A healthy 24-year-old woman presented with a recurrent, raised, red rash consisting of targetoid lesions, one to two centimetres in diameter, on the backs of both hands (Figure 1A). The patient reported that the plaques became blisters (Figure 1B) and faded to circular violet patches, healing within one week with residual hyperpigmentation. The patient reported nine similar episodes in the same location, and two such episodes on her face. A typical episode began with a burning sensation, followed by itching and headache lasting 30 minutes to 24 hours. The patient thought that the eruptions might be related to the premenstrual phase of her cycle. She had no history of infection with the herpes simplex virus.

Our differential diagnosis included fixed drug eruption and erythema multiforme. Although fixed drug eruption is primarily a clinical diagnosis, we biopsied a bullous lesion and asked the patient to construct a medication diary. Histopathology of the biopsied tissue was consistent with fixed drug eruption. The patient’s diary showed that the eruptions might be related to the premenstrual phase of her cycle. She had no history of infection with the herpes simplex virus.

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Fixed drug eruption accounts for about 11%–30% of all adverse drug reactions.1 Our patient had a classic fixed drug eruption. The hallmark of this reaction is the occurrence of eruptions in the same location during each episode, with additional lesions appearing upon drug rechallenge. Typical locations for the lesions include the genitals, face, hands and feet. Although fixed drug eruption is frequently confused with erythema multiforme, the histopathologies are different. Common causes of fixed drug eruption include sulphonamide drugs, tetracycline and nonsteroidal anti-inflammatory drugs. Fluconazole is an uncommon cause, with only 18 published cases.1,2 Cross-reactions may occur with structurally related agents, such as itraconazole.3

Fluconazole is a commonly used antifungal agent, and fixed drug eruption should be considered a possible adverse effect of its use. Our patient’s case highlights the importance of taking a comprehensive history of medication use when examining cutaneous adverse reactions.

References