A Case of Acute Telogen Effluvium After SARS-CoV-2 Infection

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Abstract: As the number of COVID-19 cases increasing, more and more patients are concerning about alopecia, a sequela after SARS-CoV-2 infection. We here report a case of a 38-year-old woman with a typical acute telogen effluvium (ATE) after recovery from COVID-19.

Keywords: COVID-19, SARS-CoV-2, acute telogen effluvium, alopecia

Case Presentation

On January 29, 2020, a woman aged 38 years old presented fever and cough but she did not visit a doctor. When she visited Beijing Haidian Hospital on February 8, she was diagnosed with COVID-19 and admitted to the hospital. On February 21, she recovered and was discharged from the hospital. Up to April 30, she visited our department with complaint about severe hair loss in the past week (>150 hairs/day), as well as oily scalp and trichodynia. Dermatology examination revealed diffuse hair loss involving the entire scalp. There was no obvious patchy hair loss area, and the forehead hairline was not significantly receded (Figure 1A). The pull test was positive in the whole head. Dermoscopy showed scalp inflammation, capillarectasia, dandruff, increased density of telogen hairs, and relatively uniform diameter of the terminal hair shaft. There was no broken hair, black dots, exclamation point hairs, etc (Figure 2A). The pigmentation at the proximal end of hair root was reduced, and the end of hair root was clubbed (Figure 3). Blood tests excluded vitamin B12, trace elements deficiencies, autoimmunity, and thyroid dysfunction. Therefore, the patient was diagnosed as acute telogen effluvium. Topical 5% minoxidil, mixed with halcinonide solution, was sprayed evenly on the hair loss area 1–2 times a day for treatment. Meanwhile, selenium sulfide lotion mixed with shampoo 2–3 mL was used to wash hair 2–3 times a week. After 3 months of treatment, the patient felt hair loss was significantly alleviated; symptoms of oily scalp and trichodynia disappeared. The hair pull test turned negative, and many new hairs grew out (Figures 1B and 2B). The study protocol was approved by the Medical Ethical Committee of China-Japan Friendship Hospital, and written informed consents were obtained from the patient to publish the case details.

Discussion

Acute telogen effluvium (ATE) is clinically manifested as diffuse alopecia of the whole head with a sudden increase in hair loss. The clinical manifestation of this “the pain in the hair” are named trichodynia. The onset and progression of hair loss

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Figure 1 Head hair at initial diagnosis (A) and 3 months later (B).

Figure 2 Trichoscopic graph at initial diagnosis (A) and 3 months later (B).

Figure 3 Trichoscopy presentation of hair roots.

is rapid, usually with positive result of hair pull test. ATE usually lasts less than 6 months.

The typical trichoscopic features of ATE include some hair follicles without hair shaft, increased density of telogen hairs, and relatively uniform diameter of the terminal hair

shaft. The proximal end of loosen hairs' roots are clubbed, and the pigmentation of them are reduced.¹ In this case, the patient's trichoscopy showed scalp inflammation, capillar ectasia, dandruff, many telogen hairs and relatively uniform diameter of the terminal hair shaft. The patient's hair loss was diffuse and did not restricted to the top of the head. Thus, this case is distinguished from diffuse alopecia areata and female-pattern androgenetic alopecia.

First reported by Kligman² in 1961, ATE is a non-cicatricial alopecia disease characterized by the simultaneous loss of many telogen hairs. Based on the duration of the disease, it can be divided into acute telogen effluvium (ATE) and chronic telogen effluvium (CTE). ATE usually occurs 2 to 3 months after a trigger event, which includes postpartum, discontinuation of birth control pills, hyperthermia, malnutrition, bulimia, blood loss, shock, surgery, and severe psychological factors, etc. Previous studies had observed telogen effluvium (TE) occurred after dengue virus and rickettsia conorii infection.^{3,4} Recent studies also showed that some patients developed hair loss after SARS-CoV-2 infection. In an investigation of clinical sequelae of COVID-19 survivors in Wuhan, China,⁵ the prevalence of alopecia as a sequela was as high as 48.5%. However, the exact mechanism between COVID-19 and TE is unclear. A prospective study⁶ enrolled 214 patients with ATE from March 2020 to August 2020, of whom 89.7% (191 patients) had a confirmed diagnosis of prior SARS-CoV-2 infection. The case we reported here suggests that SARS-CoV-2 infection may also be one of the causes of ATE.

Hoffmann et al⁷ demonstrated that SARS-CoV-2 used the SARS-CoV receptor ACE2 for entry and the serine protease TMPRSS2 for S protein priming. Therefore, any cell that carries the ACE2 receptor is a potential target for SARS-CoV-2, including the skin. A recent study⁸ has detected SARS-CoV-2 in the sweat glands and sweat ducts of COVID-19 patients. To further clarify the specific mechanism of TE following COVID-19 infection, additional testing of hair follicle tissue is required.

What makes this TE case different from the previous ones is that in the era of COVID-19 pandemic, TE has become a common sequela, which seriously affects the psychosocial condition of patients recovered from COVID-19. Also, the COVID-19 epidemic has an impact on people's social life, and some are more prone to anxiety and depression, from which TE may occur. Turkmen et al⁹ investigated the effect of COVID-19 pandemic on hair disorders through online questionnaire, which showed that 27.9% of participants experienced TE during the pandemic. A retrospective study

in Turkey¹⁰ showed that the percentage of TE in dermatology clinics increased 5.51 times during the pandemic by comparing with the same season of the previous year. Therefore, we need to link TE with this emerging and highly impactful disease and further investigate the specific mechanisms by which COVID-19 may cause TE.

Consent

The patient in this manuscript has given written informed consent to the publication of her case details.

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Disclosure

The authors report no conflicts of interest in this work.

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