Primary Cutaneous Aspergillosis in an Immunocompetent Patient: A Case Report from a Tertiary Care Hospital in Chennai

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Abstract
Aspergillosis is a systemic fungal infection that commonly affects immunocompromised individuals and, less frequently, immunocompetent individuals. It is the most common opportunistic fungal disease after candidiasis. This is primarily a pulmonary infection and can also involve other body sites like paranasal sinuses and cutaneous tissues. Aspergillus fumigatus, Aspergillus niger, and Aspergillus flavus are the common species infecting humans. Primary cutaneous aspergillosis (PCA) is usually caused by A. flavus and A. fumigatus. It is commonly seen in immunocompromised patients such as those suffering from diabetes, malignancies, tuberculosis, human immunodeficiency virus, or patients on long-term steroids and antibiotics. In this article, we report a case of PCA, in the immediate postoperative period, following a road traffic accident, in an immunocompetent patient. This posed a diagnostic challenge to the treating physicians. A. flavus was confirmed with 10% potassium hydroxide mount, lactophenol cotton blue, and growth on Sabouraud dextrose agar from tissue culture sample. Antifungal treatment was initiated with oral itraconazole 200 mg after performing antifungal susceptibility testing based on Clinical and Laboratory Standards Institute guidelines. The patient’s condition improved and was discharged. Thus, early detection of PCA combined with medical and surgical intervention can successfully eradicate infection and help in preventing disseminated aspergillosis.

Keywords
► Aspergillus flavus
► immunocompetent
► primary cutaneous aspergillosis
► systemic fungal infection

Introduction
Aspergillosis is a systemic fungal infection that commonly affects immunocompromised individuals and less frequently immunocompetent individuals. It is the commonest opportunistic fungal disease after Candidiasis.¹ This is primarily a pulmonary infection and can also involve other body sites, like para nasal sinuses and cutaneous tissues. It is caused by...
several species of genus *Aspergillus*, which are predominantly found in the environment, in decaying organic matter.\(^2\) *Aspergillus fumigatus, Aspergillus niger*, and *Aspergillus flavus* are the common species infecting humans. Primary cutaneous aspergillosis (PCA) is usually caused by *A. flavus* and *A. fumigatus*.\(^3\) It is commonly seen in patients with compromised immune system such as those suffering from diabetes, chronic diseases like tuberculosis, those on long-term steroids and antibiotics, patients with malignancies on immunosuppressive therapy, and human immunodeficiency virus. Organ transplant and autoimmune disease also increase the risk.\(^4\) Patients of PCA can present with molluscum like papules, nodules, ulcers, and plaques.\(^1\)

In this article, a case of PCA, in the immediate postoperative period, following a road traffic accident, in an immunocompetent patient is reported. This posed a diagnostic challenge to the treating physicians.

**Case Report**

A 50-year-old female patient presented with multiple phalangeal fracture and second metatarsal undisplaced fracture of the left foot caused by a motorcycle accident (►Fig. 1). She had no other comorbidities. Her blood sugar levels were normal, was normotensive, and was not on steroids. Her renal parameters were normal and she did not have any previous history of antifungal use. All other blood investigations performed were within the normal range. Fracture treatment included Kirschner wire fixation (►Fig. 2). Repeated surgical debridement of the wound was also done since it had large tissue defects (►Figs. 3 and 4). No samples were sent for culture at this point of time. There was no intensive care unit admission during this period. The patient was discharged with the advice to report regularly for wound dressing on alternate days, for which the patient did not comply. A week later, the patient reported to the outpatient department with pain and on examination, the wound looked unhealthy. She was, therefore, admitted for wound debridement and regular saline dressing. Her wound was dressed regularly and tissue bit was sent for culture in view of presence of necrotic tissue suggestive of infection. Ten percent potassium hydroxide (KOH) mount culture and lactophenol cotton blue (LPCB) mount were performed. KOH mount showed hyaline septate hyphae. Light green velvety colonies of *A. flavus* was isolated in culture (►Figs. 5, 6, 7, 8, 9). LPCB mount demonstrated conidiophores of variable length, rough, pitted and spiny along with biseriate phialides covering the entire vesicle and pointing out in all directions (►Figs. 10, 11, 12).

Additional two tissue bit samples were obtained on consecutive days. This helped to distinguish between colonization and infection. KOH mount was positive for septate hyphae and the growth was similar to previously reported cultures. Antifungal susceptibility testing was performed.
according to Clinical and Laboratory Standards Institute guidelines (►Fig. 13). It showed resistance to fluconazole, ketoconazole, amphotericin-B, and nystatin and was sensitive to clotrimazole, voriconazole, and itraconazole. Infectious necrotic tissue areas persisted despite surgical debridement. Further, *A. flavus* growth was observed in all three consecutive cultures. Hence, antifungal treatment was initiated and patient was started on oral itraconazole 200 mg twice daily for 10 days.

There was no other factor contributing to immunocompromised status of the patient. There were no findings suggestive of onychomycosis. No other bacterial organism was isolated from the samples sent. All three cultures yielded *A. flavus* proving it to be a true pathogen.

Regular saline dressing on a daily basis and surgical debridement were continued. Tissue bit cultures sent after the end of antifungal therapy yielded no growth. Saline dressing was continued till the wound healed (►Fig. 14). Patient's condition improved. She was then discharged on oral analgesics (SOS) alone. She was advised to come regularly on a weekly basis for change of dressing.

### Discussion

*A. flavus*, an important causative agent of invasive aspergillosis, can also commonly cause superficial infections, even of those affecting the skin and oral mucosa.\(^5\) *A. flavus* grows optimally at 37°C. Growth is also observed at temperatures ranging from 12 to 48°C.\(^5\) This is an important reason contributing to its pathogenicity in humans.

The cutaneous infections are generally classified as primary and secondary. Primary infection can be triggered by several causes including direct instillation, like traumatic inoculation, intravenous catheters, occlusive dressings, burns, and surgical procedures. Secondary infection happens mostly from hematogenous spread.\(^7\)

In our case, *A. flavus* growth was seen in an immunocompetent patient following a road traffic accident. All other parameters were normal, the only source of infection could be traumatic inoculation. The patient reported to the emergency department initially with acute trauma, following which she was operated and managed by the orthopaedic team. No samples were sent for culture at this point of time, as per the protocol, keeping in mind the early surgical treatment needed for the patient. After surgery and discharge, patient was asked to come for regular dressings on alternate days. But the patient reported only after a week, with unhealthy looking superficial wound and complaints of pain. The wound being superficial rules out the possibility of an iatrogenic inoculation. The possible ways by which healthy hosts can develop cutaneous aspergillosis are traumatic inoculation, surgical wound operations, or exposure to high levels of sporulating organisms in occupations such as farming. This patient had no history of farming activities. A
Fig. 3  Preoperative image showing gross injury and tissue damage with necrosis.

Fig. 4  Preoperative picture showing extensive tissue damage and necrosis of foot.

Fig. 5  Initial moldy growth observed on chocolate agar.

Fig. 6  Initial mouldy growth observed on blood agar.

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similar growth of *Aspergillus* spps. has been noted in immunocompetent patients in the studies of Tahir et al, Camus et al, Avkan-Oğuz et al, and Neki et al.²⁻¹⁰

The treatment of PCA in immunocompromised patients is well understood but the management of PCA in immunocompetent patients is less comprehensible and poses a challenge to the treating physicians.¹¹ In our case, the patient
Fig. 10 LPCB mount showing conidia, vesicles, conidiophores, and septate hyphae suggestive of *Aspergillus flavus*, under low power magnification.

Fig. 11 LPCB mount showing conidia, vesicles, and conidiophores of *Aspergillus flavus*, under high power magnification, taken from sample sent for tissue bit culture.

Fig. 12 LPCB mount showing conidia, vesicles, and conidiophores of *Aspergillus flavus*, under high power magnification, taken from sample sent for repeat tissue bit culture.

Fig. 13 Antifungal susceptibility test performed, showing resistance to multiple drugs, including Amphotericin-B and Nystatin.

Fig. 14 Postoperative picture showing healed areas with improved tissue function, taken after surgical debridement and antifungal therapy.
responded very well to oral antifungal therapy with itraconazole. She showed no adverse side effects and improved well. Amphotericin B is the drug of choice for disseminated aspergillosis. Oral itraconazole, a triazole antifungal with activity against *Aspergillus species*, is an alternative to amphotericin B for all types of aspergillosis. Voriconazole is given for the most advanced cases.\(^\text{12}\)

Thus, there has been limited case reports on aspergillosis reported in immunocompetent patients. The accurate identification and appropriate treatment give better wound healing and prognosis. The treatment of cutaneous aspergillosis causes no harm to healthcare workers who are immunocompetent.\(^\text{13}\) Early detection, combined with medical and surgical intervention, can successfully eradicate infection. This will help in treating the locally destructive disease and thereby prevents disseminated aspergillosis.\(^\text{14}\)

**Conclusion**

To conclude, the most beneficial management plan would be to make decisions based on individual history and clinical findings. Appropriate antifungal treatment combined with regular dressing and repeated surgical debridement till culture negativity should be followed.\(^\text{15}\)

**Authors’ Contribution**

SR, AB, and PS collected and analyzed the data. SR and AB prepared the manuscript and it was reviewed and modifications were made by PS. SR participated in the manuscript revision under the guidance of AB and PS. All authors of the manuscript gave their complete approval.

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**Conflict of Interest**

None declared.

**References**