

## Validation of Psoriasis Diagnoses in the Danish National Patient Register

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In recent years, administrative registry data has been used with increasing frequency in dermato-epidemiological studies (1, 2). A common approach to the identification of patients is through use of the International Classification of Diseases, 10<sup>th</sup> revision (ICD-10) codes (2–4). This study aimed to validate the diagnostic code for psoriasis (ICD-10 L40.0) recorded in the Danish National Patient Register (5).

The study was approved by the Danish Data Agency (ref. 2012-58-0004, j.no. VD-2018-286, I-Suite no.: 6528) and the Danish Patient Safety Authority (j.no. 3-3013-2654/1, ref. EMGW).

From among all patients seen in our department (the Department of Dermatology and Allergy, Herlev and Gentofte Hospital, Hellerup, Denmark), we randomly selected 100 adults and all children with a recorded diagnosis of psoriasis vulgaris in the National Patient Register ICD-10 code L40.0 between 1 January 2008 and 23 October 2018. In case of multiple diagnoses of psoriasis during this period, the first of these was used as the index date. The criterion for validity was the use of terms consistent with psoriasis vulgaris with no other competing dermatological diagnosis (e.g. dermatitis or pustular psoriasis) in the patient's medical chart on the same date that the diagnosis was recorded. Data extracted from the chart included age at recorded diagnosis, age of psoriasis onset, sex, psoriasis type, Dermatology Life Quality Index (DLQI), body surface area (BSA), Psoriasis Area Severity Index (PASI), location, potential triggers, first location, and familiar disposition.

The study comprised a total of 137 patients, of whom 100 were adults and 37 were children (**Table I**). Few patients had an identified trigger ( $n=3$ ), and BSA was reported in only 2 patients. The psoriasis diagnosis could not be verified in a total of 4 patients, 2 adults and 2 children, from the medical chart on the same date as the diagnosis was recorded. In the medical chart 1 of the 2 children was noted to definitively have psoriasis on a later date, whereas the 3 remaining patients were reported to have psoriasisform dermatitis. Consequently, the overall positive predictive value (PPV) of the study was 97.1% (95% confidence interval (95% CI) 95.5–98.1). In adults, the PPV was 98.0% (95% CI 95.7–99.1), and in children the PPV was 94.6% (95% CI 88.7–97.5).

This study found a high PPV of the ICD-10 code for psoriasis in the Danish National Patient register. Recently, a medical chart review found a similarly high PPV

**Table I. Clinical and demographic characteristics of patients with L40 diagnosis**

	Adults (n=100)	Children (n=37)
Psoriasis diagnosis, n (%)		
At index date	98 (98)	35 (95)
At a later time-point	0	1 (3)
Psoriasisform dermatitis, n (%)	2 (2)	1 (3)
Psoriasis type, n (%)		
Vulgaris <sup>a</sup>	80 (80)	25 (71)
Guttate <sup>b</sup>	7 (7)	2 (2)
Pustular <sup>c</sup>	0	0
Sex, n (%)		
Male	58 (58)	18 (49)
Female	42 (42)	19 (51)
Age, years, mean ± SD	47.9 ± 15.9	12.6 ± 3.5
Age of onset, years, mean ± SD	28.0 ± 16.2	9.1 ± 4.0
Disease duration, years, mean ± SD	19.7 ± 16.7	3.0 ± 2.9
PASI, mean ± SD [range]	10.3 ± 6.1 [0.6–34.9] (n=61)	7.8 ± 3.8 [0–13.6]* (n=18)
DLQI, mean ± SD [range]	10.8 ± 6.7 [1–27] (n=43)	11.2 ± 11.2 [1–29] (n=6)
Patients with reported familial history, n (%)	80 (80)	33 (89)
Any	45 (45)	23 (62)
Father	22 (22)	8 (22)
Mother	11 (11)	7 (19)
Siblings	8 (8)	4 (11)
Grandparents	6 (6)	5 (14)
Patients with reported anatomical involvement, n (%)	67 (67)	31 (84)
Hands	18 (18)	2 (5)
Feet	10 (10)	2 (5)
Inverse	9 (9)	1 (3)
Genital	4 (4)	5 (14)
Scalp	34 (34)	27 (73)
Knee	14 (14)	2 (5)
Elbows	24 (24)	7 (19)
Face	10 (10)	11 (30)
Patients with reported first involvement, n (%)	11 (11)	14 (38)
Hands	1 (1)	2 (5)
Scalp	6 (6)	7 (19)
Elbows	1 (1)	1 (3)
Knee	1 (1)	0 (0)
Other	3 (3)	5 (14)

\*One patient was well-treated with a biologic at referral with a PASI of 0.

<sup>a</sup>The term(s) psoriasis vulgaris and/or plaque psoriasis are used in the chart. <sup>b</sup>The term guttate has been used in the chart. <sup>c</sup>The terms pustular and/or palmoplantar pustulosis are used in the chart.

PASI: Psoriasis Area and Severity Index; DLQI: Dermatology Life Quality Index; SD: standard deviation.

for diagnosis of atopic dermatitis in the same register (6). This demonstrates consistency in the use of ICD-10 codes in skin diseases in Denmark. However, as diagnoses were made by a dermatologist, the validity of diagnoses from other departments, e.g. emergency departments, might not be similar. Nevertheless, the given estimate of the PPV is considered conservative, as only patients with the diagnosis on the index date were included as cases.

The observed mean PASI corresponds to moderate-to-severe psoriasis (7) consistent with that seen in patients receiving systemic (mean  $\pm$  standard deviation (SD), PASI of  $8.3 \pm 6.4$ ) (8) and biologics (PASI of  $13.2 \pm 8.1$ ) (9), which is consistent with this being the first time most of the patients were seen in a university hospital. However, the recorded PASI ranged from negligible to very severe disease, which might be attributed to the fact that some patients were already receiving systemic treatments at referral. The high proportion of children with a familial history of psoriasis is consistent with type 1 psoriasis showing a strong genetic predisposition (10).

This study was limited by the single-centre design; however, since all hospital dermatology departments in Denmark are academic university departments, we suspect that similar findings would apply at other departments in the country.

In summary, this study suggests that the use of the ICD-10 code L40.0 to identify patients with psoriasis is a valid approach in registry-based studies.

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Leo Pharma and is a member of the advisory boards of Abbvie, Pfizer, Janssen-Cilag, Sanofi, Eli Lilly, Celgene and Novartis. CA and ASHO report no conflicts of interest.

## REFERENCES

1. Hennessy S, Bilker WB, Knauss JS, Margolis DJ, Kimmel SE, Reynolds RF, et al. Cardiac arrest and ventricular arrhythmia in patients taking antipsychotic drugs: cohort study using administrative data. *BMJ* 2002; 325: 1070.
2. Solomon DH, Massarotti E, Garg R, Liu J, Canning C, Schneeweiss S. Association between disease-modifying antirheumatic drugs and diabetes risk in patients with rheumatoid arthritis and psoriasis. *JAMA* 2011; 305: 2525–2531.
3. Egeberg A, Mallbris L, Gislason G, Hansen P, Mrowietz U. Risk of periodontitis in patients with psoriasis and psoriatic arthritis. *J Eur Acad Dermatol Venereol* 2017; 31: 288–293.
4. Egeberg A, Mallbris L, Gislason GH, Skov L, Hansen PR. Risk of multiple sclerosis in patients with psoriasis: a Danish nationwide cohort study. *J Invest Dermatol* 2016; 136: 93–98.
5. Schmidt M, Schmidt SAJ, Sandegaard JL, Ehrenstein V, Pedersen L, Sørensen HT. The Danish National Patient Registry: a review of content, data quality, and research potential. *Clin Epidemiol* 2015; 7: 449–490.
6. Andersen YM, Egeberg A, Skov L, Thyssen JP. Demographics, health care utilization, and drug use in children and adults with atopic dermatitis in Denmark: a population-based cross-sectional study. *J Eur Acad Dermatol Venereol* 2019; 33: 1133–1142.
7. 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: 194–199.
8. Egeberg A, Gyldenløve M, Zachariae C, Skov L. Validation of psoriasis severity classification based on use of topical or systemic treatment. *J Eur Acad Dermatol Venereol* 2018; 32: e4–e5.
9. Loft N, Skov L, Bryld L, Gislason G, Egeberg A. Treatment history of patients receiving biologic therapy for psoriasis – a Danish nationwide study. *J Eur Acad Dermatol Venereol* 2017; 31: e362–e363.
10. Henseler T, Christophers E. Psoriasis of early and late onset: characterization of two types of psoriasis vulgaris. *J Eur Acad Dermatol Venereol* 1985; 13: 450–456.