

Use of technology for the objective evaluation of scratching behavior: A systematic review



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Introduction: Pruritus is a common symptom across various dermatologic conditions, with a negative impact on quality of life. Devices to quantify itch objectively primarily use scratch as a proxy. This review compares and evaluates the performance of technologies aimed at objectively measuring scratch behavior.

Methods: Articles identified from literature searches performed in October 2020 were reviewed and those that did not report a primary statistical performance measure (eg, sensitivity, specificity) were excluded. The articles were independently reviewed by 2 authors.

Results: The literature search resulted in 6231 articles, of which 24 met eligibility criteria. Studies were categorized by technology, with actigraphy being the most studied (n = 21). Wrist actigraphy's performance is poorer in pruritic patients and inherently limited in finger-dominant scratch detection. It has moderate correlations with objective measures (Eczema and Area Severity Index/Investigator's Global Assessment: $r_s(\rho) = 0.70-0.76$), but correlations with subjective measures are poor ($r^2 = 0.06$, $r_s(\rho) = 0.18-0.40$ for itch measured using a visual analog scale). This may be due to varied subjective perception of itch or actigraphy's underestimation of scratch.

Conclusion: Actigraphy's large variability in performance and limited understanding of its specificity for scratch merits larger studies looking at validation of data analysis algorithms and device performance, particularly within target patient populations. (JAAD Int 2021;5:19-32.)

Key words: algorithm; atopic dermatitis; disease management; drug development; eczema; general dermatology; itch; machine learning; pediatric dermatology; pruritus; technology.

INTRODUCTION

Pruritus is a common symptom of systemic and dermatologic disorders, and scratching is the innate reflex.¹ The itch-scratch cycle is a hallmark symptom of atopic dermatitis (AD) and perpetuates skin barrier dysfunction. Notably more severe during sleep, itch in AD has been shown to impact sleep quality.²⁻⁵ Historically, itch has been assessed subjectively through visual analog scales (VAS) and numeric rating scales.⁶ However, these measures

often do not correlate to visually observed scratch, especially in children.⁷⁻⁹ More recently, studies have explored device-driven methods to objectively measure scratch as a proxy for itch.

Actigraphy is the most commonly tested method and entails the use of accelerometers to monitor wrist movements, a proxy for scratching. Other technologies include acoustic devices,^{10,11} strain gauges,^{12,13} pressure sensors,^{12,14} and vibratory sensors.^{12,13,15-18} The commonly accepted gold

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standard is video recording of scratching with manual coding by an observer, which is time-consuming and impractical in clinical settings.¹⁹⁻²¹ The purpose of this systematic review is to assess the performance and algorithms of technological methods currently available to evaluate scratching behavior objectively.

METHODS

Search strategy

We queried PubMed, MEDLINE, Embase (Elsevier), Cochrane Library and Cochrane Central Register of Controlled Trials (CENTRAL), Scopus (Elsevier), Web of Science (Clarivate Analytics), and IEEE Xplore Digital Library in October 2020 without limits on publication date. The search strategy is fully detailed in the Supplemental Materials under “Search Strategy” (available via Mendeley at <https://data.mendeley.com/datasets/ryg97c26t6/2>).

Study selection

Eligibility assessment was performed independently by 2 authors. Included articles must feature critical assessment of a technology designed to measure itch objectively and report at least 1 of the primary outcomes described below. Exclusion criteria included studies of nonhuman subjects, articles without original data, and studies describing technology without assessing its performance.

Quality assessment

Study quality was assessed using a rating scheme (1-5), which was modified from the Oxford Centre for Evidence-Based Medicine²² for rating levels of evidence. The individual studies assessed are described in [Tables I and II](#) and assessment was performed by at least 2 authors.

Data extraction and outcomes

Performance values were extracted using a standardized survey. Primary outcomes included sensitivity, specificity, and positive and negative predictive values of scratch detection methods. Secondary outcomes included correlations of

detection methods to other technologies and subjective assessments.

Performance metrics

Sensitivity is defined as the ability to detect the number of true positives (eg, true scratching) and specificity is the ability to detect the number of true negatives (eg, nonscratching movements). Positive predictive value (PPV, precision) is the proportion of positives that are true positives (eg, movements labeled as scratch that are true scratches). The F1 score encompasses both sensitivity and precision. Root mean square error (RMSE) is the standard deviation of residuals and is effectively an estimation of how well an algorithm predicts the observed data (ie, accuracy).

CAPSULE SUMMARY

- We assessed ways to quantify itch by measuring scratching behavior via various technological modalities (eg, actigraphy, smartwatch applications, acoustic sensors).
- The overall performance of current objective tools for quantifying itch suffers from low accuracy and variable performance. Further development will allow for more-objective evaluation of disease management and treatment.

Algorithms

To efficiently extract and analyze device data, algorithms capable of distinguishing scratch from nonscratch movements are essential. Linear regression modeling is generated from the number of activity counts above a frequency threshold and total scratch time; however, this model is limited by confounding movements (eg, walking, restlessness).²³ Logistic regression modeling is a simple approach to binary classification (eg, scratch vs nonscratch) and analogous to linear regression. Bidirectional recurrent neural networks are a form of machine learning whereby the network can detect patterns directly (eg, scratch waveforms) from raw input data, thereby eliminating precursory extraction of patterns required for other models.²⁴ The k-means clustering analysis is another approach that involves clustering a set number of subgroups within a data set. The algorithm then allocates device signals into their respective subgroups based on frequency, waveform, or other qualities.²³

RESULTS

Of the 6231 articles identified, 72 were assessed based on exclusion criteria and 24 fully met eligibility criteria. Most articles looked at AD, although other conditions were also examined (eg, urticaria). Articles reporting performance and correlation measures are summarized in [Tables I and II](#).

Abbreviations used:

| | |
|-------|----------------------------------|
| AD: | Atopic dermatitis |
| PPV: | Positive predictive value |
| RMSE: | Root mean square error |
| VAS: | Visual analog scale |
| TST%: | total scratching time percentage |

Sensitivity and specificity ranges of technologies compared to video recording are summarized in Table III. An overview of benefits and limitations is seen in Table IV.

Actigraphy

Performance. Actigraphy is the most studied technology.^{5,21,25-27} Twenty-one articles investigated actigraphy devices and data extraction algorithms, with 7 compared to video recording.^{12,16,21,24,25,28,29} While all 7 articles looked at healthy subjects, only 2 reported sensitivity values (0.00-0.96; zero values indicate no true positives).^{24,29} Specificity was reported by 1 article (0.92).²⁹ Four articles explored actigraphy in AD subjects, with 1 reporting sensitivity values (0.00-0.89) and PPV values (0.00-0.57).²⁴ Specificity was not reported in this population. The large ranges likely stem from the various extraction algorithms and actigraphs (eg, PAM-RL,^{26,29} Actiwatch Plus,^{8,26} DigiTrac^{30,31}).

Each algorithm has its limitations. The k-means clustering analysis algorithm of Feuerstein et al²³ yielded high performance values, but required all anticipated movements to be determined a priori. While logistic regression approach from Petersen et al²⁹ for detecting total nocturnal scratch time yielded comparable performance to the algorithm from Feuerstein et al,²³ the model had significantly decreased performance when tested with a separate data set.²⁴ The bidirectional recurrent neural networks algorithm proposed by Moreau et al²⁴ yielded higher sensitivity, PPV, and F1-scores than the logistic regression model; however, it has not been tested in further datasets. Correlation between actigraphy data and video recording was evaluated by Moreau et al,²⁴ reporting Spearman rank correlation coefficients ($r_s(\rho)$) of 0.95-0.96.²⁸ Other studies report correlations between actigraphy and video recording for total scratching time percentage (TST%) calculation ($r_s(\rho) = 0.91$),²⁵ and correlation values between actigraphy and video recording of sleep efficiency were all reported to be greater than 0.92 by Benjamin et al.²¹

Correlations with other objective and subjective measures. Several articles explored correlations between actigraphy and subjective sleep

measures, disease severity, AD-associated serum markers, and subjective itch measures. Ten articles compared actigraphy to subjective sleep measures, with 4 reporting correlations. VAS sleep, a patient-reported measure of sleep quality, was examined in 1 article, reporting correlation coefficients of -0.44 in adults and 0.48 in children when compared to average hourly activity scores.²⁶ The total scoring AD index, which includes both subjective (eg, itch and sleep) and objective (eg, disease severity) measures, had moderate correlations with various activity measures ranging from 0.53 - 0.64 in adults ($P < .05$) and 0.42 - 0.62 in children ($P < .05$).^{26,27,30} While total and objective total scoring AD indexes both resulted in $r_s(\rho) = 0.52$ ($P < .001$) in children ($n = 24$) compared to wrist activity, correlations with pruritus and sleep subscores were not significant.³⁰

Two articles evaluated other disease severity indices in children and adults, with moderate correlation for objective measures (Eczema and Area Severity Index and Investigator's Global Assessment) compared to actigraphic wake after sleep onset, ranging from 0.70 - 0.76 ($P < .02$, $n = 10$).³² Six area six sign AD was found to have a weak correlation with average nocturnal movement ($r_s(\rho) = 0.15$, $P = .02$, $n = 235$).³³

Four articles investigated -serum markers associated with AD. Statistically significant correlations with actigraphy measurements ranged from 0.51 - 0.93 .^{17,27,30,31} These studies were not compared to video recording, however, and thus conclusions specifically related to scratch are difficult to make.

While there seems to be a moderate correlation between actigraphy and objective measures, this is not the case with subjective measures. Fourteen articles compared actigraphy to subjective itch, with 2 articles reporting correlation coefficients. Comparison between VAS itch and mean actigraphy scores yielded coefficients of determination (r^2) of 0.06 in children and adults with various pruritic conditions ($n = 118$) and 0.08 in adult AD subjects ($n = 20$).⁸ VAS itch and hourly activity scores yielded $r_s(\rho) = 0.40$ ($P = .049$) in children and 0.18 ($P = .9$) in adults.²⁶

Actigraphy-based scratch measurements correlate poorly to VAS itch scores, sleep quality, and other subjective patient-reported outcomes.^{8,21,26,27,30} The reasons for this are likely multifactorial. In pediatric populations, proxy measures may be under or overestimated by caregivers. More likely, there are inherent differences between a subject's perception of itch and the objective actions of scratching. An individual may report a high level of subjective itch

Table I. Summary table for studies exploring wrist actigraphs and smartwatch applications

| Device types | Study | Sample size and population | Study focus | Video recording? (Yes/No) | Sensitivity | Specificity | Correlation | Accuracy | Study quality (1-5)* |
|--------------|--------------------------|--|--|---------------------------|---|---|--|---|----------------------|
| Actigraphy | Feuerstein ²³ | Healthy adults (n = 12) | Testing k-means cluster algorithm | No | 0.90 ± 0.10 | 0.98 ± 0.05 (walking) 0.88 ± 0.06 (restlessness) | | 0.92 (scratch) 0.92 (walking) 0.97 (restless sleep) | 3 |
| | Petersen ²⁹ | Healthy adults (n = 12) | Testing logistic regression algorithm | Yes | 0.96 (all data) 0.96 (cross-validation, mean) | 0.92 (all data) 0.92 (cross-validation, mean) | | | 3 |
| | Almazan ²⁸ | Healthy adults (n = 3), AD adults (n = 9) | Testing BRNN algorithm | Yes | | | _{r_s} (ρ) = 0.96 (actigraphy and video scoring) _{r_s} (ρ) = 0.90 (number of scratching events at home and polysomnography) | | 3 |
| | Moreau ²⁴ | Healthy adults (n = 6), AD adults (n = 18) | Testing BRNN algorithm compared to logistic regression | Yes | <u>AD:</u> 0.45-0.91 (BRNN) 0.00-0.10 (logistic regression) <u>Healthy:</u> 0.00-0.75 (BRNN) 0.00-0.50 (logistic regression) <u>Total:</u> 0.66 (BRNN) 0.06 (logistic regression) | | r ² = 0.98 r _s (ρ) = 0.95 (BRNN and video recording) | F1 scores: <u>AD:</u> 0.27-0.90 (BRNN) 0.00-0.14 (logistic regression) <u>Healthy:</u> 0.00-0.29 (BRNN) 0.00-0.08 (logistic regression) <u>Total:</u> 0.68 (BRNN) 0.09 (logistic regression) | |
| | Kurihara ¹² | Healthy adults (n = 10) | Actigraphy vs video recording and other devices for TST% calculation | Yes | | | RMSE = 5.32%-8.12% | | 2 |

| | | | | | | |
|-----------------------|--|---|-----|---|---|---|
| Murray ⁸ | <u>Study 1</u> : healthy subjects (n = 24; 12 adults, 12 children), pruritic subjects (n = 118; 68 adults, 50 children) <u>Study 2</u> : AD adults (n = 20) | Actigraphy vs VAS itch | No | | <u>Study 1</u> : $r^2 = 0.06$ <u>Study 2</u> : $r^2 = 0.08$ | 3 |
| Shino ³⁷ | Healthy adults (n = 1) | Actigraphy vs video recording and other devices for TST% extraction via novel algorithm | Yes | RMSE = 0.83s (0.64s) TST% error = +5.02% (+4.33%) (parentheses are from visually scoring outputs) | | 3 |
| Wootton ³³ | AD children (n = 336) | Actigraphy vs AD severity (SASSAD, POEM) | No | | r_s (ρ): SASSAD = 0.15 ($P = .02$) POEM = .10 ($P = .13$) | 3 |
| Hon ³⁰ | AD children (n = 24 for subjective surveys, n = 20 chemokines) | Actigraphy vs SCORAD scores and AD-associated chemokines | No | | r_s (ρ): †Total SCORAD = 0.52 ‡Objective SCORAD = 0.52 SCORAD pruritus = 0.23 SCORAD sleep loss = 0.36 †CTACK = 0.56 §MDC = 0.63 †TARC = 0.54 | 3 |
| Hon ³¹ | AD children (n = 28) | Actigraphy vs BDNF and substance P | No | | r_s (ρ): ‖BDNF = 0.83-0.91 ‖Substance P = .83-.87 | 3 |

Continued

Table I. Cont'd

| Device types | Study | Sample size and population | Study focus | Video recording? (Yes/No) | Sensitivity | Specificity | Correlation | Accuracy | Study quality (1-5)* |
|--------------|--------------------------|---|---|---------------------------|-------------|-------------|--|----------|----------------------|
| | Fujita ²⁷ | AD adults (n = 15) | Actigraphy vs SCORAD, VAS itch, serum cytokines | No | | | $r_s (\rho)$: \dagger VAS daytime itch = 0.58 \dagger SCORAD = 0.54 \dagger TARC = 0.51 \dagger LDH = 0.65 | | 3 |
| | Bender ⁴³ | Healthy adults (n = 14), AD adults (n = 14) | Actigraphic sleep measures vs VAS itch | No | | | $r_s (\rho)$: \dagger WASO = 0.35 \dagger Sleep efficiency = 0.38 \dagger Average sleep = 0.46 | | 3 |
| | Benjamin ²¹ | Healthy children (n = 7), AD children (n = 14) | Video recording (sleep time, scratch time, restlessness) vs actigraphy and VAS itch | Yes | | | $r_s (\rho)$: \ddagger Actigraphy, all > 0.92 VAS itch = 0.16-0.30 ($P > .05$) | | 3 |
| | Bringhurst ²⁶ | Pruritic subjects (n = 33 adults, n = 25 children), healthy subjects (n = 30 adults, n = 17 children) | Actigraphy vs subjective scores (VAS sleep, VAS itch, VAS skin disease), and SCORAD | No | | | $r_s (\rho)$: <u>Children:</u> \dagger VAS sleep = 0.48 \dagger VAS itch = 0.40 \dagger VAS skin disease = 0.49 \ddagger SCORAD = 0.62 <u>Adults:</u> \dagger VAS sleep = -0.44 VAS itch = 0.18 \dagger VAS skin disease = 0.15 \dagger SCORAD = 0.53 | | 3 |
| | Ebata ²⁵ | Healthy adults (n = 5), AD adults (n = 29) | Actigraphy vs video recording in TST% calculation | Yes | | | $\parallel r_s (\rho) = 0.91$ | | 3 |

| | | | | | | | | | | |
|----------------------------|------------------------|-------------------------------|--|-----|---|--|--|--|---|---|
| | Sandoval ³² | AD adults (n = 10) | Actigraphic WASO vs IGA and EASI at baseline and after 5-day fluocinonide 0.1% cream | No | | | | | r_s (ρ): †baseline EASI = 0.75 †baseline IGA = 0.76 ‡end treatment EASI = 0.70 †end treatment IGA = 0.73 | 3 |
| | Kaburagi ¹⁶ | Healthy adults (n = 12) | TST% estimation algorithm for various devices | Yes | | | | | RMSE = 4.29% (4.85%) (parentheses are from visual scoring of outputs for TST%) | 4 |
| Smartwatch applications | Ikoma ³⁶ | AD adults (n = 5) | “ItchTracker” (now “DermaTrack”) testing for scratch detection | Yes | 0.85 ± 0.10 | | | | R = 0.85-0.90 | 4 |
| | Lee ³⁴ | Healthy adults (n = 3) | “Itchtector” prototype testing | Yes | dominant hand = 0.98-1.00 nondominant hand = 0.63- 0.82 | dominant hand = 0.98-1.00 nondominant hand = 0.99 | | dominant hand = 0.985-0.99 nondominant hand = 0.933- 0.976 | | 3 |
| | Lee ³⁵ | Pruritic subjects (n = 13) | “Itchtector” testing in pruritic subjects | Yes | 0.75 | | | 0.90 | | 3 |

AD, Atopic dermatitis; BDNF, brain-derived neurotrophic factor; BRNN, bidirectional recurrent neural network; CTACK, cutaneous T-cell-attracting chemokine; EASI, Eczema Area and Severity Index; IGA, Investigator’s Global Assessment; r_s (ρ), Spearman’s rank correlation coefficient; LDH, lactate dehydrogenase; MDC, macrophage-derived chemokine; r^2 , coefficient of determination; RMSE, root mean square error; POEM, Patient-Oriented Eczema Measure; SASSAD, Six Area, Six Sign Atopic Dermatitis; SCORAD, SCORing Atopic Dermatitis; TARC, thymus and activation-regulated chemokine; TST%, total sleep time percentage; VAS, visual analog scale; WASO, wake after sleep onset.

*Study quality was assessed using a rating scheme modified from the Oxford Centre for Evidence-Based Medicine for ratings of individual studies: (1) properly powered and conducted randomized clinical trial or systematic review with meta-analysis; (2) well-designed controlled trial without randomization or prospective comparative cohort trial; (3) case-control study or retrospective cohort study; (4) case series with or without intervention or cross-sectional study; and (5) opinion of respected authorities or case reports.²²

† $p < .05$.

‡ $p < .01$.

§ $p < .005$.

|| $p < .001$.

Table II. Summary table for studies exploring acoustic, vibratory, pressure, and strain gauge devices. Note that no specificity values are reported for any of the studies listed

| Device type | Study | Sample size and population | Study focus | Video recording? (Yes/No) | Sensitivity | Correlation | Accuracy | Study quality (1-5)* |
|-------------|------------------------|---|---|---------------------------|-------------|---|--|----------------------|
| Acoustic | Kurihara ¹² | Healthy adults (n = 10) | Finger-mounted microphone vs video recording and other devices for TST% calculation | Yes | | | RMSE = 1.09% | 2 |
| | Noro ¹⁰ | Healthy adults (n = 8), AD adults (n = 4) | Wristwatch-type piezoelectric device for scratching rate compared to video recording | Yes | | r ² = 0.98 (nocturnal scratching rate by acoustic device vs video recording) | | 3 |
| Vibratory | Kurihara ¹⁸ | Healthy adults (n = 12) | Validation of piezoceramic disk devices placed under bed legs vs video recording for scratch and nonscratch | Yes | | | RMSE (staying calmly) = 0.35-0.72s RMSE (moving hand, turning over, moving foot) = 0.94-1.26s RMSE (scratching) = 0.56-1.29s | 3 |
| | Kurihara ¹² | Healthy adults (n = 10) | Piezoceramic disk bed devices placed under bed legs vs video recording and other devices for TST% calculation | Yes | | | RMSE = 0.87 = 6.31% | 3 |
| | Shino ³⁷ | Healthy adults (n = 1) | Piezoceramic bed devices vs video recording and other devices for TST% extraction via novel algorithm | Yes | | | RMSE = 0.68-0.79s (0.40-0.94s) TST% error = 2.13-4.11% (-6.51-0.82%) (parentheses are from visually scoring outputs) | 3 |
| | Kaburagi ¹⁶ | Healthy adults (n = 12) | TST% estimation algorithm for various devices | Yes | | | RMSE (left bed head) = 1.51% (1.84%) RMSE (right bed head) = 0.92% (1.86%) RMSE (left bed foot) = 6.58% (6.27%) | 4 |

| | Kogure ¹⁷ | AD subjects (n = 20) | Evaluation of sheet-shaped body vibrometer vs wrist actigraphy for measurement of scratching, activity count, and sleep efficiency | No | | r_s (ρ): activity count per minute = 0.63-0.82 [†] sleep efficiency = 0.82-0.91 [‡] | RMSE (right foot bed) = 3.97% (6.83%) (parentheses are from visual scoring of outputs for TST%) 3 |
|-----------------|------------------------|---|--|-----|--|--|---|
| Pressure Sensor | Endo ¹⁴ | Healthy adults (n = 10), AD adults (n = 20 total; 10 male, 10 female) | Evaluation of "Scratch Monitor" device on dorsal hand | No | 0.74 (overall) 0.65 (male) 0.83 (female) | | 3 |
| | Kurihara ¹² | Healthy adults (n = 10) | Ceramic sheet placed on dorsal hand vs video recording and other devices for TST% calculation | Yes | | RMSE = 0.73% | 3 |
| Strain Gauge | Kurihara ¹² | Healthy adults (n = 10) | Strain gauge on index finger vs video recording and other devices for TST% calculation | Yes | | RMSE = 2.41% | 3 |
| | Shino ³⁷ | Healthy adults (n = 1) | Strain gauge on index finger vs video recording and other devices for TST% extraction via novel algorithm | Yes | | RMSE = 0.53s (0.37s) TST% error = +1.38% (-1.54%) (parentheses are from visually scoring outputs) | 3 |
| | Kaburagi ¹⁶ | Healthy adults (n = 12) | TST% estimation algorithm for various devices | Yes | | RMSE = 1.29% (1.63%) (parentheses are from visual scoring of outputs for TST%) | 4 |

AD, Atopic dermatitis; r^2 , coefficient of determination; RMSE, root mean square error; r_s (ρ), Spearman's rank correlation coefficient; TST%, total sleep time percentage.

*Study quality was assessed using a rating scheme modified from the Oxford Centre for Evidence-Based Medicine for ratings of individual studies: (1) properly powered and conducted randomized clinical trial or systematic review with meta-analysis; (2) well-designed controlled trial without randomization or prospective comparative cohort trial; (3) case-control study or retrospective cohort study; (4) case series with or without intervention or cross-sectional study; and (5) opinion of respected authorities or case reports.²²

[†] $P < .005$.

[‡] $P < .001$.

Table III. Reported sensitivity of algorithms for scratch detection in studies focused on subjects with atopic dermatitis, which used video recording as comparison

| Performance metric | Actigraphy | Smartwatch applications |
|---------------------|---|----------------------------|
| Sensitivity (range) | 0.45-0.91 (BRNN) ²⁴ 0.00-0.10 (logistic regression) ²⁴ | 0.75-0.85 ^{34,36} |

BRNN, Bidirectional recurrent neural network.

but exhibit an equally high level of scratching restraint. In contrast, some individuals with chronic itch are habituated to it and report low scores despite frequent scratching. Ultimately, scratch measurements with objective tools and subject-reported outcomes are interrelated outputs that provide complementary information.

Smartwatch applications

Applications leveraging smartwatches and their accelerometers show comparable performance in detecting scratch when compared to actigraphs. Three articles examined smartwatch applications compared to video recording. In preliminary testing of their “Itchector” app, Lee et al³⁴ reported sensitivity (0.63-1.00), specificity (0.98-1.00), PPV (0.83-0.98), negative predictive value (0.93-1.00), and accuracy (93.3%-99.0%) in healthy adults (n = 3). When cross-validated in pruritic subjects (n = 13), the app yielded lower sensitivity (0.75), PPV (0.74), and accuracy (90%), which may be due to the small initial sample size, different subject populations, and different smartwatches.³⁵

Ikoma et al³⁶ also tested the “ItchTracker” app in adult AD subjects (n = 5) and reported a sensitivity of 0.85 and PPV of 0.90. They reported a correlation between the app and video recording for an hourly scratch duration of $r_s(\rho) = 0.851-0.901$ ($P < .001$). The authors further compared scratching duration percentage to current and 7-day itch in healthy and AD adults, reporting $r_s(\rho) = 0.36-0.43$ ($P < .001$). Similar findings were reported regarding self-reported sleep disturbance ($r_s(\rho) = 0.45$) and daytime disturbance ($r_s(\rho) = 0.42$). Disease severity measured by the Eczema and Area Severity Index was significantly correlated to scratching duration percentage ($r_s(\rho) = 0.60$).³⁶ However, they excluded finger-only scratching movements. Additionally, the small sample size should be taken into consideration. Although smartwatch applications show good sensitivity, there are no reported specificity ranges for pruritic subjects,

making it difficult to assess their ability to distinguish between scratch and nonscratch movements.

Acoustic

Acoustic devices detect sound waves generated from scratching. Two articles studied healthy subjects and compared the performance of their respective devices to that of video recording. No sensitivity or specificity values were reported. The finger-mounted microphone presented by Kurihara et al¹² yielded an RMSE of 1.09% for TST% calculation when compared to video recording. Noro et al¹⁰ reported $r^2 = 0.98$ when comparing scratching rate captured by their acoustic sensor and scratching rate obtained from video observation. While the devices show strong accuracy in detecting fine finger movements, the technology is not widely available and follow-up studies have not been conducted since first reported in 2014.

Vibratory

Vibratory devices allow for noninvasive monitoring of body movements and mitigate lesion exacerbation by devices that require skin contact. Four articles studied bed vibratory sensors compared to video recording.^{12,13,18,37} Accuracy was measured by RMSE, ranging 0.56-1.29s for scratching time¹⁸ and 0.87%-6.31% for TST% calculation.¹² Shino et al³⁷ reported comparable RMSE values for their TST% algorithm (0.68-0.79s) when compared to visually scored device outputs (0.40-0.94s) (n = 1). For both studies, the vibratory RMSE values were among the lowest when compared to other technologies. While vibratory devices have comparable accuracy to actigraphy and are largely burden-free once installed, their cost and setup may be deterrents.

Pressure sensors

Pressure sensors placed on the dorsal hand detect pressure changes with hand movements. Only 1 of 2 articles was compared to video recording. Kurihara et al¹² compared a ceramic sheet to other devices in healthy subjects, and reported a RMSE of 0.72% for TST% calculation when compared to video recording, the lowest among the devices tested. Although not compared to video recording, the Scratch Monitor pressure sensor presented by Endo et al¹⁴ was tested in healthy adults and yielded sensitivity ranging from 0.65-0.83.

Strain gauge

Strain gauges placed on the index finger to measure finger bending were evaluated in 2 studies, both of which were compared to video recording

Table IV. Comparison of various technologies used to detect scratching

| Device type | Benefits/pros | Limitations/cons | Algorithms for scratch detection |
|-------------------------|--|--|---|
| Actigraphy | <ul style="list-style-type: none"> • Most studied, has a large literature base • Validated against video recording • High sensitivity for wrist-dominant scratching movements in healthy subjects • Statistically significant moderate correlation with other objective measures (eg, SCORAD, IGA, EASI) | <ul style="list-style-type: none"> • Very poor correlation with subjective assessment tools for itch • Varied performance regarding scratch detection • Poor sensitivity for finger-dominant scratching movements • Deterioration of performance in pruritic subjects • Poor specificity given difficulty distinguishing wrist movements from scratching • False positives with similar waveforms (eg, walking) • Larger studies in target populations (eg, AD subjects) needed for algorithm development | <ul style="list-style-type: none"> • The k-means cluster analysis algorithm has good performance, but impractical in clinical setting given required determination of all movements a priori²³ • The BRNN model has good performance in pruritic subjects (albeit poorer than healthy subjects) and moderate F1-scores²⁴ • Logistic regression model in the study by Petersen et al²⁹ has comparable performance to k-means cluster analysis, but poorer performance in separate data set by Moreau et al²⁴ • Note that all of the aforementioned algorithms are for determination of TST |
| Smartwatch applications | <ul style="list-style-type: none"> • Similar to actigraphy in that it utilizes the smartwatch's built-in accelerometer, more convenient for current smartwatch owners • Bluetooth and cloud capabilities make accessing data easy for both patients and health care providers | <ul style="list-style-type: none"> • Few applications available • Some applications (eg, "DermaTrack", formerly called "ItchTracker") do not show raw data output • Smartwatches may be cumbersome for pediatric subjects, with no currently reported pediatric data | <ul style="list-style-type: none"> • Algorithm proposed by Lee et al^{34,35} reveals good accuracy in pruritic subjects, but authors report false negatives (eg, nonperiodic scratching) and false positives (nonscratching periodic movements such as arm shaking) |
| Acoustic | <ul style="list-style-type: none"> • Greater specificity with detection of scratch-generated sounds; will have different pattern than restlessness or turning over • Able to detect both finger and wrist scratching | <ul style="list-style-type: none"> • Limited research • Privacy concerns/risk • Unable to use in patients who do not sleep alone or have OSA | <ul style="list-style-type: none"> • Able to estimate TST% in healthy subjects with high accuracy (low RMSE compared to video recording) in healthy subjects |
| Vibratory | <ul style="list-style-type: none"> • Noninvasive • Able to localize scratching based on different waveforms | <ul style="list-style-type: none"> • Subject must use specific bed and unable to be used in patients who do not sleep alone | <ul style="list-style-type: none"> • Able to estimate TST% with variable accuracy depending on distance between the sensor and scratch site in healthy subjects |
| Pressure sensor | <ul style="list-style-type: none"> • Able to detect finger scratching if placed on dorsal hand along metatarsal bone (ceramic sheet) | <ul style="list-style-type: none"> • Performance dependent on technology • Eg, false positives from any hand movement that causes changes in pressure • Limited research | <ul style="list-style-type: none"> • Able to estimate TST% with high accuracy (low RMSE) due to distinct waveforms in healthy subjects |
| Strain gauge | <ul style="list-style-type: none"> • Higher sensitivity for finger-dominant scratching when placed on index finger compared to actigraphy | <ul style="list-style-type: none"> • False positives with non-scratch finger bending movements • Limited research | <ul style="list-style-type: none"> • Able to estimate TST% with good accuracy in healthy subjects |

AD, Atopic dermatitis; BRNN, bidirectional recurrent neural network; EASI, Eczema Area and Severity Index; IGA, Investigator's Global Assessment; N/A, not available; OSA, obstructive sleep apnea; RMSE, root mean square error; SCORAD, SCORing Atopic Dermatitis; TST, total scratch time; TST%, total sleep time percentage.

and tested in healthy subjects. The devices yielded an RMSE of 2.41% for TST% calculation, half that of wrist actigraphy.¹² The devices also yielded a TST% error of 1.38% when automatically extracted via an algorithm proposed by Shino et al,³⁷ which was compared to a TST% error of -1.54% when the data were visually scored. No sensitivity or specificity values were reported. It should be noted that strain gauges may be more susceptible to false positives (eg, nonscratching finger movements).

DISCUSSION

While the development of existing and novel devices has progressed tremendously, their performances reveal large areas in need of improvement. Actigraphy-based algorithms appear to have good sensitivity and specificity in healthy subjects; however, their performance deteriorates considerably when applied to pruritic subjects. This may be due to a lack of algorithm generalizability and failure to capture finger scratching. Additionally, most data used for establishing scratch parameters were obtained from small healthy samples. While there have been cross-validation studies with data from small AD samples, testing in larger samples of pruritic patients has not been performed. The same principle applies to newer scratch technologies, whereby further testing in both populations is needed for robust algorithms. While certain devices have demonstrated greater sensitivity for detecting finger scratching, the studies do not explicitly mention their abilities to detect rubbing or use of other scratching tools (eg, back scratchers). Rubbing, like scratching, is a natural reaction to itch; if devices are unable to distinguish rubbing or use of scratching tools from other motions, they may be underestimating itch. Further development of these technologies may help provide a more comprehensive picture of itch.

Performance metrics and algorithms

With advances in machine learning, data-driven approaches for objective scratch monitoring have gained significant interest. Various metrics have been employed to evaluate performance. While specificity and accuracy are useful, they need to be used with caution as they can be prone to class imbalances. Under typical situations, scratching arises sporadically, each over a brief period, ranging from a few seconds to several minutes depending on symptom severity. Thus, the majority of data collected features nonscratching behaviors; only a small amount of data feature scratching, resulting in a significant class imbalance. For example, a poor classification algorithm that predicts nonscratch all the time will, most likely, produce excellent accuracy and

specificity. Given this problem, other metrics, such as sensitivity, precision, and F1-score are deemed more appropriate to quantify performance.

Future considerations

While patient history and examination remain important tools in assessing itch, there remains an ongoing need for adjunctive objective and precise, tools to quantify itch, such as in the case of subconscious habitual scratching. Many technologies and algorithmic strategies have been studied, though their performances are highly variable, with validation studies rarely extending beyond small samples. In addition, most studies focus on nocturnal scratching. Given that the perception of itch varies during the day, daytime scratching remains an important behavior that is largely unstudied.

In this review, very few studies reported specificity values. While this is understandable in nocturnal scratching, during which the targeted behavior scratching is rare overall, daytime wear introduces other confounders, such as texting or walking. Thus, specificity may hold greater relevance in daytime wear, during which the wristwatch-based systems may struggle to differentiate scratching from other movements. Our group has introduced a novel mechano-acoustic skin device that incorporates actigraphy and acoustic detection of scratching by conforming to the dorsal hand and sampling at higher frequencies (~1600 Hz) compared to actigraphy (20-100 Hz). Scratch algorithm development performed in healthy subjects yielded high sensitivity and specificity with comparable performance among AD datasets using an IR camera gold standard, even with confounders.^{38,39} A comparison of data outputs for scratch from actigraphy, smartwatch application, and mechano-acoustic device is shown in Supplemental Fig 1.

CONCLUSION

While actigraphy remains the most frequently studied modality in clinical studies, performance is variable with no assessment of daytime performance. Further testing of these technologies will be needed before used in the clinical setting. A reliable technological modality would allow for objective support of drug development outcomes,⁴⁰⁻⁴² guide disease management, and assess treatment response.

Conflicts of interest

Drs Yang, Nguyen, Li, Lee, Chun, Wu, Fishbein, and Paller have no conflicts of interest to declare. Dr Xu has equity in a private company with a commercial interest in

scratch sensors and inventorship interest in patents related to a scratch sensor.

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