Effect of Hidradenitis Suppurativa Disease Duration on Psychiatric Comorbidity

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Hidradenitis suppurativa (HS) is a chronic inflammatory disease characterized by painful, often discharging, lesions of intertriginous areas of the skin. HS is more common in females than males and it occurs most often after puberty (1). HS is associated with several somatic and psychiatric comorbidities (2–4), depression and anxiety being the most common in both adult and adolescents (4, 5). HS is often unrecognized, and patients with HS may not seek medical help. Diagnosis may be delayed for more than a decade (6). The long diagnostic delay makes it impossible to reliably study the temporal relationship of HS and its comorbidities via registry data. Therefore, we performed a retrospective hospital-based study to analyse the effect of disease duration on psychiatric comorbidity.

MATERIALS AND METHODS

The Oulu University Hospital (OUH) database was queried to obtain the records of all patients diagnosed with HS at the hospital between 1996 and 2015. Each diagnosis was identified by the presence in the record of International Classification of Diseases, Tenth Revision (ICD-10) code L73.2. All patients with at least one registered L73.2 code were included. The retrieved patient records were reviewed by the authors for demographics, disease duration, smoking status, and pre-selected comorbidities, as well as the diagnosis date of any psychiatric comorbidity. Statistical analyses were conducted using the SAS software package (version 9.4, SAS Institute Inc., Cary, NC, USA). All results are presented as proportions and means. Because the study was based on a retrospective review of records, the agreement of the ethics committee was not required. However, the study methods were approved by the Medical Director of OUH.

RESULTS AND DISCUSSION

The database search returned the records of 304 patients; 187 females (61.5%) and 117 males (38.5%), mean age 38.5 years (range 12–79 years). At the time of diagnosis over 60% (n = 191) had had symptoms of HS for longer than 1 year. Smoking status was reported for 60.1% (n = 178/304) of cases and of these 75.8% (n = 135/178) were current smokers (Table I).

Sixty percent of patients had at least 1 comorbid disease, and over half of these (51.3%) had at least 2 comorbid diseases. All the investigated comorbidities are shown in Fig. 1. The most common comorbidities were cardiovascular diseases (CVD) (35.9%), diabetes (23.0%), obesity (20.4%) and depression (15.5%).

Regarding psychiatric comorbidities and their relationship with duration of HS symptoms, those with <1 year of HS symptoms were most likely to have a diagnosis of depression or another psychiatric comorbidity (Table II). Those who had had HS symptoms for longer than 5 years had a lower rate of psychiatric comorbidities. Patients with depression or other psychiatric comorbidities did not differ from those without in terms of age at diagnosis, diagnostic delay or other comorbidities studied.
This study found that 15.5% of patients with HS had a diagnosis of depression. This reflects both our previous findings based on Finnish registry data (4) and similar findings from Israel (3). A diagnosis of at least 1 psychiatric disorder other than depression was also found in 15% of cases. However, the study design prevented us from analysing these diagnoses in detail.

In the current study, depression and other psychiatric comorbidities were most often seen in patients whose HS had been diagnosed recently. The prevalence of depression and other psychiatric comorbidities decreased after onset of HS symptoms. Over the past decade there has been an increase in awareness of HS comorbidities (7). However, this does not entirely explain our findings, because the cut-off date for our study period was 2015. One possible explanation could be that patients who experienced a long diagnostic delay learned to cope with their HS symptoms before diagnosis, and were no longer experiencing any psychiatric consequences by the time of their HS diagnosis.

Most patients in the current study had at least 2 HS comorbidities. Cardiovascular diseases (CVD) and diabetes were the most common comorbidities. CVD were present in approximately 40% of the study population. This is notable because the reported prevalence of CVD in overweight middle-aged Finns is approximately 8%; much lower than that found in our comparatively young population (mean age <40 years) (8). However, more than 1 in 5 of our study subjects was obese. Previously, a large Danish registry-based study found that HS is a remarkable and independent risk factor for cardiovascular diseases (9), and an Israeli study reported that patients with HS have an elevated risk of metabolic syndrome, diabetes, hypertension and hyperlipidaemia (10). In addition, most of the subjects were previous or current smokers; smoking is known to be strongly associated with HS (11).

Although metabolic diseases are the most common comorbidities of HS, psychiatric disorders greatly diminish patients’ quality of life. To further reduce the psychiatric morbidity of HS patients, it is important that dermatologists are aware of the possibility of psychiatric problems when diagnosing HS. This would help to guide patients with HS to seek social support and psychiatric care if needed. These procedures may bolster resilience and coping in patients with HS, which, in turn, may help to prevent mental illness from becoming chronic (12).

A strength of this study is that it includes all patients with HS treated in the OUH over a 19-year period. All medical records of patients included in the study were reviewed carefully and used to verify HS diagnoses. The individual follow-up time varied from less than 1 year to 2 decades, and this can be considered as a limitation of this study. Furthermore, since this study was based on hospital medical records, it was unable to capture diagnoses made in the primary care setting.

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REFERENCES

1. Jemec GB, Kimball AB. Hidradenitis suppurativa: epidemiology and scope of the problem. J Am Acad Dermatol 2015; 73: 4.
2. Shlyankevich J, Chen AJ, Kim GE, Kimball AB. Hidradenitis suppurativa is a systemic disease with substantial comorbidity burden: a chart-verified case-control analysis. J Am Acad Dermatol 2014; 71: 1144–1150.
3. Shavit E, Dreijer J, Freud T, Halevy S, Vinker S, Cohen AD. Psychiatric comorbidities in 3207 patients with hidradenitis suppurativa. J Eur Acad Dermatol Venereol 2015; 29: 371–376.
4. Hulilja L, Tiri H, Jokelainen J, Timonen M, Tasanen K. Patients with hidradenitis suppurativa have a high psychiatric disease burden: a Finnish nationwide registry study. J Invest Dermatol 2018; 138: 46–51.
5. Tiri H, Jokelainen J, Timonen M, Tasanen K, Hulilja L. Somatic and psychiatric comorbidities of hidradenitis suppurativa in children and adolescents. J Am Acad Dermatol 2018; 79: 514–519.
6. Garg A, Neuren E, Cha D, Kirby JS, Ingram JR, Jemec GBE, et al. Evaluating patients’ unmet needs in hidradenitis suppurativa: results from the Global Survey of Impact and Healthcare Needs (VOICE) Project. J Am Acad Dermatol 2020; 82: 366–376.
7. Naik HB, Lowes MA. A call to accelerate hidradenitis suppurativa research and improve care-moving beyond burden. JAMA Dermatol 2019 Jul 10 [E-pub ahead of print].
8. Uusitupa M, Peltonen M, Lindström J, Aunola S, Ilanne-Parikka P, Keinänen-Kiukaanniemi S, et al. Ten-year mortality and cardiovascular morbidity in the Finnish Diabetes Prevention Study – secondary analysis of the randomized trial. PLoS One 2009; 4: e5656.
9. Egeberg A, Gislason GH, Hansen PR. Risk of Major adverse cardiovascular events and all-cause mortality in patients with hidradenitis suppurativa. JAMA Dermatol 2016; 152: 429–434.
10. Shalom G, Freud T, Harman-Boehm I, Polischuk I, Cohen AD. Hidradenitis suppurativa and metabolic syndrome: a comparative cross-sectional study of 3207 patients. Br J Dermatol 2015; 173: 464–470.
11. Kohorst JJ, Kimball AB, Davis MD. Systemic associations of hidradenitis suppurativa. J Am Acad Dermatol 2015; 73: 27.
12. Kirby JS, Butt M, Esmann S, Jemec GBE. Association of resilience with depression and health-related quality of life for patients with hidradenitis suppurativa. JAMA Dermatol 2017; 153: 1263–1269.