Two Cases of Paradoxical Hidradenitis Suppurativa while on Adalimumab

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Abstract
Hidradenitis suppurativa (HS) is a chronic, inflammatory skin disease characterized by recurring abscesses, nodules, and fistulas predominantly in the groin and axillae. The association between HS and Crohn’s disease (CD) has been well documented. Tumor necrosis factor (TNF) inhibitors have shown to be effective in treating both HS and CD. We report 2 patients who developed HS while on TNF inhibitor treatment for CD.

Introduction
Crohn’s disease (CD) is an idiopathic chronic inflammatory bowel disease that can affect any part of the gastrointestinal tract. Hidradenitis suppurativa (HS) is a chronic inflammatory skin disease characterized by recurrent deep, painful nodules, often of the axilla, groin and gluteal areas, that can lead to abscesses and chronic draining sinus tracts [1]. The association between HS and CD is well documented [2]. Patients with inflammatory bowel disease were approximately nine times more likely to develop HS than the general population in a population-based cohort study [3]. Tumor necrosis factor (TNF) inhibitors have been shown to be effective in treating recalcitrant HS [4] and moderate to severe CD. [5] Parado-

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ical psoriatic skin lesions while on anti-TNF therapy have also been well described [6]. We present 2 patients who developed HS while receiving treatment with adalimumab for CD. We describe some of the immunologic similarities between HS and the psoriasiform rash that has been noted on anti-TNF therapy, and explain why HS should be considered as one of the rare skin rashes that can develop while a patient is on anti-TNF therapy.

Case 1

A 57-year-old African American female presented to our emergency department with a 6-month history of diarrhea, urgency and tenesmus. She was found to have a perirectal abscess. CT of the abdomen was significant for thickening of the rectum and sigmoid colon. On rectal examination, she was noted to have an anal fistula. The perirectal abscess was drained, and a seton was placed for the anal fistula. A colonoscopy was performed showing erythema and friability consistent with colitis extending to the transverse colon, although most pronounced in the sigmoid colon and rectum. The remainder of the colon and the terminal ileum were normal. Biopsies from the involved portion of the colon including the rectum showed acute cryptitis with crypt abscess formation and crypt distortion and were consistent with CD. The patient was started on adalimumab and mesalamine suppositories with significant clinical improvement. A follow-up sigmoidoscopy after 8 months showed resolution of the colitis with only mild proctitis remaining. The patient continues to smoke despite counseling.

Twelve months into treatment, she developed recurrent boils and nodules in her groin. Her white blood cell count was normal (6,900/μl). She was started on amoxicillin 875 mg and clavulanic acid 125 mg orally twice a day for 10 days. With the suspicion of HS, she was referred to the dermatology clinic, and the diagnosis was confirmed. Her skin lesions improved only transiently on the antibiotics. Her disease progressed with multiple boils and draining sinus tracts in her groin bilaterally with involvement of her right posterior thigh (fig. 1, HS of the right posterior thigh; fig. 2, HS of the left groin) but no axillary involvement. She had no prior history of HS or psoriasis. She was started on doxycycline 100 mg orally twice a day for 10 days and subsequently also with 10-day courses of-cephalexin followed by clindamycin with minimal improvement. She was started on suppressive therapy with doxycycline 100 mg orally twice a day for 2 months with some improvement. She was transitioned off the antibiotics but the disease recurred. She was treated with several additional 10-day courses of amoxicillin/clavulanic acid followed by clindamycin. She was placed back on suppressive therapy with doxycycline 100 mg orally twice a day for the next 3 months followed by minocycline for one month. Over the past 2 years, she received intralesional triamcinolone injections on five occasions with little improvement. Currently, she is being evaluated by plastic surgery for surgical excision.

Thirty months into treatment, the patient developed erythematous scaly plaques on her back (fig. 3) and at the site of adalimumab injections (fig. 4) and was diagnosed with psoriasis. Meanwhile, her CD was well controlled on adalimumab with mild disease confined to the rectum with no new fistula formation. It was believed that the adalimumab was causing the HS, and the drug was discontinued despite good control of her CD. The patient was switched to azathioprine.
Case 2

A 24 year-old African American male non-smoker presented to the emergency room with a 1-year history of intermittent bright red blood per rectum with no associated abdominal pain or tenesmus. A colonoscopy was performed showing erythema and friability from the rectum to the descending colon. The biopsies showed chronic colitis with granulomas consistent with Crohn’s colitis. He was started on oral Lialda 2.4 g daily, but the disease progressed with worsening abdominal pain, diarrhea and the development of an anal fistula. Therapy was changed to adalimumab for moderate-severe CD. Nine months into adalimumab therapy, he developed a right groin abscess with spontaneous drainage. Initially, it was thought that this was folliculitis, and he was treated with Bactrim for 2 weeks. He went on to develop multiple nodules in his groin bilaterally with the development of draining sinuses but no axillary lesions. He was diagnosed with HS; he had no personal or family history of HS. He was treated by dermatology with several 10- to 14-day courses of Bactrim and amoxicillin 875 mg/clavulanic acid 125 mg separately with transient improvement. He also had surgical excision of these areas and was started on chronic suppressive therapy with doxycycline 100 mg orally twice a day. On follow-up, his CD was not well controlled with worsening anal fistula, and azathioprine was added to adalimumab. HS also progressed to involve his axillary region.

Discussion

The association of CD with HS has been determined [2], and TNF antagonists to include infliximab have been shown to be helpful in treating patients with CD and HS [2, 7]. TNF plays an important role in both CD and HS [7]. CD is characterized by the predominance of increased T helper 1 (Th1) lymphocytes with Th17 cytokine expression [8]. TNF antagonists such as infliximab and adalimumab have been used to treat diseases such as CD, ulcerative colitis and psoriasis.

Recent studies have implicated anti-TNF antibodies as a cause of psoriasiform skin lesions in a small subset of patients with inflammatory bowel disease treated with these agents. Historically, this risk has been believed to be very small. However, several studies have indicated a higher risk than previously thought. In a recent study of 434 anti-TNF treated patients with inflammatory bowel disease, 4.8% developed psoriasiform skin lesions [9]. Smoking has been shown to be a risk factor for anti-TNF-induced skin lesions [9]. In the study by Tillack et al., 76% of the patients that developed psoriasiform skin lesions on anti-TNF were smokers or ex-smokers [9]. The skin lesions developed on average 13 months after initiation of the drug in multiple studies [6, 10], and most of the cases were confined to the palmoplantar and scalp areas [10]. In many cases, the drug had to be withdrawn due to the rash (43% in the study by Cullen et al. [11]). The psoriasiform rash is believed to be likely a class effect. Of the patients switched to an alternative anti-TNF in the review by Cullen et al. [11], 52% were unable to continue the drug due to recurrence of the rash, and only 15% were successful in controlling the rash by switching to an alternative anti-TNF.

In normal skin, TNF-α acts to inhibit the overproduction of interferon-γ [10]. It has been theorized that anti-TNF drugs allow unopposed interferon-γ production leading to activation of T helper cells producing IL-17 [10]. Increased concentrations of IL-17A and IL-23 have been shown in CD patients who developed skin lesions while on anti-TNF-α therapy [12]. It has also been shown that anti-TNF-induced psoriasiform skin lesions have infiltrates
of Th17 and Th1 cells, which are similar to the cytokine profile of psoriasis [9], and as noted below are similar to the profiles seen in HS. It is believed that the TNF blockade may cause the disruption of the cytokine balance in patients with psoriasiform skin lesions leading to unopposed interferon-α production by plasmacytoid dendritic cells in these genetically predisposed patients [13].

Studies have highlighted skin lesions such as psoriasiform lesions and eczema that can develop on anti-TNF therapy, but there is no mention of HS [6, 10, 12]. There was a recent case series of several patients that developed HS during treatment with adalimumab [14]. This is interesting, as adalimumab has been shown to be effective in treating HS in some small studies [4], though not all [15]. There was a range of several weeks to 24 months to the onset or exacerbation of HS after starting adalimumab in this series [14]. Two of the 4 patients in this series did have CD. The development of HS has also been described on etanercept [16]. Also, in a study looking at adalimumab for treatment of HS, 40% of the patients withdrew due to worsening of HS [15]. Due to very few reported cases of HS developing while on anti-TNF therapy, a conclusion as to whether this is a class effect of TNF antagonists cannot be determined. However, the occurrence of anti-TNF-induced psoriasiform lesions is believed to be a class effect [9], and this opens up the possibility that anti-TNF-induced HS might also be a class effect.

In the case series by Delobeau et al., 3 of the 4 patients were able to continue on the anti-TNF, while 1 patient required a switch to ustekinumab [14]. One of the 2 patients in our series had to be switched to another agent due to poorly controlled HS despite multiple therapeutic agents. This is only the second case in which we are aware of the development of HS associated with psoriasis [14]. Our series and the series by Delobeau et al. are unique in that the patients developed HS while already on adalimumab rather than being started as a potential treatment for HS.

Increased concentrations of TNF-α has been seen in the serum of patients with HS [17] and in HS skin [18]. Adalimumab treatment has been shown to cause a large reduction in the number of CD11c+ dendritic cells which are involved in HS pathophysiology [18]. Abundant expression of IL-12 and IL-23 was seen in the lesional skin of patients with HS [19]. IL-23 has an important role in Th17 development, and Th17 cells were seen to infiltrate the lesional skin in a study of HS patients [19].

The cytokine profiles of HS, psoriasis and anti-TNF induced psoriasiform lesions as noted above are very similar. This opens the possibility that like psoriasis, HS is a rare skin lesion that can develop while on anti-TNF therapy and is perhaps due to a cytokine dysregulation as has been seen with TNF antagonists and psoriasiform skin lesions. Ustekinumab is an anti-IL-12/IL-23 antibody, which has been used for anti-TNF-induced psoriasiform skin lesions [9] and has shown some benefit in CD [20]. This opens the way for a possible use of Ustekinumab to treat both CD and HS.

In conclusion, HS should be considered as a potential rare paradoxical side effect of anti-TNF therapy.

Statement of Ethics

The authors have no ethical conflicts to disclose.
Disclosure Statement

There are no conflicts of interest for either author.

References

Fig. 1. HS of the posterior thigh.

Fig. 2. HS of the left groin.
Fig. 3. Psoriasiform rash that developed on her back.

Fig. 4. Psoriasiform skin lesion at the adalimumab injection site.