



ORIGINAL ARTICLE

The Clinical Profile of Patients with Psoriasis in Korea: A Nationwide Cross-Sectional Study (EPI-PSODE)

Hae Jun Song, Chul Jong Park¹, Tae Yoon Kim², Yong Beom Choe³, Seok-Jong Lee⁴, Nack In Kim⁵, Jae We Cho⁶, Jie Hyun Jeon, Min Soo Jang⁷, Jai Il Youn⁸, Myung Hwa Kim⁹, Joonsoo Park¹⁰, Ki Ho Kim¹¹, Byung Soo Kim¹², Sang Woong Youn¹³, Joo-Heung Lee¹⁴, Min-Geol Lee¹⁵, Sung Ku Ahn¹⁶, Young Ho Won¹⁷, Seok Kweon Yun¹⁸, Bong Seok Shin¹⁹, Seong Jun Seo²⁰, Ji Yeoun Lee²¹, Kwang Joong Kim²², Young Suck Ro²³, Youngdoe Kim²⁴, Dae Young Yu²⁴, Jee-Ho Choi²⁵

Department of Dermatology, Korea University Guro Hospital, Seoul, ¹Department of Dermatology, The Catholic University of Korea, Bucheon St. Mary's Hospital, Bucheon, ²Department of Dermatology, The Catholic University of Korea, Seoul St. Mary's Hospital, ³Department of Dermatology, Konkuk University School of Medicine, Seoul, ⁴Department of Dermatology, Kyungpook National University Hospital, Daegu, ⁵Department of Dermatology, Kyung Hee University Medical Center, Seoul, ⁶Department of Dermatology, Keimyung University Dongsan Medical Center, Daegu, ⁷Department of Dermatology, Kosin University College of Medicine, Busan, ⁸Department of Dermatology, National Medical Center, Seoul, ⁹Department of Dermatology, Dankook University Hospital, Cheonan, ¹⁰Department of Dermatology, Catholic University of Daegu School of Medicine, Daegu, ¹¹Department of Dermatology, Dong-A University Hospital, ¹²Department of Dermatology, Pusan National University School of Medicine, Busan, ¹³Department of Dermatology, Seoul National University Bundang Hospital, Seoul National University College of Medicine, Seongnam, ¹⁴Department of Dermatology, Samsung Medical Center, Sungkyunkwan University School of Medicine, ¹⁵Department of Dermatology, Severance Hospital, Yonsei University College of Medicine, Seoul, ¹⁶Department of Dermatology, Wonju Severance Christian Hospital, Wonju, ¹⁷Department of Dermatology, Chonnam National University Hospital, Gwangju, ¹⁸Department of Dermatology, Chonbuk National University Hospital, Jeonju, ¹⁹Department of Dermatology, Chosun University Hospital, Gwangju, ²⁰Department of Dermatology, Chung-Ang University Hospital, Seoul, ²¹Department of Dermatology, Chungbuk National University College of Medicine, Cheongju, ²²Department of Dermatology, Hallym University Sacred Heart Hospital, Anyang, ²³Department of Dermatology, Hanyang University Medical Center, ²⁴Medical Affairs, Janssen Korea, ²⁵Department of Dermatology, Asan Medical Center, University of Ulsan College of Medicine, Seoul, Korea

Background: Psoriasis is an immune-mediated, chronic inflammatory disease affecting multiple aspects of patients' lives. Its epidemiology varies regionally; however, nationwide epidemiologic data on psoriasis depicting profile of Korean patients has not been available to date. **Objective:** To understand nationwide epidemiologic characteristics and clinical features of adult patients with psoriasis visited university hospitals in Korea. **Methods:** This multicenter, non-in-

terventional, cross-sectional study recruited 1,278 adult patients with psoriasis across 25 centers in Korea in 2013. Various clinical data including PASI, BSA, DLQI, SF-36 and PASE were collected. **Results:** A total of 1,260 patients completed the study (male:female = 1.47:1). The mean age was 47.0 years with a distribution mostly in the 50s (24.9%). Early onset (<40 years) of psoriasis accounted for 53.9% of patients. The mean disease duration was 109.2 months; mean body mass index was 23.9 kg/m²; and 12.7% of patients had a family history of psoriasis. Plaque and guttate types of psoriasis accounted for 85.8% and 8.4%, respectively. Patients with PASI ≥ 10 accounted for 24.9%; patients with body surface area ≥ 10 were 45.9%. Patients with DLQI ≥ 6 accounted for 78.8%. Between PASI < 10 and PASI ≥ 10 groups, significant difference was noted in age at diagnosis, disease duration, blood pressure, waist circumference of female, and treatment experiences with phototherapy, systemic agents, and biologics. **Conclusion:** This was the first nationwide epidemiologic study of patients with psoriasis in Korea and provides an overview of the epidemio-

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Corresponding author: Jee-Ho Choi, Department of Dermatology, Asan Medical Center, University of Ulsan College of Medicine, 88 Olympic-ro 43-gil, Songpa-gu, Seoul 05505, Korea. Tel: 82-2-3010-3463, Fax: 82-2-486-783, E-mail: jhchoy@amc.seoul.kr

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logic characteristics and clinical profiles of this patient population. (*Ann Dermatol* 29(4) 462~470, 2017)

-Keywords-

Epidemiologic studies, Korea, Psoriasis

INTRODUCTION

Psoriasis is an immune-mediated, chronic inflammatory disease affecting various aspects of patients' lives far beyond the skin and challenging clinicians in its management¹. While psoriasis is prevailing worldwide, there are regional differences in its epidemiological characteristics. Local differences in genetic, environmental, cultural, and geographic characteristics may be related. Although outstanding progress have achieved in various fields of psoriasis research, understanding epidemiologic profiles of the disease is still important. A more elaborate understanding of regional epidemiology is essential to refining the management of psoriasis. Unlike many western countries, where data from many large-scale epidemiologic studies are available, epidemiologic study data of nationwide scale were unavailable in Korea. In 2013, the Korean Society for Psoriasis (KSP) conducted a cross-sectional, multicenter study to delineate the clinical profile of adult patients with psoriasis visiting university hospitals in Korea. Here we report the results of this study.

MATERIALS AND METHODS

Study design

This was a multicenter, non-interventional, cross-sectional study. Patients were recruited for approximately 4 months, from February to June 2013. The 25 centers participating in the study were mainly university hospitals, located across the country (with the exception of the Jeju island region). This study was reviewed and approved by the Institutional Review Board of each center. To extract a representative sample reflecting the distribution of psoriasis in Korea, numbers and locations of study centers were selected on the basis of population distribution and health insurance reimbursement data of the patients with psoriasis in Korea.

Because half of the total population of Korea lives near the capital, 13 of 25 centers were selected from the capital area. To avoid selection bias, study patients were examined and enrolled consecutively by visiting orders. After obtaining informed consent, all procedures of the study were performed on 1 day. The study patients were adult

patients aged 20 or older diagnosed with psoriasis.

Clinical data were collected by interviewing the patients on demographic factors (age, sex, height, weight, waist circumference, disease duration, age at diagnosis, family history in first-degree relative, drinking/smoking history, and both past and ongoing concomitant therapies). Physical examination for disease-severity measures (psoriasis area and severity index [PASI], body surface area [BSA]), morphological characteristics (clinical types and location) of psoriasis and blood pressure were carried out for all enrolled patients. Quality of life was evaluated by the dermatology life quality index (DLQI) and MOS 36-item short-form health survey (SF-36). Screening for psoriatic arthritis was done using the psoriatic arthritis screening and evaluation (PASE) tool. In addition, data on treatment-related factors (past and current treatment history, patient satisfaction for treatment) were collected. In order to evaluate patient satisfaction for various therapies, questions from the Medication Satisfaction Questionnaire (MSQ)², which is a validated tool for evaluating treatment satisfaction among patients with schizophrenia, were used in this study. Patients scored their response to the question, "In general, how satisfied are you with your current treatment?" on a 7-point scale ranging from 1 to 7 (a score of 4 means neutral).

Statistical analysis

Descriptive statistics for continuous variables are presented as means with standard deviation, and dichotomous variables are presented as frequencies with percentages in parentheses. Types of psoriasis, location of involved lesions, and clinical/demographic characteristics were compared to assess the statistical significance of any differences among the groups by disease severity (PASI <10, PASI ≥10). We used, where appropriate, the Wilcoxon rank-sum and t tests for continuous variables, and analyses for dichotomous variables that were based on Pearson's chi-square test and Fisher's exact test, without missing data imputation. In addition, a multiple logistic regression analysis was used to assess the adjusted effect of characteristics on PASI ≥10. Multicollinearity between independent variables had been checked by computing correlation coefficient and variance inflation factors (VIF) before the variables to be included in the regression model were selected. All statistical tests were performed using two-sided and *p*-values less than 0.05 were considered statistically significant. Analyses were performed using the statistical software package SAS 9.2 (SAS Institute, Cary, NC, USA).

RESULTS

Demographics

A total of 1,278 patients were enrolled in the study, and 1,260 (98.6%) completed the study. Eighteen patients (1.4%) were excluded from analysis, 16 (1.3%) for violation of the inclusion/exclusion criteria, and two (0.2%) for lack of data for the primary assessment. Demographic data from the 1,260 patients are presented in Table 1. A total of 749 patients (59.4%) were male and 511 (40.6%) were female; the male to female ratio was 1.47:1. The mean age of the study patients was 47.0 ± 14.5 years (range, 20~89 years). Most study patients were in their 50s (24.9%), with a gradual increase from 20s to 50s and a sharp decrease after 60s. The distribution pattern of age at diagnosis revealed dual peaks at 21~25 years and 46~50 years (Fig. 1). A ratio of early onset (<40 years) versus late onset (≥ 40 years) at diagnosis was 53.9% versus 46.1% (33.8% versus 66.2% when divided by age of 30 years old). The mean disease duration was 109.2 ± 122.0 months. Mean height and weight were 166.5 ± 8.4 cm and 66.4 ± 12.4

kg, respectively, and mean body mass index (BMI) was 23.9 ± 3.5 kg/m². At the time of the study, 48.3% (n=609) of the total patients had a history of smoking and 64.0% (n=806) had a history of drinking. Male showed higher rate in smoking/drinking history than female. Patients who had a family history of psoriasis in a first-degree relative accounted for 12.7% (n=160) of the total.

Clinical type of psoriasis

Plaque-type psoriasis was the most prevalent, occurring in 1,081 patients (85.8%), followed by guttate type in 106 (8.4%), pustular type (including localized form) in 66 (5.2%), erythrodermic type in 44 (3.5%), annular type in 24 (1.9%), and inverse type in 4 (0.3%) (Table 2).

Location of lesions

Psoriasis lesions were most frequently found in the lower leg (72.6%, n=913), followed by the back (60.9%, n=766), upper leg (60.8%, n=764), elbow (58.0%, n=729), knee (57.1%, n=718), lower arm (56.8%, n=714), abdomen (55.7%, n=700), upper arm (53.5%, n=672), and scalp

Table 1. Demographics

Characteristics	Total	Overall	PASI < 10	PASI \geq 10	<i>p</i> -value
Gender	1,260				0.215
Male		749 (59.4)	553 (58.5)	196 (62.4)	
Female		511 (40.6)	393 (41.5)	118 (37.6)	
Age (yr)	1,260	47.0 ± 14.5	47.3 ± 14.3	46.3 ± 15.0	0.310
Age at diagnosis (yr)	1,257	38.6 ± 16.5	39.3 ± 16.2	36.4 ± 17.0	0.008*
< 40		677 (53.9)	492 (52.1)	185 (59.1)	
≥ 40		580 (46.1)	452 (47.9)	128 (40.9)	0.032*
Disease duration (mo)	1,257	109.2 ± 122.0	103.3 ± 118.2	126.9 ± 131.5	0.003*
Family history	1,260	160 (12.7)	128 (13.5)	32 (10.2)	0.124
Height (cm)	1,259	166.5 ± 8.4	166.5 ± 8.5	166.5 ± 8.3	0.925
Weight (kg)	1,259	66.4 ± 12.4	66.1 ± 12.2	67.4 ± 12.9	0.116
BMI (kg/m ²)	1,259	23.9 ± 3.5	23.8 ± 3.4	24.2 ± 3.8	0.059
Male	748	24.5 ± 3.2	24.5 ± 3.1	24.6 ± 3.5	0.699
Female	511	23.0 ± 3.7	22.8 ± 3.5	23.6 ± 4.2	0.046*
Waist circumference (cm)	1,113	84.3 ± 11.1	83.8 ± 11.0	85.6 ± 11.4	0.022*
Male (≥ 90 cm)	659	254 (38.5)	193 (38.6)	61 (38.4)	0.958
Female (≥ 80 cm)	454	199 (43.8)	138 (40.0)	61 (56.0)	0.003*
SBP ≥ 140 mmHg or DBP ≥ 90 mmHg	1,164	231 (19.8)	158 (17.8)	73 (26.5)	0.001*
Smoking history	1,260	609 (48.3)	449 (47.5)	160 (51.0)	0.283
Male	749	538 (71.8)	394 (71.2)	144 (73.5)	0.552
Female	511	71 (13.9)	55 (14.0)	16 (13.6)	0.905
Drinking history	1,260	806 (64.0)	591 (62.5)	215 (68.5)	0.055
Male	749	595 (79.4)	431 (77.9)	164 (83.7)	0.088
Female	511	211 (41.3)	160 (40.7)	51 (43.2)	0.628

Values are presented as total number only, number (%), or mean \pm standard deviation. Total number does not always represent 100% of the study population because of missing values. Percentages of subgroup (by gender) were based on the total number of subjects in each PASI group per subgroup. PASI: psoriasis area and severity index, BMI: body mass index, SBP: systolic blood pressure, DBP: diastolic blood pressure. **p*-value < 0.05.

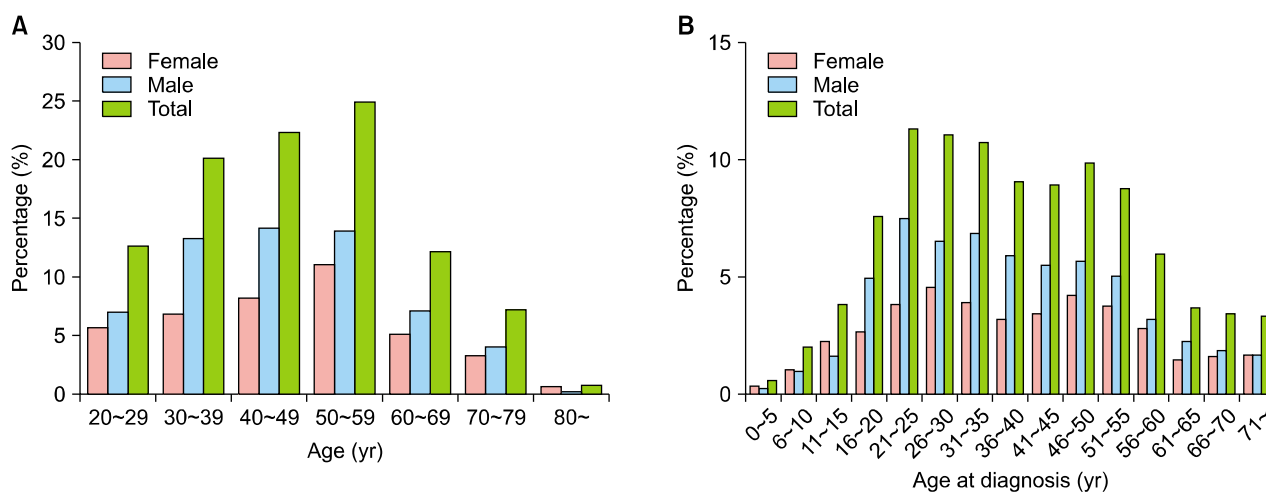


Fig. 1. Age distribution of patient (A) and distribution of age at diagnosis (B).

Table 2. Types of psoriasis

Classification	Overall (n=1,260)
Plaque	1,081 (85.8)
Guttate	106 (8.4)
Pustular	66 (5.2)
Erythrodermic	44 (3.5)
Annular	24 (1.9)
Inverse	4 (0.3)

Values are presented as number (%).

(53.1%, n=668). Nail involvement was noted in 12.3% of patients and genitalia involvement in 8.8% of patients. Details of lesion location are listed in Table 3.

Disease severity

Of the total study patients, 75.1% (n=946) had PASI < 10 and 24.9% (n=314) scored PASI ≥ 10 (19.0% had 10 ≤ PASI < 20, 5.9% had PASI ≥ 20). The mean PASI score was 7.5 ± 7.1. The mean BSA (%) measurement was 13.1 ± 14.6. 54.1% (n=681) of the total patients had BSA < 10, 45.9% (n=577) had BSA ≥ 10 (14.3% had BSA < 3%, 39.8% had 3 ≤ BSA < 10, 25.4% had 10 ≤ BSA < 20, and 20.4% had BSA ≥ 20). On the other hand, with regards to the DLQI, results showed that of the 1,256 study patients whose DLQI data were available, 21.2% (n=266) scored DLQI ≤ 5 (no/small effect), 24.7% (n=310) scored 6 ≤ DLQI ≤ 10 (moderate effect), and 54.1% (n=680) scored DLQI ≥ 11 (very/extremely large effect). The mean value of DLQI was 12.0 ± 7.2 (Fig. 2). The mean SF-36 results were 48.8 ± 8.0 for the physical component summary score and 42.6 ± 11.2 for the mental component summary score (data not shown). PASE data were available for 1,255 patients. The mean PASE score was 33.7 ± 13.1;

Table 3. Location of involved lesions

Classification	Overall (n=1,257)*	PASI < 10 (n=943)	PASI ≥ 10 (n=314)
Scalp	668 (53.1)	453 (48.0)	215 (68.5)
Face	349 (27.8)	218 (23.1)	131 (41.7)
Neck	205 (16.3)	104 (11.0)	101 (32.2)
Chest	536 (42.6)	334 (35.4)	202 (64.3)
Abdomen	700 (55.7)	460 (48.8)	240 (76.4)
Back	766 (60.9)	497 (52.7)	269 (85.7)
Upper arm	672 (53.5)	425 (45.1)	247 (78.7)
Elbow	729 (58.0)	484 (51.3)	245 (78.0)
Lower arm	714 (56.8)	462 (49.0)	252 (80.3)
Palm [†]	164 (13.0)	115 (12.2)	49 (15.6)
Back of hand	295 (23.5)	178 (18.9)	117 (37.3)
Nail	155 (12.3)	89 (9.4)	66 (21.0)
Upper leg	764 (60.8)	499 (52.9)	265 (84.4)
Knee	718 (57.1)	472 (50.1)	246 (78.3)
Lower leg	913 (72.6)	620 (65.7)	293 (93.3)
Foot sole [†]	146 (11.6)	103 (10.9)	43 (13.7)
Back of foot	217 (17.3)	127 (13.5)	90 (28.7)
Buttock	471 (37.5)	280 (29.7)	191 (60.8)
Genitalia	111 (8.8)	59 (6.3)	52 (16.6)
Other area [†]	8 (0.6)	4 (0.4)	4 (1.3)

Values are presented as number (%). PASI: psoriasis area and severity index. *3 patients are missing. [†]No significant p-value (other involved areas showed p-value < 0.0001).

301 (24.0%) patients had a PASE score of 44 points or higher (Fig. 3).

Past and concomitant therapies

Of all past therapies utilized by the study patients (n=1,260), topical agents were the most common (93.6%, n=1,179), followed by systemic agents (61.2%, n=771), phototherapy (50.8%, n=640), and biologics (5.7%, n=72).

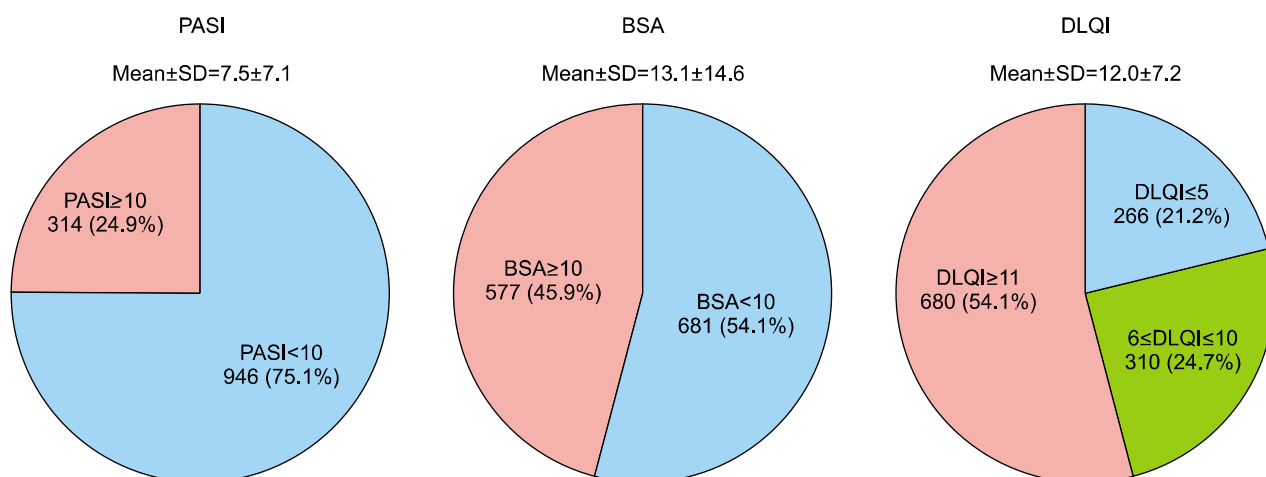


Fig. 2. Disease severity classified by psoriasis area and severity index (PASI), body surface area (BSA), and dermatology life quality index (DLQI). Overall mean ± standard deviation (SD) are presented below the title. Each n (%) of the classified group is presented.

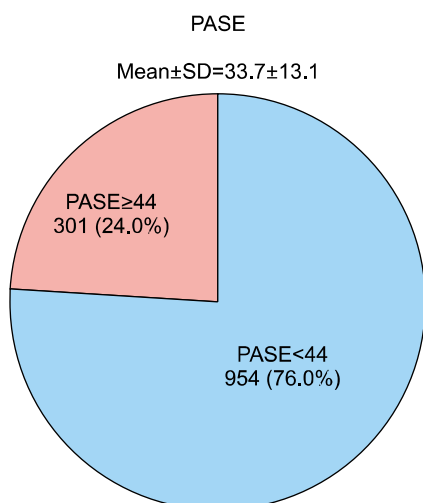


Fig. 3. Psoriatic arthritis screening and evaluation (PASE). SD: standard deviation.

The order of current therapy maintained at the time of enrollment was topical agent (86.0%, n=1,083), oral systemic agents (45.9%, n=578), phototherapy (30.6%, n=385), and biologics (4.6%, n=58). Although herbal medicine was used at some point by a considerable number of patients, it was seldom maintained (Table 4).

Comparison of factors related to disease based on disease severity

The patients were divided into two groups of PASI < 10 and PASI ≥ 10 and factors that could be related with disease severity were compared. Gender, age, family history, height and weight, smoking/drinking history, and male gender's waist circumference and BMI did not showed any significant difference between two groups. However,

there was significant difference between two groups regarding BMI in female, age at diagnosis, disease duration, waist circumference of female, blood pressure. The group with PASI ≥ 10 had longer disease duration ($p=0.003$) and earlier age of diagnosis ($p=0.008$) than the group with PASI < 10 and comprised a significantly higher proportion of patients with systolic blood pressure ≥ 140 mmHg or diastolic blood pressure ≥ 90 mmHg ($p=0.001$). BMI in female was significantly higher ($p=0.046$) and the proportion of waist circumference with ≥ 80 cm in female was higher in PASI ≥ 10 group ($p=0.003$, Table 1). Most of body parts were involved more as disease severity increased except for palm, sole (Table 3). After adjusting for variables in demographic characteristics except age at diagnosis, height and weight due to multicollinearity, younger age, longer disease duration, absence of family history, longer waist circumference and higher blood pressure were significantly associated with severe disease (PASI ≥ 10, Table 5). More patients with PASI ≥ 10 had experienced treatment with phototherapy, systemic agents, and biologics compared to patients with PASI < 10 (Table 4).

Treatment satisfaction

Patients were divided into three treatment satisfaction groups: dissatisfied (from extremely dissatisfied to somewhat dissatisfied), neutral, or satisfied (from somewhat satisfied to extremely satisfied). The satisfied group included 733 patients (58.5%); neutral group, 317 (25.3%); and dissatisfied group, 202 (16.1%). In the PASI < 10 group, satisfied proportion was significantly higher than the PASI ≥ 10 group ($p=0.002$, Table 6).

Table 4. Past and current therapies

Category	Overall (n=1,260)		PASI<10 (n=946)	PASI≥10 (n=314)	p-value
	Past and current	Current only			
Topical	1,179 (93.6)	1,083 (86.0)	884 (93.4)	295 (93.9)	0.753
Glucocorticosteroid	937 (74.4)	774 (61.4)	697 (73.7)	240 (76.4)	0.333
Calcipotriol/tacalcitol	986 (78.3)	893 (70.9)	746 (78.9)	240 (76.4)	0.367
Phototherapy	640 (50.8)	385 (30.6)	463 (48.9)	177 (56.4)	0.023*
UVA/UVB	35 (2.8)	19 (1.5)	32 (3.4)	3 (1.0)	0.023*
Narrowband UVB	564 (44.8)	338 (26.8)	404 (42.7)	160 (51.0)	0.011*
Broadspectrum UVB	10 (0.8)	6 (0.5)	7 (0.7)	3 (1.0)	0.717
PUVA (systemic)	42 (3.3)	26 (2.1)	31 (3.3)	11 (3.5)	0.847
Systemic agents	771 (61.2)	578 (45.9)	561 (59.3)	210 (66.9)	0.017*
Fumaric acid esters	4 (0.3)	0 (0.0)	2 (0.2)	2 (0.6)	0.260
Methotrexate	280 (22.2)	174 (13.8)	205 (21.7)	75 (23.9)	0.413
Acitretin	300 (23.8)	185 (14.7)	206 (21.8)	94 (29.9)	0.003*
Cyclosporine	366 (29.1)	221 (17.5)	253 (26.7)	113 (36.0)	0.002*
Oral corticosteroid	100 (7.9)	26 (2.1)	79 (8.4)	21 (6.7)	0.345
Biologics	72 (5.7)	58 (4.6)	45 (4.8)	27 (8.6)	0.011*
Adalimumab	6 (0.5)	4 (0.3)	5 (0.5)	1 (0.3)	1.000
Etanercept	13 (1.0)	2 (0.2)	7 (0.7)	6 (1.9)	0.102
Infliximab	7 (0.6)	5 (0.4)	5 (0.5)	2 (0.6)	0.687
Ustekinumab	53 (4.2)	47 (3.7)	32 (3.4)	21 (6.7)	0.012*
Others	181 (14.4)	42 (3.3)	137 (14.5)	44 (14.0)	0.837
Diet therapy	28 (2.2)	19 (1.5)	27 (2.9)	1 (0.3)	0.008*
Herbal medicine	106 (8.4)	9 (0.7)	81 (8.6)	25 (8.0)	0.740
Exercise therapy	8 (0.6)	8 (0.6)	7 (0.7)	1 (0.3)	0.688
Spa	14 (1.1)	11 (0.9)	11 (1.2)	3 (1.0)	1.000
Supplements	13 (1.0)	9 (0.7)	10 (1.1)	3 (1.0)	1.000
None	19 (1.5)	-	18 (1.9)	1 (0.3)	0.058

Values are presented as number (%). PASI: psoriasis area and severity index, UV: ultraviolet, PUVA: psoralen and ultraviolet A. *p-value<0.05.

Table 5. Adjusted effect of characteristics on PASI ≥10 based on multiple logistic regression model (n=1,086)

Characteristics	Ref.	OR	95% CI	p-value
Gender	Male	1.182	0.810~1.725	0.386
Age		0.987	0.977~0.998	0.020*
Disease duration		1.002	1.001~1.003	0.002*
Family history	No	0.571	0.361~0.905	0.017*
BMI		0.992	0.943~1.044	0.763
Waist circumference		1.018	1.001~1.036	0.036*
SBP ≥140 mmHg or DBP ≥90 mmHg	No	1.535	1.090~2.162	0.014*
Smoking history	No	1.006	0.705~1.435	0.975
Drinking history	No	1.061	0.758~1.485	0.731

One hundred seventy-four patients with missing data were excluded from the logistic regression analysis. PASI: psoriasis area and severity index, Ref.: reference for odds ratio, OR: odds ratio, CI: confidence interval, BMI: body mass index, SBP: systolic blood pressure, DBP: diastolic blood pressure. *p-value<0.05.

DISCUSSION

Prior to this study, epidemiologic data for the patients with psoriasis in Korea were limited to small-scale or single-center studies using accumulated patient pools over

long periods of time³. For this reason, it was difficult to obtain an overview of nationwide epidemiologic profile of psoriasis that is necessary for further research and better policy making for the patients with psoriasis in Korea. Hence, the KSP designed and conducted a nationwide

Table 6. Comparing the degree of satisfaction with treatment

Category	Overall (n=1,252)*	PASI < 10 (n=939)	PASI ≥ 10 (n=313)	p-value
Degree of satisfaction				0.002 [†]
Extremely satisfied	42 (3.4)	34 (3.6)	8 (2.6)	
Very satisfied	229 (18.3)	192 (20.4)	37 (11.8)	
Somewhat satisfied	462 (36.9)	350 (37.3)	112 (35.8)	
Neutral	317 (25.3)	226 (24.1)	91 (29.1)	
Somewhat dissatisfied	125 (10.0)	89 (9.5)	36 (11.5)	
Very dissatisfied	58 (4.6)	36 (3.8)	22 (7.0)	
Extremely dissatisfied	19 (1.5)	12 (1.3)	7 (2.2)	

Values are presented as number (%). PASI: psoriasis area and severity index. *8 patients are missing. [†]p-value < 0.05.

study in 2013 to delineate the epidemiologic profile of Korean patients with psoriasis. To minimize regional sampling bias and improve temporal focusing, we matched the number and location of study centers with general population distribution data. As a result, the distributed proportion of recruited patients was very similar to that of the distributed proportion of claims for psoriasis in the Health Insurance Review and Assessment (HIRA) database of 2013.

The mean age of patients in our study (47.0 years) was higher than that observed in hospital based Japanese studies (45.5 years⁴ and 44.7 years⁵) as well as a Chinese population-based study (41.8 years⁶), but was similar to a Taiwanese cohort study (46.4 years⁷). Unlike classical bimodal peaks in distribution patterns of age at diagnosis in western studies, which showed peaks located in the late teens and late 50s, Asian studies^{4,6}, including ours, showed dual peaks located in the 20s and 40s. Additionally, early-onset (type 1, < 40 years) psoriasis accounted for more than 75% of patients in western studies⁸ and 67.6% of patients in the Chinese study⁶, but only 53.9% in our study.

The prevalence of psoriasis has been reported to be equal between men and women in western studies⁹⁻¹¹. However, in Asian studies^{4,5,7,12}, male predominance has been reported repeatedly. Our study also revealed the predominance of this disease in males (male to female ratio 1.47:1). This tendency was also noted in data from the Korean health insurance database in 2012 (1.36:1). But the degree of male predominance was somewhat lower than studies from other Asian countries such as Japan and Taiwan, which reported a male to female ratio of 1.6 ~ 2.1:1^{5,7}. Although this phenomenon may be unique to the study population, the cause of this interesting observation needs to be elucidated.

Although it is well known that obesity is closely related to psoriasis in western populations^{8,13,14}, the mean BMI of 23.9 ± 3.5 kg/m² noted in our study was considerably lower than the mean BMI of 28.0 kg/m² observed in western

populations¹³. In addition, Kim and Lee¹⁵ reported no statistical difference in BMI between the patients with psoriasis and a control group in one study of a Korean population. This implies that further research is necessary to clarify the association between obesity and psoriasis in the Korean population.

Plaque-type psoriasis was the predominant clinical type in Korea, consistent with observations in Caucasian and other Asian populations^{5,16-18}. Interestingly, although the proportion of plaque psoriasis is similar, the proportion of guttate psoriasis is relatively high in Korean patients compared with Japanese patients (8.4% versus 2.8%/3.9%)^{4,5}. It was difficult to compare the involved site with other published studies, as each uses different criteria to distinguish the location in the body. For future studies, it is necessary to define the boundary of anatomical regions in a standardized manner and use common protocol to express involved areas. Hernandez et al.¹⁹ reported that patients with more severe psoriasis had greater skin involvement in most of area including the palms and soles. However, our data showed no significant differences in the involvement of palms and soles by disease severity divided by PASI score 10.

When comparing disease severity as measured by PASI and BSA with quality of life as measured by DLQI, it was found that the impact of psoriasis on quality of life was not fully explained by the severity or extent of skin disease. Therefore, an independent assessment of quality of life should be considered in the management of psoriasis. Some studies^{20,21} reported a higher disease severity in male patients than in female patients. Although there was a higher proportion of males in the severe group, it was not statistically significant in our study ($p=0.215$).

Smoking has been known to be associated with the clinical severity of psoriasis²² but the ratio of smoking history between two groups by disease severity did not differ statistically. Some studies have reported that patients with more severe disease had an earlier age of onset and a lon-

ger disease duration^{19,23}. Those results are in line with our findings. The more severe group presented with significantly higher blood pressure, as reported by Cohen et al.²⁴.

Interestingly, the patients with family history showed significantly milder disease severity than the patients without family history after adjusting for relevant demographic characteristics. Although there are some studies^{25,26} that family history is associated with early onset age, but it was difficult to find a study to explain a relationship between family history and disease severity. However, López-Esteban et al.²⁷ recently presented that the severity of psoriasis was not affected by the presence of a family history. Therefore, further studies are necessary to determine the association between family history and disease severity.

Finally, it was remarkable that significantly more patients in the PASI ≥ 10 group were dissatisfied with treatment results than the PASI < 10 group. This result indicates that more effective and satisfactory treatment is needed for patients with PASI ≥ 10 .

Although we attempted to minimize the bias from regional and temporal factors and reveal multilateral aspects of the clinical profiles of Korean patients with psoriasis, due to the recruitment of adult patients mostly from university hospitals, the representation of the general psoriasis population in Korea is limited. Further, because this study was cross-sectional, it is necessary to use caution when interpreting results for causal relationships between disease-related factors and outcomes over time. Therefore, additional research in the form of prospective longitudinal cohort studies is needed to further understanding of psoriasis in the Korean population.

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CONFLICTS OF INTEREST

Dr. Hae Jun Song has participated as an advisory board member and has conducted clinical trials for Janssen and has received speaking fees or research support from Janssen. Dr. Chul Jong Park, Dr. Tae Yoon Kim, Dr. Yong Beom Choe, Dr. Seok-Jong Lee, Dr. Nack In Kim, Dr. Jae We Cho, Dr. Jie Hyun Jeon, Dr. Min Soo Jang have no conflict of interest to declare. Dr. Jai Il Youn has conducted clinical trial for Pfizer, Janssen, Lilly, Leo. Dr. Myung Hwa Kim, Dr. Joonsoo Park, Dr. Ki Ho Kim, Dr. Byung Soo Kim have no conflict of interest to declare. Dr. Sang Woong Youn has served as an advisor, received speaker honoraria, and participated in clinical trials for

Janssen. Dr. Joo-Heung Lee has no conflict of interest to declare. Dr. Min-Geol Lee has conducted clinical trials for Pfizer, Novartis, Eli Lilly, and Janssen and has received speaking fees from Janssen. Dr. Sung Ku Ahn, Dr. Young Ho Won, Dr. Seok Kweon Yun, Dr. Bong Seok Shin have no conflict of interest to declare. Dr. Seong Jun Seo has conducted clinical trials for Amore, Regeneron, Novartis, Roche. Dr. Ji Yeoun Lee, Dr. Kwang Joong Kim have no conflict of interest to declare. Dr. Young Suck Ro has conducted post-marketing surveillance for Janssen. Mr. Dae Young Yu and Mr. Youngdoe Kim are employee of Janssen Korea. Dr. Jee-Ho Choi has conducted clinical trials for Janssen.

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