ORIGINAL RESEARCH



Demographic and Clinical Factors Associated with Patient-Reported Remission in Psoriatic Arthritis

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ABSTRACT

Introduction: Achievement of remission in psoriatic arthritis is a key goal for patients and clinicians, yet definitions of remission may vary. Previous efforts have utilized multidomain measures such as minimal disease activity that assess the status of joints, skin, and function to determine current level of psoriatic arthritis (PsA) disease activity. The goal of this study is to identify factors associated with patient-reported psoriatic arthritis remission.

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Methods: The National Psoriasis Foundation conducted a cross-sectional study using an online survey of a random stratified sample of 1570 individuals with psoriatic disease in the USA. Participants were asked about a provider diagnosis of psoriasis and/or psoriatic arthritis, comorbid conditions, and psoriatic arthritis impact and disease activity, and demographic questions. All participants reporting a physician-given diagnosis of psoriatic arthritis were asked if they felt their psoriatic arthritis was in remission ("Do you feel your psoriatic arthritis is in remission?" Yes/No/Unsure) and, if so, length of remission. Individuals with psoriasis and psoriatic arthritis reporting a body surface area impacted by psoriasis 3% or less were asked if they felt their psoriasis was in remission. Psoriatic arthritis disease activity and impact was assessed using the nine-question Psoriatic Arthritis Impact of Disease (PsAID-9) instrument and a global PsA-related quality of life question. PsAID-9 scores ≤ 4 were used to indicate acceptable disease state. Multivariate logistic regression was used to identify factors associated with patient-perceived PsA remission.

Results: Of 834 participants with PsA, including 76 (4.8%) with PsA without skin involvement ever, 144 (17.3%) felt their psoriatic arthritis was in remission, with an average remission duration of 43 months. Of those in remission, 116 (78.4%) reported currently using a treatment for their PsA, with most (75.7%)

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reporting using a biologic therapy for their PsA in the past 12 months. Multivariate logistic regression revealed that patient-perceived psoriatic arthritis remission was independently associated with experiencing acceptable disease state (PsAID-9 \leq 4), perception of psoriasis remission, lower impact of PsA on global quality of life, and non-white race. Age, sex, body mass index, or biologic use in the last 12 months were not associated with patient-reported PsA remission.

Conclusion: Overall, patient perception of PsA remission was most strongly associated with patient-reported psoriasis remission.

Keywords: Psoriatic arthritis; Psoriasis; Remission; Treat-to-target; Quality of life

Key Summary Points

Defining remission for psoriatic arthritis is challenging.

Understanding how patients with psoriatic arthritis define remission can inform efforts to define remission for psoriatic arthritis.

This study sought to understand patientdefined remission for psoriatic arthritis and identify factors associated with patient-reported psoriatic arthritis remission.

Study results suggest that patient perceived psoriatic arthritis remission is experiencing acceptable disease state (PsAID-9 \leq 4), perception of psoriasis remission, lower impact of PsA on global quality of life, and non-white race.

Our study highlights that psoriatic arthritis remission is multifactorial and patient-dependent.

Patient-defined psoriatic arthritis remission is linked with patient perception of skin disease and psoriatic arthritis impact on quality of life.

INTRODUCTION

Psoriatic arthritis (PsA) is a chronic inflammatory joint disease associated with psoriasis [1]. The clinical presentation of PsA is heterogeneous and may involve peripheral joint arthritis, dactylitis, enthesitis, and skin and nail disease [2]. Unlike rheumatoid arthritis. patients with PsA commonly have asymmetric disease. The wide variety of impacted health domains and the multidimensional nature of PsA makes defining remission status complex [1]. Although some have proposed clinical guidelines to facilitate treatment goals and define treatment success, there is currently no single, universally accepted definition of PsA remission. One "treat-to-target" guideline suggests that the goal for PsA treatment is to achieve minimal disease activity (MDA), which is defined as low disease activity across several domains of joints, skin, and other patient-reported outcomes, including quality of life [3]. However, the difference between remission and low disease activity remains ill defined.

Given the complex nature of PsA, patients should also be involved in defining their treatment goals, as every patient may have different perspectives on the target for therapy and the means to achieve it. [3]

Evaluating the course of the patient's disease activity can lead to objective measures to assess PsA remission. However, since patients often perceive remission in terms of the impact on their quality of life and physical limitations, objective measures have poor correlation with patient-reported remission [4]. Furthermore, it is difficult to expect the same level of remission across patients with different disease manifestations. For example, a complete absence of disease activity may be achievable in patients with early signs of PsA and limited joint damage but may not be realistic in patients with a more established disease with irreversible damage. There may also be a substantial pain component that does not track with improvement in inflammation and that activates central sensitization pathways [5]. Thus, allowing patients to define remission of their disease becomes more important.

Question: Has a doctor ever told you that you have any of the following conditions?					
Cardiovascular disease	9.6% (71)	Dyslipidemia	1.5% (11)	COPD	5.1% (38)
Stroke	2.0% (15)	Depression	41.2% (305)	Uveitis	3.6% (27)
Hypertension	21.8% (161)	Anxiety	42.6% (315)	Liver disease	1.9% (14)
High blood pressure	48.0% (355)	Sexual dysfunction	7.7% (57)	Fatty liver disease	13.1% (97)
Heart attack	3.6% (27)	Crohn's disease	2.0% (15)	Rheumatoid arthritis	9.3% (69)
Adult-onset diabetes (type 2 diabetes)	17.3% (128)	Ulcerative colitis	2.2% (16)	Gout	7.4% (55)
Metabolic syndrome	5.8% (43)	Cancer	10.7% (79)	Osteoarthritis	28.6% (212)
High triglyceride levels	19.3% (143)	Cutaneous T-cell lymphoma	0.3% (2)	Fibromyalgia	17.3% (128)
High cholesterol	39.6% (293)	Chronic kidney disease	3.6% (27)	Thyroid disease	19.1% (141)
Coronary artery disease	4.6% (34)	Sleep apnea	26.2% (194)	Other (specify)	13.9% (116)
Atherosclerosis	3.0% (22)	Hardening/narrowing of arteries	3.1% (23)	None of the above	8.8% (73)

Table 1 Comorbidities among participants with PsA

To further explore remission from a patientcentered view, the National Psoriasis Foundation aimed to identify demographic and clinical factors associated with patient-reported remission of psoriatic arthritis.

METHODS

The National Psoriasis Foundation conducted a cross-sectional study using an online and telephone-based survey of a random, stratified sample of individuals with psoriasis in October and November 2019. The survey was approved by the Genetic Alliance Institutional Review Board. Participants included individuals living in the USA with a physician-given diagnoses of psoriasis and/or psoriatic arthritis, 18 years of age or older, who contacted the foundation between 2017 and 2019. Participants reported physician-diagnosed psoriasis and/or psoriatic arthritis. All participants who reported having psoriatic arthritis were asked questions about their perspective on whether their psoriatic arthritis is in remission. Quality of life was assessed using a single-question global PsA-related quality of life measure and the ninequestion Psoriasis Impact of Disease (PsAID-9), a validated patient-reported outcome measure that assesses impact of PsA on patients' lives [6]. PsAID-9 score < 4 is considered to represent patient-acceptable symptom state (PASS) and PsAID-9 score > 4 represents non-PASS. The global PsA-related quality of life question asked participants the following: "[i]n all the ways your psoriatic arthritis affects you, how would you rate the way your felt over the past week" on a scale of 0 to 10, with 0 being "excellent" and 10 being "poor." The survey included questions about current and past treatments, physician-diagnosed comorbidities (Table 1), and participant demographics, including race, biologic sex, height, and weight. Participantreported height and weight were used to

calculate body mass index (BMI) using the standard formula: weight (lb)/height $(in)^2 \times 703$.

Participants with psoriasis were asked to selfreport severity of their psoriasis at the time of data collection using the Patient Report of Extent of Psoriasis Involvement (PREPI) [7]. The PREPI is a validated patient-reported outcome measure to assess percentage of body surface area affected by psoriasis. Participants who reported a body surface area (BSA) of 3% or less were asked questions about their perspective on whether their psoriasis is in remission. All participants with PsA were asked if the question, "Do you feel your psoriatic arthritis is in remission?" To more closely reflect patient perspective on remission, participants were not given a definition of remission for psoriasis or PsA or any explanation of potential indicators for remission. Individuals who perceived their PsA to be in remission were asked if they were currently taking therapies to treat their PsA. Separately, participants were asked if they used any of the following therapy types in the last 12 months: biologics, phototherapy, topical medications, prescribed oral systemic medications, or over-the-counter products, as well as if they are not currently on any treatment or a treatment other than those listed.

Descriptive statistics were used to report number and percentage of respondents who feel like their psoriatic arthritis is in remission. For univariate analyses, chi-square tests of independence were conducted to explore associations with patient perception of psoriatic arthritis remission. Independent-samples t-test was used to assess if global psoriatic arthritis quality of life differed on the basis of perception of remission. Results from univariate analyses informed logistic regression models, which were used to assess factors related to patient perception of psoriatic arthritis remission. For the logistic regression models, the continuous variable for age was used, race was collapsed into a dichotomous "white" and "non-white" variable owing to the lack of racial diversity among participants, and a continuous variable for BSA was used for disease severity.

Table 2 Participant demographics

Characteristic	n (%)		
Disease			
Psoriatic arthritis	76	(9.1%)	
Psoriatic arthritis and psoriasis	758	(90.9%)	
Race			
White or Caucasian	732	(89.3%)	
Asian or Asian American	19	(2.3%)	
Black or African American	20	(2.4%)	
American Indian or Alaskan Native	6	(0.7%)	
Native Hawaiian or other Pacific Island	2	(.2%)	
Two or more races	20	(2.4%)	
Other	13	(1.6%)	
Unsure	8	(1.0%)	
Biologic sex			
Male	310	(37.4%)	
Female	520	(62.6%)	
Age			
18–35 years	59	(7.3%)	
36–50 years	219	(27.1%)	
51–65 years	377	(46.7%)	
Older than 65 years	152	(18.8%)	
BMI categories			
Underweight	10	(1.3%)	
Normal weight	174	(22.4%)	
Overweight	251	(32.3%)	
Obese	342	(44.0%)	
Mean number of comorbidities	3.70	(± 2.64)	

RESULTS

A total of 834 individuals with psoriatic arthritis completed the survey (Table 2) and were asked if they felt their PsA was in remission. Of these, 144 (21.6%), felt that their PsA was in remission

Response	n (%)	
Perception of remission		
PsA not in remission	521 (78.4%)	
PsA in remission	144 (21.6%)	
Length of remission		
Number of months (mean, SD)	25.6 (± 23.8)	
PsA activity and impact		
Unacceptable (PsAID > 4)	477 (59.5%)	
Acceptable (PsAID ≤ 4)	324 (40.5%)	
Biologic use		
No	250 (30.6%)	
Yes	567 (69.4%)	

Table 3Descriptive statistics

(Table 3). Of participants who felt their PsA was
in remission, most are currently taking a treat-
ment for the PsA (76.7%) (Table 4) and the
average length of remission was 25.3 months
(Table 3). Patient perception of PsA remission
did not differ on the basis of presence of psori-
asis (17.7%-without skin involvement ever;
22.4% PsA and psoriasis) (Table 3). Overall,
most participants with PsA (59.5%) reported not
experiencing PASS (Table 3).

Univariate analysis results suggest that PASS, non-white race, BMI < 25.0, using one therapy to treat psoriatic disease, and perceiving psoriasis to be in remission were independently associated with patient-reported psoriatic arthritis remission (Table 4). Compared with individuals with non-PASS (5.1%), a higher percentage of participants with PASS (47.8%) felt their psoriatic arthritis was in remission (p < 0.001). Patient-reported psoriatic arthritis remission was higher among non-white participants (32.8%) compared with white participants (10.6%; p < 0.05) and among nonoverweight versus obese participants (31.5% versus 18.9%, p = 0.001). A higher percentage of those who felt their psoriasis was in remission reported using only one therapy (26.5%) compared with those using more than one therapy (18.1%, p < 0.05) to treat their psoriatic disease.

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Response	Psoriatic arthritis not in remission		
Disease diagnose	d		
Psoriatic arthritis without skin involvement	51 (82.3%)	11 (17.7%)	<i>p</i> = 0.431
Psoriatic arthritis and psoriasis	470 (90.2%)	133 (22.4%)	
PsAID-9 score			
PsAID-9 > 4	375 (95.9%)	20 (5.1%)	<i>p</i> < 0.001
$PsAID-9 \le 4$	128 (52.2%)	117 (47.8%)	
Biologic use			
Not biologic user	159 (82.4%)	34 (17.6%)	<i>p</i> = 0.108
Biologic user	349 (76.7%)	106 (23.3%)	
Race*			
Non-white	45 (67.2%)	22 (32.8%)	<i>p</i> < 0.05
White	465 (79.3%)	121 (10.6%)	
Biologic sex			
Female	330 (80.5%)	80 (19.5%)	p = 0.096
Male	189 (75.0%)	63 (25.0%)	
Obese or overwe	ight**		
Normal or underweight	102 (68.5%)	47 (31.5%)	p = 0.001
Obese or overweight	383 (78.1%)	89 (18.9%)	
Psoriasis in remi	ssion***		
No	137 (90.1%)	15 (9.9%)	<i>p</i> < 0.001
Yes	123 (59.4%)	84 (40.6%)	

Table 4 Chi-square tests of independence

Response	Psoriatic arthritis not in remission		p value
Currently using t	herapies to trea	t psoriatic art	hritis
Yes	-	102 (76.7%)	<i>p</i> = 0.378
No	_	31 (23.3%)	
Number of thera	pies**		
Only one therapy	161 (73.5%)	58 (26.5%)	<i>p</i> < 0.05
More than one therapy	299 (81.9%)	66 (18.1%)	
Number of comorbidities			
None	39 (7.6%)	13 (9.4%)	<i>p</i> < 0.05
1	55 (10.7%)	27 (19.6%)	
2	73 (14.2%)	28 (20.3%)	
3	89 (17.3%)	18 (13.0%)	
4	79 (15.4%)	20 (14.5%)	
5	58 (11.3%)	10 (7.3%)	
6 or more	121 (23.5%	22 (15.9%)	

 Table 4 continued

Patient-reported psoriatic arthritis remission was higher among individuals who felt their psoriasis was in remission (40.6%) compared with those who did not feel their psoriasis is in remission (9.9%). Individuals who do not perceive their PsA as being in remission experience higher impact of PsA on their quality of life than those who feel their PsA is in remission (M = 5.90, SD 2.53 versus M = 1.92, SD 1.65) (Fig. 1).

Factors associated with patient-reported PsA remission were explored using logistic regression. Results from multivariate logistic regression, controlling for BMI, sex, age, and type and number of therapies used, suggest that a PsAID-9 score ≤ 4 [odds ratio (OR) 4.58, 95% confidence interval (CI) 2.65–13.43, p < 0.05],

perceiving psoriasis to be in remission (OR 4.51, 95% CI 1.84–11.02, p < 0.05), global quality of life score (OR 0.55, 95% CI 0.43-0.69, p < 0.001), and white race (OR 0.28, 95% CI 0.28–1.00, p < 0.05) are associated with patientreported PsA remission (Table 5). Individuals achieving PASS were 358% more likely to report their PsA is in remission (p < 0.05). As impact of PsA on quality of life increased, the likelihood of perceiving PsA as in remission decreased by 45%. Individuals who felt that their psoriasis was in remission were 351% more likely to report their PsA was in remission (p < 0.05). White race was associated with an 82% decrease in likelihood of reporting PsA was in remission compared with non-white race (p < 0.05).

DISCUSSION

This study highlights several clinical and demographic factors associated with PsA remission from the patient perspective.

Patients with low psoriatic arthritis disease activity and impact, as determined by a PsAID-9 < 4, were more likely to report remission. The PsAID-9 was developed by Gossec et al. to specifically assess quality of life and physical function in PsA patients [6]. The PsAID-9 uses a series of numeric rating scale (NRS) items that range from 0 to 10 and integrates up to 12 different domains of health. PsAID-12 provides questions that are most relevant to clinical management and patient care over a long period of time, whereas PsAID-9 uses questions that are more relevant for clinical trials [6]. Several studies have shown that the PsAID is a reliable method of measuring patient-reported outcomes that can differentiate PsA disease activities across patients [6, 8–10]. The results in our study strengthen the relationship between patient-reported outcome measures and remission status. It also highlights the importance of understanding the patient's perspective on not only the physical burden, but also the psychological impact of PsA when defining remission.

Patients who perceived their psoriasis to be in remission were also more likely to report PsA remission. This highlights that, although PsA involves numerous health domains, skin



Fig. 1 Boxplot-PsA Global QoL by Perception of Remission

involvement is an important factor associated with patient-perceived remission, or that reduced skin activity tracks closely with the use of especially efficacious therapies that also alleviate PsA symptoms. A recent study demonstrated that patients with PsA involving both joint and skin symptoms have worse patientreported outcomes and more severe disease [11]. Moreover, a second study demonstrated that improvement in health-related quality of life in patients with psoriatic arthritis was dependent on successfully treating both skin and joint symptoms [12]. While studies have shown that skin symptoms largely impact the course of psoriatic arthritis, our study further highlights that skin involvement independently conpatients' perception of PsA tributes to remission.

Lower global PsA quality of life score was also found to be significantly associated with remission status even when joint disease activity was controlled. This questionnaire was measured on a scale of 0–10, with 0 indicating the least impact on the overall quality of life due to psoriatic arthritis. Patients with PsA were found to have significantly reduced healthrelated quality of life compared with the general population even after 5 years from disease onset [13]. This suggests that long-term reductions in quality of life may be associated with PsA disease, which explains why improvement in quality of life is associated with patient-reported disease remission.

Non-white patients were also more likely to report remission than white patients. The role of race in psoriatic arthritis remission is unclear; however, studies have demonstrated the prevalence and disease characteristics of PsA may vary between races. A cross-sectional US claims database study found that PsA prevalence is highest among white patients compared with other racial and ethnic groups [14]. Another study found that PsA was also estimated to occur twice as frequently in white patients compared with Black patients [15]. The severity of psoriatic arthritis was found to be higher in individuals of Indian ethnicity compared with Chinese ethnicity, suggesting genetic or environmental factors may play a role [16]. Future studies are needed to evaluate the role of race in psoriatic arthritis severity, disease characteristics, and treatment to understand their impact

		Dermatol Th

	Psoriatic arthritis in remission			
	Odds ratio	SE	95% CI	p value
Model 1				
PsA acceptable disease activity	17.13	4.50	10.24-28.68	< 0.0001
Model 2				
PsA acceptable disease activity	3.56	1.17	1.86-6.80	< 0.0001
PsA Global QoL	0.55	0.40	0.48-0.64	< 0.0001
Model 3				
PsA acceptable disease activity	4.68	2.04	1.99–11.00	< 0.0001
PsA Global QoL	0.58	0.05	0.48-0.70	< 0.0001
Psoriasis in remission	7.07	2.78	3.27-15.27	< 0.0001
Model 4				
PsA acceptable disease activity	4.70	2.10	1.96–11.24	0.01
PsA Global QoL	0.57	0.05	0.47-0.69	< 0.0001
Psoriasis in remission	6.80	2.68	3.14-14.74	< 0.0001
White	0.28	0.17	0.08-0.92	0.05
Model 5				
PsA acceptable disease activity	5.10	2.74	1.78–14.61	0.01
PsA Global QoL	0.54	0.06	0.43-0.69	< 0.0001
Psoriasis in remission	4.84	2.18	2.00-11.69	< 0.0001
White	0.28	0.18	0.08-1.00	0.051
Multiple therapy use	0.62	0.24	0.28-1.34	0.225
Obese/overweight	1.06	0.47	0.45-2.52	0.888
Male	0.78	0.33	0.34-1.78	0.561
Biologic user	0.97	0.50	0.35-2.69	0.953
Model 6				
PsA acceptable disease activity	4.58	2.51	1.56-13.43	0.01
PsA Global QoL	0.55	0.06	0.43-0.69	< 0.0001
Psoriasis in remission	4.51	2.05	1.84-11.02	0.01
White	0.28	0.18	0.08-1.00	0.05
Multiple therapy use	0.61	0.24	0.28-1.33	0.215
Obese/overweight	1.04	0.46	0.44-2.46	0.935
Male	0.82	0.36	0.35-1.93	0.665
Biologic user	0.91	0.48	0.32-2.57	0.853

Table 5 Logistic regression models

	Psoriatic arthritis in remission			
	Odds ratio	SE	95% CI	p value
Age	0.99	0.01	0.96-1.02	0.725
Number of comorbidities	1.05	0.09	0.88-1.25	0.567

Table 5 continued

on PsA remission. A notable limitation of our study is the predominance of white race subjects.

This study has some limitations. First, using an online survey is a potential weakness of the study. Online surveys require access to the internet and a certain level of computer skills to navigate. Additionally, age and race also affect response to web-based surveys. Individuals who contact the National Psoriasis Foundation differ from the general population with psoriasis in several ways that could influence study results. Members of the National Psoriasis Foundation tend to be older and more affluent, have more significant disease, and be more knowledgeable about treating the psoriatic disease compared with the general public.

CONCLUSIONS

PsA is a heterogeneous, inflammatory disease with multiple domains of importance. We evaluated the clinical and demographic factors associated with patient-perceived PsA remission through the use of a cross-sectional survey. Low disease activity and impact (low PSAID score) was associated with remission status. Furthermore, perceived psoriasis remission, quality of life, and non-white race were also significantly associated with patient-reported PsA remission. Ultimately, our study highlights that PsA remission status is multifactorial and patient-dependent.

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Compliance with Ethics Guidelines. This study was approved by the Genetic Alliance Institutional Review Board (Protocol # NPF-ASP-2019) and performed in accordance with the Helsinki Declaration of 1964 and its later amendments. Study participants consented to the study electronically.

Data Availability. The datasets generated during and/or analyzed during the current study are available from the corresponding author on reasonable request.

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