Folliculitis Decalvans
Long-lasting Response to Combined Therapy with Fusidic Acid and Zinc

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In 3 patients, the diagnosis of folliculitis decalvans was based on clinical, histopathological and laboratory criteria. All patients responded to a combination therapy consisting of oral and topical fusidic acid and oral zinc sulphate. The follow-up period exceeded more than one year. Key words: Staphylococci; Antimicrobial chemotherapy.

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Folliculitis decalvans is defined as a rare but characteristic circumscribed chronically progressive purulent folliculitis causing follicular atrophy and subsequent hair loss (1, 2). Other terms used in the literature to describe the disease are “acne decalvante” and “folliculite epilante et destructrice de regions velues”. Skin manifestations which do not cause subjective symptoms, such as itching, affect in most cases both scalp and face, but involvement of axillae and the pubic region as well as spreading to the trunk are possible (1, 3). Numerous rounded or irregular bald atrophic areas (pseudopelade state), usually surrounded by inflammatory folliculitis or purulent milial abscesses, form the clinical picture. Larger plaque-like areas can form due to confluence.

The etiology is still unknown, although Staphylococcus aureus (S. aureus) seems to act as a co-factor, since in most cases cultivation has revealed the presence of this bacterium.

Folliculitis decalvans is known for its resistance to treatment – hence the unfavourable prognosis. Most patients suffering from the disease have been confronted with a wide variety of treatment protocols. Scribe, for example, reports on a patient who was successively treated with Soy-Dome Cleanser and Neosporin Ointment topically, superficial X-ray therapy and lincocin and Pentids orally (4). Only a few publications reporting successful treatment can be found in the literature. One of these is by Bogg (1) who described a positive effect of fusidic acid treatment, without giving information about the

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<th>Table 1. Findings in patients with folliculitis decalvans</th>
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<td><strong>Initials</strong></td>
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<td><strong>Blood sedimentation</strong></td>
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<td><strong>Complete blood count</strong></td>
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<td>(normal range: ≤2.0)</td>
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<td><strong>Antinuclear antibodies</strong></td>
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was reduced to one tablet daily. Treatment with fusidic acid and zinc sulphate was started simultaneously. 1.5% fusidic acid in a cream base (Fuzidine® Creme) was applied topically for the first 2 weeks, in addition.

Follow-up
Each patient was seen for follow-up after 10 days, after 4 weeks and subsequently at 3-month intervals.

Response to therapy
For all 3 patients, a clearing of purulent folliculitis eruptions was noted at the first follow-up on day 10, with areas of reddening still to be seen on the previously active sites of disease. Cultivation for S. aureus proved negative at that time. The size of the reddened areas was noticeably smaller in all 3 cases at the 4-week follow-up. During the next year, no active folliculitis has been observed in patients B. O. and M. F.

Patient H. P. had taken a lengthy holiday in the Sahara 2 months after starting treatment, during a disease-free interval. Unfortunately, the supply of zinc sulphate was running out. Within 2 weeks after ending zinc sulphate therapy, a recurrence of folliculitis decalvans was noticed. On returning home, S. aureus was again isolated from affected sites. By repeating the same treatment regimen, complete cure was achieved.

Treatment for all 3 patients consists at the moment of one tablet of zinc sulphate each day.

DISCUSSION
Diagnosis of folliculitis decalvans in the 3 patients was based on a combination of the following findings: the characteristic clinical picture of follicular and pustular lesions in circumscribed areas resulting subsequently in hair loss and atrophy (psuedopelade state); the histopathology findings, with suppurative folliculitis and perifolliculitis, numerous plasma cells during the acute stage of the disease, and the bacteriological culture results revealing the presence of S. aureus.

The efficacy of fusidic acid in the treatment of folliculitis decalvans has been reported previously (1, 2). Fusidic acid, isolated from a strain of Fusidicum coccineum (6), is a steroid-like antibiotic. It is known primarily as an anti-staphyloccocal drug, with practically all strains of S. aureus being sensitive, whether or not they are resistant to methicillin and related

CASE REPORTS
Between June 1988 and February 1989, 3 patients suffering from folliculitis decalvans could be assessed. Table I summarizes the important clinical, histopathologic and laboratory findings separately for each patient.

Treatment protocol
Each patient received a 3-week oral course of fusidic acid (Fuzidine®; 3×500 mg daily) and a 6-month course of zinc sulphate (Solvezink® Brausetablette; 2 × 200 mg zinc sulphate daily) after which the dose

Fig. 1. Folliculitis decalvans (Patient B. O.). Clinical picture before treatment; top of the head: central atrophy surrounded by a zone of erythema with dense folliculitis.

protocol used. Suter (2) also successfully treated folliculitis decalvans with oral and topical fusidic acid. Like Bogg, he failed to report for what period of time his patients were observed for follow-up after concluding the treatment. Recently, Brozena et al. (5) reported on a case of folliculitis decalvans in a patient who remained free of disease for more than one year after having received rifampicin in a dose of 600 mg/day for 10 weeks. Although all reports published so far concern single cases and only provide partial information about the period of remission following discontinuation of therapy, antibiotics, topically or orally used, seem to be the preferred drugs.

In the following, we present a treatment regimen based on oral and topical use of fusidic acid in combination with zinc sulphate that has proved successful on 3 patients who could be followed-up for more than one year.

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Fig. 2. Folliculitis decalvans (Patient B. O.). Clinical picture one year after treatment initiation; top of the head: central atrophy without signs of active purulent folliculitis and a very small reddened area.
penicillins (7). Currently, however, fusidic acid is being subjected to two clinical trials in AIDS patients in Europe, since a marked clinical improvement in a Danish AIDS patient was noted after treatment with the drug (8). Although the clinical efficacy of fusidic acid in the treatment of folliculitis decalvans is established and can be supported by the drug's antimicrobial profile, no information is given in the publications regarding what period of time patients had remained free of disease after ending fusidic acid treatment.

Maintenance of the clinical state of absence of inflammatory reactions in the patients presented here was, in our opinion, most likely due to continuous zinc sulphate therapy. This trace metal has proved useful against other chronic and inflammatory diseases such as chronic folliculitis (9), erosive pustular dermatitis of the scalp (10), and perifolliculitis capitis abscedens et suffodiens (Hoffman) (11). Zinc supplementation is the preferred treatment for acrodermatitis enteropathica in which there is a zinc deficiency (12). However, supplementation of zinc in diseases in which a zinc deficiency cannot be found, such as acne vulgaris, alopecia areata, and leg ulcers (13) is a controversial topic (14). The exact scientific basis for the efficacy of zinc is not known as yet, but it is possible that zinc has per se, in some way, an anti-inflammatory effect (15, 16) and can modulate the immune response (17). Since an influence of the immune system in vitro could also be shown for fusidic acid (18), the combination of these two drugs could result in the demonstrated long-lasting therapeutic success. Complete withdrawal of zinc sulphate has not been carried out so far, in view of the experience of the possible effects, as was seen when one patient had to stop involuntarily when the supply of the drug ran out whilst on holiday. To learn more about the possible involvement of these two drugs in chronic inflammatory skin diseases, further studies are needed.

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REFERENCES