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Case Series

Successful Treatment of Chronic *Staphylococcus aureus*-Related Dermatoses with the Topical Endolysin Staphefekt SA.100: A Report of 3 Cases

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Keywords

Staphefekt · *Staphylococcus aureus* · Endolysin · Folliculitis · Atopic dermatitis · Antibiotic resistance

Abstract

Staphylococcus aureus plays an important role in skin and soft tissue infections and contributes to the pathophysiology of complex skin disorders such as atopic dermatitis. Bacterial resistance against commonly used antibiotics has increased considerably in the last decades demanding alternative treatment approaches. We present 3 cases where patients with chronic and recurrent *S. aureus*-related dermatoses were successfully treated with Staphefekt SA.100. Staphefekt SA.100 is a recombinant phage endolysin for topical skin application that specifically targets both methicillin-sensitive and methicillin-resistant *S. aureus*. As a conse-

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quence of its specific mechanism of action, bacterial resistance is unlikely to develop. In our 3 cases, resistance induction was not observed. Our results indicate that targeted treatment with Staphfek might be an attractive alternative for (long-term) classical antibiotic therapy, and confirmatory randomized controlled trials are warranted to evaluate its clinical efficacy and safety.

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Introduction

Staphylococcus aureus is the most common microbe causing skin and soft tissue infections and contributes to the pathophysiology of complex skin disorders such as atopic dermatitis [1–3]. As antibiotic resistance against classical antibiotics is rapidly emerging, alternative treatment options are desperately needed [4–6]. Staphfek SA.100 is a recombinant phage endolysin for topical skin application that specifically binds to the cell wall of *S. aureus* and cleaves bonds in the peptidoglycan wall, resulting in bacterial cell death [7]. In vitro studies showed that Staphfek exclusively targets *S. aureus* without affecting other (commensal) skin bacteria. Induction of resistance was not observed and is also not expected due to its specific mechanism of action [7–9]. Staphfek is currently registered as a (class 1) medical device in Europe and is available in a cetomacrogol-based cream and in a gel as over-the-counter treatment. Evidence for its clinical efficacy is currently based on questionnaire studies and patient reviews only (Table 1) [8–10]. In this report we describe 3 cases with difficult-to-treat chronic or recurrent *S. aureus*-related dermatoses requiring antibiotic therapy that were successfully treated with Staphfek SA.100 in our clinic.

Case Presentation

Case 1

The first patient is a 23-year-old male with a history of osteosarcoma at the age of 16 and acne vulgaris of the face and trunk. He presented with an extensive papulopustular eruption on the trunk consistent with bacterial folliculitis. Consecutive empirical treatments with clarithromycin 500 mg once daily combined with Betadine scrub resulted in moderate improvement. However, after discontinuing treatment the symptoms recurred within 3 days. Taking into consideration the patient's reluctance to use continuous oral antibiotics and after discussing the alternative (on- and off-label) treatment options, our patient decided to try a short course of Staphfek. He applied the cream twice daily on the affected skin for 2 weeks. At his return visit he reported a strong decrease of inflammatory symptoms that started within a few days. This triggered us to further investigate the clinical effect of Staphfek using clinical photographs and bacterial cultures in this and 2 other cases. One week after he had stopped treatment with Staphfek, the papules and pustules recurred and were similar to those at the baseline (visit 1; Fig. 1a). Staphfek was restarted twice daily, again resulting in a clear decrease in the number of the papules and pustules observed 4 weeks later (visit 2). This improvement continued up to visit 3 after another 4 weeks (Fig. 1b). Nasal cultures showed persistent carriage of *S. aureus*. Skin cultures were only taken at visits 2

and 3 with a positive culture at visit 3. An extra skin culture was taken 2 months later (visit 4) to determine minimal inhibitory concentrations of Staphfek. The minimal inhibitory concentrations of the cultured *S. aureus* strain at visits 3 and 4 was stable, indicating the absence of resistance during long-term daily therapy.

Case 2

The second patient, a 63-year-old male, was diagnosed with a secondary bacterial infection (impetigo) of his nummular eczema, mainly located at the face and the extremities. The diagnosis was confirmed by histopathology and *S. aureus* cultured from a lesion. Treatment with clarithromycin 250 mg twice daily during 7 days, combined with topical fusidic acid cream, did not result in clinical improvement. Oral prednisolone 30 mg once daily for 2 weeks combined with flucloxacillin 500 mg 4 times daily during 7 days resulted in clinical improvement. However, 2 days after the treatment was stopped, the symptoms recurred (Fig. 2a). After discussing the alternative treatment options, Staphfek cream was started twice daily on the affected skin of the face and extremities, and the extremities were additionally treated with betamethasone ointment once daily. The patient reported improvement of the superinfection within 1 week. His symptoms did not recur over a treatment period of more than 12 weeks (Fig. 2b), and he did not report any adverse effects. Although the superinfection had cleared, the underlying eczema at the extremities remained. His eczema gradually improved after the treatment with betamethasone ointment was replaced with tacrolimus 0.1% ointment in tar 10% (solutio carbonis detergens) 1 month later. Cultures from the facial skin lesions and nose collected at baseline, and after 6 and 12 weeks of treatment were all positive for *S. aureus*.

Case 3

The third case is a 23-year-old man who presented with a bacterial folliculitis of the lower legs. *S. aureus* was cultured from a pustule but also from his nose and perineum. The patient was diagnosed with nummular eczema mainly at the extremities, which was active at the time of presentation. His medical history comprised biliary atresia followed by a liver transplant and chronic use of oral immunosuppressive medication. Class 2–3 topical corticosteroids, topical tacrolimus, and an emollient were prescribed to treat the eczema. The folliculitis was treated with topical fucidic acid cream, oral flucloxacillin for 1 week, and Beta-dine scrub, resulting in an improvement of the symptoms. An exacerbation 6 months later with follicular pustules at the lower legs was treated again with flucloxacillin but 2 months later the folliculitis recurred again, including extended perifollicular pustules that evolved to large confluent plaques at the lower legs (visit 1). After discussing the alternative treatment options, we started topical Staphfek twice daily. Three weeks later (visit 2) he reported a strong improvement of his symptoms starting 4 days after initiation of the treatment. Clinical examination at this time showed the remaining extensive eczematous lesions, but pustules were clearly absent and the redness of the lesions was significantly reduced. With continued treatment, further improvement of the eczema and absence of pustules were reported after 1 month. At visit 3, 2 months later, an increase of the eczema was reported. This was probably related to a lack of adherence since our patient used the corticosteroid cream, tacrolimus ointment, and emollient inconsistently. The legs showed dry eczematous lesions and

postinflammatory hyperpigmentation. However, pustules were still absent. Cultures of the skin taken at visits 1, 2, and 3 were all positive for *S. aureus*.

Discussion

This case series describes the successful treatment of *S. aureus*-related dermatoses with an antibacterial endolysin, which is currently available as an over-the-counter medical device. All 3 patients had recurrent *S. aureus* infections despite adequate antibiotic treatment. Atopic predisposition and nasal carriage of *S. aureus* probably contributed to the chronicity of their symptoms [2, 3, 11]. Despite the improvement of symptoms in these patients, *S. aureus* was still cultured from their skin during and after treatment with Staphfek. We believe that Staphfek induces a clinically relevant reduction of *S. aureus* rather than a total eradication. In addition, it is well-known that the skin lesions can get recolonized quickly due to constant exposure to *S. aureus* from the nose and environment. Apparently, complete and sustained eradication is not necessary to achieve clinically relevant improvement with an endolysin-based topical treatment. Since the results from our cultures are only qualitative in nature, future studies should further investigate the effect of Staphfek on the *S. aureus* load and composition of the microbiome, using quantitative methods and sequencing technologies.

Clinical symptoms quickly reappeared both after stopping antibiotics (cases 1 and 2) and Staphfek (case 1), likely due to rapid *S. aureus* recolonization to a clinically relevant state. The positive clinical effect after restarting *S. aureus* suppression with Staphfek in case 1 supports this hypothesis. Concurrent treatment of the nostrils combined with treatment of the complete skin surface could potentially further reduce recolonization.

It should be noted that both impetiginized eczema and folliculitis have a naturally fluctuating disease course, which complicates the interpretation of our data. However, all 3 patients visited our outpatient clinic frequently with recurring and continuous symptoms (despite adequate treatment) for long periods already before the start of Staphfek treatment (10, 3.5, and 13 months). This suggests that remission is likely due to the use of Staphfek rather than to chance.

Although limited by its uncontrolled design, this case series illustrates that topical therapy with Staphfek SA.100 to continuously suppress *S. aureus* can lead to the reduction and prevention of clinical symptoms in 3 cases of *S. aureus*-related skin conditions, folliculitis and impetiginized dermatitis. Importantly, Staphfek targets both methicillin-sensitive and methicillin-resistant *S. aureus* but does not interfere with the commensal skin microbes and is also not expected to induce bacterial resistance due to its specific mechanism of actions [8, 9]. It could therefore be considered as a feasible alternative for *S. aureus*-related superficial skin infections, since classical (long-term) antibiotic therapy is clearly associated with the induction of bacterial resistance and other adverse effects [4, 5]. The observed beneficial clinical effect of Staphfek in these cases is encouraging, and further study of its efficacy and safety is warranted in randomized controlled clinical trials.

Statement of Ethics

The patients described in this case report provided informed consent.

Disclosure Statement

The Department of Dermatology of the Erasmus MC University Medical Center Rotterdam received an unrestricted grant from Microcos Human Health, the Netherlands, to perform clinical studies with Staphhefekt SA.100. Microcos Human Health provided Staphhefekt SA.100 for the patients.

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Fig. 1. Folliculitis. **a** Baseline; extensive follicular papules and pustules. **b** After 8 weeks; few follicular papules, postinflammatory hyperpigmentation, and scarring.



Fig. 2. Nummular eczema with secondary infection. **a** Baseline; erythematous nummular lesions forming confluent plaques covered with crustae and swelling of the ear. **b** Six weeks later; very mild erythematous macula with minimal scaling.

Table 1. Overview of in vitro and clinical studies with Staphfek

Authors, year	Published in	Summary of findings
Herpers et al., 2014 [8]	Abstract R144 at the 24th European Congress of Clinical Microbiology and Infectious Diseases (ECCMID), Barcelona	Lysis of <i>S. aureus</i> by Staphfek is dose dependent, specific, and efficient, compared to control strains of coagulase-negative staphylococci (in vitro) MSSA and MRSA proved to be equally susceptible to the endolysin Induction of resistance against Staphfek could not be achieved in vitro
Herpers et al., 2014 [9]	Summary of presentation at EuroSciCon meeting, London	In 3 out of 7 patients with rosacea, <i>S. aureus</i> was cultured from a lesion; after the local application of Staphfek for 1 week, <i>S. aureus</i> was eradicated from the lesion in all 3 patients In 6 out of 8 patients with recurring dermatitis (3 constitutional eczema, 2 chronic contact dermatitis, and 1 perioral dermatitis), <i>S. aureus</i> was found in skin cultures before treatment; in 5 of 6 patients, symptoms decreased during treatment with Staphfek, and patients reported less need of corticosteroids
Totté et al., 2015 [10]	Abstract P1212 at 24th EADV Congress, Copenhagen	Patients with rosacea ($n = 112$) and acne ($n = 56$) reported an improvement of inflammation-related symptoms and quality of life (Skindex-29) after 6 weeks of topical treatment with Staphfek SA.100 in a prospective questionnaire study
Pasmans et al., 2016 [12]	Study protocol randomized controlled trial, registered at ClinicalTrials.gov, NCT02840955	Ongoing double-blinded randomized controlled trial aiming to assess the difference in the need for corticosteroid cotherapy between the Staphfek and a placebo group over a 12-week treatment period Secondary objectives include differences in clinical efficacy, quality of life, microbial composition, and safety and tolerability