A Recalcitrant Acrodermatitis Continua of Hallopeau Successfully Treated with Etanercept

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Acrodermatitis continua of Hallopeau (ACH) is considered as a localized form of pustular psoriasis that is usually refractory to the treatment. The condition is characterized by sterile pustules, onychodystrophy, anonychia, and osteolysis of distal phalanges. The authors report a case of recalcitrant ACH who previously failed to topical corticosteroid, methotrexate, acitretin, and phototherapy and rapidly responded to etanercept in combination with acitretin. The primary varicella infection occurred during the therapy. The patient was able to discontinue etanercept within four months after starting the treatment. The lesion was then under-controlled by acitretin and topical clobetasol ointment with an 8-month clinical remission.

Keywords: Acrodermatitis continua of Hallopeau, Pustular psoriasis, Etanercept, Varicella infection

Case Report

A 24-year-old patient was referred to the department of Dermatology with a 5-year history of persistent erythematous plaques with pustules on distal phalanges of all fingers, some toes and both palms (Fig. 1). All the fingernails and some of her toenails had totally shed, causing a limitation in fine movement. None of the lesions was detected at other parts of the body. She was previously healthy and denied tobacco smoking or alcohol consumption. She had no family history of psoriasis. Gram’s stain and potassium hydroxide (KOH) preparation from the pustules showed no organism. Radiological examination of the hands revealed periarticular osteoporosis of distal interphalangeal (DIP) joint of both hands. Laboratory investigation included complete blood count (CBC), blood chemistry (liver function test, blood urea nitrogen, creatinine), hepatitis B and C viruses, Anti-HIV, chest x-ray and tuberculin skin test were within normal limits or negative. Acrodermatitis continua of Hallopeau was diagnosed. She was treated with various treatment modalities including topical high potency steroids, 3 months of methotrexate (15 mg/week), 2 years of acitretin (0.7 mg/kg/day) and local PUVA (psoralen and ultraviolet A) without significant response. Because of the lacking of efficacy of previous treatments, the symptoms were getting worse. Eventually, the authors started etanercept 50 mg subcutaneous (SC) twice weekly combined with daily acitretin 0.5 mg/kg for a week, and then decreased etanercept to 25 mg SC twice weekly combined with daily acitretin 0.5 mg/kg.

After the second doses of 50 mg etanercept, the lesions and joint symptom were dramatically improved. After six doses of etanercept, the lesions were completely clear (Fig. 2). Simultaneously, the patient developed generalized papulovesicular lesions on the face, trunk, and arms without dermatomal distribution. She also developed coryza and low-grade fever. No mucosal lesion was detected. She had never had chickenpox before. Tzanck’s smear form the vesicle was positive for multinucleated giant cell. Primary varicella infection was diagnosed and oral acyclovir had been prescribed for a week. The authors decided to discontinue etanercept and the pustules on phalanges recurred. After one week, the varicella lesions turned to small crusts and in two weeks, the lesions had completely disappeared. After a 2-week discontinuation, the authors restarted etanercept 50 mg weekly combined with 1 mg/kg/day of acitretin without any problems. The disease responded well within 4 weeks. During the course of therapy, no new
lesion recurred. Hence, etanercept was discontinued after four months of therapy. The authors continued acitretin 0.7 mg/kg/day and topical clobetasol ointment for controlling her lesions. During the 8-month follow-up period, no disease recurrence and her nails started to grow back.

Discussion

Acrodermatitis continua of Hallopeau (ACH) is a localized form of pustular psoriasis that is usually refractory to treatment. It is more common in middle-aged women but can occur in any age group. Typically, lesions contain pustules that coalesce to form lakes of pus on a scaly erythematous base. The lesions start from distal part of digits and can spread proximally to hands and dorsum of forearms and feet. Lesions on nail bed and nail matrix generally lead to onychodystrophy and anonychia. The affected part can cause osteolysis of distal phalanges, as seen in x-ray. At present, management of ACH is a challenging problem. Systemic acitretin is considered as the first-line treatment. Other therapies that have been used include corticosteroids, methotrexate, phototherapy (psoralen plus ultraviolet A, ultraviolet B), calcipotriol, topical tacrolimus and thalidomide. However, these treatments have been tried with uncertain results.

Etanercept (Enbrel®) is a recombinant human receptor fusion protein that competitively inhibits the interaction of TNF-α with cell surface receptor. TNF-α is a pro-inflammatory cytokine involved in pathogenesis of ACH and other forms of psoriasis. Its concentration was detected in the joints and skin of psoriasis patients. In 2005, etanercept was initially reported successful in ACH treatment. Subsequently, many reports demonstrated the benefits of this drug in ACH, sometimes added to other therapies. Nevertheless, few cases of ACH had been reported failure to etanercept. In Table 1 the authors summarize ACH cases treated with etanercept during 2005-2010. Duration varied from three months to 30 months. Overall etanercept was effective in ACH except for the two studies. Etanercept was usually combined with other treatments such as methotrexate and acitretin. Other anti-TNF-α, including infliximab and adalimumab, were reported as successful for ACH. The adverse effect of anti-TNF-α is related to reactivation of many infections such as tuberculosis, fungi, and viruses due to immunomodulating properties. Varicella zoster virus (VZV) infection (primary varicella, recurrent varicella, herpes zoster infection) had been increasingly reported in patients who received anti-TNF-α. Recently there was a report of recurrent varicella infection in a psoriasis patient receiving etanercept for one month. It was proposed that anti-TNF-α agents might aggravate the VZV replication and VZV antigen expression. The present patient developed primary varicella infection during etanercept possibly due to the change in the immune status. However, she responded well to acyclovir treatment.

In conclusion, the authors demonstrated the recalcitrant ACH patient who had a dramatic response to etanercept combined with acitretin. The authors recommend etanercept should be considered as an option for recalcitrant ACH.

Potential conflicts of interest

None.

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Table 1. ACH cases treated with etanercept

<table>
<thead>
<tr>
<th>Authors</th>
<th>Yrs</th>
<th>Age/sex</th>
<th>Dose of etanercept</th>
<th>Combined treatment</th>
<th>Duration</th>
<th>Results</th>
</tr>
</thead>
<tbody>
<tr>
<td>Puig et al (9)</td>
<td>2010</td>
<td>78/F</td>
<td>50 mg twice weekly</td>
<td>Methotrexate 10-15 mg/week</td>
<td>30 months</td>
<td>Effective but relapsed after discontinuation, switch to adalimumab</td>
</tr>
<tr>
<td>Gokdemir et al (11)</td>
<td>2009</td>
<td>47/M</td>
<td>50 mg twice weekly</td>
<td>-</td>
<td>3 months</td>
<td>Ineffective</td>
</tr>
<tr>
<td>Thielen et al (8)</td>
<td>2008</td>
<td>64/M</td>
<td>25 mg twice weekly</td>
<td>-</td>
<td>9 months</td>
<td>Effective</td>
</tr>
<tr>
<td>Adisen et al (10)</td>
<td>2007</td>
<td>40/M</td>
<td>25-50 mg twice weekly</td>
<td>Topical steroid</td>
<td>3 months</td>
<td>Ineffective</td>
</tr>
<tr>
<td>Weisshaar and Diepgen (16)</td>
<td>2007</td>
<td>50/M</td>
<td>25 mg twice weekly</td>
<td>-</td>
<td>9 months</td>
<td>Effective</td>
</tr>
<tr>
<td>Ahmad and Rogers (6)</td>
<td>2007</td>
<td>Not stated</td>
<td>Not stated</td>
<td>Not stated</td>
<td>Not stated</td>
<td>Effective</td>
</tr>
<tr>
<td>Bonish et al (17)</td>
<td>2006</td>
<td>76/M</td>
<td>50 mg biweekly</td>
<td>-</td>
<td>Not stated</td>
<td>Effective</td>
</tr>
<tr>
<td>Nikkels and Pierard (18)</td>
<td>2006</td>
<td>74</td>
<td>50 mg twice weekly</td>
<td>Acitretin 50 mg/day</td>
<td>6 months</td>
<td>Effective</td>
</tr>
<tr>
<td>Kazinski et al (7)</td>
<td>2005</td>
<td>65/M</td>
<td>25-50 mg twice weekly</td>
<td>Acitretin 75 mg/day</td>
<td>3 months</td>
<td>Effective</td>
</tr>
<tr>
<td>This patient</td>
<td>2010</td>
<td>24/F</td>
<td>25-50 mg twice weekly</td>
<td>Acitretin 0.5-0.7 mg/kg/d</td>
<td>4 months</td>
<td>Effective</td>
</tr>
</tbody>
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ผู้ป่วยอโครเดอร์มาไตตีส คอนตินูอา ออฟ ฮาโลโพ ที่ดื้อต่อการรักษา 1 ราย ตอบสนองดีมากต่อการรักษาด้วยยาอะซิเทรตติน

นพมล ศิลปอาชา, ชนิษฎา วงษ์ประภารัตน์