of elevated autistic traits.

Across all participants, autistic traits and sensory features positively correlated with severity, duration and impact of spontaneous and provoked itch. This fits with our predictions made on the basis of higher rates of baseline scratching found as part of a study of itch contagion.³⁰ These significant correlations also persisted within groups. Together our studies present early corroborative objective evidence that itch is heightened in autistic populations. Helt et al.'s (2021) study also serves to highlight that scratching in autism, a phenomenon commonly encompassed under self-injury and 'stimming' behaviours, may sometimes be itch-related behaviour. Our study is unique in its probing of itch experience from self-report, rather than via a surrogate of scratching behaviour. However, we did not explore the possible overlap between itching and repetitive motor behaviours. Also, some respondents may have been drawing on their experiences of itch-related 'stimming' behaviours as much as of the underlying experience of itch sensation. The relationship between itch sensation, scratching, 'stimming' and self-injurious behaviour will be an important avenue for future research.

Epidemiology of Itch

Itch in the general population is a widespread phenomenon, but one with a poorly defined epidemiology.⁵⁵⁻⁵⁷ Our study was not epidemiological in its design, however some approximated prevalence estimates can be made for the purpose of comparison. Within our non-autistic participants, 55% experienced itch within the last week and 39% experienced current 'chronic' itch. These are both higher than most literature estimates at 8.5% and between 8-26%, respectively,⁵⁷⁻⁶¹ but are comparable to a study where efforts were made to focus on a general patient population.² However, the previous investigations of itch within the last week excluded "yes, a little" as a response option,⁵⁸ and the studies

that explored chronicity used longer durations (>6 weeks versus >4 weeks in ours). Not only might these methodological differences account for disparities, but our counts may also be greater because of the non-clinical nature of our study: itch asked about as a phenomenon not a 'problem' or 'complaint' might encourage people to report itch otherwise considered trivial. Indeed, a very high percentage of our participants (95%), both autistic and non-autistic, reported itch at some point during their lives. Of note, a condition termed sensitive skin syndrome (SSS) has a worldwide prevalence of at least 38%, close to our chronic itch prevalence estimate for our non-autistic sample.⁶² Its high prevalence may reflect the distress that unwanted skin sensations can cause. SSS is defined as "stinging, burning, pain, pruritus [itch] and tingling sensations ... in response to stimuli that normally should not provoke such sensations..." (p.5)⁶³ and is a 'diagnosis of exclusion', only considered once other causes of itch have been ruled-out.⁶⁴ SSS presents a syndrome worthy of exploration within autism. Beyond SSS, prevalence studies of 'incidental' itch of the type that we have measured (e.g., without medical cause and captured by our categories of provoked and spontaneous itch), are not readily found. However, the possible inflation of itch rates described above is unlikely to fully account for the negative impacts of itch that were endorsed by our participants.

Itch, Autism and Interoception

Many hypothetical models of autism predict that heightened interoceptive experiences contribute to sensory differences.¹⁹ Itch can be considered an interoceptive sensation as it provides the organism with information about the state of its own structures or tissues. Models propose that attention biased towards interoceptive information, occurring as a result of any of overamplification, reduced attenuation, or under-prediction, might variously lead to greater reliance on the immediate interoceptive signal. This is sometimes at the expense of integrating other sensations, including socially relevant exteroceptive cues.^{19,20} Thus, what for non-autistic people might be a trivial itch sensation, for autistic people might become a highly salient event that interferes with social interactions.

Itch and Anxiety

The itch literature shows a clear interplay between itch experience and anxiety.³⁹ Given that autistic people are more at risk of anxiety, and that our autistic participants reported heightened itch experience, it is interesting to consider whether anxiety might contribute part of the explanation. Within our demographics section, we asked about diagnosed anxiety. As expected, AUT-D had higher proportions of diagnosed anxiety and anxiety was significantly associated with the presence of all three categories of itch. Severity was also significantly higher for those with an anxiety diagnosis for spontaneous and provoked itch. Anxiety was not formally measured however, and this highlights an important area that warrants follow-up.

Limitations

In interpreting our findings, it is important to recognise that our participants were a self-selected cohort; personal experiences with itch or autism may have attracted them to the study and 'volunteer bias' may have inflated the self-report of itch experience. Also, as recruitment was uncontrolled there was a disproportionate response by UK nationals compared to Australian, which raises the potential for socio-cultural confounders. Those with an autism diagnosis had relatively late diagnoses, were early middle-aged, and predominantly female. Although age and sex did not differ significantly across groups and therefore does not invalidate our group comparisons, late diagnosed autistic people are not necessarily representative of the wider spectrum of autism.⁶⁵⁻⁶⁸ Further, completion of the survey necessitated a certain intellectual capacity, which also limited our ability to capture the heterogeneity of autism. The experience of itch for autistic children is also another important area for further study which will additionally provide insights into the development of unusual itch experiences in autistic people.

Our survey was built on a validated clinical itch scale.³¹ However, our study was explorative, and our survey was adapted and developed within this frame. The scope of our survey was necessarily constrained by considerations about completion time and participant fatigue (the primary motivation for using the abbreviated AQ). For example, we did not explore the contexts within which itch occurs. It might be that autistic people are more likely to experience itch, or experience itch as distressing, in particular environments. For example, we know that chronic itch and SSS are exacerbated by stress, negative emotions^{63,69} and anxiety.³⁹ Additionally, autistic children show increased scratching when viewing others scratching, which indicates sensitivity to the social context in the form of contagious itch.³⁰ There is the possibility that social context was a causative factor in 'spontaneous' itch reported in our study. Further research into context could be important for helping autistic people better manage itch in the future. Another unexplored avenue that warrants follow-up is how ethnicity affects itch experience⁷⁰ and how this might intersect with autism.

We also focussed on itch as a problem. However, it is known that some autistic people find components of their intense sensory experience pleasurable,^{71,72} and scratching an itch is often highly pleasurable.⁷³ Therefore, asking about itch-related pleasure will be important moving forwards.

Finally, itch features and their relationships with autistic traits (AQ-10) and sensory features (GSQ) were explored via iterative regression modelling, a technique that can artificially inflate the predictive power of the final model.⁴⁴ The robustness of inferences that can be drawn from the AQ-10 data are also hampered by ceiling effects. It is important to note however that these models were designed to be exploratory and the findings require replication and extension in future work. Recent concerns regarding the psychometric properties of the AQ-10 need also to be taken into account when evaluating the model outcomes.⁷⁴

Implications

Our evidence suggesting heightened itch is a relevant and impactful feature of the autistic experience has implications for supporting the health and well-being of autistic people. For example, existing psychoeducational information on sensory differences in autism could be expanded to include discussion of itch. Clinically, the experience of itch in autism is not well documented or recognised. It may be important going forward to consider ways of disseminating information about the autistic experience of itch to clinical services, including both autism services and medical services that treat itch. Related to this, existing interventions for the management of forms of chronic itch may be amenable to adaptation for the problematic itch experienced by autistic people. For example, web-based tools for managing itch in eczema and psoriasis include information about coping techniques and itch triggers that could be applicable.⁷⁵ Indeed, working with the autistic community and clinicians to make existing therapies more autism-specific may be an endeavour that autistic people consider an important future step. A challenge of any research on a subjective experience that cannot be objectively measured is that personal perceptions may vary. We did not include a definition of itch, or examples of itch, due to concern that a description that did not mirror a person's own definition may limit or bias their responses. It is important to note, therefore, that our data is underpinned by participants drawing on personal definitions and understanding of itch, and not on a singular definition. Currently, there has been no exploration of how autistic people describe itch and whether these descriptions can be considered similar or different to non-autistic people. These types of detailed insights and understanding would be best supported by qualitative methodology, such as interpretive phenomenological analysis.⁷⁶

Conclusion

Itch is a largely overlooked aspect of the autistic sensory experience. Our online survey has revealed that autistic adults report more itch, which is of both greater severity and longer duration, compared to non-autistic adults. Importantly, itch also had a greater impact on key elements of daily life, including leisure, housework, work and sleep. The group differences were more pronounced for provoked itch and spontaneous itch compared to medical itch. However, and importantly, differences in medical itch did not appear to drive the experiences of other types of itch. Our group of participants who self-identified as autistic generally fell between the other two groups. These novel findings are a clear indication of a need to better understand the experience of itch for autistic people and to translate this research into support for autistic people who are being negatively impacted by itch.

Authorship Contribution Statement

All Authors were involved in Conceptualising the study and developing its Methodology. GT undertook the initial Formal analysis, with subsequent input and advice from All Authors. The Original Draft was written by GT and has been extensively Reviewed & Edited by All Authors.

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Tables

Table 1

Participant Characteristics

		AUT-D	AUT-S	N-AUT	All groups
n		158	37	85	280
Age, Mean (SD)		43 (12.6)	44 (12.9)	41.4 (13.6)	42.6 (12.9)
Age at diagnosis, Mean (SD)*		36 (14.5)	NA	NA	NA
% female		71.5	64.9	70.6	70.4
AQ10 score†, Mean (SD)		8.1 (1.8)	6.6 (2.4)	2.8 (2.2)	6.3 (3.1)
GSQ score†, Mean (SD)		81 (23.4)	65.3 (24.1)	37.8 (21.2)	65.7 (29.8)
Location at time of participation§:	UK, n	127	26	54	207
	Aus., n	19	9	30	58
	Other, n	12	2	1	15

AQ-10, The Short Autism Quotient; GSQ, Glasgow Sensory Questionnaire; UK, United Kingdom; Aus., Australia.

AUT-D, diagnosed autistic; AUT-S, self-identified autism; N-AUT, not autistic.

* Slightly right-skewed (Shapiro-Wilk $p < 0.01$)

† ANOVA, $p < 0.01$; Post-hoc Tukey, all-comparisons, $p < 0.01$.

§ Chi-square, $p < 0.01$; Post-hoc comparison, $p < 0.01$ for AUT-D vs. N-AUT, including when limited to UK and Aus.

Table 2.

Medical and Provoked Itch subtypes, participant counts, and additional information

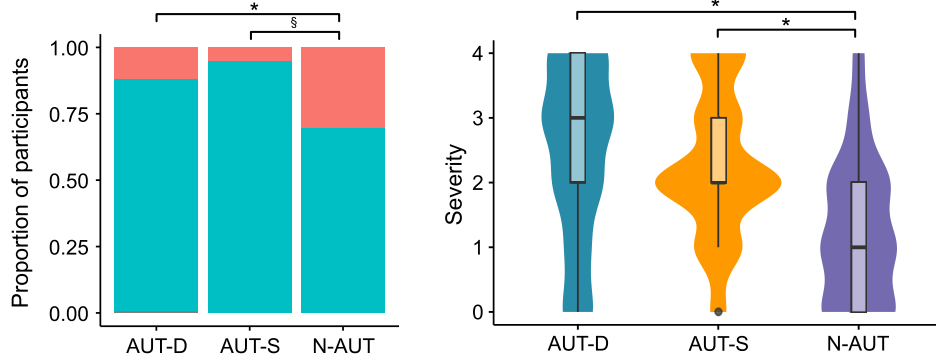
Itch type	Itch Subtype	n	Additional information
Provoked Itch			Subtype description in the survey
	Material/clothing	169	When certain materials or clothes touch your skin
	Objects	88	When your skin makes contact with certain objects
	Other's touch	103	When someone else touches you
	Temperature	42	When something hot/cold touches your skin
	Instead of pain	62	Do you sometimes feel itch instead of pain?
	Other	24	Another cause(s)
Medical Itch			Subcategories
	Eczema	83	Dermatological
	Psoriasis	21	Dermatological
	Contact dermatitis	39	Dermatological
	Dandruff	43	Dermatological
	Chronic pain	14	Systemic
	Athletes foot	34	Infection / Infestation
	Jaundice	0	Systemic
	Liver disease	3	Systemic
	Renal disease	0	Systemic
	Anaemia	8	Systemic
	Diabetes	9	Systemic
	Thyroid problems	2	Systemic
	Thrush	18	Infection / Infestation
	Sunburn	47	Dermatological
	Dry skin	68	Dermatological
	Urticaria / Hives	26	Dermatological
	Lice or scabies	9	Infection / Infestation
	Other	26	N/A

N = participant counts for itch subtypes

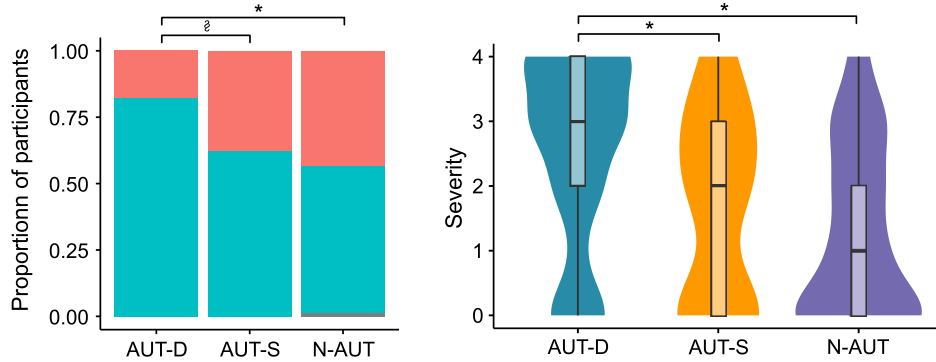
Figures

Figure 1. Proportion of participants reporting each itch type (participant counts) and severity of itch types by group

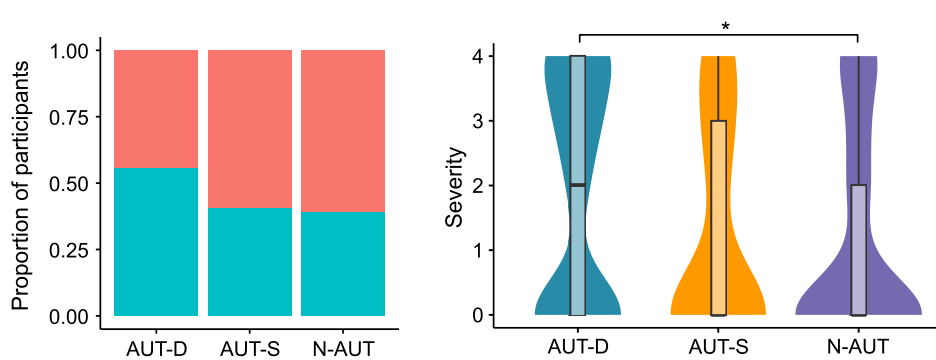
A. Spontaneous Itch



B. Provoked Itch



C. Medical Itch



Key: Yes (cyan), No (red), N/A (grey)

*, $p < 0.01$; §, $p < 0.05$; (both post-hoc)

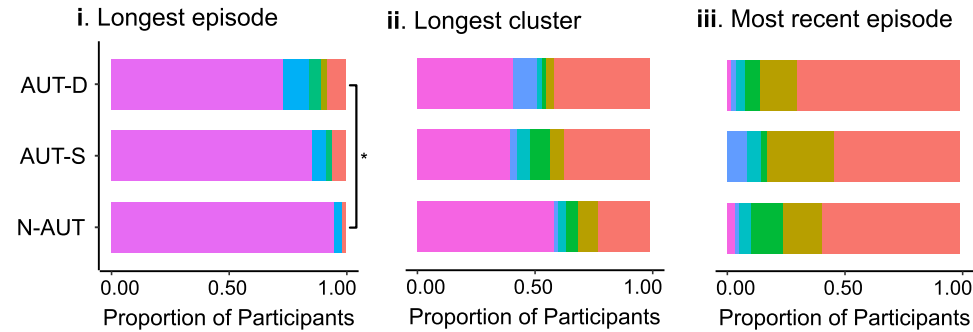
[Legend:]

AUT-D, diagnosed autistic; AUT-S, self-identified autism; N-AUT, not autistic.

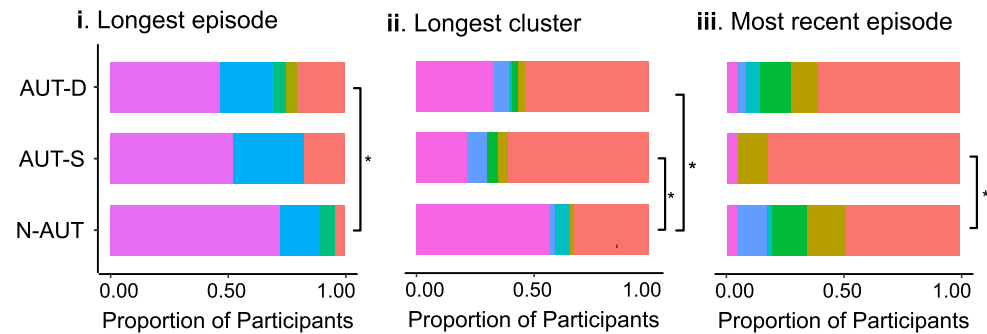
Severity was assessed by a 5-point scale with the following wording: 0 = no itching, 1 = mild, 2 = moderate, 3 = severe, 4 = very severe.

Figure 2. Duration and timing of itch categories by group

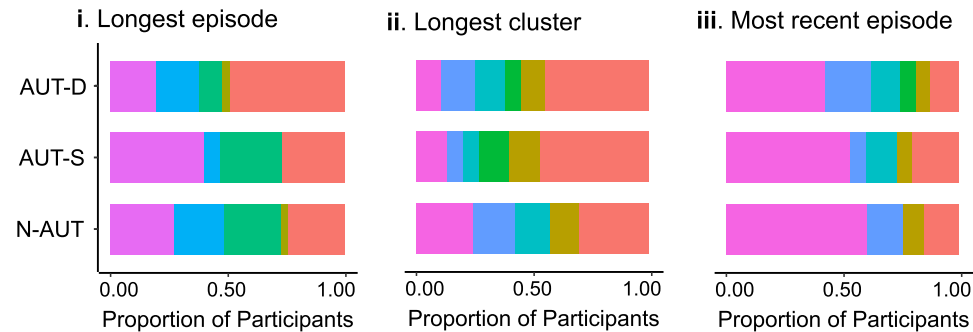
A. Spontaneous



B. Provoked



C. Medical



Key: Longest episode

- Red: All day (or more)
- Green: Between 18-23 hours per day
- Blue: Between 12-18 hours per day
- Yellow: Between 6-12 hours per day
- Pink: Less than 6 hours per day

Key: Longest cluster

- Red: More than 5 years
- Green: Up to 5 years
- Blue: Up to 1 year
- Yellow: Up to 6 months
- Pink: Up to 1 month
- Light Pink: Up to 1 week

Key: Most recent episode

- Red: Within the last week
- Green: Within the last month
- Blue: Within the last 6 months
- Yellow: Within the last year
- Pink: Within the last 5 years
- Light Pink: Longer than 5 years ago

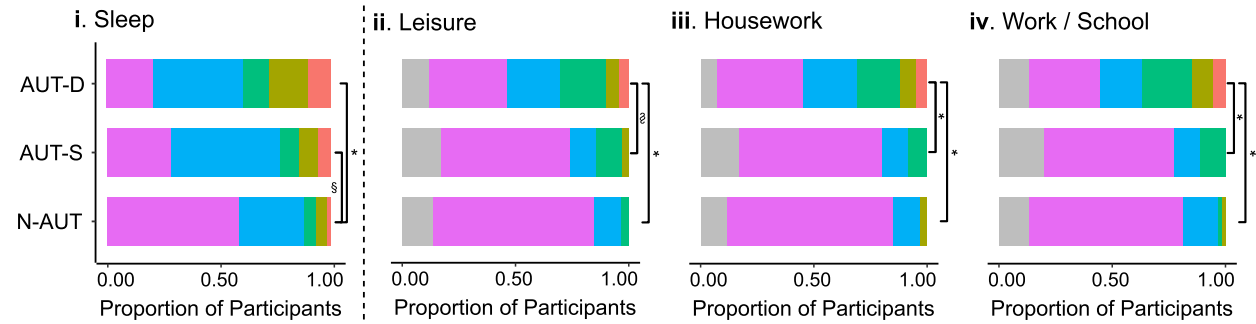
*, $p < 0.01$ (post-hoc)

[Legend:]

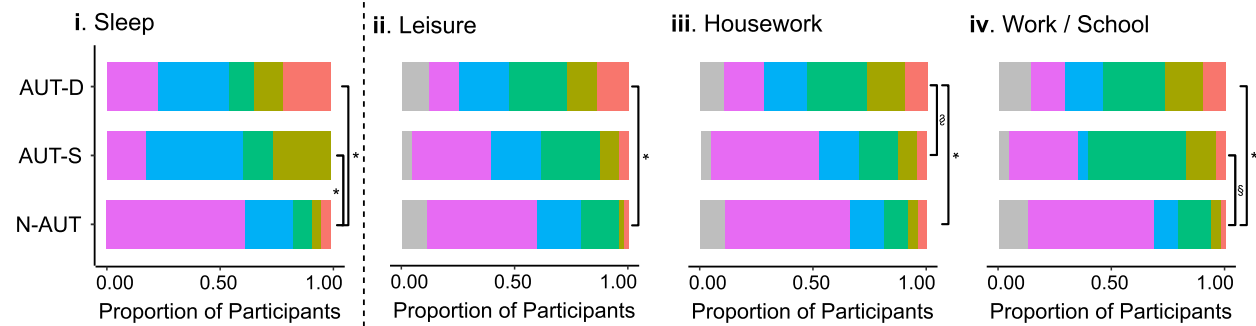
AUT-D, diagnosed autistic; AUT-S, self-identified autism; N-AUT, not autistic.

Figure 3. Impact of categories of itch on different aspects of everyday life by group

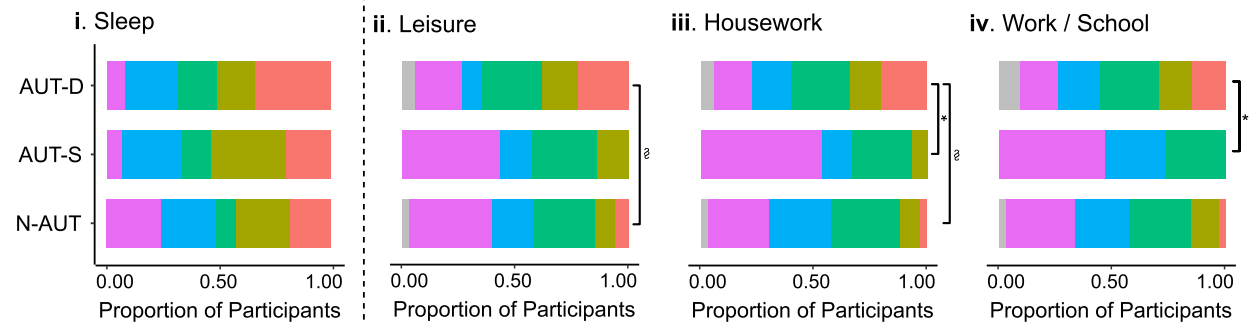
A. Spontaneous



B. Provoked



C. Medical



Key: Sleep

- Delayed falling asleep and frequently wakes me up at night
- Delayed falling asleep and occasionally wakes me up at night
- Frequently delayed falling asleep
- Occasionally delayed falling asleep
- Never affected sleep

Key: Leisure / Housework / Work / School

- Always affects this activity
- Frequently affects this activity
- Occasionally affects this activity
- Rarely affects this activity
- Never affects this activity
- N/A

*, $p < 0.01$; §, $p < 0.05$; (both post-hoc)

[Legend:]

AUT-D, diagnosed autistic; AUT-S, self-identified autism; N-AUT, not autistic.

Supplemental Material

Result tables: Parent itch categories & demographics

Table S1a.

Results summary table for itch participant counts across groups

Itch Category	χ^2	DF	V	p-value	Post-hoc comparisons (p-values)		
					AUT-D vs AUT-S	AUT-D vs N-AUT	AUT-S vs N-AUT
Spon.	17.44	2, 279	0.24	<0.001	1	0.002	0.015
Prov.	20.63	2, 279	0.26	<0.001	0.042	<0.001	1
Med.	7.40	2, 280	0.14	0.025	0.417	0.053	1

p < 0.05 in **bold**.

AUT-D, diagnosed autistic; AUT-S, self-suspected autism; N-AUT, not autistic.

Spon., spontaneous itch; Prov., provoked itch of any cause; Med., itch of any medical cause.

DF, degrees of freedom; V, Cramer's V.

Table S1b.

Counts of participants reporting itch parent categories by group

Itch category	AUT-D, n (%*)	AUT-S, n (%*)	N-AUT, n (%*)	All groups, n (%*)
Spontaneous	138 (88%)	35 (95%)	59 (69%)	232 (83%)
Provoked	130 (82%)	23 (62%)	47 (56%)	200 (71%)
Medical	88 (56%)	15 (41%)	33 (39%)	136 (49%)
<i>All itch categories</i>	<i>153 (97%)</i>	<i>35 (95%)</i>	<i>77 (91%)</i>	<i>265 (95%)</i>

* Percentage of group reporting type of itch.

AUT-D, diagnosed autistic; AUT-S, self-suspected autism; N-AUT, not autistic

Table S2.

Itch parent category severity across groups

Itch Category	AUT-D, M	AUT-S, M	N-AUT, M	H	DF	η^2	p-value	Post-hoc comparisons (p-values)		
								AUT-D vs AUT-S	AUT-D vs N-AUT	AUT-S vs N-AUT
Spon.	2.44	2.27	1.26	44.39	2	0.16	<0.001	0.777	<0.001	<0.001
Prov.	2.56	1.68	1.25	42.56	2	0.15	<0.001	0.002	<0.001	0.494
Med.	1.87	1.24	1.14	10.25	2	0.03	0.006	0.167	0.01	1

p < 0.05 in **bold**.

AUT-D, diagnosed autistic; AUT-S, self-suspected autism; N-AUT, not autistic.

Spon., spontaneous itch; Prov., provoked itch of any cause; Med., itch of any medical cause.

M, mean; DF, degrees of freedom; H, Kruskal-Wallis H statistic; η^2 , rank Eta-squared.

Table S3.

Itch parent category timings & durations across groups

Temporal Measure	Itch Category	AUT-D, M	AUT-S, M	N-AUT, M	H	DF	η^2	p-value	Post-hoc comparisons (p-values)		
									AUT-D vs AUT-S	AUT-D vs N-AUT	AUT-S vs N-AUT
Longest Episode	Spon.	0.61	0.34	0.10	13.07	2	0.05	0.001	0.402	0.002	0.362
	Prov.	1.29	1	0.47	10.72	2	0.04	0.005	1	0.005	0.278
	Med.	2.44	1.67	1.76	5.57	2	0.03	0.061	0.289	0.141	1
Longest Cluster	Spon.	2.4	2.49	1.71	5.77	2	0.02	0.056	1	0.060	0.260
	Prov.	2.95	3.43	1.83	10.24	2	0.04	0.006	1	0.015	0.021
	Med.	3.26	3.47	2.48	4.29	2	0.02	0.117	1	0.151	0.435
Most Recent Episode	Spon.	0.56	0.86	0.83	4.18	2	0.00	0.123	0.270	0.330	1
	Prov.	0.92	0.35	1.26	7.76	2	0.03	0.021	0.105	0.490	0.018
Episode	Med.	3.48	3.4	3.73	1.68	2	0.00	0.431	1	0.57	1

$p < 0.05$ in **bold**.

AUT-D, diagnosed autistic; AUT-S, self-suspected autism; N-AUT, not autistic.

Spon., spontaneous itch; Prov., provoked itch of any cause; Med., itch of any medical cause.

M, mean; DF, degrees of freedom; H, Kruskal-Wallis H statistic; η^2 , rank Eta-squared.

Table S4.

Itch parent category impact across groups

Impact	Itch Category	AUT-D, M	AUT-S, M	N-AUT, M	H	DF	η^2	p-value	Post-hoc comparisons (p-values)		
									AUT-D vs AUT-S	AUT-D vs N-AUT	AUT-S vs N-AUT
Sleep	Spon.	1.56	1.14	0.61	29.46	2	0.12	<0.001	0.242	<0.001	0.015
	Prov.	1.78	1.48	0.68	24.04	2	0.11	<0.001	1	<0.001	0.003
	Med.	2.45	2.33	1.88	4.08	2	0.02	0.130	1	0.136	0.946
Leisure	Spon.	1.12	0.52	0.22	31.09	2	0.15	<0.001	0.019	<0.001	0.363
	Prov.	1.92	1.23	0.76	27.10	2	0.14	<0.001	0.067	<0.001	0.327
	Med.	2.12	1.14	1.28	11.12	2	0.07	0.004	0.060	0.016	1
Housework	Spon.	1.11	0.34	0.25	33.11	2	0.15	<0.001	0.002	<0.001	1
	Prov.	1.80	1.00	0.74	25.50	2	0.13	<0.001	0.020	<0.001	1
	Med.	2.04	0.87	1.31	12.81	2	0.09	0.002	0.009	0.037	0.567
Work / School	Spon.	1.31	0.43	0.27	36.13	2	0.18	<0.001	0.001	<0.001	1
	Prov.	1.88	1.55	0.71	25.62	2	0.14	<0.001	0.896	<0.001	0.021
	Med.	1.91	0.80	1.31	11.48	2	0.08	0.003	0.009	0.099	0.488

p < 0.05 in **bold**.

AUT-D, diagnosed autistic; AUT-S, self-suspected autism; N-AUT, not autistic.

Spon., spontaneous itch; Prov., provoked itch of any cause; Med., itch of any medical cause.

M, mean; DF, degrees of freedom; H, Kruskal-Wallis H statistic; η^2 , rank Eta-squared.

Table S5a.

Further demographics: ethnicity

Ethnicity [§]	AUT-D, n (%*)	AUT-S, n (%*)	N-AUT, n (%*)
Asian	1 (1%)	3 (8%)	10 (12%)
Black	2 (1%)	1 (3%)	0 (0%)
Mixed	7 (4%)	2 (5%)	1 (1%)
White	148 (94%)	31 (84%)	72 (85%)
Other	0 (0%)	0 (0%)	2 (2%)

* Percentage of group.

AUT-D, diagnosed autistic; AUT-S, self-suspected autism; N-AUT, not autistic.

§, group comparison: $\chi^2(2) = 23.69$, p-value = 0.003.**Table S5b.**

Further demographics: education attainment

Education attainment	AUT-D, n (%*)	AUT-S, n (%*)	N-AUT, n (%*)
Some Primary Education	0 (0%)	0 (0%)	1 (1%)
Completed Primary Education	1 (1%)	0 (0%)	0 (0%)
Some Secondary Education	6 (4%)	0 (0%)	0 (0%)
Completed Secondary Education	32 (20%)	8 (22%)	10 (12%)
Bachelor's degree	58 (37%)	15 (41%)	26 (31%)
Master's degree	38 (24%)	9 (24%)	21 (25%)
PhD	7 (4%)	1 (3%)	17 (20%)
Other	16 (10%)	4 (11%)	10 (12%)

* Percentage of group.

AUT-D, diagnosed autistic; AUT-S, self-suspected autism; N-AUT, not autistic.

Table S5c.

Further demographics: medical, psychiatric and neurodevelopmental history

Condition	AUT-D, n (%*)	AUT-S, n (%*)	N-AUT, n (%*)	All groups
Allergy	44 (27.8%)	11 (29.7%)	24 (28.2%)	79
Anxiety	112 (70.9%)	15 (40.5%)	25 (29.4%)	152
Social anxiety	42 (26.6%)	6 (16.2%)	11 (12.9%)	59
Personality disorder	10 (6.3%)	0 (0%)	1 (1.2%)	11
Eating disorder	13 (8.2%)	2 (5.4%)	2 (2.4%)	17
Asthma	32 (20.3%)	6 (16.2%)	4 (4.7%)	42
ADHD / ADD	18 (11.4%)	4 (10.8%)	3 (3.5%)	25
Bipolar disorder	12 (7.6%)	1 (2.7%)	3 (3.5%)	16
Cerebral palsy	2 (1.3%)	0 (0%)	0 (0%)	2
Depression	104 (65.8%)	13 (35.1%)	20 (23.5%)	137
Developmental Delay	2 (1.3%)	0 (0%)	0 (0%)	2
Dyspraxia	16 (10.1%)	0 (0%)	1 (1.2%)	17
Dyslexia	9 (5.7%)	1 (2.7%)	2 (2.4%)	12
Hearing impairment	8 (5.1%)	2 (5.4%)	3 (3.5%)	13
Intellectual Disability	3 (1.9%)	0 (0%)	0 (0%)	3
OCD	16 (10.1%)	2 (5.4%)	2 (2.4%)	20
Orthopedic impairment	8 (5.1%)	0 (0%)	2 (2.4%)	10
Schizophrenia	2 (1.3%)	0 (0%)	0 (0%)	2
Seizure disorder, epilepsy	4 (2.5%)	1 (2.7%)	0 (0%)	5
Speech impairment	3 (1.9%)	0 (0%)	0 (0%)	3
Tourette's syndrome	0 (0%)	0 (0%)	0 (0%)	0
Visual Impairment	9 (5.7%)	2 (5.4%)	3 (3.5%)	14
Other	29 (18.4%)	6 (16.2%)	7 (8.2%)	42
None	7 (4.4%)	8 (21.6%)	31 (36.5%)	46

*percentage of group

Comparison of Itch Metrics Across Itch Categories

Comparisons of severity, timing and duration, and impact measures across the three itch categories were performed using chi-square comparison of proportions and Kruskal-Wallis chi-squares. These are conservative methods that do not benefit from the paired nature of the data (e.g. a single participant with both spontaneous and provoked itch metrics). However, the method is valid and was used for simplicity and comprehensibility (Looney & Jones, 2003).

Comparison of Participant Counts Across Itch Categories

Across all participants, 265 (95%) reported at least one form of itch. The proportions of respondents reporting spontaneous, provoked and medical itch were significantly different (83%, 71% and 49%, respectively; χ -square = 77.95, df = 2, p-value < 0.001).

Comparison of Severity Across Itch Categories

Severity scores were significantly different across the three itch categories ($H(2) = 43.82$, $\eta^2=0.07$, $p<0.001$) with significantly higher severity for medical versus provoked ($p=0.002$) and spontaneous itch ($p<0.001$), and significantly higher severity for provoked versus spontaneous itch ($p<0.001$). Significant differences remained when the sample was limited to AUT-D only, however post-hoc comparison of medical and provoked itch was no-longer significantly different.

Comparison of Timing and Duration Across Itch Categories

Longest episode length was significantly different across itch categories ($H(2) = 125.01$, $\eta^2=0.22$, $p<0.001$) with significantly shorter episodes for spontaneous versus provoked, and provoked versus medical (all pairs, $p<0.001$). Longest cluster length was also significantly different across itch categories ($H(2) = 14.73$, $\eta^2=0.02$, $p<0.001$) with a significantly longer clusters for medical itch versus spontaneous

itch ($p < 0.001$). Timing of most recent episode was also significantly different across itch categories ($H(2) = 179.69$, $\eta^2 = 0.32$, $p < 0.001$) with more recent episodes for spontaneous and provoked forms of itch compared to medical itch (both $p < 0.001$). The same pattern of significant differences remained when analyses were restricted to only AUT-D.

Comparison of Impact Across Itch Categories

Impacts on sleep, leisure, housework and school/work were all significantly different across the three itch categories ($H(2) = 47.077$, $\eta^2 = 0.08$; $H(2) = 54.146$, $\eta^2 = 0.10$; $H(2) = 49.643$, $\eta^2 = 0.09$; $H(2) = 35.38$, $\eta^2 = 0.07$; respectively; p (all) < 0.001), with significantly greater impact on sleep for medical itch versus spontaneous and provoked itch (both $p < 0.001$). There was also significantly less impact on leisure, housework and work / school for spontaneous itch versus provoked itch and medical itch (all $p < 0.001$), but no significant difference between spontaneous itch and medical itch. When limited to only AUT-D participants, the patterns of significant differences remain the same.

Results tables: Exploded subtypes**Table S6a.**

Counts of participants reporting itch: Provoked and medical subtypes

Parent itch category	Itch subtype	AUT-D, n (%*)	AUT-S, n (%*)	N-AUT, n (%*)	<i>All groups,</i> <i>n (%*)</i>
Provoked	Materials	116 (34%)	21 (35%)	32 (37%)	169 (60%)
	Objects	65 (19%)	11 (18%)	12 (14%)	103 (37%)
	Other's touch	74 (22%)	9 (15%)	20 (23%)	42 (15%)
	Temperature	27 (8%)	7 (12%)	8 (9%)	62 (22%)
	Instead of pain	43 (13%)	9 (15%)	10 (11%)	24 (9%)
	Other	16 (5%)	3 (5%)	5 (6%)	83 (30%)
	Medical	Eczema	58 (19%)	12 (24%)	13 (13%)
Psoriasis		15 (5%)	2 (4%)	4 (4%)	39 (14%)
Dermatitis		26 (9%)	2 (4%)	11 (11%)	43 (15%)
Dandruff		26 (9%)	8 (16%)	9 (9%)	14 (5%)
Chronic Pain		9 (3%)	2 (4%)	3 (3%)	34 (12%)
Tinea		20 (7%)	5 (10%)	9 (9%)	0 (0%)
Jaundice		0 (0%)	0 (0%)	0 (0%)	3 (1%)
Liver		3 (1%)	0 (0%)	0 (0%)	0 (0%)
Renal		0 (0%)	0 (0%)	0 (0%)	8 (3%)
Anaemia		7 (2%)	1 (2%)	0 (0%)	9 (3%)
Diabetes		5 (2%)	1 (2%)	3 (3%)	2 (1%)
Thyroid		1 (0%)	0 (0%)	1 (1%)	18 (6%)
Candida		13 (4%)	0 (0%)	5 (5%)	47 (17%)
Sunburn		34 (11%)	6 (12%)	7 (7%)	68 (24%)
Dry Skin		45 (15%)	8 (16%)	15 (15%)	26 (9%)
Urticaria		17 (6%)	2 (4%)	7 (7%)	9 (3%)
Lice		4 (1%)	1 (2%)	4 (4%)	26 (9%)
Other		19 (6%)	0 (0%)	7 (7%)	0 (0%)

Table S6b.

Counts of participants reporting itch: Medical sub-categories

Medical itch sub-category	AUT-D, n (%*)	AUT-S, n (%*)	N-AUT, n (%*)	<i>All groups,</i> <i>n (%*)</i>
Dermatological	83 (53%)	15 (41%)	29 (34%)	127 (46%)
Infection / Infestation	32 (20%)	5 (14%)	12 (14%)	49 (17%)
Systemic	21 (13%)	4 (11%)	6 (7%)	31 (11%)

* Percentage of group reporting type of itch.

AUT-D, diagnosed autistic; AUT-S, self-suspected autism; N-AUT, not autistic

Table S7a.

Provoked and medical itch subtype participant counts across groups: Summary of statistics

Parent itch category	Itch subtype	χ^2	DF	V	p-value	Post-hoc comparisons (p-values)		
						AUT-D vs AUT-S	AUT-D vs N-AUT	AUT-S vs N-AUT
Provoked								
	Material	28.91	2,279	0.31	<0.001	0.218	<0.001	0.263
	Objects	18.38	2,279	0.24	<0.001	0.822	<0.001	0.244
	Other's touch	15.39	2,279	0.22	<0.001	0.063	0.002	1.000
	Temperature	2.95	2,279	0.06	0.228	2.944	0.484	0.756
	Instead of pain	7.55	2,279	0.14	0.023	2.639	0.030	0.434
	Other	1.23	2,279	0.00	0.541	2.845	1.172	1.000
Medical								
	Eczema	12.31	2,280	0.19	0.002	1.000	0.002	0.168
	Psoriasis	2.10	2,280	0.02	0.351	1.000	0.847	1.000
	Dermatitis	3.15	2,280	0.06	0.207	0.429	1.000	1.000
	Dandruff	2.75	2,280	0.05	0.253	1.000	0.880	0.547
	Chronic_Pain	0.56	2,280	0.00	0.755	1.000	1.000	1.000
	Tinea	0.30	2,280	0.00	0.862	1.000	1.000	1.000
	Jaundice	NA						
	Liver	2.34	2,280	0.03	0.310	1.000	1.000	0.000
	Renal	NA						
	Anaemia	3.91	2,280	0.08	0.141	1.000	0.351	1.000
	Diabetes	0.06	2,280	0.00	0.971	1.000	1.000	1.000
	Thyroid	0.54	2,280	0.00	0.765	1.000	1.000	1.000
	Candida	3.43	2,280	0.07	0.180	0.450	1.000	0.938
	Sunburn	6.99	2,280	0.13	0.030	1.000	0.042	0.960
	Dry_Skin	3.69	2,280	0.08	0.158	1.000	0.261	1.000
	Urticaria	1.18	2,280	0.00	0.554	1.000	1.000	1.000
	Lice	0.88	2,280	0.00	0.645	1.000	1.000	1.000
	Other	5.31	2,280	0.11	0.070	0.167	1.000	0.508

Table S7b.

Medical itch sub-category participant counts across groups: Summary of statistics

Parent itch category	χ^2	DF	V	p-value	Post-hoc comparisons (p-values)		
					AUT-D vs AUT-S	AUT-D vs N-AUT	AUT-S vs N-AUT
Dermatological	7.96	2,280	0.15	0.019	0.775	0.027	1.000
Infection / Infestation	1.91	2,280	0.00	0.385	1.000	0.938	1.000
Systemic	2.18	2,280	0.03	0.336	1.000	0.623	1.000

p < 0.05 in **bold**.

AUT-D, diagnosed autistic; AUT-S, self-suspected autism; N-AUT, not autistic.

Spon., spontaneous itch; Prov., provoked itch of any cause; Med., itch of any medical cause.

DF, degrees of freedom; V, Cramer's V.

Table S8a.

Provoked and medical itch subtype severities across groups: Summary of statistics

Parent itch category	Itch subtype	AUT-D, M	AUT-S, M	N-AUT, M	H	DF	η^2	p-value	Post-hoc comparisons (p-values)		
									AUT-D vs AUT-S	AUT-D vs N-AUT	AUT-S vs N-AUT
Provoked											
	Material	0.86	1.49	2.24	42.06	2	0.15	<0.001	0.022	<0.001	0.07
	Objects	0.3	0.54	1.03	18.62	2	0.06	<0.001	0.304	<0.001	0.185
	Other's touch	0.4	0.49	1.18	19.07	2	0.06	<0.001	0.026	<0.001	1
	Temperature	0.21	0.35	0.46	2.57	2	0	0.277	1	0.424	0.484
	Instead of pain	0.24	0.57	0.74	8.09	2	0.02	0.017	1	0.014	0.231
	Other	0.13	0.16	0.25	0.54	2	0	0.762	1	1	1
Medical											
	Eczema	0.41	0.97	1.18	13.1	2	0.04	0.001	1	0.001	0.082
	Psoriasis	0.12	0.11	0.23	2.1	2	0	0.351	1	0.573	1
	Dermatitis	0.33	0.11	0.49	3.55	2	0.01	0.17	0.209	1	0.624
	Dandruff	0.18	0.49	0.39	3.57	2	0.01	0.168	1	0.434	0.193
	Chronic_Pain	0.06	0.19	0.1	0.29	2	0	0.866	1	1	1
	Tinea	0.25	0.38	0.35	0.31	2	0	0.857	1	1	1
	Jaundice	NA									
	Liver	0	0	0.06	2.33	2	0	0.311	0.808	0.408	NA
	Renal	NA									
	Anaemia	0	0.05	0.1	3.29	2	0	0.193	1	0.207	0.403
	Diabetes	0.07	0	0.08	1.24	2	0	0.538	0.851	1	0.78
	Thyroid	0.01	0	0.01	0.53	2	0	0.767	1	1	1
	Candida	0.18	0	0.24	3.07	2	0	0.216	0.253	1	0.407
	Sunburn	0.15	0.22	0.49	7.19	2	0.02	0.028	1	0.026	0.659
	Dry_Skin	0.26	0.41	0.68	4.55	2	0.01	0.103	1	0.113	1
	Urticaria	0.25	0.16	0.36	1.03	2	0	0.596	1	1	1
	Lice	0.15	0.08	0.06	1.53	2	0	0.464	1	0.661	1
	Other	0.29	0	0.25	3.55	2	0.01	0.17	0.172	1	0.224

Table S8b.

Medical itch sub-category severities across groups: Summary of statistics

Medical itch sub-category	AUT-D, M	AUT-S, M	N-AUT, M	H	DF	η^2	p-value	Post-hoc comparisons (p-values)		
								AUT-D vs AUT-S	AUT-D vs N-AUT	AUT-S vs N-AUT
Dermatological	1.73	1.14	0.91	12.79	2	0.04	0.002	0.205	0.002	1
Infection / Infestation	0.59	0.41	0.41	1.64	2	0	0.441	1	0.814	1
Systemic	0.3	0.24	0.13	1.45	2	0	0.483	1	0.737	1

p < 0.05 in **bold**.

AUT-D, diagnosed autistic; AUT-S, self-suspected autism; N-AUT, not autistic.

Spon., spontaneous itch; Prov., provoked itch of any cause; Med., itch of any medical cause.

M, mean; DF, degrees of freedom; H, Kruskal-Wallis H statistic; η^2 , rank Eta-squared.

Table S9a.

Provoked and medical itch subtype timings & durations across groups: Summary of statistics

Temporal Measure	Parent category	Itch subtype	AUT-D, M	AUT-S, M	N-AUT, M	H	DF	η^2	p-value	Post-hoc comparisons (p-values)		
										AUT-D vs AUT-S	AUT-D vs N-AUT	AUT-S vs N-AUT
Episode length	Prov.	Material Objects	0.97	0.67	0.56	2.61	2	0	0.271	1	0.455	1
		Other's touch	0.69	0.27	0	4.35	2	0.03	0.114	1	0.123	0.195
		Temperature	0.35	0	0	5.41	2	0.03	0.067	0.386	0.106	NaN
		Instead of pain	0.72	0	0.5	2.88	2	0.02	0.237	0.296	1	0.611
		Other	1.34	1.22	0.4	3.26	2	0.02	0.196	1	0.216	0.751
		Eczema	0.86	0.33	0	2.32	2	0.02	0.313	1	0.454	0.905
		Psoriasis	2.2	1.25	1.62	3.48	2	0.02	0.175	0.258	0.98	1
		Dermatitis	1.27	0	1	1.2	2	0	0.547	1	1	1
		Dandruff	1.2	1	0.64	0.42	2	0	0.809	1	1	1
		Chronic_Pain	1.4	1	0.67	1.33	2	0	0.513	1	1	1
	Tinea	1	1	0.67	0.14	2	0	0.932	1	1	1	
	Jaundice	1.47	1	1.44	0.16	2	0	0.921	1	1	1	
	Liver	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	
	Renal	2.67	NA	NA	NA	NA	NA	NA	NA	NA	NA	
	Anaemia	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	
	Diabetes	1.17	1	NA	NA	NA	NA	NA	NA	NA	NA	
	Thyroid	1.2	NA	0.33	0.67	1	0	0.414	NA	NA	NA	
	Candida	1	NA	0	NA	NA	NA	NA	NA	NA	NA	
	Sunburn	1.92	NA	3	1.25	1	0	0.263	NA	NA	NA	
	Dry_Skin	2.03	0.67	1.29	4.35	2	0.05	0.114	0.158	1	0.864	
Urticaria	1.4	0.75	0.53	4.64	2	0.04	0.098	0.819	0.136	1		
Lice	2.07	1	0.86	2.83	2	0.04	0.243	1	0.356	1		
Other	2	2	3	1.56	2	0	0.459	1	1	1		
Cluster length	Prov.	Material Objects	1.69	NA	1.71	0.01	1	0	0.934	NA	NA	NA
		Material Objects	2.79	3.05	1.91	3.96	2	0.01	0.138	1	0.224	0.251
		Other's touch	2.52	2.8	0.42	8.86	2	0.09	0.012	1	0.01	0.053
		Temperature	2.08	3.89	1.65	5.19	2	0.03	0.075	0.112	1	0.098
		Instead of pain	1.84	3	2	1.73	2	0	0.421	0.621	1	1
	Med.	Other	2.43	2.78	1.3	2.79	2	0.01	0.248	1	0.359	0.562
		Eczema	2.36	1	0	5.85	2	0.2	0.054	0.944	0.074	0.905
		Psoriasis	2.85	1.92	3	2.6	2	0.01	0.273	0.396	1	0.491
		Dermatitis	2.53	0.5	2	2.24	2	0.01	0.326	0.503	1	1
		Other	2.24	0	0.55	10.19	2	0.23	0.006	0.198	0.019	0.907

		Dandruff	1.92	2.62	1.56	1.78	2	0	0.411	0.716	1	0.751
		Chronic_Pain	2.67	3.5	1.67	0.56	2	0	0.755	1	1	1
		Tinea	1.84	1.8	1.22	0.14	2	0	0.93	1	1	1
		Jaundice	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
		Liver	1	NA	NA	NA	NA	NA	NA	NA	NA	NA
		Renal	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
		Anaemia	1.83	1	NA	NA	NA	NA	NA	NA	NA	NA
		Diabetes	2.8	NA	1	0.22	1	0	0.643	NA	NA	NA
		Thyroid	0	NA	0	NA	NA	NA	NA	NA	NA	NA
		Candida	2.25	NA	0	6.3	1	0.31	0.012	NA	NA	NA
		Sunburn	0.76	0	0.14	4.11	2	0.05	0.128	0.267	0.733	1
		Dry_Skin	2.65	2.38	0.67	10.95	2	0.15	0.004	1	0.002	0.632
		Urticaria	2.13	0	1.71	2.78	2	0.04	0.249	0.322	1	1
		Lice	0.67	0	1.25	1.72	2	0	0.424	1	1	1
		Other	2.38	NA	2.71	0.11	1	0	0.746	NA	NA	NA
Most recent	Prov.	Material	1.11	0.81	2	10.06	2	0.05	0.007	0.579	0.023	0.024
		Objects	1.2	1.09	1.67	0.84	2	0	0.656	1	1	1
		Other's touch	1.58	1.78	1.6	0.01	2	0	0.997	1	1	1
		Temperature	1.24	1.29	1.25	0.23	2	0	0.891	1	1	1
		Instead of pain	1.76	1.67	1.2	1.3	2	0	0.523	1	0.797	1
	Med.	Other	2.21	0.33	2	1.73	2	0	0.421	0.634	1	1
		Eczema	2.27	2.58	2.85	0.99	2	0	0.61	1	1	1
		Psoriasis	1.93	0	2.5	2.05	2	0	0.359	0.535	1	1
		Dermatitis	2.31	2.5	2.36	0.06	2	0	0.97	1	1	1
		Dandruff	1.28	0.62	0.67	1.47	2	0	0.479	1	1	1
		Chronic_Pain	1.57	1.5	2.33	1.79	2	0	0.408	1	0.459	1
		Tinea	2.53	4	2.44	3.11	2	0.04	0.211	0.285	1	0.451
		Jaundice	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
		Liver	3.67	NA	NA	NA	NA	NA	NA	NA	NA	NA
		Renal	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
		Anaemia	2.5	5	NA	NA	NA	NA	NA	NA	NA	NA
		Diabetes	2.4	NA	2	0.02	1	0	0.879	NA	NA	NA
		Thyroid	5	NA	0	NA	NA	NA	NA	NA	NA	NA
		Candida	2.67	NA	4.2	2.56	1	0.04	0.11	NA	NA	NA
		Sunburn	4.06	4.67	4	3.23	2	0.03	0.199	0.272	1	1
	Dry_Skin	1.07	1.5	1.4	0.51	2	0	0.775	1	1	1	
	Urticaria	3.12	4.5	2.71	1.39	2	0	0.499	0.657	1	1	
	Lice	4.67	5	5	1.67	2	0	0.435	1	0.773	NaN	
	Other	1.86	NA	1.43	0.88	1	0	0.348	NA	NA	NA	

p < 0.05 in **bold**.

AUT-D, diagnosed autistic; AUT-S, self-suspected autism; N-AUT, not autistic.

Spon., spontaneous itch; Prov., provoked itch of any cause; Med., itch of any medical cause.

M, mean; DF, degrees of freedom; H, Kruskal-Wallis H statistic; η^2 , rank Eta-squared.

Table S9b.

Medical itch sub-category timings & durations across groups: Summary of statistics

Temporal Measure	Medical itch sub-category	AUT-D, M	AUT-S, M	N-AUT, M	H	DF	η^2	p-value	Post-hoc comparisons (p-values)		
									AUT-D vs AUT-S	AUT-D vs N-AUT	AUT-S vs N-AUT
Episode length	Dermatological	2.37	1.53	1.41	7.74	2	0.05	0.021	0.25	0.038	1
	Infection / Infestation	1.71	1.2	2.17	1.24	2	0	0.537	1	1	0.816
	Systemic	1.59	1	0.5	2.35	2	0.02	0.308	1	0.413	1
Cluster length	Dermatological	3	3.07	2.21	3.29	2	0.01	0.193	1	0.217	0.835
	Infection / Infestation	1.74	1.8	1	1.07	2	0	0.586	1	1	1
	Systemic	2.41	2.67	1.33	1.37	2	0	0.503	1	1	0.836
Most recent	Dermatological	3.45	3.2	3.52	0.51	2	0	0.775	1	1	1
	Infection / Infestation	2.84	4	3.92	5.51	2	0.08	0.064	0.466	0.116	1
	Systemic	2.56	2.67	2.17	0.18	2	0	0.915	1	1	1

Table S10a.

Provoked and medical itch subtype impacts across groups: Summary of statistics

Impact	Parent category	Itch subtype	AUT-D, M	AUT-S, M	N-AUT, M	H	DF	η^2	p-value	Post-hoc comparisons (p-values)		
										AUT-D vs AUT-S	AUT-D vs N-AUT	AUT-S vs N-AUT
Sleep	Provoked	Material	1.61	1.29	0.62	17.33	2	0.09	<0.001	1	<0.001	0.012
		Objects	1.28	0.64	0.33	7.65	2	0.07	0.022	0.236	0.059	1
		Other's touch	1.1	0.44	0.45	6.12	2	0.04	0.047	0.267	0.144	1
		Temperature	1.92	0.57	1	5.7	2	0.1	0.058	0.1	0.47	1
		Instead of pain	2.39	1.44	0.9	9.74	2	0.14	0.008	0.206	0.016	1
	Medical	Other	1	0.67	0.4	1.38	2	0	0.5	1	0.809	1
		Eczema	2.34	2.08	1.62	2.8	2	0.01	0.247	1	0.289	1
		Psoriasis	1	0.5	0.25	2.39	2	0.02	0.303	1	0.454	1
		Dermatitis	1.62	0.5	0.73	5.68	2	0.1	0.058	0.681	0.09	1
		Dandruff	1.32	0.75	0.67	2.44	2	0.01	0.295	0.99	0.518	1
		Chronic_Pain	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
		Tinea	1.26	0.8	0.78	1.42	2	0	0.492	1	0.915	1
		Jaundice	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
		Liver	3	NA	NA	NA	NA	NA	NA	NA	NA	NA
		Renal	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
		Anaemia	1.83	0	NA	NA	NA	NA	NA	NA	NA	NA
		Diabetes	1.4	NA	1.33	0.26	1	0	0.608	NA	NA	NA
		Thyroid	0	NA	1	NA	NA	NA	NA	NA	NA	NA
		Candida	2.42	NA	1.2	2.31	1	0.02	0.129	NA	NA	NA
		Sunburn	1.67	1.17	1.71	0.35	2	0	0.838	1	1	1
Dry_Skin	1.5	0.62	0.6	9.68	2	0.12	0.008	0.216	0.016	1		
Urticaria	2.47	1	1.14	4.82	2	0.13	0.09	0.552	0.175	1		
Lice	2	1	2.25	0.46	2	0	0.795	1	1	1		
Other	2	NA	1.57	0.59	1	0	0.444	NA	NA	NA		
Leisure	Provoked	Material	1.74	1.11	0.72	17.83	2	0.11	<0.001	0.147	<0.001	0.874
		Objects	1.47	0.33	0.78	7.09	2	0.08	0.029	0.047	0.465	1
		Other's touch	1.39	1.12	0.44	8.82	2	0.09	0.012	1	0.01	0.549
		Temperature	1.78	0.71	0.88	3.86	2	0.06	0.145	0.354	0.397	1
		Instead of pain	1.64	0.57	0.67	7.15	2	0.13	0.028	0.122	0.12	1
	Medical	Other	1.3	1	0.6	1.19	2	0	0.552	1	0.907	1
		Eczema	1.92	0.67	1.09	7.58	2	0.09	0.023	0.05	0.317	0.673
		Psoriasis	0.92	0	0.67	1.37	2	0	0.504	0.868	1	1
		Dermatitis	1.38	0.5	0.6	2.5	2	0.02	0.287	1	0.436	1
		Dandruff	1.32	0.62	0.89	1.16	2	0	0.56	0.994	1	1

		Chronic_Pain	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	
		Tinea	0.78	0.8	1	0	2	0	0.999	1	1	1	
		Jaundice	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	
		Liver	4	NA	NA	NA	NA	NA	NA	NA	NA	NA	
		Renal	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	
		Anaemia	2	0	NA	NA	NA	NA	NA	NA	NA	NA	
		Diabetes	1.67	NA	0.67	0.89	1	0	0.346	NA	NA	NA	
		Thyroid	0	NA	0	NA	NA	NA	NA	NA	NA	NA	
		Candida	2.2	NA	0.6	3.21	1	0.1	0.073	NA	NA	NA	
		Sunburn	1.31	0.67	0.33	3.22	2	0.03	0.2	1	0.325	1	
		Dry_Skin	0.82	0.38	0.36	2.06	2	0	0.358	1	0.636	1	
		Urticaria	1.5	2	0.71	2.66	2	0.03	0.264	1	0.717	0.441	
		Lice	2.33	2	2	0.07	2	0	0.964	1	1	1	
		Other	1.21	NA	1.43	0.15	1	0	0.696	NA	NA	NA	
Housework	Provoked	Material	1.6	0.79	0.64	19.43	2	0.12	<0.001	0.022	<0.001	1	
		Objects	1.65	0.67	0.33	10.6	2	0.13	0.005	0.133	0.017	1	
		Other's touch	0.53	0.57	0.27	2.13	2	0	0.344	1	0.548	1	
		Temperature	1.95	0.4	0.86	7.08	2	0.18	0.029	0.087	0.215	1	
		Instead of pain	1.48	0.4	0.89	3.96	2	0.05	0.138	0.243	0.655	1	
	Medical	Other	1.3	0.5	0.2	2.81	2	0.06	0.246	1	0.374	1	
		Eczema	1.92	1.1	1.09	5.67	2	0.05	0.059	0.251	0.176	1	
		Psoriasis	1	0	0.67	1.77	2	0	0.414	0.682	1	1	
		Dermatitis	1.88	0.5	1	5.47	2	0.1	0.065	0.384	0.166	1	
		Dandruff	0.52	0.25	0.33	0.43	2	0	0.806	1	1	1	
		Chronic_Pain	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
		Tinea	0.59	0	0.5	1.72	2	0	0.423	0.763	1	0.656	
		Jaundice	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
		Liver	4	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
		Renal	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
		Anaemia	1	0	NA	NA	NA	NA	NA	NA	NA	NA	NA
		Diabetes	0.33	NA	0.67	0.07	1	0	0.796	NA	NA	NA	NA
		Thyroid	0	NA	0	NA	NA	NA	NA	NA	NA	NA	NA
		Candida	1.8	NA	0.4	3.11	1	0.09	0.078	NA	NA	NA	NA
		Sunburn	1	0.67	0.33	1.56	2	0	0.46	1	0.702	1	
Dry_Skin	1.15	0.38	0.5	4.07	2	0.04	0.131	0.351	0.392	1			
Urticaria	1.46	2	1.29	0.74	2	0	0.691	1	1	1			
Lice	1	2	1	0.4	2	0	0.818	1	1	1			
Other	0.85	NA	0.83	0.01	1	0	0.924	NA	NA	NA			
Work	Provoked	Material	1.87	1.4	0.68	21.23	2	0.13	<0.001	0.461	<0.001	0.083	
		Objects	1.4	0.44	0.33	8.18	2	0.1	0.017	0.147	0.068	1	
		Other's touch	1.07	1.12	0.44	5.56	2	0.05	0.062	1	0.052	0.602	
		Temperature	1.4	0.5	1.25	1.9	2	0	0.388	0.533	1	1	
		Instead of pain	1.67	0.86	0.62	5.44	2	0.09	0.066	0.409	0.132	1	
		Other	1.4	0.5	0	5.95	2	0.28	0.051	1	0.07	0.618	

Medical	Eczema	1.83	0.64	0.82	11.86	2	0.15	0.003	0.015	0.046	1
	Psoriasis	1.08	0	0.5	1.99	2	0	0.369	0.779	1	1
	Dermatitis	1.29	0	0.44	4.33	2	0.08	0.115	0.857	0.217	1
	Dandruff	0.86	0.62	0.56	0.37	2	0	0.831	1	1	1
	Chronic_Pain	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
	Tinea	0.75	0	0.62	2.89	2	0.04	0.236	0.277	1	0.656
	Jaundice	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
	Liver	4	NA	NA	NA	NA	NA	NA	NA	NA	NA
	Renal	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
	Anaemia	2.33	0	NA	NA	NA	NA	NA	NA	NA	NA
	Diabetes	1.67	NA	0.33	1.34	1	0	0.246	NA	NA	NA
	Thyroid	0	NA	1	NA	NA	NA	NA	NA	NA	NA
	Candida	2.11	NA	0.6	3.19	1	0.11	0.074	NA	NA	NA
	Sunburn	0.76	0.5	0.17	1.35	2	0	0.508	1	0.789	1

Table S10b.

Medical itch sub-category impacts across groups: Summary of statistics

Impact	Itch subtype	AUT-D, M	AUT-S, M	N-AUT, M	H	DF	η^2	p-value	Post-hoc comparisons (p-values)		
									AUT-D vs AUT-S	AUT-D vs N-AUT	AUT-S vs N-AUT
Sleep	Dermatological	2.42	2.13	1.55	7.87	2	0.05	0.02	1	0.017	0.585
	Infection / Infestation	1.65	0.8	1.5	1.77	2	0	0.414	0.579	1	1
	Systemic	2.17	2.33	1.33	1.38	2	0	0.503	1	0.815	1
Leisure	Dermatological	1.96	0.79	1.11	12.45	2	0.09	0.002	0.018	0.024	1
	Infection / Infestation	1.39	0.8	1.18	0.81	2	0	0.665	1	1	1
	Systemic	2	1	0.4	4.23	2	0.12	0.121	1	0.171	1
Housework	Dermatological	2	0.87	1.25	12.96	2	0.09	0.002	0.01	0.03	0.769
	Infection / Infestation	1	0.4	0.64	0.97	2	0	0.615	1	1	1
	Systemic	1.53	1.5	0.8	0.69	2	0	0.707	1	1	1
Work	Dermatological	1.77	0.8	1	11.75	2	0.09	0.003	0.026	0.021	1
	Infection / Infestation	1.31	0.4	1.09	2.33	2	0.01	0.312	0.419	1	0.776
	Systemic	2.08	0	0.8	4.11	2	0.13	0.128	0.753	0.317	1

p < 0.05 in **bold**.

AUT-D, diagnosed autistic; AUT-S, self-suspected autism; N-AUT, not autistic.

Spon., spontaneous itch; Prov., provoked itch of any cause; Med., itch of any medical cause.

M, mean; DF, degrees of freedom; H, Kruskal-Wallis H statistic; η^2 , rank Eta-squared.

Regression model outputs**Table S11.**

Results from hierarchical multiple regression analysis with AQ-10 as the outcome.

Step	Model summary	Predictor	st.			
			beta	error	95% CI	p
1A	R ² =0.28; adj. R ² =0.27, p<0.001	Spontaneous itch severity	0.64	0.13	0.4, 0.9	<0.001
		Provoked itch severity	0.66	0.11	0.4, 0.9	<0.001
		Medical itch severity	0.12	0.09	-0.1, 0.3	0.215
1B	R ² =0.26; adj. R ² =0.24, p<0.001	Spontaneous itch severity	0.66	0.14	0.4, 0.9	<0.001
		Provoked itch severity: Material	0.43	0.14	0.2, 0.7	0.003
		Provoked itch severity: Other's touch	0.05	0.16	-0.3, 0.4	0.767
		Provoked itch severity: Objects	0.29	0.17	0, 0.6	0.094
		Provoked itch severity: Instead of pain	0.29	0.16	0, 0.6	0.082
		Provoked itch severity: Temperature	-0.07	0.19	-0.4, 0.3	0.715
		Provoked itch severity: Other	0.03	0.24	-0.4, 0.5	0.895
1C	R ² =0.25; adj. R ² =0.25, p<0.001	Spontaneous itch severity	0.7	0.13	0.4, 1	<0.001
		Provoked itch severity: Material	0.6	0.11	0.4, 0.8	<0.001
1D	R ² =0.27; adj. R ² =0.27, p<0.001	Spontaneous itch severity	0.65	0.13	0.4, 0.9	<0.001
		Provoked itch severity	0.68	0.11	0.5, 0.9	<0.001
2A	R ² =0.13; adj. R ² =0.1, p<0.001	Spontaneous longest episode	0.18	0.2	-0.2, 0.6	0.374
		Provoked longest episode	0.43	0.17	0.1, 0.8	0.009
		Medical longest episode	0.01	0.17	-0.3, 0.3	0.968
		Spontaneous longest cluster	0.16	0.09	0, 0.3	0.075
		Provoked longest cluster	0.12	0.09	-0.1, 0.3	0.203
		Medical longest cluster	0.02	0.12	-0.2, 0.3	0.867
		Spontaneous most recent episode	-0.03	0.16	-0.4, 0.3	0.857
		Provoked most recent episode	0.21	0.14	-0.1, 0.5	0.131
		Medical most recent episode	0.13	0.11	-0.1, 0.3	0.241
2B	R ² =0.08; adj. R ² =0.06, p<0.001	Provoked longest episode: Material	0.31	0.19	-0.1, 0.7	0.103
		Provoked longest episode: Objects	0.57	0.31	0, 1.2	0.068
		Provoked longest episode: Other's touch	0.54	0.4	-0.2, 1.3	0.178
		Provoked longest episode: Instead of pain	0.45	0.24	0, 0.9	0.063
		Provoked longest episode: Temperature	0.1	0.41	-0.7, 0.9	0.803
		Provoked longest episode: Other	-0.33	0.55	-1.4, 0.8	0.544
3A	R ² =0.19; adj. R ² =0.18, p<0.001	Spontaneous total impact	0.67	0.19	0.3, 1	<0.001
		Provoked total impact	0.65	0.17	0.3, 1	<0.001
		Medical total impact	0.14	0.15	-0.1, 0.4	0.339
3B	R ² =0.19; adj. R ² =0.16, p<0.001					

		Spontaneous total impact	0.77	0.2	0.4, 1.2	<0.001
		Provoked total impact: Material	0.69	0.22	0.3, 1.1	0.002
		Provoked total impact: Objects	0.07	0.27	-0.5, 0.6	0.805
		Provoked total impact: Other's touch	0.07	0.3	-0.5, 0.7	0.827
		Provoked total impact: Temperature	0.16	0.25	-0.3, 0.6	0.508
		Provoked total impact: Instead of pain	-0.34	0.3	-0.9, 0.2	0.251
		Provoked total impact: Other	-0.27	0.49	-1.2, 0.7	0.582
3C	R ² =0.18; adj. R ² =0.17, p<0.001					
		Spontaneous total impact	0.76	0.19	0.4, 1.1	<0.001
		Provoked total impact: Material	0.69	0.17	0.4, 1	<0.001
4A	R ² =0.28; adj. R ² =0.26, p<0.001					
		Spontaneous itch severity	0.26	0.23	-0.2, 0.7	0.272
		Provoked itch severity	0.13	0.23	-0.3, 0.6	0.576
		Provoked longest episode	-0.04	0.16	-0.4, 0.3	0.789
		Spontaneous longest cluster	-0.01	0.08	-0.2, 0.1	0.894
		Spontaneous total impact	0.55	0.16	0.2, 0.9	<0.001
		Provoked total impact: Material	0.62	0.15	0.3, 0.9	<0.001
4B	R ² =0.27; adj. R ² =0.27, p<0.001					
		Spontaneous itch severity	0.65	0.13	0.4, 0.9	<0.001
		Provoked itch severity	0.68	0.11	0.5, 0.9	<0.001
4C	R ² =0.63; adj. R ² =0.63, p<0.001					
		Dummy: AUT-S	3.42	0.38	2.7, 4.2	<0.001
		Dummy: AUT-D	4.64	0.29	4.1, 5.2	<0.001
		Spontaneous itch severity	0.2	0.1	0, 0.4	0.045
		Provoked itch severity	0.35	0.08	0.2, 0.5	<0.001
4D	R ² =0.5; adj. R ² =0.49, p<0.001					
		GSQ score	0.06	0.01	0.1, 0.1	<0.001
		Spontaneous itch severity	0.15	0.12	-0.1, 0.4	0.212
		Provoked itch severity	0.26	0.1	0, 0.5	0.016
4E	R ² =0.59; adj. R ² =0.58, p<0.001					
		Dummy: AUT-S	3.75	0.39	3, 4.5	<0.001
		Dummy: AUT-D	5.29	0.27	4.8, 5.8	<0.001
4F	R ² =0.49; adj. R ² =0.48, p<0.001					
		GSQ score	0.07	0	0.1, 0.1	<0.001

Table S12.

Results from hierarchical multiple regression analysis with GSQ as the outcome.

Step	Model summary	Predictor	st.			p
			beta	error	95% CI	
1A	R ² =0.39; adj. R ² =0.39, p<0.001	Spontaneous itch severity	7.86	1.15	5.6, 10.1	<0.001
		Provoked itch severity	7.26	1.02	5.2, 9.3	<0.001
		Medical itch severity	1.12	0.84	-0.5, 2.8	0.185
1B	R ² =0.44; adj. R ² =0.43, p<0.001	Spontaneous itch severity	6.83	1.14	4.6, 9.1	<0.001
		Provoked itch severity: Material	3.18	1.15	0.9, 5.4	0.006
		Provoked itch severity: Other's touch	2.81	1.28	0.3, 5.3	0.029
		Provoked itch severity: Objects	3.07	1.4	0.3, 5.8	0.029
		Provoked itch severity: Instead of pain	5.52	1.34	2.9, 8.2	<0.001
		Provoked itch severity: Temperature	2.55	1.56	-0.5, 5.6	0.103
		Provoked itch severity: Other	-1.11	1.93	-4.9, 2.7	0.566
1C	R ² =0.44; adj. R ² =0.43, p<0.001	Spontaneous itch severity	6.79	1.12	4.6, 9	<0.001
		Provoked itch severity: Material	3.82	1.11	1.6, 6	<0.001
		Provoked itch severity: Other's touch	2.88	1.26	0.4, 5.4	0.023
		Provoked itch severity: Objects	3.1	1.36	0.4, 5.8	0.023
		Provoked itch severity: Instead of pain	5.33	1.31	2.8, 7.9	<0.001
2A	R ² =0.27; adj. R ² =0.24, p<0.001	Spontaneous longest episode	2.35	1.83	-1.2, 5.9	0.200
		Provoked longest episode	5.56	1.47	2.7, 8.5	<0.001
		Medical longest episode	1.71	1.52	-1.3, 4.7	0.262
		Spontaneous longest cluster	1.83	0.8	0.3, 3.4	0.022
		Provoked longest cluster	2.57	0.83	0.9, 4.2	0.002
		Medical longest cluster	-0.47	1.07	-2.6, 1.6	0.660
		Spontaneous most recent episode	0	1.46	-2.9, 2.9	0.998
		Provoked most recent episode	-1.49	1.24	-3.9, 1	0.232
		Medical most recent episode	1.04	0.99	-0.9, 3	0.297
2B	R ² =0.24; adj. R ² =0.23, p<0.001	Provoked longest episode	6.89	1.33	4.3, 9.5	<0.001
		Spontaneous longest cluster	1.97	0.78	0.4, 3.5	0.012
		Provoked longest cluster	2.39	0.82	0.8, 4	0.004
2C	R ² =0.27; adj. R ² =0.23, p<0.001	Provoked longest episode: Material	2.83	1.72	-0.6, 6.2	0.102
		Provoked longest episode: Objects	3.49	3.28	-3, 9.9	0.288
		Provoked longest episode: Other's touch	8.45	3.55	1.5, 15.4	0.018
		Provoked longest episode: Instead of pain	3.23	2.49	-1.7, 8.1	0.195
		Provoked longest episode: Temperature	4.93	4	-2.9, 12.8	0.218
		Provoked longest episode: Other	-2.17	5.57	-13.1, 8.8	0.698
		Spontaneous longest cluster	1.87	0.8	0.3, 3.4	0.019
		Provoked longest cluster: Material	0.85	0.92	-1, 2.7	0.357
		Provoked longest cluster: Objects	2.36	1.48	-0.5, 5.3	0.112

	Provoked longest cluster: Other's touch	1.02	1.13	-1.2, 3.2	0.367
	Provoked longest cluster: Instead of pain	1.71	1.58	-1.4, 4.8	0.279
	Provoked longest cluster: Temperature	-0.22	1.71	-3.6, 3.1	0.898
	Provoked longest cluster: Other	-0.79	2.49	-5.7, 4.1	0.752
2D	R2=0.14; adj. R2=0.13, p<0.001				
	Provoked longest episode: Other's touch	14.37	3.36	7.8, 21	<0.001
	Spontaneous longest cluster	3.59	0.75	2.1, 5.1	<0.001
3A	R2=0.43; adj. R2=0.42, p<0.001				
	Spontaneous total impact	9.97	1.56	6.9, 13	<0.001
	Provoked total impact	10.29	1.37	7.6, 13	<0.001
	Medical total impact	0.33	1.18	-2, 2.7	0.781
3B	R2=0.42; adj. R2=0.4, p<0.001				
	Spontaneous total impact	9.92	1.66	6.6, 13.2	<0.001
	Provoked total impact: Material	6.87	1.78	3.4, 10.4	<0.001
	Provoked total impact: Objects	3.24	2.22	-1.1, 7.6	0.146
	Provoked total impact: Other's touch	4.91	2.48	0, 9.8	0.049
	Provoked total impact: Temperature	2.29	2.01	-1.7, 6.3	0.257
	Provoked total impact: Instead of pain	-0.48	2.43	-5.2, 4.3	0.845
	Provoked total impact: Other	-4.55	4.03	-12.5, 3.4	0.260
3C	R2=0.41; adj. R2=0.4, p<0.001				
	Spontaneous total impact	10.37	1.61	7.2, 13.6	<0.001
	Provoked total impact: Material	8.36	1.52	5.4, 11.3	<0.001
	Provoked total impact: Other's touch	5.33	2.38	0.6, 10	0.026
4A	R2=0.48; adj. R2=0.46, p<0.001				
	Spontaneous total impact	5.61	2.02	1.6, 9.6	0.006
	Provoked total impact: Material	2.79	2.15	-1.5, 7	0.197
	Provoked total impact: Other's touch	3.33	3.21	-3, 9.7	0.301
	Provoked longest episode: Other's touch	-1.59	1.42	-4.4, 1.2	0.266
	Spontaneous longest cluster	-0.39	0.74	-1.8, 1.1	0.603
	Provoked longest cluster	1.19	0.77	-0.3, 2.7	0.124
	Spontaneous itch severity	4.73	1.34	2.1, 7.4	<0.001
	Provoked itch severity: Material	2.32	1.41	-0.4, 5.1	0.100
	Provoked itch severity: Other's touch	0.68	1.65	-2.6, 3.9	0.681
	Provoked itch severity: Objects	2.81	1.35	0.1, 5.5	0.039
	Provoked itch severity: Instead of pain	4.02	1.38	1.3, 6.7	0.004
4B	R2=0.44; adj. R2=0.43, p<0.001				
	Spontaneous total impact	7.33	1.84	3.7, 11	<0.001
	Spontaneous itch severity	5.51	1.3	3, 8.1	<0.001
	Provoked itch severity: Objects	4.95	1.23	2.5, 7.4	<0.001
	Provoked itch severity: Instead of pain	5.51	1.28	3, 8	<0.001
4C	R2=0.6; adj. R2=0.59, p<0.001				
	Spontaneous total impact	5.14	1.59	2, 8.3	0.001
	Spontaneous itch severity	3.21	1.13	1, 5.4	0.005
	Provoked itch severity: Objects	3.66	1.05	1.6, 5.7	<0.001
	Provoked itch severity: Instead of pain	4.97	1.09	2.8, 7.1	<0.001
	Dummy: S-AUT	19.45	3.9	11.8, 27.1	<0.001
	Dummy: D-AUT	29.22	2.89	23.5, 34.9	<0.001
4D	R2=0.42; adj. R2=0.41, p<0.001				

		Dummy: S-AUT	27.49	4.55	18.5, 36.4	<0.001
		Dummy: D-AUT	43.17	3.08	37.1, 49.2	<0.001
4E	R ² =0.63; adj. R ² =0.62, p<0.001	Spontaneous total impact	6.38	1.51	3.4, 9.4	<0.001
		Spontaneous itch severity	2.23	1.1	0.1, 4.4	0.044
		Provoked itch severity: Objects	2.82	1.03	0.8, 4.8	0.006
		Provoked itch severity: Instead of pain	3.91	1.06	1.8, 6	<0.001
		AQ-10 score	4.69	0.42	3.9, 5.5	<0.001
4F	R ² =0.49; adj. R ² =0.48, p<0.001	AQ-10 score	6.76	0.42	5.9, 7.6	<0.001

Table S13.

Prevalence estimates of current and chronic itch.

Medical itch sub-category	AUT-D, n (%*)	AUT-S, n (%*)	N-AUT, n (%*)	<i>All groups,</i> <i>n (%*)</i>
Current (within last week)	123 (78%)	27 (73%)	47 (55%)	197 (70%)
Chronic + current (>4 weeks duration)	103 (65%)	26 (70%)	33 (39%)	162 (58%)

* Percentage of group reporting type of itch.

AUT-D, diagnosed autistic; AUT-S, self-suspected autism; N-AUT, not autistic