Immuneological Pathways in Hidradenitis Suppurativa: Current Concepts and Innovative Therapies

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Abstract

Hidradenitis Suppurativa is a chronic and disabling disease characterized by pustules, nodules, abscesses, and scarring. It affects natural body folds (inguinal, axillary, inframammary), perianal region and genitalia. Several comorbidities are related to this disease, such as obesity, smoking, acne, folliculitis, and hyperandrogenism. Disorders in keratinization, systemic upregulation of complement (C5a) activation and changes in cutaneous microbiome stimulate the development of an autoimmune state, promoting the increase of CD4+ T-help lymphocytes (Th-17 and Th-1). These changes stimulate the expression of toll-like receptors type 2, enabling the NLRP3 inflammasome and triggering of the caspase-dependent inflammatory cascade. Intralesional increase of dendritic cells, macrophage and neutrophil influx stimulate the production of pus, promoting necrosis and apoptosis of inflammatory cells undergoing a special type of cell death program, releasing proinflammatory cytokines (pyroptosis). In addition, patients affected with this condition presents an imbalance of IL-36 proinflammatory cytokines and decreased levels of its antagonist receptor IL-36Ra. Treatment is frustrating and includes the use of broad-spectrum antibiotics, surgical procedures, and oral retinoids. The first and only one biologic therapy approved is a TNF-α inhibitor (Adalimumab); other alternatives are under development, including IL-17 (Secukinumab, Bimekizumab and CJM112) IL-12 and IL-23 (Ustekinumab); and IL-1 (Anakinra, Canakinumab). Recent articles showed that liraglutide, a glucagon-like-1 peptide (GLP-1) agonist, could be used to treat one of the comorbidities associated with HS (insulin resistance). Finally, inhibition of IL-36 and C5a could be an option to treat this recalcitrant and exhausting disease.

Highlights: Pathogenesis of HS involves genetic and environmental factors that trigger the innate immune response.

Several cytokines participate in the inflammatory response of the skin.

HS is chronic and relapsing causing a negative impact on patient’s quality of life, expensive medical approaches, and huge number of comorbidities.

New biological drugs, former medical treatments and cutaneous surgical interventions can be combined to better assist of patients suffering of HS.

Keywords: Hidradenitis suppurativa; Biologics; IL-17; IL-1; IL-23; C5a; Autoinflammatory diseases;

Introduction

Hidradenitis Suppurativa (HS), also known as “ace inversa” or “maladie de Verneuil,” is a rare inflammatory skin disorder, about 0.035% to 4% of the world population suffer from this condition, mostly affecting adults and adolescents with few reported cases in prepubertal children \([1,2]\). Disease course is recurrent or intermittent and chronic, with multiple clinical variants, a 3:1 female-male ratio, and apparently without ethnic preference \([2]\). However, Afro-descendants seem to develop more exuberant lesions than Caucasians patients. Usually the onset is puberty-related, suggesting an endocrine component and associated with increased levels of androgenic hormones \([2]\).

Due to a delay in diagnosis and lack of awareness by physicians, final diagnostic is achieved after an average of 12 years \([2]\). Also, HS patients score high values in the Dermatology Quality of Life Index (DLQI), with higher values than the average found in psoriasis \([3]\). High values in this score are related to a greater impact in patient’s life \([3]\).

Heterogeneous mutation in the intramembranous protease complex γ-secretase and its subunits genes NCSTN, PSENEN, and PSEN1 has been associated with familial inheritance of HS. These mutations have been associated with impaired inhibition of the innate immunity by the Notch-MKP-1 signaling, promoting cutaneous proinflammatory microenvironment, inducing follicular keratinization and epidermal hyperplasia. Furthermore, mutations of γ-secretase complex have also been implicated in Alzheimer’s Disease (AD); however, no link between HS and AD has been documented. About 42% of patients report family history, early onset at puberty and more severe disease manifestations \([4]\). In addition, mutations in PSENEN are found to be present in patients with Dowling-Degos Disease (DDD). Around half of patients with DDD develop HS and share similar trigger factors \([5]\).

Discussion

Clinical Features

HS is characterized by papules, nodules, abscesses, and painful
suppurating sinus, these lesions can progress into fibrosclerotic plaques, keloids, or atrophic scars, decreasing and disabling the patient movement [6]. Inguinal, axillary, inter gluteal and inframammary folds are predominantly affected, however, ectopic locations such as the head, torso, genitals, dorsum, and limbs are uncommonly found in some patients [6]. Chronic and advanced cases may develop lower limb and genitalia lymphedema, as well as, hinder social and sexual relationships [7]. The disease has a recurrent course with an average of 2 to 3 crises every 6 months [2].

Diagnosis of HS is often clinical. Cutaneous biopsies are not required in most cases [2]. There are several scoring systems for assessing disease severity: Hurley Staging System (HSS), HS-Physician’s Global Assessment, Modified Sartorius Score (MSS), and Hidradenitis Suppurativa Severity Index. Unfortunately, all of them have some limitations in daily practice, with no consensus among the outcome measures used for HS [2].

HSS and MSS are the most widely applied disease activity scores [2]. Hurley scoring is usually stratified into three stages [2]. Stage I: single or multiple abscess, without scars and sinus tracts formation. Stage II: recurrent single or multiple abscesses with sinus tracts and scars formation, lesions are widely separated. Stage III: diffuse involvement or multiple interconnected tracts and abscesses formation across entire area.

Smoking, obesity, hyperandrogenism and women with Polycystic Ovarian Syndrome (PCOS) seem related to HS, worsening the disease evolution [8].

Remarkably, the presence of other associated diseases such as acne, pyoderma gangrenosum, neutrophilic diseases, Crohn’s disease, pilonidal cyst, spondyloarthropathies and psoriatic arthritis are commonly seen in some patients. These conditions could be overlapping and develop syndromes such as the SAPHO (Synovitis, Acne, Pustulosis, Hyperostosis and Osteitis), PASH (Pyoderma gangrenosum, Acne and Hidradenitis suppurativa), PAPA (Pyogenic Arthritis, Acne, and Hidradenitis suppurativa) and PAPASH (Arthritis, Pyoderma, Acne, and Hidradenitis suppurativa) [8].

Due to its polymorphism, multiple differential diagnoses are included, such as lymphogranuloma venereum, donovanosis, tertiary syphilis, paracoccidioidomycosis, furunculosis, atypical mycobacterial and tuberculosis infection with abscesses, metastatic Crohn’s disease and anal fistulas. Delayed diagnosis is common and usually done by an experienced physician [8].

Pathophysiology

The pathogenesis of HS until recent years was still poorly understood. Because lesions are localized in body folds areas such as the axilla, groin, perineum, and genitalia, it was long thought to be a primary disorder of the apocrine glands [9]. During the last decade, several evidences clarified that the primary defect in HS pathophysiology involves occlusion and subsequent inflammation of the hair follicle [9]. Morphological changes in the pilosebaceous unit plays a critical role in the development of HS, these follicular changes are characterized by an hyperkeratinization of the follicle infundibulum associated to defects in keratin proteins production (decreased number of cytokeratin CK17, with upregulated CK5 and CK6) [10,11]. This suggests fragility of the draining sinus epithelium, which may allow rupture to occur more easily, thus resulting in a subcutaneous abscess formation. Blocked follicles modify the skin microbiome and increases bacterial proliferation (S. epidermidis and P. acnes), generating a film-like substance that remain fixed to the follicular unit (bacterial biofilm), and stimulating skin overexpressed Toll-like receptors type 2 (TLR2) [12-15].

TLR2 is a vital component of the innate immunity, present in polymorphonuclear cells, keratinocytes, and dendritic cells. Fragments of bacterial antigens called PAMPs (Molecular patterns associated with pathogens) activate this receptor and initiate skin inflammation.

Activation of TLR2, increase intralesional expression of endogenous antibiotics called antimicrobial Peptides (AMPs), such as is human beta defensin-2 (HBD2) and cathelicidin LL37 (LL37) [15]. These peptides interfere further in the inflammatory process and induce hyperkeratinization of the hair follicle, triggering the formation of the NLRP3 inflammasome and of the pro inflammatory cytokines (IL-1β, IL-12, IL-23, IL-17, and TNF-α).

NLRP3 inflammasome activates the caspase inflammatory cascade, increasing caspase-1 expression, this enzyme plays an important role in phagocytes tissue migration, promoting polymorphonuclear cells death in presence of newly formed interleukin-1beta (IL-1β) by a process called pyroptosis (cell death with release of inflammatory cytokines) [16].

This process leads to cellular DNA damage and rapid pore formation in dendritic and macrophages cells membrane, resulting in cellular lysis and release of intracellular inflammatory contents, including Interleukin-1beta and alpha (IL-1β, IL-1α) [17]. Excess release of IL-1α from the cytosol of necrotic and stressed-cells leads to the recruitment of hematopoietic cells that can further sustain perilessional inflammation through endothelial activation and disruption of vascular walls [17].

Neutrophils migrate into the damaged skin and undergo phagocytosis and cellular death, forming networks of extracellular fibers mainly composed by neutrophilic nuclear and granular contents such as elastase, myeloperoxidase, and DNA, binding to pathogens fragments (NETosis) [18]. This process further stimulates the expression of interleukins, especially IL-8, TNF-α and IL-17; promoting chemotaxis of new neutrophils into the affected skin, pus formation and continuing the inflammatory loop [19].

Recent studies indicate that intralesional and serological increase of Th-17 T-helper lymphocytes and its cytokine (IL-17 and IL-22) may be a significant risk factor for the development of HS [20]. Patients with metabolic syndrome, smokers and Polycystic Ovary Syndrome (PCOS) have a considerable serological augmentation in both IL-17 and TNF-α [21-24]. These systemic alterations could explain the link between these comorbidities with HS [19].
Hyperandrogenism, puberty and PCOS stimulate the production of androgen hormones, increasing follicular hyperkeratosis and sebum production; these alterations clog the pore, generating more inflammation [24,25]. Besides the systemic inflammatory effect, obesity generates friction in the affected body folds, worsening the follicular obstruction [26]. Remarkably, smoking also increase inflammatory cytokines and stimulate skin cholinergic receptors, these alterations will increase follicular keratinization and neutrophil migration [22].

Recent publications have shown an increased expression of proinflammatory interleukins IL-36 (IL-36α, IL-36β and IL-36γ) and decreased antagonist cytokines (IL-36Ra, IL-37, IL-38) in the lesional and perilesional skin of patients with HS [27].

Pro-inflammatory IL-36 are members of the interleukin-1 (IL-1) superfAMILY, involved in the inflammassome activation, and inducing pro-inflammatory signaling pathways through the activation of the Nuclear Factor-kB (NF-kB) and Mitogen Activated Protein Kinase (MAPK) [27]. These cytokines may have an important role in the onset and regulation of the inflammatory response, especially in those patients with autoimmune and autoinflammatory diseases, including pustular psoriasis, rheumatoid arthritis, and inflammatory bowel diseases [28-30].

In HS, IL-36 increase Dendritic Cells (DC) activation, neutrophils influx, keratinocytes proliferation, and secretion of proinflammatory factors, (IL-1β, TNF-α, IL-6, IL-8), stimulating the production of T-helper lymphocytes type 1 (Th1) and Th17, and its cytokines (IL-17, IL-22 and IL-23). This loop perpetuates the autoinflammatory process [31].

Patients suffering of HS present decreased expression of the IL-36 receptor antagonist (IL-36Ra). This antagonist is commonly expressed in dendritic cells, lymphocytes, and keratinocytes, inhibiting the union of IL-36 with its peripheral receptor (IL-36R), regulating the inflammatory process in healthy patients [28]. Also, immunosuppressive cytokine (IL-10) seen to be upregulated in affected skin as a compensatory response to the pro inflammatory process. However, this augmentation suppresses only IL-12 with no modification in IL-17 lesional levels. Lower levels of IL-22 induce a decreased formation of AMP compared to psoriasis; maybe this could explain the defected skin antimicrobial function in HS [8].

Recently, Kanni et al. demonstrated that systemic complement activation is up-regulated in HS patients [32]. Elevated C5a and membrane attack complex (C5b-9) induce the expression of TNF-α by circulating mononuclear cells, contributing to the lesional chemotaxis and infiltration of neutrophils [32]. (Figure 1)
Therapeutic Approaches

Hidradenitis suppurativa is a difficult-to-treat disease. Due to smoking and obesity show the strongest correlation with disease severity, there is a consensus that patients suffering for HS must be advised and supported in the cessation of tobacco use and in reducing their body weight [2].

For long time, patients with mild and intermediate stages (Hurley I-II) undergo clinical treatment consisted of oral broad-spectrum antibiotics (Doxycycline, Trimethoprim-Sulfamethoxazole, Clindamycin), oral retinoids (Acitretin, Isotretinoin), dapsone, 15% topical resorcinol and small surgical procedures (Deroofing, laser, and local excision) [2].

In severe stages (Hurley III) the treatment is based on wide excision surgeries, forming deformed scars, with high rate of recurrence after procedures [2,33].

On table 1, we summarized the actual and new promising biological drugs studied to treat HS in distinct points of its pathophysiology.

<table>
<thead>
<tr>
<th>Name</th>
<th>Target</th>
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<th>Administration</th>
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<tr>
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<td>Subcutaneous</td>
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<td>TNF-α</td>
<td>II</td>
<td>Subcutaneous</td>
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<td>Infliximab</td>
<td>TNF-α</td>
<td>II</td>
<td>Intra venous</td>
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<td>IL-1 antagonist receptor</td>
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<td>MABp1</td>
<td>IL-1α</td>
<td>II</td>
<td>Intra venous</td>
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<td>Ustekinumab</td>
<td>IL-12 e IL-23 from p40 subunits</td>
<td>II</td>
<td>Subcutaneous</td>
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<td>Secukinub</td>
<td>IL-17A</td>
<td>I</td>
<td>Subcutaneous</td>
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<td>Bimekizumab</td>
<td>IL-17, IL-17F</td>
<td>II</td>
<td>Subcutaneous and Intra venous</td>
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<td>CJM112</td>
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<td>II</td>
<td>Subcutaneous</td>
</tr>
<tr>
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<td>GLP-1 agonist</td>
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<tr>
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<td>IL-36R</td>
<td>I, in a study for generalized pustular psoriasis</td>
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<td>1FX-1</td>
<td>C5a</td>
<td>II</td>
<td>Intra venous</td>
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TNF-α Inhibitors

With the development of monoclonal antibody-based therapy, the first approved immunobiological for HS was launched, adalimumab, a human monoclonal antibody capable of blocking TNF-α improving lesions and patient's quality of life after 12 weeks of treatment [34]. In some patients, early initiation of this therapy combined with surgery may result in a decrease in outcome and deforming lesions [35]. Adalimumab is given with an initial subcutaneous dose of 160mg on day 0 of treatment, then 80mg on day 14, and lastly 40mg per week from day 29 of treatment [34]. Other anti-TNF-α agents such as infliximab or etanercept have only reached Phase II clinical studies and failed to prove to be more effective than adalimumab [36-38].

IL-12 and IL-23 Inhibitors

Ustekinumab is a human monoclonal antibody that acts against the p40 subunit of interleukins IL-12 and IL-23. Approved for use in psoriasis and Crohn's disease, it was used in an open study where 17 HS patients participated, achieving 12 of them a good response after 40 weeks of treatment [39]. However, more clinical studies are needed to prove the efficacy and safety of this drug in HS. Three new selective p19 subunit inhibitors are currently in phase III clinical trials, guselkumab, risankizumab, and tildrakizumab, preliminary results have shown greater efficacy than ustekinumab in psoriatic patients [40]. However, the use of them in HS is still not being studied; moreover, it would be interesting to do a clinical study focused on this condition.

IL-1 Inhibitors

Anakinra is a recombinant IL-1 receptor antagonist (IL-1Ra) used in patients with rheumatoid arthritis. In a study involving 12 HS patients, 10 of them achieved considerable results after 12 weeks of treatment [41]. However, there are reports of isolated cases with no therapeutic benefits [42].

Canakinumab is a monoclonal antibody against IL-1β, useful in autoimmune diseases [43]. Several reports have shown promising results in patients with HS. However, there is no extensive study proving its efficacy.

MABp1 is a recombinant human IgG1 monoclonal antibody specific for human interleukin-1α. The mechanism of action is thought to be the inhibition of neovascularization and modulation of the production of IL-8 and human β-defensin-2. A randomized study with 20 patients, has recently viewed this drug as a promising agent for HS, including those not eligible for treatment with adalimumab [44].
IL-17 Inhibitors

Secukinumab is a monoclonal antibody against IL-17A, with approved use for moderate to severe psoriasis, there are 2 cases with HS treated with satisfactory results and currently the originator is recruiting HS patients for new clinical trials [45-47]. Two new monoclonal antibodies against IL-17, bimekizumab and CJM11243, are being evaluated as possible treatment in patients with HS, yet without preliminary results [48,49].

IL-36 Inhibitors

AnaptysBio Inc. announced positive results in a phase I trial evaluating a new drug (ANB019), an anti-interleukin-36 (IL-36R) antibody [51]. Obtained data demonstrated pharmacokinetic and pharmacodynamic safety and may now continue phase II clinical research in patients with palmoplantar and generalized pustular psoriasis [51]. As previously described, in HS there is an increase of this interleukin. Although these drugs are still in development, soon they may be a promising therapeutic option against this chronic and recalcitrant condition.

C5a Inhibitors

In an open label clinical phase II Study, a novel C5a-specific monoclonal antibody (IFX-1) effectively neutralized the overproduced C5a in HS patients. IFX-1 was tested in 12 patients with moderate to severe HS. All patients received nine intravenous doses of 800 mg IFX-1 on days 1, 4, 8, 15, 22, 29, 36, 43 and 50. Plasma level of C5a was determined before the trial, at days 22 and 50 during the treatment with IFX-1, with a significant decrease of C5a levels the end of study [52].

Hidradenitis Suppurativa Clinical Response (HiSCR) was applied to evaluate clinical improvement to treatment, achieving up to 83% response rate at the endpoint of the study. This novel drug may represent an effective treatment for HS patients [52].

Conclusion

In summary, HS is a recurrent, disabling, and difficult-to-treat disease. Several biological events play to develop an autoinflammatory milieu in the skin.

This condition is related with other comorbidities, as insulin resistance and hyperandrogenic states. With the development of new therapies, better knowledge of the pathophysiology, we hope that more physicians will be able to improve the quality of life of patients suffering from HS.

References


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