Evaluating Factors Impacting Hidradenitis Suppurativa Disease Severity in Patients With Darker Skin Types

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PRACTICE POINTS

- Anxiety and depression are highly prevalent among Black patients with hidradenitis suppurativa (HS).
 Implementing self-report questionnaires at medical office visits are crucial to identifying patients who could benefit from additional psychiatric resources.
- Hidradenitis suppurativa patients with skin of color may have a higher incidence of comorbid gastrointestinal disease than other HS patients.
- Investigating the impact of weight-management practices on disease severity would be helpful in outlining nonpharmacologic treatments for patients with HS.
- The patient cohort described here had a high prevalence of marijuana use but lower prevalence of current cigarette use compared to studies conducted in predominantly White HS patient populations, emphasizing the multifactorial nature of HS pathophysiology.

Hidradenitis suppurativa (HS) is a debilitating chronic skin disease that has a considerable psychosocial impact on patient quality of life. Despite HS disproportionately affecting individuals with darker skin types, most published HS research features patient cohorts predominantly consisting of White patients. In this retrospective chart review, we aimed to evaluate demographics, comorbidities, and pharmacologic treatments in a predominantly Black patient population that underwent surgical treatment for HS during the period from April 2014 to April 2023 at a regional verified burn center. Our results suggest that Black patients with HS may be impacted by anxiety, depression, and comorbid gastrointestinal disease at higher rates than White patients with HS. Also, the low rates of cigarette smoking yet high rates of marijuana use in our patient population suggest a multifactorial disease pathology, warranting further research. Studies investigating disease pathophysiology, risk factors, and outcomes

in HS patients with skin of color (SOC) are imperative to minimizing health care disparity gaps, improving disease outcomes, and providing more equitable health care for all patients.

idradenitis suppurativa (HS) is a debilitating chronic skin disease that often affects apocrine-bearing regions of the skin such as the axillae, perineum, and groin.¹ Although current research on the etiology and pathogenesis of HS is limited, the disease is known to have a considerable psychosocial impact on patient quality of life.

Clinically, HS lesions manifest as tender subcutaneous nodules that rupture to form painful and deep dermal abscesses.² These lesions typically develop due to hair follicle occlusion, followed by a cyclic process of inflammation, healing, re-inflammation, and scarring. Often, they are mistaken for cysts or a simple abscess in the early stages of the disease, leading to a delay in diagnosis.¹ Disease severity is categorized based on Hurley staging: stage 1 involves abscess formation without scarring; stage 2 involves limited sinus tracts and recurrent abscesses with scarring and/or multiple separated lesions; and stage 3 is the most advanced stage, with diffuse involvement or multiple interconnected sinus tracts across an area with scarring. The condition primarily is medically managed with antibiotics and immunomodulators, but patients who have refractory disease can benefit from surgical excision.^{1,2}

The prevalence of HS in the United States ranges from 0.77% to 1.19%, and individuals who self-identify as Black have 3-fold higher odds of having this condition compared with all other racial groups.³⁻⁵ Black patients also

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are thought to have a greater number and size of apocrine glands compared with patients who self-identify as White, suggesting an anatomic predisposition to developing HS and greater disease severity.⁶ However, despite HS disproportionately impacting individuals with skin of color (SOC), the majority of published HS research includes predominantly White patient cohorts.⁵ There is insufficient research assessing HS epidemiology, comorbidities, and treatment responses in patients with SOC.

A 2020 review reported the notable lack of clinical trials that sufficiently examine systemic medication treatment response in HS patients with SOC.7 Of the 15 HS treatment trials published from 2000 to 2019, only 16.4% (138/840) of the patient population were of African descent.7 Clinical trials investigating the efficacy of adalimumab in reducing HS burden also did not adequately evaluate clinical response in patients with SOC. One clinical trial did not include any Black patients as part of the cohort, 8 and in 3 other studies, 80% to 85% of the study participants self-identified as White.9 The current literature does not reflect the patient populations most affected by HS, as several studies have reported that 65% of patients diagnosed with HS in the United States annually are Black.^{5,7} These results emphasize the underrepresentation of SOC populations in the current HS literature and the need for more research that investigates the disease processes, comorbidities, and treatment outcomes of the diverse patient population impacted by HS.

Methods

Study Population and Data Extraction—Following a protocol reviewed and approved by the MedStar Health/ Georgetown University institutional review board (IRB #00006783), a retrospective chart review of 31 adult patients with HS who underwent surgery at a regional verified burn center from April 2014 to April 2023 was conducted. The following variables were collected from the electronic medical record (EMR): baseline demographics including age, sex, body mass index (BMI), obesity status, race, ethnicity, Fitzpatrick skin type, smoking status, substance use, employment status, and family history of HS; HS-specific details including Hurley staging, affected areas, and age at initial diagnosis; comorbidities such as dermatologic conditions, autoimmune disorders, infectious diseases, cardiovascular and associated diseases, ovarian disorders, gastrointestinal diseases, and other common chronic comorbidities (psychiatric illness, kidney disease, type 2 diabetes [T2D], asthma, allergies, lymphedema, and inflammatory eye disease); and use of pharmacologics such as topical medications, oral antibiotics, immunomodulators, and steroids.

Study Definitions—Obesity was defined as both a continuous and categorical variable. Each patient's BMI at the surgery date was recorded from the EMR as a continuous variable. Patients with obesity also had this condition listed under their complaints and problem list in the EMR, which was recorded as a categorical variable. Race and ethnicity

were self-reported by patients. Comorbidity data, including T2D and hyperlipidemia, were defined by previously diagnosed diseases listed in the EMR. Pharmacologic medication data were included in the study if a patient was recommended/prescribed a medication and they had confirmed use of the medication in a subsequent office visit.

Statistical Analysis—Descriptive statistics were calculated for demographics, HS characteristics (eg, location, Hurley stage), and comorbidities. Continuous variables were presented as mean and standard deviation or median and interquartile range and were evaluated using a t test or Mann-Whitney U test when appropriate. Categorical variables were presented as frequencies and percentages and tested for associations using the χ^2 or Fisher exact test. Data analyses were performed using SAS software version 9.4 (SAS Institute Inc.).

Results

Thirty-one patients (17 females, 14 males; mean age, 40.9 years) were included in the study. Twenty-nine (93.5%) patients identified as Black. All study patients had at least 1 comorbidity. Obesity was diagnosed in 22 (71.0%) patients (mean BMI, 35.5 kg/m²). A total of 16 (51.6%) patients were current smokers, 3 (9.7%) were past smokers, 22 (71%) reported alcohol use, and 17 (54.8%) were active marijuana users. Only 3 (9.7%) patients had a family history of HS (Table 1).

Other common comorbidities associated with HS were anemia (64.5% [20/31]), a non-inflammatory bowel disease gastrointestinal disease (61.3% [19/31]), allergies (54.8% [17/31]), hypertension (41.9% [13/31]), cardiovascular disease (41.9% [13/31]), T2D (32.3% [10/31]), asthma (32.3% [10/31]), kidney disease (29.0% [9/31]), and atopic dermatitis (25.8% [8/31]). More than half (54.8% [17/31]) of patients were diagnosed with psychiatric illnesses, including depression, anxiety, bipolar depression, psychosis, anorexia, impulsive anger, hallucinations, delusion, attention deficit-hyperactivity disorder, and panic disorder (Table 2). Depression was diagnosed in 38.7% (12/31) of patients, and 22.6% (7/31) were diagnosed with anxiety.

The most common anatomic locations for HS were the right axilla (74.2% [23/31]), left axilla (74.2% [23/31]), groin (71% [22/31]), perineum (61.3% [19/31]), buttocks (41.9% [13/31]), and thigh (41.9% [13/31]). Other locations included the breast, lower back, posterior neck, dorsal foot, and scalp (all 3.2% [1/31])(Table 3). Twenty (64.5%) patients had Hurley staging recorded in the EMR. Seventeen (54.8%) were categorized as Hurley stage 3, and 3 (9.7%) were categorized as Hurley stage 2.

Twenty-nine (93.5%) patients were prescribed an oral antibiotic regimen. The most common oral antibiotics were clindamycin (35.5% [11/31]), doxycycline (35.5% [11/31]), rifampin (29% [9/31]), trimethoprim/sulfamethoxazole (22.6% [7/31]), and cephalexin (22.6% [7/31]). Of the patients who were prescribed rifampin, 87.5% (8/9) also were prescribed an adjunct oral clindamycin regimen. Twenty-nine percent (9/31) of patients were prescribed

TABLE 1. Patient Demographics, Characteristics, and Substance Use (N=31)

Characteristic	Reported value
Sex, n (%)	
Male	14 (45.2)
Female	17 (54.8)
Mean age, y (SD)	40.9 (14.9)
Race/ethnicity	
Black	29 (93.5)
Other ^a	2 (6.5)
Mean BMI, kg/m² (SD)	35.5 (10.7)
Diagnosed with obesity, n (%)	22 (71.0)
Smoking status, n (%)	
Current smoker	16 (51.6)
Past smoker	3 (9.7)
Alcohol user, n (%)	22 (71.0)
Drug user, ^b n (%)	19 (61.3)
Marijuana user, n (%)	17 (54.8)
Positive family history of HS, n (%)	3 (9.7)

Abbreviations: BMI, body mass index; HS, hidradenitis suppurativa. ^aDid not have race information available in their chart or had "other" listed as their race.

a biologic regimen; 22.6% (7/31) were prescribed adalimumab, 3.2% (1/31) were prescribed secukinumab, and 3.2% (1/31) were prescribed ustekinumab (Table 4).

Twenty-five (80.6%) patients were prescribed a topical treatment regimen, the most common being topical clindamycin (45.2% [14/31]). Other topical medications included triamcinolone (12.9% [4/31]), chlorhexidine gluconate wash (9.7% [3/31]), clobetasol (3.2% [1/31]), hydrocortisone (3.2% [1/31]), and hydroquinone (3.2% [1/31]) (Table 4).

Other medical treatments for HS included metformin (25.8% [8/31]), spironolactone (16.1% [5/31]), and zinc supplements (12.9% [4/31]). Four patients (12.9%) were prescribed clindamycin plus rifampin as well as a combination of metformin, spironolactone, and/or zinc (Table 4).

Twenty-two (71.0%) patients had a history of receiving incision and drainage procedures as treatment for HS. All 31 patients underwent excisional surgery followed by appropriate reconstruction. The total number of excisional surgeries a single patient underwent for HS treatment ranged from 1 to 9, with a mean of 2 excisional surgeries per patient.

Comment

Our regional verified burn center in Washington, DC, serves a large population of patients with SOC, making it a unique and important sample to study for HS. Our results suggest that

TABLE 2. Most Common HS Comorbidities and Prevalence Rates Within the Study Population (N=31)

Comorbidity	No. of patients (%)
Anemia	20 (64.5)
Gastrointestinal disease ^a	19 (61.3)
Allergies	17 (54.8)
Psychiatric illness	17 (54.8)
Hypertension	13 (41.9)
Cardiovascular disease	13 (41.9)
T2D	10 (32.3)
Asthma	10 (32.3)
Kidney disease	9 (29.0)
Atopic dermatitis	8 (25.8)
Lymphedema	8 (25.8)
Hyperlipidemia	8 (25.8)
High cholesterol	7 (22.6)
Inflammatory eye disease	7 (22.6)
Female gynecologic disease	7 (22.6)
Acne	4 (12.9)
Folliculitis	4 (12.9)
Inflammatory bowel disease	3 (9.7)
MRSA infection	3 (9.7)
Rheumatoid arthritis	3 (9.7)
History of herpes zoster	2 (6.5)

Abbreviations: HS, hidradenitis suppurativa; T2D, type 2 diabetes; MRSA, methicillin-resistant *Staphylococcus aureus*.

Black patients with HS may be at a higher risk for depression and anxiety. Twelve (38.7%) of our patients were diagnosed with depression, which is substantially higher than the 17% to 21% depression prevalence rate among all HS patients reported in meta-analyses. 10,11 Additionally, 22.6% (7/31) of our patients were diagnosed with anxiety, which is higher than the 5% to 12% prevalence rate of anxiety among HS patients reported in meta-analyses.^{10,11} The stress of chronic disease management, psychosocial impact of living with HS, social stigma, sexual dysfunction, pain, and financial concerns make mental illness a debilitating yet common comorbidity for patients with HS. The results of our study suggest that anxiety and depression are highly prevalent among Black patients with HS. It is important to identify if this finding is due to the interplay of health care disparities and social determinants of health; the cause likely is multifactorial, as race and ethnicity may be potential predictors for increased disease severity. Hidradenitis suppurativa is known to be a major economic burden on patients, and race-dependent

blncludes marijuana, heroin, cocaine, and methamphetamines.

^aExcluding inflammatory bowel disease.

structural and societal inequalities may be influencing the increased prevalence of anxiety and depression among Black patients with HS.¹² Therefore, clinicians must be vigilant for the signs and symptoms of mental illnesses to refer patients for psychiatric treatment when appropriate. Implementing self-report Patient Health Questionnaire-9, General Anxiety Disorder-7 depression and anxiety screening tools, and Dermatology Life Quality Index questionnaires at primary care and dermatology office visits may be a beneficial step toward identifying patients who could benefit from additional mental health resources.¹³

The patients included in our study predominantly self-identified as Black, and the current smoker prevalence rate was 51.6% (16/31). This percentage is lower than the smoking rates of other published HS studies conducted in predominantly White patient populations, which report up to a 76.5% smoking prevalence rate. 14-16 One review article published in 2022 reported that approximately 90% of HS patients are current or former smokers.¹⁷ Additionally, a retrospective cohort analysis identifying HS cases among 3,924,310 tobacco smokers in the United States reported that tobacco smokers diagnosed with HS most commonly racially self-identified as White (66.2%). 18 Tobacco chemicals and smoke can increase inflammatory cytokine levels, and the activation of nicotinic acetylcholine receptors surrounding pilosebaceous-apocrine units can increase follicular occlusion. 14 While several studies 1-3,14,19,20 support the strong correlation between tobacco smoking and HS, there are very few that specifically investigate the association between smoking and HS disease in SOC populations. It is possible that smoking rates may be lower in Black patients with HS compared with White patients with HS, which would suggest a multifactorial nature of HS disease pathophysiology. Future large, multicenter studies are needed that investigate smoking rates and HS disease severity in patients across various racial groups.

Prior research has shown a strong correlation between cigarette smoking and HS, but there is minimal data on the role of use of marijuana and other illicit drugs in HS disease pathophysiology.²¹ A total of 54.8% of our patients were

TABLE 3. Most Common Anatomic Sites Impacted by HS Within the Study Population (N=31)

Anatomic site	No. of patients (%)	
Right axilla	23 (74.2)	
Left axilla	23 (74.2)	
Groin	22 (71.0)	
Perineum	19 (61.3)	
Buttock	13 (41.9)	
Thigh	13 (41.9)	
Abbreviations: HS, hidradenitis suppurativa.		

active marijuana users with daily or weekly usage. Further research is needed to investigate whether marijuana use is linked with HS disease pathophysiology and severity or if patients with HS may be using marijuana to relieve pain, anxiety, and depression. Additional studies that survey the method of marijuana use (eg, joint, vape devices, or edibles) would clarify the relationship between not only HS and marijuana but also a potential link between disease severity and the process of inhaling large amounts of smoke vs a link with the active ingredients in the marijuana plant itself.

Approximately 61% (19/31) of our patients were diagnosed with a gastrointestinal disease in addition to HS.

TABLE 4. Pharmacologic Treatment Interventions Prescribed in Study Population (N=31)

Treatment	No. of patients (%)
Oral antibiotics ^a	
Clindamycin	11 (35.5)
Doxycycline	11 (35.5)
Rifampin	9 (29.0)
Trimethoprim/sulfamethoxazole	7 (22.6)
Cephalexin	7 (22.6)
Total	29 (93.5)
Biologics/immunomodulators	
Adalimumab	7 (22.6)
Secukinumab	1 (3.2)
Ustekinumab	1 (3.2)
Total	9 (29.0)
Topical pharmacologics ^a	
Clindamycin	14 (45.2)
Triamcinolone	4 (12.9)
Chlorhexidine gluconate wash	3 (9.7)
Clobetasol	1 (3.2)
Hydrocortisone	1 (3.2)
Hydroquinone	1 (3.2)
Total	25 (80.6)
Other treatments ^b	
Metformin	8 (25.8)
Spironolactone	5 (16.1)
Zinc	4 (12.9)
Total	13 (35.5)

^aSeveral patients underwent combination oral and topical treatment consisting of multiple oral and/or multiple topical agents.

^bSeveral patients were taking a combination of one or more of these medications.

Current research reports the link between HS and inflammatory bowel disease, but few studies have investigated if a relationship exists between the gut microbiome and HS, as well as the incidence of general gastrointestinal disease among Black patients with HS.14,22 Our patients were diagnosed with gastrointestinal conditions such as colonic polyps, gastroesophageal reflux disease, benign neoplasms of the cecum and sigmoid colons, small bowel obstruction and perforation, biliary tract diseases, ileus, abdominal hernia, peritonitis, and diverticulosis. Further research is warranted to identify if there is a true relationship between gastrointestinal disease, the gut microbiome, and skin conditions such as HS.22 Biochemical research on the common genetic and inflammatory cytokine pathways involved in HS and gastrointestinal manifestations could help predict disease severity and management in HS patients with SOC.

Several research studies have reported the association between obesity and HS, likely due to adipose cells producing increased estrogen and leading to an estrogendominant hormone profile and increased local androgen production in adipose tissue. 14,23,24 Antiandrogenic drugs such as finasteride and spironolactone lead to positive results in HS treatment compared to oral antibiotics alone.²⁴ While 71.9% (22/31) of our patients were diagnosed with obesity, only 16.1% (5/31) were prescribed antiandrogen therapy such as spironolactone. It is unclear if this result reflects a health disparity due to insufficient insurance coverage and low prescribing rates or if there is patient hesitancy to taking antiandrogen medications. Additional clinical trials are needed to investigate the efficacy of antiandrogen therapies for HS. If proven to be efficacious, providers should consider adding these medications to the pharmacologic regimen of HS patients with SOC prior to recommending wide-excision surgeries. Furthermore, in addition to antiandrogen medication, weight-management interventions may be helpful in reducing HS disease. The results of a survey conducted in 35 HS patients who underwent bariatric surgery reported 48.6% (17/35) experienced complete disease remission after more than a 15% weight reduction. 25,26 Investigating the impact of weight-management practices on disease severity would be helpful in outlining nonpharmacologic treatments for patients with HS.

Limitations

Our study was limited by the constraints of a retrospective chart review and small sample size. Retrospective chart reviews are susceptible to recall bias, variability in providers' charting practices, and human error from data collectors. We acknowledge that a control group of non-HS patients should be the next step in furthering our research on HS disease comorbidities. Also, since 35.5% (11/31) of our patients did not have Hurley staging recorded in the EMR, it would be beneficial to conduct a future study comprehensive of all 3 Hurley stages. Since 93.5% (29/31) of the patients in our study racially

identified as Black, having a control group of racially diverse HS patients would help further our understanding of HS pathophysiology. Lastly, since the inclusion criteria required patients to have undergone excisional surgery for HS, future studies that consider comorbidities among both surgical and nonsurgical patients with HS will aid in our understanding of HS patients with SOC.

Conclusion

The results of our study demonstrate a descriptive analysis of the demographics, most common comorbidities, lesion sites, pharmacologic treatments, and surgical profiles in patients with SOC who underwent surgical treatment for HS. Our data show that HS patients with SOC may be more likely to experience anxiety, depression, and gastrointestinal disease than other HS patients. Additionally, our patients had a high prevalence of marijuana use but lower prevalence of current cigarette use compared to studies conducted in predominantly White HS patient populations, emphasizing the multifactorial nature of HS pathophysiology. Furthermore, despite published research on the efficacy of immunomodulator therapy for HS, most of our HS patients with SOC underwent surgical intervention without first attempting biologic treatment regimens, indicating possible gaps in health care access for minority patients that may be impacting disease severity and outcomes. Studies such as this one that investigate disease pathophysiology and risk factors in SOC patient populations with HS are imperative in minimizing the health care disparity gap, improving disease outcomes, and providing more equitable health care for all patients.

REFERENCES

- Wieczorek M, Walecka I. Hidradenitis suppurativa—known and unknown disease. Reumatologia. 2018;56:337-339. doi:10.5114/reum.2018.80709
- Alikhan A, Lynch PJ, Eisen DB. Hidradenitis suppurativa: a comprehensive review. J Am Acad Dermatol. 2009;60:539-563. doi:10.1016/j. jaad.2008.11.911
- Garg A, Lavian J, Lin G, et al. Incidence of hidradenitis suppurativa in the United States: a sex- and age-adjusted population analysis. J Am Acad Dermatol. 2017;77:118-122. doi:10.1016/j.jaad.2017.02.005
- Ingram JR, Jenkins-Jones S, Knipe DW, et al. Population-based Clinical Practice Research Datalink study using algorithm modelling to identify the true burden of hidradenitis suppurativa. *Br J Dermatol*. 2018;178:917-924. doi:10.1111/bjd.16101
- Lee DE, Clark AK, Shi VY. Hidradenitis suppurativa: disease burden and etiology in skin of color. *Dermatology*. 2017;233:456-461. doi:10.1159/000486741
- Brown-Korsah JB, McKenzie S, Omar D, et al. Variations in genetics, biology, and phenotype of cutaneous disorders in skin of color—part I: genetic, biologic, and structural differences in skin of color. *J Am Acad Dermatol*. 2022;87:1239-1258. doi:10.1016/j.jaad.2022.06.1193
- Narla S, Lyons AB, Hamzavi IH. The most recent advances in understanding and managing hidradenitis suppurativa. F1000Res. 2020;9:F1000 Faculty Rev-1049. doi:10.12688/f1000research.26083.1
- Arenbergerova M, Gkalpakiotis S, Arenberger P. Effective long-term control of refractory hidradenitis suppurativa with adalimumab after failure of conventional therapy. *Int J Dermatol.* 2010;49:1445-1449. doi:10.1111/j.1365-4632.2010.04638.x
- Kimball AB, Okun MM, Williams DA, et al. Two phase 3 trials of adalimumab for hidradenitis suppurativa. N Engl J Med. 2016;375:422-434. doi:10.1056/NEJMoa1504370

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- Jalenques I, Ciortianu L, Pereira B, et al. The prevalence and odds of anxiety and depression in children and adults with hidradenitis suppurativa: systematic review and meta-analysis. *J Am Acad Dermatol*. 2020;83:542-553. doi:10.1016/j.jaad.2020.03.041
- Machado MO, Stergiopoulos V, Maes M, et al. Depression and anxiety in adults with hidradenitis suppurativa: a systematic review and meta-analysis. *JAMA Dermatol*. 2019;155:939-945. doi:10.1001/jamadermatol.2019.0759
- Kilgour JM, Li S, Sarin KY. Hidradenitis suppurativa in patients of color is associated with increased disease severity and healthcare utilization: a retrospective analysis of 2 U.S. cohorts. *JAAD Int*. 2021;3:42-52. doi:10.1016/j.jdin.2021.01.007
- Rymaszewska JE, Krajewski PK, Szczęch J, et al. Depression and anxiety in hidradenitis suppurativa patients: a cross-sectional study among Polish patients. Postep Dermatol Alergol. 2023;40:35-39. doi:10.5114ada.2022.119080
- Johnston LA, Alhusayen R, Bourcier M, et al. Practical guidelines for managing patients with hidradenitis suppurativa: an update. J Cutan Med Surg. 2022;26(2 suppl):2S-24S. doi:10.1177/12034754221116115
- Vazquez BG, Alikhan A, Weaver AL, et al. Incidence of hidradenitis suppurativa and associated factors: a population-based study of Olmsted County, Minnesota. J Invest Dermatol. 2013;133:97-103. doi:10.1038/jid.2012.255
- Seyed Jafari SM, Knüsel E, Cazzaniga S, et al. A retrospective cohort study on patients with hidradenitis suppurativa. *Dermatology*. 2018;234:71-78. doi:10.1159/000488344
- Lewandowski M, Świerczewska Z, Barańska-Rybak W. Hidradenitis suppurativa: a review of current treatment options. *Int J Dermatol.* 2022;61:1152-1164. doi:10.1111/ijd.16115
- Garg A, Papagermanos V, Midura M, et al. Incidence of hidradenitis suppurativa among tobacco smokers: a population-based

- retrospective analysis in the U.S.A. *Br J Dermatol.* 2018;178:709-714. doi:10.1111/bid.15939
- Garg A, Malviya N, Strunk A, et al. Comorbidity screening in hidradenitis suppurativa: evidence-based recommendations from the US and Canadian Hidradenitis Suppurativa Foundations. *J Am Acad Dermatol*. 2022;86:1092-1101. doi:10.1016/j.jaad.2021.01.059
- Tzellos T, Zouboulis CC. Which hidradenitis suppurativa comorbidities should I take into account? Exp Dermatol. 2022;31(suppl 1):29-32. doi:10.1111/exd.14633
- Metko D, Mehta S, Piguet V. Cannabis usage among patients with hidradenitis suppurativa: a scoping review. J Cutan Med Surg. 2024;28:307-308. doi:10.1177/12034754241238719
- Mahmud MR, Akter S, Tamanna SK, et al. Impact of gut microbiome on skin health: gut-skin axis observed through the lenses of therapeutics and skin diseases. *Gut Microbes*. 2022;14:2096995. doi:10.1080/194 90976.2022.2096995
- Mair KM, Gaw R, MacLean MR. Obesity, estrogens and adipose tissue dysfunction—implications for pulmonary arterial hypertension. *Pulm Circ*. 2020;10:2045894020952019. doi:10.1177/2045894020952023
- Abu Rached N, Gambichler T, Dietrich JW, et al. The role of hormones in hidradenitis suppurativa: a systematic review. *Int J Mol Sci.* 2022;23:15250. doi:10.3390/ijms232315250
- Alikhan A, Sayed C, Alavi A, et al. North American clinical management guidelines for hidradenitis suppurativa: a publication from the United States and Canadian Hidradenitis Suppurativa Foundations: part I: diagnosis, evaluation, and the use of complementary and procedural management. J Am Acad Dermatol. 2019;81:76-90. doi:10.1016/j.jaad.2019.02.067
- Choi ECE, Phan PHC, Oon HH. Hidradenitis suppurativa: racial and socioeconomic considerations in management. *Int J Dermatol*. 2022;61:1452-1457. doi:10.1111/ijd.16163