

Original Paper

Adherence to Psoriasis Remote Monitoring and Its Associations With Demographic, Clinical, and Health Care Use Characteristics: Multicenter Observational Study

Maarja Lember^{1*}, BSc; Priit Kruus^{1,2*}, MSc; Katrin Kaarna^{3,4*}, MD; Tanel Traks^{3,4}, MSc; Laura Prett⁵, MD; Liisi Raam⁴, MD, PhD; Oliver Taul⁴, MD; Liis Ilves⁴, MD, PhD; Kaisa Viljar⁴, MD; Pille Konno⁶, MD; Secil Matasova¹, BSc, MEng; Peeter Ross¹, MD, PhD; Külli Kingo^{3,4}, MD, PhD

¹Tallinn University of Technology, Tallinn, Harjumaa, Estonia

²Dermtest OÜ, Tallinn, Harjumaa, Estonia

³University of Tartu, Tartu, Tartu, Estonia

⁴Tartu University Hospital, Tartu, Tartu, Estonia

⁵OÜ Linnamõisa Perearstikeskus, Tallinn, Harjumaa, Estonia

⁶Confido Medical Centre, Tallinn, Harjumaa, Estonia

*these authors contributed equally

Corresponding Author:

Katrin Kaarna, MD

University of Tartu

L.Puusepa 8

Tartu, Tartu, 50406

Estonia

Phone: 372 737 4119

Email: katrin.kaarna@ut.ee

Abstract

Background: Psoriasis is a lifelong, relapsing, quick-to-flare skin disease that requires personalized, continuous management. Due to its unpredictable nature, there is an increasing interest in using electronic patient-reported outcome measures (ePROMs) for remote monitoring of the condition. However, adherence to ePROMs and predictors of adherence have not been sufficiently explored among patients with psoriasis.

Objective: This study aimed to assess the adherence to ePROMs and the factors associated with adherence among patients with psoriasis.

Methods: We conducted an observational study that spanned multiple centers to measure adherence to ePROMs among 55 patients managed in primary care and 55 patients managed in specialized care over 12 months. We analyzed the data together with longitudinal health care data recorded in Estonia's countrywide health insurance database—a unique dataset previously unavailable to researchers studying remote monitoring. While continuing their routine clinical care, patients were periodically prompted to complete ePROMs. Adherence to remote monitoring was evaluated in relation to a range of demographic, clinical, and health care use factors using ordinal logistic regression.

Results: A total of 102 patients (male: n=49, 48%; female: n=53, 52%) were included in the adherence calculation, including 46 (45.1%) patients in the high-adherence group, 22 (21.6%) in the medium-adherence group, and 34 (33.3%) in the low-adherence group. The total mean adherence ratio was 63.4% (SD 30.8%). The mean adherence ratio was 60.9% (SD 32.8%) among patients in primary care and 66.0% (SD 28.9%) among patients in specialized care. In multivariable ordinal logistic regression, older age was associated with higher adherence ($P=.03$). No significant associations were observed for the other investigated factors.

Conclusions: This study adds to existing research on the use of ePROMs in chronic diseases by demonstrating the feasibility of using ePROMs for remote monitoring of chronic skin diseases across different levels of care. The findings provide insights into factors associated with adherence, which may inform the development of tailored approaches to improve the effectiveness of remote monitoring in psoriasis. Importantly, adherence remained strong among older adults, supporting broader implementation of ePROM-based remote care.

KEYWORDS

psoriasis; remote monitoring; electronic patient-reported outcome; ePROM; adherence; eHealth; insurance database

Introduction

Remote monitoring with routine patient-reported outcome measures (PROMs) is an effective strategy for capturing vital disease information for efficient patient-physician communication [1]. Remote monitoring makes timely tracking of symptom progression possible [2] and raises physicians' awareness of patients' perceived health status [3], resulting in improved patient-centered care [4].

Remote monitoring with electronic PROMs (ePROMs) is also being considered for implementation in the care process for psoriasis [5-7], a chronic, remitting, and relapsing immune-mediated skin disease affecting more than 60 million people of all ages worldwide [8,9]. Psoriasis is associated with a range of comorbidities of varying severity, including psoriatic arthritis [10-15]. Psychological distress in patients is caused both by the visible presence of lesions on the skin and by the subjective experience of lesion-related pain and itching [16]. The severity of psoriasis fluctuates throughout a person's lifetime, and health care needs vary depending on disease severity [17]. The disease requires personalized ongoing monitoring and management to increase the duration of remission, improve quality of life [9,17], and reduce the risk of comorbidities [18-20]. Monitoring psoriasis with ePROMs shows promise in equipping physicians with better information [21].

The chronic and variable progression of psoriasis poses significant challenges to developing effective remote monitoring programs for its management [22-24]. eHealth solutions are potentially susceptible to low adherence [25], and the rate of patient correspondence "with agreed recommendations from a health care provider," in the words of a World Health Organization report [26], can be even lower in psoriasis monitoring [27]. Reaching sufficiently high adherence, which Donkin et al [28] defined as "the degree to which the user followed the program as it was designed," assumes that the program needs proper operational design and fits the health care context [29].

Understanding the possible factors that can hinder or support adherence is therefore important to ensure successful program design and sufficient adherence. Adherence to eHealth solutions is increasingly investigated, as inconsistent adherence limits the potential benefits of these solutions and could negatively impact treatment efficacy [25]. However, there are limited studies about which quantitative factors affect adherence to remote monitoring with ePROMs [30], specifically in chronic inflammatory skin disorders.

Previous research on adherence to remote monitoring systems or the predictors of high adherence among patients with psoriasis is limited. This study design is unique compared to existing remote monitoring programs in terms of its complexity (2 intervention groups, one at the primary care level and the other

at the specialized care level, across 9 locations) and the range of factors investigated. In this paper, we focused on remote psoriasis management with ePROMs to study adherence and adherence factors based on countrywide health insurance fund claims data. The data from all patient insurance claims across Estonia were linked with the adherence results of patients in the program. The recruitment period for remote monitoring was from January 2022 to June 2022, followed by 12 months of monitoring for each patient. Adherence to remote monitoring with ePROMs among patients with psoriasis treated in either primary care or a dermatology outpatient clinic over 12 months was analyzed, and possible predictive factors for adherence were identified.

Methods

Study Design

An observational study used multisite monitoring data collected from January 2022 to June 2023. The publicly reimbursed monitoring was conducted across 7 primary care centers and 1 dermatology outpatient department at Tartu University Hospital, all in Estonia. While patients continued with routine clinical care, they were requested to complete validated ePROMs commonly used in psoriasis management, including the Psoriasis Symptoms and Signs Diary (PSSD) [21], the Dermatology Life Quality Index (DLQI) [31], and the Early Arthritis for Psoriatic Patients (EARP) screening questionnaire [18]. Patients also had the opportunity to add photos to the questionnaires.

The web application could be used with any browser, either on a computer or a mobile device (iOS and Android). Patients were notified through email when a new ePROM became available for answering and were provided with a link to the web application. On the first day of every month, the PSSD was requested; on the 15th day of every month, the DLQI was requested; and the EARP was requested on days 90 and 270. The web application transmitted patient-generated data to a secure online platform. Physicians were notified and could then access the results, review them, and use the information as part of standard care provision.

A previous publication reported results from the same cohort and evaluated the effectiveness of remote psoriasis monitoring across primary and specialist care levels, demonstrating that specialist-led care was associated with significantly greater improvements in quality of life [32]. In contrast, the present study focuses on adherence to remote monitoring and aims to identify factors associated with successful ePROM completion.

Participants and Recruitment Procedure

In total, 110 patients were initially registered—55 (50%) from the dermatology outpatient department and 55 (50%) from 7 primary care centers. Eligible individuals were identified during an outpatient or primary care visit with their regular physician responsible for managing their psoriasis. Inclusion criteria were

as follows: (1) male and female outpatients aged 18 to 75 years, (2) plaque psoriasis diagnosed by a physician, (3) access to a computer or smartphone with internet access and a digital camera, and (4) an understanding of the study time commitment and provision of informed consent. Exclusion criteria were as follows: (1) a psoriasis subtype other than plaque psoriasis, (2) a history of alcohol or drug abuse within the 6 months prior to the study [33], (3) inability to comply with study requirements, and (4) other unspecified reasons that made the patient unsuitable for inclusion based on the discretion of the investigator.

For onboarding, the following activities were performed for each patient: completion of the physician-assessed Psoriasis Area and Severity Index (PASI) [34] questionnaire by the physician and initiation of the patient's remote monitoring protocol on the application platform. During the initial face-to-face onboarding visit, the physician was required to ensure that the patient could complete an ePROM via the web application. This was accomplished by sending an ePROM prompt to the patient's email and instructing them to complete it.

Determining Adherence Ratio

The adherence ratio was presented as a percentage and was determined by calculating the ratio of reported electronic patient-reported outcomes (ePROs) to the total number of ePROs that could have been reported [25]. Adherence was operationalized as "the number of necessary ePROMs completed and submitted within 30 days of receiving notification" [29]. This completion window was selected to evaluate overall engagement with remote monitoring over the 12-month period rather than strict compliance with scheduled dates. Considering the nature of the questionnaires, delays within this time frame were not expected to substantially affect the reported outcomes.

PSSD, DLQI, and EARP results were collected from the eHealth company database. Data were cleaned to eliminate any duplicate or incomplete entries for PSSD, DLQI, and EARP responses. The response dates of the ePROMs were assessed to ensure that responses were submitted within a 30-day period of receiving notification. ePROs completed during face-to-face onboarding sessions were excluded from the adherence calculations as these

did not accurately reflect patient-initiated adherence behavior. Adherence was assessed for each participant and summarized across the sample. Adherence was categorized into 3 categories: low (<50%), medium (50%-75%), and high (>75%).

Predictors of Adherence and Statistical Analysis

The characteristics of the study cohort are presented in Table 1. The investigated demographic data comprised sex and age. Psoriasis-specific factors comprised disease duration, treatment status, and baseline PASI scores. Investigated health care factors included the number of psoriasis-related comorbidities, patients' care level (primary and specialized care), and the total number of unique health care services used during the study period (primary care contact visits or primary care remote consultations and specialist contact visits or specialist remote consultations). The Estonian Health Insurance Fund (EHIF) data (ie, health care service type and service dates) were cross-checked with the patients' individual study dates. All comorbidity data pertaining to the year 2022 were comprehensively incorporated into the investigated dataset to minimize the risk of overlooking any diagnoses. Treatment data used in this study were collected from prescriptions fulfilled by patients during their individual study dates and the preceding 3-month period. The psoriasis duration for each patient was calculated from the first recorded psoriasis diagnosis code in the EHIF database.

Factors potentially associated with adherence were evaluated using multivariable ordinal logistic regression. Multicollinearity was assessed using variance inflation factors (VIFs) calculated from a linear regression model including all predictors. The proportional odds assumption was evaluated for each variable using a parallel lines test. PASI data were available for a subset of participants ($n=86$, 84.3%). Therefore, PASI was not included in the primary regression model to preserve statistical power. As a sensitivity analysis, the ordinal logistic regression was repeated among participants with available PASI data, with PASI included as an additional predictor. The stability of the regression coefficients and odds ratios (ORs) was compared between the primary and sensitivity models to ensure the robustness of the conclusions. All statistical analyses were conducted using Jamovi (version 2.6.44) with the GAMLj module for advanced generalized linear modeling diagnostics. A $P<.05$ was considered statistically significant.

Table 1. Characteristics of the study population (N=102).

Characteristics	Values
Sex, n (%)	
Male	49 (48)
Female	53 (52)
Age (years), mean (SD)	44.4 (10.4)
Psoriasis duration (years), mean (SD)	10.8 (6.3)
Treatment, n (%)	
Topical only	64 (62.7)
Systemic or biological	7 (6.9)
Systemic and topical	13 (12.7)
No treatment	18 (17.6)
Baseline Psoriasis Area and Severity Index score, mean (SD)	4.7 (6.3) ^a
Comorbidities, n (%)	
Hypertension	14 (13.7)
Cardiovascular disease	10 (9.8)
Dyslipidemia	12 (11.8)
Psoriatic arthritis	12 (11.8)
Mood disorders (depression and anxiety)	9 (8.8)
Obesity	8 (7.8)
Type 2 diabetes	2 (2)
Autoimmune thyroiditis	3 (2.9)
Care level, n (%)	
Primary care	51 (50)
Specialized care	51 (50)
Primary care clinic consultations, mean (SD)	2.5 (3.1)
Primary care remote consultations, mean (SD)	4.8 (3.5)
Specialist clinic consultations, mean (SD)	1.7 (1.6)
Specialist remote consultations, mean (SD)	0.2 (0.5)

^aPsoriasis Area and Severity Index data were available for 86 participants.

Ethical Considerations

The project was approved by the research ethics committee of the University of Tartu, Estonia (350/T-14). Eligible patients who understood the study time commitment were asked to provide informed consent. To protect privacy and confidentiality, the study used only anonymized data that had undergone a process of anonymization by authorized personnel involved in the study. Data were initially pseudonymized during queries of the respective databases, after which they were merged into a single database and then anonymized by assigning a new unique identifier to each patient. No incentives or compensation were offered to the patients.

Results

Among the 110 patients initially registered for the study, 8 (7.3%) did not complete any ePROM questionnaires.

Consequently, the adherence analysis included 102 (92.7%) participants who initiated the use of the remote monitoring program and contributed adherence data. The remaining cohort had an almost equal distribution of male (n=49, 48%) and female (n=53, 52%) participants. On the basis of the PASI scores at the start of the study, 68 (66.7%) patients had no to mild psoriasis (PASI <7) [35]. Prevalent comorbidities found in our sample population were hypertension and other cardiovascular diseases, dyslipidemia, psoriatic arthritis, mood disorders, obesity, type 2 diabetes, and autoimmune thyroiditis (Table 1). Patient characteristics did not differ significantly between primary care and specialist care, except for the higher proportion of patients in specialized care who received topical, systemic, or biological therapy, as shown in Table S1 in Multimedia Appendix 1.

The number of patients in each adherence group was as follows: 46 (45.1%) patients in the high-adherence group, 22 (21.6%)

in the medium-adherence group, and 34 (33.3%) in the low-adherence group. The overall mean adherence ratio was 63.4% (SD 30.8%). The mean adherence ratio was 60.9% (SD 32.8%) among patients in primary care and 66.0% (SD 28.9%) among patients in specialized care. There was a gradual decrease in the mean adherence ratio during the 12-month monitoring period, from 74.0% (SD 27.2%) in the first quarter to 62.8% (SD 39.3%) in the final quarter. Average adherence to the monthly ePROMs was 65.3% (SD 31.7%) for the DLQI and 64.2% (SD 32.7%) for the PSSD, whereas average adherence to the less frequently administered EARP was 47.5% (SD 39.4%). In the primary ordinal logistic regression analysis (N=102), increasing age was independently associated with higher ePROM adherence (OR 1.05, 95% CI 1.01-1.09; $P=.03$; Table 2). No other predictors reached statistical significance, although a greater number of primary care remote consultations and receipt of specialist care showed a trend toward significance

($P=.09$ and $P=.09$, respectively). Multicollinearity among the included predictors was negligible, with all VIF values remaining <2 . Parallel lines tests indicated that the proportional odds assumption was generally satisfied, with only primary care clinic consultations showing evidence of deviation ($P=.02$). PASI data were available for 86 (84.3%) participants and therefore were not included in the primary model. A secondary sensitivity analysis was restricted to participants with available PASI data and included PASI as an additional predictor (Table 2). The baseline PASI score was not associated with ePROM adherence ($P=.81$). Importantly, the inclusion of PASI did not meaningfully alter the model coefficients, and age remained a statistically significant predictor with an identical effect size (OR 1.05, 95% CI 1.00-1.10; $P=.04$). This finding supports the robustness of the primary findings and demonstrates their independence from baseline disease severity.

Table 2. Multivariable ordinal logistic regression analysis of factors associated with adherence to electronic patient-reported outcome measures (ePROMs).

Adherence factors	Primary model (N=102)		Sensitivity model (N=86)	
	Odds ratio (95% CI)	P value	Odds ratio (95% CI)	P value
Sex (female vs male ^a)	1.15 (0.52-2.57)	.73	0.99 (0.42-2.34)	.99
Age (years)	1.05 (1.01-1.09)	.03 ^b	1.05 (1.00-1.10)	.04 ^b
Psoriasis duration (years)	0.98 (0.92-1.04)	.48	0.98 (0.91-1.04)	.48
Treatment status (treatment vs no treatment ^a)	0.65 (0.21-2.00)	.46	0.89 (0.25-3.17)	.85
Baseline Psoriasis Area and Severity Index score	— ^c	—	0.99 (0.92-1.07)	.81
Number of comorbidities	0.97 (0.67-1.44)	.89	1.03 (0.67-1.58)	.91
Care level (specialized vs primary ^a)	2.39 (0.88-6.74)	.09	1.99 (0.71-5.70)	.19
Number of primary care clinic consultations	1.04 (0.90-1.23)	.59	0.98 (0.80-1.18)	.79
Number of primary care remote consultations	1.13 (0.99-1.29)	.09	1.04 (0.90-1.19)	.63
Number of specialist clinic consultations	1.01 (0.78-1.32)	.97	0.97 (0.72-1.31)	.84
Number of specialist remote consultations	0.60 (0.24-1.37)	.25	0.53 (0.19-1.30)	.20

^aReference category.

^bSignificant difference ($P<.05$).

^cNot estimated because the variable was not included in the model.

Discussion

Principal Results

The overall mean adherence ratio of 63.4% (SD 30.8%; N=102) is comparable to adherence rates of 49% to 96% reported in other studies that used repetitive ePROMs in noncommunicable disease management [36-42], although ePROMs have not been studied in the context of psoriasis. These findings suggest that patients with psoriasis can successfully engage in regular remote monitoring with ePROMs in both primary and specialized care settings. Average adherence decreased from 74.0% (SD 27.2%) during the first quarter to 62.8% (SD 39.3%) during the final quarter, suggesting reduced engagement over time. Lower overall completion rates were observed for the EARP, which was requested twice during the monitoring period, than for the

monthly DLQI and PSSD. This may indicate that participants perceived it as less relevant if they did not experience musculoskeletal symptoms related to psoriatic arthritis.

Most of the factors examined in the countrywide insurance claims registry were not associated with adherence, which is broadly consistent with prior studies that evaluated similar factors when analyzing adherence to PROs [36-42]. However, the results showed an association between increasing age and adherence ($P=.03$).

Previous studies have reported mixed findings regarding the relationship between older age and higher adherence compared to younger individuals. Our finding is therefore not entirely consistent with previous literature, although a systematic review of 5 studies on telemonitoring adherence identified 1 (by Colls et al [41]) showing “that patients over 65 years old had higher

adherence compared to patients <45 years old” [30]. The other studies covered in the mentioned systematic review did not show age as a predictor of adherence. Possibly relevant, another study detected an age bias in email-based PROM recruitment, reporting that older patients were more likely to respond [43]. Although age was not significantly associated with adherence in that sample, the older participants who joined may represent a more motivated and responsive subgroup, indicating survivor or adopter bias.

Other investigated factors (sex, disease duration, baseline psoriasis severity, treatment status, number of comorbidities, care level [primary or specialized], and number of health care consultations [across primary and specialist care, both clinic based and remote]) were not associated with adherence, which is consistent with existing literature on adherence to ePROM-based monitoring of chronic diseases, although some discrepancies exist. For example, in patients with rheumatoid arthritis, older age [41,42] and lower symptom severity [41] have been related to better adherence, whereas higher symptom severity has also been associated with better adherence when coupled with chronic pain [40]. Similarly, primary care management was identified as an adherence facilitator in patients with heart failure [37], while our study did not identify any significant differences in adherence between care levels. A publication using the same dataset demonstrated clear differences in outcomes between primary and specialist care groups [32], while the current paper did not find significant differences in patient adherence between the 2 care levels.

Previous adherence studies have often had an overrepresentation of female participants [38-41], but this study had an equal sex composition. Despite the finding that female patients had a higher average adherence ratio than male patients (mean 64.1%, SD 33.3% vs mean 62.7%, SD 28.6%), sex was not a statistically significant predictor of adherence, consistent with prior studies [37-41]. The findings imply that future research should concentrate on comprehensive data collection incorporating additional categories of factors that can significantly influence adherence, such as patient-level demographics, including marital status, educational level, employment status, and income [44]. Moreover, contextualizing nonadherence through interviews and personalizing remote monitoring with psychosocial or educational strategies could be useful for developing more effective, tailored interventions for chronic conditions such as psoriasis.

Limitations

The adherence factors investigated were limited to those accessible from the EHIF database, providing a unique set of potential predictors gathered at a national level that lacks some factors often examined in adherence studies. Additionally, potential attrition and volunteer biases due to purposive sampling, data cleaning, and nonrandomization were not investigated. However, we acknowledge the possibility that such biases might exist, as volunteering patients may have had

different attitudes toward completing the ePROMs than nonvolunteering patients. Furthermore, the requirement for smartphone or computer access may have selected more digitally capable patients, including older adults. In addition, the study population was largely homogeneous, which meant that the generally underexplored sociocultural and racial determinants of health could not be assessed. Therefore, this may limit the generalizability of our findings to more diverse populations. Literature indicates that eHealth solutions incorporating psychosocial or educational components have demonstrated increased quality of life [7,45,46], and these elements were not included as part of our intervention. Thus, a key limitation is the lack of data on education, income, marital status, digital literacy, work status, motivation, satisfaction, and technology access, all of which may influence adherence and should be examined in future research. Another limitation is the absence of baseline PASI data for 15.7% (16/102) of participants. Consequently, disease severity could only be evaluated in a sensitivity analysis conducted on a reduced sample, although inclusion of PASI did not materially alter the findings. The regression analyses included a relatively large number of potential predictors in relation to the sample size. Although this may have reduced the precision of individual coefficient estimates and increased the risk of overparameterization, the variables were selected to examine their associations with adherence rather than to develop a predictive model. The observed association between age and adherence remained consistent in the sensitivity analysis, including PASI, supporting the robustness of the primary finding.

Conclusions

This study contributes to the unique niche of improving the understanding of remote monitoring with ePROMs in patients with psoriasis, providing insights for selecting factors and designing quality interventions. Although some studies have evaluated remote monitoring as a long-term, sustainable management option for different chronic conditions, there is a general lack of research on the penetration, acceptability, and longitudinal sustainability of remote monitoring in psoriasis. In this study, 2 equal-sized intervention groups with very similar characteristics across primary and specialist care levels showed similar adherence. Of the 11 factors investigated, only 1 factor was found to be associated with adherence. Thus, the study builds on existing research while highlighting the massive need for future studies to expand the scope of possible adherence predictors, including psychosocial and socioeconomic factors.

Remote monitoring is an underused method of disease management that could potentially lead to improved clinical outcomes. However, the effect of the delivery methods for ePROMs, as well as the sociological factors that may play into the disease perception, is understudied. Multifaceted research that investigates adherence in conjunction with individual disease perception, education, and clinical outcomes is necessary to realize the full potential of remote monitoring.

Acknowledgments

The authors thank Anneli Vuks, product manager at Dermtest OÜ, for her help with data curation as well as for her tremendous support to the team. The authors declare that they used AI tools (ChatGPT [version 5.5; OpenAI] and Gemini [version 3.5;

Alphabet Inc]) to improve the English language and obtain feedback on selected draft statements. The authors retained full responsibility for the manuscript's content and independently evaluated, revised, and verified all text.

Funding

The study was funded by the Estonian Health Insurance Fund Model Project Competition (grant 161; April 28, 2021), Enterprise Estonia Program for Applied Research (grant 1.1-5.1/21/2144; December 22, 2021), and the Estonian Research Council (grant PRG1189). The research was conducted using the research infrastructure "National Centre for Translational and Clinical Research," funded by the Estonian Research Council (grant TARISTU24-TK22).

Authors' Contributions

Conceptualization: ML, P Kruus, SM

Data analysis: K Kaarna, TT

Data curation: ML, P Kruus

Formal analysis: ML, TT, SM

Funding acquisition: P Kruus, K Kingo

Investigation: ML

Methodology: ML, P Kruus, SM, PR

Project administration: P Kruus, K Kaarna

Resources: P Kruus

Software: P Kruus

Supervision: P Kruus, SM, K Kingo

Validation: ML, P Kruus, LP, LR, OT, LI, KV, P Konno, SM, K Kingo

Writing—original draft: ML

Writing—review and editing: P Kruus, K Kaarna, TT, LP, LR, OT, LI, KV, P Konno, SM, PR, K Kingo

Conflicts of Interest

P Kruus is a management board member and owns shares in Dermtest OÜ, the eHealth company providing the solution for remote monitoring. SM was employed as the head of research at Dermtest OÜ at the time the study was conducted. PR owns shares in Dermtest OÜ. Tartu University Hospital was the lead institution for the research project, and the role of Dermtest OÜ was to provide the remote monitoring platform. Statistical analyses were performed independently by investigators without affiliations to Dermtest OÜ using the raw study data. All other authors declare no other conflicts of interest.

Multimedia Appendix 1

Adherence predictive factors among primary care and specialized care patients.

[\[DOCX File, 22 KB-Multimedia Appendix 1\]](#)

References

1. Gibbons C, Porter I, Gonçalves-Bradley DC, Stoilov S, Ricci-Cabello I, Tsangaris E, et al. Routine provision of feedback from patient-reported outcome measurements to healthcare providers and patients in clinical practice. *Cochrane Database Syst Rev*. Oct 12, 2021;10(10):CD011589. [\[FREE Full text\]](#) [doi: [10.1002/14651858.CD011589.pub2](https://doi.org/10.1002/14651858.CD011589.pub2)] [Medline: [34637526](#)]
2. Austin L, Sharp CA, van der Veer SN, Machin M, Humphreys J, Mellor P, et al. Providing 'the bigger picture': benefits and feasibility of integrating remote monitoring from smartphones into the electronic health record. *Rheumatology (Oxford)*. Feb 01, 2020;59(2):367-378. [\[FREE Full text\]](#) [doi: [10.1093/rheumatology/kez207](https://doi.org/10.1093/rheumatology/kez207)] [Medline: [31335942](#)]
3. Shaw Y, Courvoisier DS, Scherer A, Ciurea A, Lehmann T, Jaeger VK, et al. Impact of assessing patient-reported outcomes with mobile apps on patient-provider interaction. *RMD Open*. Apr 2021;7(1):e001566. [\[FREE Full text\]](#) [doi: [10.1136/rmdopen-2021-001566](https://doi.org/10.1136/rmdopen-2021-001566)] [Medline: [33811177](#)]
4. Trettin B, Danbjørg DB, Andersen F, Feldman S, Agerskov H. An mHealth app to support patients with psoriasis in relation to follow-up consultations: qualitative study. *JMIR Dermatol*. Jun 08, 2021;4(1):e28882. [\[FREE Full text\]](#) [doi: [10.2196/28882](https://doi.org/10.2196/28882)] [Medline: [37632803](#)]
5. Garzorz-Stark N, Beicht S, Baghin V, Stark SP, Biedermann T, Lauffer F. IMPROVE 1.0: individual monitoring of psoriasis activity by regular online app questionnaires and outpatient visits. *Front Med (Lausanne)*. Jun 22, 2021;8:648233. [\[FREE Full text\]](#) [doi: [10.3389/fmed.2021.648233](https://doi.org/10.3389/fmed.2021.648233)] [Medline: [34239885](#)]
6. Zhu B, Wang Y, Zhou X, Cao C, Zong Y, Zhao X, et al. A controlled study of the feasibility and efficacy of a cloud-based interactive management program between patients with psoriasis and physicians. *Med Sci Monit*. Feb 04, 2019;25:970-976. [\[FREE Full text\]](#) [doi: [10.12659/MSM.913304](https://doi.org/10.12659/MSM.913304)] [Medline: [30713334](#)]

7. Domogalla L, Beck A, Schulze-Hagen T, Herr R, Benecke J, Schmieder A. Impact of an eHealth smartphone app on the mental health of patients with psoriasis: prospective randomized controlled intervention study. *JMIR Mhealth Uhealth*. Oct 25, 2021;9(10):e28149. [[FREE Full text](#)] [doi: [10.2196/28149](https://doi.org/10.2196/28149)] [Medline: [34431478](#)]
8. Greaves MW, Weinstein GD. Treatment of psoriasis. *N Engl J Med*. Mar 02, 1995;332(9):581-588. [doi: [10.1056/NEJM199503023320907](https://doi.org/10.1056/NEJM199503023320907)] [Medline: [7838193](#)]
9. Griffiths CE, Armstrong AW, Gudjonsson JE, Barker JN. Psoriasis. *Lancet*. Apr 03, 2021;397(10281):1301-1315. [doi: [10.1016/S0140-6736\(20\)32549-6](https://doi.org/10.1016/S0140-6736(20)32549-6)] [Medline: [33812489](#)]
10. Ogdie A, Weiss P. The epidemiology of psoriatic arthritis. *Rheum Dis Clin North Am*. Nov 2015;41(4):545-568. [[FREE Full text](#)] [doi: [10.1016/j.rdc.2015.07.001](https://doi.org/10.1016/j.rdc.2015.07.001)] [Medline: [26476218](#)]
11. Choudhary S, Pradhan D, Pandey A, Khan MK, Lall R, Ramesh V, et al. The association of metabolic syndrome and psoriasis: a systematic review and meta-analysis of observational study. *Endocr Metab Immune Disord Drug Targets*. 2020;20(5):703-717. [doi: [10.2174/1871530319666191008170409](https://doi.org/10.2174/1871530319666191008170409)] [Medline: [31595859](#)]
12. Khan SR, Bano A, Wakkee M, Korevaar TI, Franco OH, Nijsten TE, et al. The association of autoimmune thyroid disease (AITD) with psoriatic disease: a prospective cohort study, systematic review and meta-analysis. *Eur J Endocrinol*. Oct 2017;177(4):347-359. [doi: [10.1530/EJE-17-0397](https://doi.org/10.1530/EJE-17-0397)] [Medline: [28747386](#)]
13. Ruffilli I, Ragusa F, Benvenga S, Vita R, Antonelli A, Fallahi P, et al. Psoriasis, psoriatic arthritis, and thyroid autoimmunity. *Front Endocrinol*. Jun 19, 2017;8:139. [[FREE Full text](#)] [doi: [10.3389/fendo.2017.00139](https://doi.org/10.3389/fendo.2017.00139)] [Medline: [28674524](#)]
14. Fu Y, Lee CH, Chi CC. Association of psoriasis with inflammatory bowel disease: a systematic review and meta-analysis. *JAMA Dermatol*. Dec 01, 2018;154(12):1417-1423. [[FREE Full text](#)] [doi: [10.1001/jamadermatol.2018.3631](https://doi.org/10.1001/jamadermatol.2018.3631)] [Medline: [30422277](#)]
15. Vaengebjerger S, Skov L, Egeberg A, Loft ND. Prevalence, incidence, and risk of cancer in patients with psoriasis and psoriatic arthritis: a systematic review and meta-analysis. *JAMA Dermatol*. Apr 01, 2020;156(4):421-429. [[FREE Full text](#)] [doi: [10.1001/jamadermatol.2020.0024](https://doi.org/10.1001/jamadermatol.2020.0024)] [Medline: [32074260](#)]
16. Dowlatabadi EA, Wakkee M, Arends LR, Nijsten T. The prevalence and odds of depressive symptoms and clinical depression in psoriasis patients: a systematic review and meta-analysis. *J Invest Dermatol*. Jun 2014;134(6):1542-1551. [[FREE Full text](#)] [doi: [10.1038/jid.2013.508](https://doi.org/10.1038/jid.2013.508)] [Medline: [24284419](#)]
17. Khoury LR, Skov L, Møller T. Facing the dilemma of patient-centred psoriasis care: a qualitative study identifying patient needs in dermatological outpatient clinics. *Br J Dermatol*. Aug 2017;177(2):436-444. [doi: [10.1111/bjd.15292](https://doi.org/10.1111/bjd.15292)] [Medline: [28032892](#)]
18. Tinazzi I, Adami S, Zanolin EM, Caimmi C, Confente S, Girolomoni G, et al. The early psoriatic arthritis screening questionnaire: a simple and fast method for the identification of arthritis in patients with psoriasis. *Rheumatology (Oxford)*. Nov 2012;51(11):2058-2063. [doi: [10.1093/rheumatology/kes187](https://doi.org/10.1093/rheumatology/kes187)] [Medline: [22879464](#)]
19. Pearlman RA, Uhlmann RF. Quality of life in chronic diseases: perceptions of elderly patients. *J Gerontol*. Mar 1988;43(2):M25-M30. [doi: [10.1093/geronj/43.2.m25](https://doi.org/10.1093/geronj/43.2.m25)] [Medline: [3346521](#)]
20. Kimball AB, Gladman D, Gelfand JM, Gordon K, Horn EJ, Korman NJ, et al. National Psoriasis Foundation clinical consensus on psoriasis comorbidities and recommendations for screening. *J Am Acad Dermatol*. Jun 2008;58(6):1031-1042. [[FREE Full text](#)] [doi: [10.1016/j.jaad.2008.01.006](https://doi.org/10.1016/j.jaad.2008.01.006)] [Medline: [18313171](#)]
21. Mathias SD, Feldman SR, Crosby RD, Colwell HH, McQuarrie K, Han C. Measurement properties of a patient-reported outcome measure assessing psoriasis severity: the psoriasis symptoms and signs diary. *J Dermatolog Treat*. Aug 2016;27(4):322-327. [[FREE Full text](#)] [doi: [10.3109/09546634.2015.1114567](https://doi.org/10.3109/09546634.2015.1114567)] [Medline: [26634943](#)]
22. Duarte GV, Calmon H, Radel G, de Fátima Paim de Oliveira M. Psoriasis and sexual dysfunction: links, risks, and management challenges. *Psoriasis (Auckl)*. Dec 10, 2018;8:93-99. [[FREE Full text](#)] [doi: [10.2147/PTT.S159916](https://doi.org/10.2147/PTT.S159916)] [Medline: [30574453](#)]
23. Feldman SR, Regnier SA, Chirilov A, Hey F, Gilloteau I, Cella D. Patient-reported outcomes are important elements of psoriasis treatment decision making: a discrete choice experiment survey of dermatologists in the United States. *J Am Acad Dermatol*. Jun 2019;80(6):1650-1657. [[FREE Full text](#)] [doi: [10.1016/j.jaad.2019.01.039](https://doi.org/10.1016/j.jaad.2019.01.039)] [Medline: [30703461](#)]
24. Oji V, Luger TA. The skin in psoriasis: assessment and challenges. *Clin Exp Rheumatol*. 2015;33(5 Suppl 93):S14-S19. [Medline: [26472560](#)]
25. Jakob R, Harperink S, Rudolf AM, Fleisch E, Haug S, Mair JL, et al. Factors influencing adherence to mHealth apps for prevention or management of noncommunicable diseases: systematic review. *J Med Internet Res*. May 25, 2022;24(5):e35371. [[FREE Full text](#)] [doi: [10.2196/35371](https://doi.org/10.2196/35371)] [Medline: [35612886](#)]
26. Adherence to long-term therapies: evidence for action. World Health Organization. 2003. URL: <https://iris.who.int/items/bf8058c0-03b2-4b47-838f-5534849927fb> [accessed 2026-07-19]
27. Sagahyroon A. Remote patients monitoring: challenges. In: *Proceedings of the 2017 IEEE 7th Annual Computing and Communication Workshop and Conference*. 2017. Presented at: CCWC 2017; Jan 9-11, 2017; Las Vegas, NV. [doi: [10.1109/ccwc.2017.7868460](https://doi.org/10.1109/ccwc.2017.7868460)]
28. Donkin L, Christensen H, Naismith SL, Neal B, Hickie IB, Glozier N. A systematic review of the impact of adherence on the effectiveness of e-therapies. *J Med Internet Res*. Aug 05, 2011;13(3):e52. [[FREE Full text](#)] [doi: [10.2196/jmir.1772](https://doi.org/10.2196/jmir.1772)] [Medline: [21821503](#)]

29. Sieverink F, Kelders SM, van Gemert-Pijnen JE. Clarifying the concept of adherence to eHealth technology: systematic review on when usage becomes adherence. *J Med Internet Res*. Dec 06, 2017;19(12):e402. [FREE Full text] [doi: [10.2196/jmir.8578](https://doi.org/10.2196/jmir.8578)] [Medline: [29212630](https://pubmed.ncbi.nlm.nih.gov/29212630/)]
30. Wiegel J, Seppen B, van der Leeden M, van der Esch M, de Vries R, Bos W. Adherence to telemonitoring by electronic patient-reported outcome measures in patients with chronic diseases: a systematic review. *Int J Environ Res Public Health*. Sep 27, 2021;18(19):10161. [FREE Full text] [doi: [10.3390/ijerph181910161](https://doi.org/10.3390/ijerph181910161)] [Medline: [34639463](https://pubmed.ncbi.nlm.nih.gov/34639463/)]
31. Finlay AY, Khan GK. Dermatology Life Quality Index (DLQI)--a simple practical measure for routine clinical use. *Clin Exp Dermatol*. May 1994;19(3):210-216. [doi: [10.1111/j.1365-2230.1994.tb01167.x](https://doi.org/10.1111/j.1365-2230.1994.tb01167.x)] [Medline: [8033378](https://pubmed.ncbi.nlm.nih.gov/8033378/)]
32. Arsenjeva J, Kruus P, Hallik R, Matasova S, Prett L, Kaarna K, et al. Remote monitoring of psoriasis: comparing care models and evaluating quality of life outcomes: mixed methods study. *J Med Internet Res*. Jun 03, 2025;27:e73664. [FREE Full text] [doi: [10.2196/73664](https://doi.org/10.2196/73664)] [Medline: [40459920](https://pubmed.ncbi.nlm.nih.gov/40459920/)]
33. Mazilu R, Ziehfrend S, Biedermann T, Zink A. Patterns of addiction in chronic skin diseases: a comparative analysis of addictions and influencing factors in atopic dermatitis and psoriasis. *Acta Derm Venereol*. Jan 03, 2025;105:adv41350. [FREE Full text] [doi: [10.2340/actadv.v105.41350](https://doi.org/10.2340/actadv.v105.41350)] [Medline: [39750041](https://pubmed.ncbi.nlm.nih.gov/39750041/)]
34. Ashcroft DM, Wan Po AL, Williams HC, Griffiths CE. Clinical measures of disease severity and outcome in psoriasis: a critical appraisal of their quality. *Br J Dermatol*. Aug 01, 1999;141(2):185-191. [doi: [10.1046/j.1365-2133.1999.02963.x](https://doi.org/10.1046/j.1365-2133.1999.02963.x)] [Medline: [10468786](https://pubmed.ncbi.nlm.nih.gov/10468786/)]
35. Schmitt J, Wozel G. The psoriasis area and severity index is the adequate criterion to define severity in chronic plaque-type psoriasis. *Dermatology*. 2005;210(3):194-199. [doi: [10.1159/000083509](https://doi.org/10.1159/000083509)] [Medline: [15785046](https://pubmed.ncbi.nlm.nih.gov/15785046/)]
36. Wiegel J, Seppen BF, Nurmohamed MT, Bos WH, Ter Wee MM. Who stop telemonitoring disease activity and who adhere: a prospective cohort study of patients with inflammatory arthritis. *BMC Rheumatol*. Nov 30, 2022;6(1):73. [FREE Full text] [doi: [10.1186/s41927-022-00303-w](https://doi.org/10.1186/s41927-022-00303-w)] [Medline: [36447263](https://pubmed.ncbi.nlm.nih.gov/36447263/)]
37. Guzman-Clark JR, van Servellen G, Chang B, Menten J, Hahn TJ. Predictors and outcomes of early adherence to the use of a home telehealth device by older veterans with heart failure. *Telemed J E Health*. Mar 2013;19(3):217-223. [doi: [10.1089/tmj.2012.0096](https://doi.org/10.1089/tmj.2012.0096)] [Medline: [23268695](https://pubmed.ncbi.nlm.nih.gov/23268695/)]
38. Jamilloux Y, Sarabi M, Kérever S, Boussey N, le Sidaner A, Valgublas V, et al. Adherence to online monitoring of patient-reported outcomes by patients with chronic inflammatory diseases: a feasibility study. *Lupus*. Nov 2015;24(13):1429-1436. [doi: [10.1177/0961203315585814](https://doi.org/10.1177/0961203315585814)] [Medline: [25966927](https://pubmed.ncbi.nlm.nih.gov/25966927/)]
39. Rosen D, McCall JD, Primack BA. Telehealth protocol to prevent readmission among high-risk patients with congestive heart failure. *Am J Med*. Nov 2017;130(11):1326-1330. [doi: [10.1016/j.amjmed.2017.07.007](https://doi.org/10.1016/j.amjmed.2017.07.007)] [Medline: [28756266](https://pubmed.ncbi.nlm.nih.gov/28756266/)]
40. Ross EL, Jamison RN, Nicholls L, Perry BM, Nolen KD. Clinical integration of a smartphone app for patients with chronic pain: retrospective analysis of predictors of benefits and patient engagement between clinic visits. *J Med Internet Res*. Apr 16, 2020;22(4):e16939. [FREE Full text] [doi: [10.2196/16939](https://doi.org/10.2196/16939)] [Medline: [32297871](https://pubmed.ncbi.nlm.nih.gov/32297871/)]
41. Colls J, Lee YC, Xu C, Corrigan C, Lu F, Marquez-Gras G, et al. Patient adherence with a smartphone app for patient-reported outcomes in rheumatoid arthritis. *Rheumatology (Oxford)*. Jan 05, 2021;60(1):108-112. [doi: [10.1093/rheumatology/keaa202](https://doi.org/10.1093/rheumatology/keaa202)] [Medline: [32572490](https://pubmed.ncbi.nlm.nih.gov/32572490/)]
42. Wiegel J, Seppen BF, Nurmohamed MT, Ter Wee MM, Bos WH. Predictors for response to electronic patient-reported outcomes in routine care in patients with rheumatoid arthritis: a retrospective cohort study. *Rheumatol Int*. Apr 2023;43(4):651-657. [FREE Full text] [doi: [10.1007/s00296-023-05278-6](https://doi.org/10.1007/s00296-023-05278-6)] [Medline: [36715728](https://pubmed.ncbi.nlm.nih.gov/36715728/)]
43. Heilemann G, Georg D, Dobiasch M, Widder J, Renner A. Automation of ePROMs in radiation oncology and its impact on patient response and bias. *Radiother Oncol*. Oct 2024;199:110427. [FREE Full text] [doi: [10.1016/j.radonc.2024.110427](https://doi.org/10.1016/j.radonc.2024.110427)] [Medline: [39002570](https://pubmed.ncbi.nlm.nih.gov/39002570/)]
44. Jacob C, Sezgin E, Sanchez-Vazquez A, Ivory C. Sociotechnical factors affecting patients' adoption of mobile health tools: systematic literature review and narrative synthesis. *JMIR Mhealth Uhealth*. May 05, 2022;10(5):e36284. [FREE Full text] [doi: [10.2196/36284](https://doi.org/10.2196/36284)] [Medline: [35318189](https://pubmed.ncbi.nlm.nih.gov/35318189/)]
45. Balato N, Megna M, Di Costanzo L, Balato A, Ayala F. Educational and motivational support service: a pilot study for mobile-phone-based interventions in patients with psoriasis. *Br J Dermatol*. Jan 2013;168(1):201-205. [doi: [10.1111/j.1365-2133.2012.11205.x](https://doi.org/10.1111/j.1365-2133.2012.11205.x)] [Medline: [23240729](https://pubmed.ncbi.nlm.nih.gov/23240729/)]
46. Bundy C, Pinder B, Bucci S, Reeves D, Griffiths CE, Tarrier N. A novel, web-based, psychological intervention for people with psoriasis: the electronic Targeted Intervention for Psoriasis (eTIPs) study. *Br J Dermatol*. Aug 2013;169(2):329-336. [doi: [10.1111/bjd.12350](https://doi.org/10.1111/bjd.12350)] [Medline: [23551271](https://pubmed.ncbi.nlm.nih.gov/23551271/)]

Abbreviations

DLQI: Dermatology Life Quality Index
EARP: Early Arthritis for Psoriatic Patients
EHIF: Estonian Health Insurance Fund
ePRO: electronic patient-reported outcome
ePROM: electronic patient-reported outcome measure

OR: odds ratio

PASI: Psoriasis Area and Severity Index

PROM: patient-reported outcome measure

PSSD: Psoriasis Symptoms and Signs Diary

VIF: variance inflation factor

Edited by J Sarvestan; submitted 17.Dec.2025; peer-reviewed by S Feldman, J Ren; comments to author 01.May.2026; revised version received 30.Jun.2026; accepted 01.Jul.2026; published 13.Aug.2026

Please cite as:

Lember M, Kruus P, Kaarna K, Traks T, Prett L, Raam L, Taul O, Ilves L, Viljar K, Konno P, Matasova S, Ross P, Kingo K
Adherence to Psoriasis Remote Monitoring and Its Associations With Demographic, Clinical, and Health Care Use Characteristics:
Multicenter Observational Study

JMIR Form Res 2026;10:e89706

URL: <https://formative.jmir.org/2026/1/e89706>

doi: [10.2196/89706](https://doi.org/10.2196/89706)

PMID:

©Maarja Lember, Priit Kruus, Katrin Kaarna, Tanel Traks, Laura Prett, Liisi Raam, Oliver Taul, Liis Ilves, Kaisa Viljar, Pille Konno, Secil Matasova, Peeter Ross, Külli Kingo. Originally published in JMIR Formative Research (<https://formative.jmir.org>), 13.Aug.2026. This is an open-access article distributed under the terms of the Creative Commons Attribution License (<https://creativecommons.org/licenses/by/4.0/>), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work, first published in JMIR Formative Research, is properly cited. The complete bibliographic information, a link to the original publication on <https://formative.jmir.org>, as well as this copyright and license information must be included.