Trichophyton tonsurans is an anthropophilic dermatophyte, with a worldwide distribution, although its prevalence varies considerably between different geographical regions. Whereas in North America infections due to this fungus are exceptionally common, on the European continent they appear relatively seldom. Although T. tonsurans is primarily associated with tinea capitis, it can also be the cause of tinea corporis and tinea unguium. The course of infection is usually only mildly symptomatic. We describe here two cases of urease-positive T. tonsurans infections with atypically extensive cutaneous lesions and severe inflammatory responses.

Key words: Trichophyton tonsurans; dermatophyte; tinea capitis; tinea corporis.

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Trichophyton tonsurans is an important anthropophilic dermatophyte that causes an endothrix type of hair invasion. The fungus is currently the leading aetiological agent of tinea capitis in North America (1, 2). In Europe, the frequency of infections due to T. tonsurans is considerably lower, albeit increasing (3–5). In Poland, the species accounts for less than 10% of all dermatophytoses reported each year (6). Although T. tonsurans is primarily associated with tinea capitis, it can also be the cause of tinea corporis and tinea unguium. Several cases of epidemic infection with T. tonsurans, presenting as tinea capitis and/or tinea corporis, have been reported recently in different countries among wrestling and judo participants (7–10). Tinea capitis caused by T. tonsurans may present with a variety of clinical findings, including seborrhoeic dermatitis-like scaling, “black-dot” alopecia (with fracture of the hair at the skin surface), and, very rarely, a kerion type of inflammatory tinea capitis (11). Cases of tinea corporis have been observed mainly among members of sporting clubs. Such cases presented with exfoliating erythematous lesions, resembling nummular eczema (7).

We describe here two cases of tinea capitis and tinea corporis, both caused by T. tonsurans, and both with unusual clinical presentations.

CASE REPORTS

Case 1

A 5-year-old girl was admitted to the Wrocław outpatient dermatology department because of exacerbated cutaneous lesions. She had no prior history of any dermatological disease or condition. There was no history of fungal infections in other family members or children being in contact with the girl. Within a short time a severe kerion had developed in the temporoparietal area of the child’s scalp. The condition was characterized by an intense oedema of the skin, erythema, hair loss, and superficial pustules. After approximately one month, acute infiltrative inflammatory lesions could be observed on both cheeks, involving scaling and pustule formation (Fig. 1). In the following month the lesions spread to the chest, abdomen, and extremities, leading to disseminated erythematous papular eruptions. The cutaneous lesions were not associated with any general symptoms. The body temperature remained normal, as did the results of the laboratory examinations of blood and urine samples. A bacteriological culture from pustule content was negative. Skin scrapings collected from the borderline of all lesions (located on the scalp, cheeks, chest, abdomen, and extremities) and a hair plucked at the periphery of the hair loss area were processed for mycological study. Direct

Fig. 1. Case 1. Acute inflammation with pustules on the face.
microscopic examination of the scales from the scalp and cheek lesions revealed the presence of hyphae, while in the hair an endothrix invasion was evidenced. Cultures on Sabouraud’s agar yielded white and powdery colonies. Microscopy evidenced the presence of micro- and macro-conidia corresponding to *T. tonsurans*; however, spiral hyphae unusual for this species were also observed (Fig. 2). Furthermore, an enhanced growth on thiamine-containing agar, a positive urease activity test, and a negative hair perforation test matched the diagnostic criteria for *T. tonsurans*. The identification of the species performed by conventional mycological examination was confirmed by means of genotypic analysis. PCR analysis for dermatophytes was performed with a HotStartTaq Plus Master Kit (Quiagen, Quiagen GmbH, Hilden, Germany) according to the manufacturer’s instructions. The following primers were used: forward “panDerm1” 5’ GAA GAA GAT TGT CGT TTG CAT CGT CTC 3’, reversed “panDerm2” 5’ CTC GAG GTC AAA AGC ACG CCA GAG 3’ (12). The weight of amplified product was 366 bp. Positive controls (*T. tonsurans* (366 bp), *T. mentagrophytes* (366 bp)) and a negative control were included (Fig. 3). Furthermore, an additional PCR was performed with primers specific for *T. tonsurans*: forward “tonsF1” 5’ CGG AGG CCG GCC CCCCAG 3’, reversed “tonsR1” 5’ CTC GAG GTC AAA AGC ACG CCA GAG 3’ (13). Its amplified product weight was identical to that of a positive control with *T. tonsurans* (173 bp) (Fig. 4).

Samples collected from lesions located on the trunk and extremities were negative for fungal infection.

**Treatment** with oral terbinafine (125 mg/day) for 8 weeks resulted in complete resolution of the lesions.

**Case 2**

The second case was a 23-year-old male with no history of chronic illness or medication. The patient had been a member of a wrestling club for 2 years, but had not exercised during the last 3 years. The initial skin lesions (two discrete erythematous patches) were localized in the right temporal area of the scalp. Without any mycological diagnostics, topical antifungal treatment had been given to the patient, resulting in a substantial improvement in the condition. One month later, however, the patient relapsed with several skin eruptions that first occurred in the temporal region as well as on the forearms, and then spread to the trunk and extremities. The patient was misdiagnosed as having an allergic reaction and was treated with corticosteroids administered topically and systemically. After 7 months from the onset of the disease he presented with numerous extensive erythematous and slightly scaling lesions located on the trunk and limbs. The lesions consisted of concentric erythematous rings forming a grain-like pattern. Scaling of the scalp and a black-dot alopecia were also observed. Apart from the skin lesions, the general condition of the patient was good. The results of blood and urine tests were normal.

Scrapings from the lesions of the scalp and trunk showed hyphae in KOH-mounts and a dermatophyte was grown in the cultures. The same mycological and genetic tests were performed as in case 1, and *T. tonsurans* was identified. For treatment, oral terbinafine at a dose of 250 mg/day and topical treatment with clotrimazole was administered. A complete cure was achieved within 4 weeks.

**DISCUSSION**

*T. tonsurans* is a ubiquitous fungus. It is particularly common in the USA being responsible for as much as 21.1–44.9% of all dermatophyte infections (1, 2). Of all European countries only in the UK and Croatia the infections are not very rare (14–16). *T. tonsurans* predominantly causes tinea capitis. In the UK and USA 50–90% (15) and 95.8% (17) of tinea capitis were infections with *T. tonsurans*, while in continental Europe, only 3–5% (18, 19).
Cases of atypical courses of *T. tonsurans* infections have been reported: tinea corporis with an acute inflammatory reaction (20), folliculitis decalvans (21, 22), and Majocchi granulomas (23–25).

We report two cases of infection with *T. tonsurans* with severe inflammatory courses.

In the first case, a 5-year-old child, sterile papules scattered over the trunk and limbs were observed in addition to the tinea lesions on the scalp and face. Such a disseminated and intensely inflammatory eruption suggests a dermatophytid reaction. Interestingly the causative *T. tonsurans* strain showed an unusual microscopic morphology with spiral hyphae. In addition, in both cases *T. tonsurans* strains were positive in a urease test, while most of the fungi of this species are urease negative. It is possible that higher than usual enzymatic activity of *T. tonsurans* strains was the cause of the intensive inflammatory reaction.

Our second patient with an acute inflammatory skin reaction in the course of *T. tonsurans* infection, had been a wrestler in the past. The first skin symptoms appeared 2 years after quitting sport. In the literature some wrestlers have been described as asymptomatic carriers (9), in addition to *T. tonsurans* being cultured from wrestling mats (9, 26). Epidemic outbreaks of *T. tonsurans* infections among members of combat sports clubs have been reported (7–9) as well as in other populations, such as paediatric healthcare workers (27).

This report emphasizes that *T. tonsurans*, a dermatophyte that is still relatively uncommon in Europe, can give rise to infections with quite diverse clinical features, ranging from the carrier-state to severe tinea corporis with heavy inflammation.

REFERENCES