

Cutaneous Aspergillosis As a First Manifestation of Systemic Infection in Patient After Kidney Transplantation

Geun-Hwi Park^{1,3}, Kihyuk Shin^{1,2}, Hoon-Soo Kim^{1,3}, Hyun-Chang Ko^{1,2}, Byung-Soo Kim^{1,3}, Moon-Bum Kim^{1,3}, Dae-Lyong Ha^{1,3,4}

¹Department of Dermatology, School of Medicine, Pusan National University, Busan, ²Department of Dermatology, Pusan National University Yangsan Hospital, Yangsan, ³Biomedical Research Institute, Pusan National University Hospital, Busan, ⁴Department of Dermatology, School of Medicine, Kyungpook National University, Daegu, Korea

Dear Editor:

Aspergillosis is a significant cause of morbidity and mortality after organ transplantation. Cutaneous aspergillosis reportedly develops primary or secondary to hematogenous dissemination. Secondary *Aspergillus* infection resulting from hematogenous spread is extremely rare (<5%)¹. Moreover, there are no reports of cutaneous lesions as the first sign of systemic aspergillosis. Herein, we present the case of a patient with cutaneous

manifestations as the first sign of systemic aspergillosis.

A 70-year-old Korean male visited the dermatologic department for evaluation of a purple-colored nodular cutaneous lesion on the knee that appeared 2 months after renal transplantation. Histopathological analysis showed massive dermal neutrophilic and granulomatous inflammation. Moreover, multiple fungal hyphae were observed following periodic acid-Schiff with diastase (D-PAS) staining, and the patient

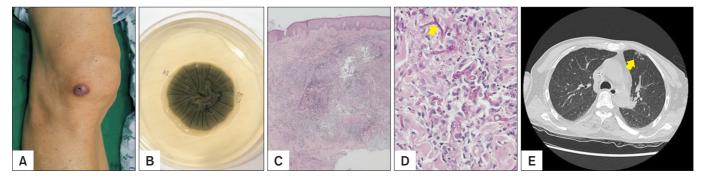


Fig. 1. (A) Purple-colored nodule on the right knee. (B) Spreading yellow-green colony. (C) Skin biopsy revealed dermal neutrophilic and granulomatous infiltration (H&E, original magnification $\times 20$). (D) Numerous septate hyphae with dichotomous branching are visible at 45° angle (D-PAS staining, original magnification $\times 200$; yellow arrow). (E) Fungal balls were observed on chest computed tomography (yellow arrow).

Received September 2, 2020 Revised November 24, 2020 Accepted December 5, 2020

Corresponding Author

Dae-Lyong Ha

Department of Dermatology, School of Medicine, Kyungpook National University, 130 Dongdeok-ro, Jung-gu, Daegu 41944, Korea Tel: +82-53-200-5838, Fax: +82-53-426-0770, E-mail: dhwl222@naver.com https://orcid.org/0000-0002-2268-4795

was diagnosed with cutaneous fungal infection (Fig. 1). Unexpectedly, before his scheduled visit with the dermatologist, the patient visited the emergency department with left-sided weakness and dysarthria. Magnetic resonance imaging of the brain revealed an abscess suspected to have a fungal origin. The patient was diagnosed with systemic aspergillosis based on the presence of a fungal ball observed on chest computed tomography and a positive result in the broncho-alveolar lavage fluid galactomannan test. Although the patient received systemic antifungal therapy, such as amphotericin B, and voriconazole, he died of systemic mycoses 6 weeks after diagnosis.

Cutaneous, subcutaneous, and systemic fungal infections are responsible for significant morbidity and mortality, particularly in immunocompromised patients². *Aspergillus* sp. infection occurs in 1%~15% of organ transplant patients and has a mortality rate of 74%~92%³. Risk factors for dissemination after initial infection include long-term use of immunosuppressants and transplant organ dysfunction. In this case, the patient took high doses of immunosuppressants—prednisolone, cyclosporin, and azathioprine—before and after kidney transplantation; the use of immunosuppressants is considered a major predisposing factor for the spread of *Aspergillus* into the skin^{4,5}.

Early diagnosis can greatly reduce mortality due to systemic aspergillosis by provision of prompt treatment; however, the condition is diagnosed after considerable disease progression in most patients. This makes treatment difficult and worsens the clinical course, leading to death, as in the present case.

The authors report a case of cutaneous aspergillosis in the form of a cutaneous nodule in a patient using immunosuppressive agents after kidney transplantation. Such cutaneous lesions in immunosuppressed patients may be an early symptom of opportunistic infection. We received the patient's consent form about publishing all photographic materials.

CONFLICTS OF INTEREST

The authors have nothing to disclose.

FUNDING SOURCE

None.

ORCID

Geun-Hwi Park, https://orcid.org/0000-0001-8371-2676
Kihyuk Shin, https://orcid.org/0000-0001-8955-9828
Hoon-Soo Kim, https://orcid.org/0000-0002-7649-1446
Hyun-Chang Ko, https://orcid.org/0000-0002-3459-5474
Byung-Soo Kim, https://orcid.org/0000-0003-0054-8570
Moon-Bum Kim, https://orcid.org/0000-0003-4837-0214
Dae-Lyong Ha, https://orcid.org/0000-0002-2268-4795

REFERENCES

- Bonduel M, Santos P, Turienzo CF, Chantada G, Paganini H. Atypical skin lesions caused by Curvularia sp. and Pseudallescheria boydii in two patients after allogeneic bone marrow transplantation. Bone Marrow Transplant 2001;27:1311-1313.
- Tessari G, Naldi L, Piaserico S, Boschiero L, Nacchia F, Forni A, et al. Incidence and clinical predictors of primary opportunistic deep cutaneous mycoses in solid organ transplant recipients: a multicenter cohort study. Clin Transplant 2010;24:328-333.
- Einollahi B, Lessan-Pezeshki M, Pourfarziani V, Nemati E, Nafar M, Pour-Reza-Gholi F, et al. Invasive fungal infections following renal transplantation: a review of 2410 recipients. Ann Transplant 2008;13:55-58.
- 4. Shoham S, Marr KA. Invasive fungal infections in solid organ transplant recipients. Future Microbiol 2012;7:639-655.
- Tsai WC, Lee CH, Wu WM, Lin SH, Yang YC, Cheng YW, et al. Cutaneous manifestations of subcutaneous and systemic fungal infections in tropical regions: a retrospective study from a referral center in southern Taiwan. Int J Dermatol 2017;56:623-629.