

A Case of Giant Basal Cell Carcinoma of the Ear Complicated by Primary Cutaneous Aspergillosis

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Abstract: A 78-year-old female patient with right ear agenesis presented with a skin manifestation of approximately 7 cm × 8 cm deep-invasive ulcer with well-defined borders and a small amount of yellow purulent discharge visible at the base, surrounded by pearl-like margins in a dyke-like elevation, covered with a small amount of necrotic tissue and black crust. The disease lasted for more than 20 years and was diagnosed as giant basal cell carcinoma complicated by primary cutaneous aspergillosis after two histopathological examinations of the skin lesions. There are similarities in the clinical manifestations of these two diseases, which need to be differentiated, and the simultaneous complications are infrequent. It has not been reported.

Keywords: giant basal cell carcinoma, primary, cutaneous aspergillosis

Case Report

A 78-year-old woman presented to the dermatology department with a skin ulcer with pain around her right ear for more than 20 years. The patient developed a nodule the size of a pea-sized behind her right ear with no apparent cause 20 years ago, accompanied by itching. Since then, the lesion gradually expanded and formed an ulcer with painful itching. She was previously in good health with no underlying disease and no similar family history.

On physical examination, a deep ulcer of about 7cm × 8cm with clear borders is visible in the right ear, a small amount of yellow purulent discharge is visible at the base, surrounded by pearl-like edges in a dyke-like elevation, covered with a small amount of necrotic tissue and black scabs, and the right ear is absent (Figure 1).

Blood tests showed haemoglobin 108 g/L, red cell count $3.65 \times 10^{12}/L$, C-reactive protein 11.83 mg/L, and Immunoglobulin E 1660 IU/mL. Mycobacteria were detected in stool 1–2. Bacterial culture of skin lesion secretions isolated *Klebsiella oxytoca* and *Proteus mirabilis*. Fungal D-glucan test, aspergillus galactomannan, the fungal smear of skin lesion secretions and culture were negative. CT scans of the head and chest showed brain atrophy, right middle ear mastoiditis, scattered foci of fibrosis in both lungs, and left pleural thickening. Histopathological examination of the lesions showed visible *Aspergillus* clusters, mostly neutrophil-dominated inflammatory necrotic tissue and keratinized epithelium (Figure 2A). Periodic Acid-Schiff stain showed fungal hyphae, which were septate and branched (Figure 2B). Alcian blue staining shows fungal mycelium and spores (Figure 2C). The initial diagnosis was primary cutaneous aspergillosis, and there was no significant improvement after 2 months of treatment with itraconazole capsules (200 mg bid). A second histopathological examination showed a basal cell-like tumor mass in the dermis with extensive infiltration between the fibers and a fenestrated arrangement of cells around the mass (Figure 2D). No fungus was seen in the lesion on this histopathological examination. A diagnosis of giant basal cell carcinoma complicated by primary cutaneous aspergillosis disease was made. Surgical excision was recommended, but the patient refused due to financial reasons and continuous follow-up.



Figure 1 Clinical feature. The lesion on the right ear.

Discussion

Basal cell carcinoma (BCC) is the most common skin tumor in the human body, which is more common in older people and more common in men than women, with a male-to-female ratio of about 2:1.¹ Ultraviolet radiation and PTCH1 gene mutations are its main causative factors, and inactivating mutations of PTCH1 are found in 90% of patients with sporadic BCC.² Surgical resection is effective for most primary BCC, with a less than 2–8% recurrence rate at 5 years after surgery.³ In addition to conventional surgical therapy, Mohs microsurgery can be used, which has the advantage of completely removing tumor cells by histopathological examination and maximum preservation of normal tissue. Giant basal cell carcinoma (GBCC) refers to BCC with a diameter of ≥ 5 cm, and this clinical subtype is rare, accounting for about 0.5–1% of BCCs.⁴ GBCC is aggressive and can infiltrate deep tissues involving muscle, cartilage, and bone, and may metastasize, so the prognosis is often poor.^{5,6}

Aspergillus is widely present in cereals, soil, air, and other nature and is a common opportunistic pathogenic fungus. Primary Cutaneous Aspergillosis is the direct invasion of pathogenic fungi into damaged skin, such as trauma sites, burns, intravenous catheters, surgical wounds, and closed dressing impregnated areas. Clinical lesions present in a variety of ways, including erythema, ulcers, papules, and nodules. The characteristic pathological manifestation of cutaneous aspergillosis can be seen as clear, separated mycelium branched at 45°. Treatment includes amphotericin B, itraconazole, or terbinafine.

Conclusion

Our patient suffered from BCC, which caused the ear ulcers for a long time, and the skin barrier was destroyed during the period without regular and standardized treatment, and coupled with long-term agricultural activities, the huge ulcerated surface was exposed to the air, leading to opportunistic infection by Aspergillus, which further aggravated and concealed

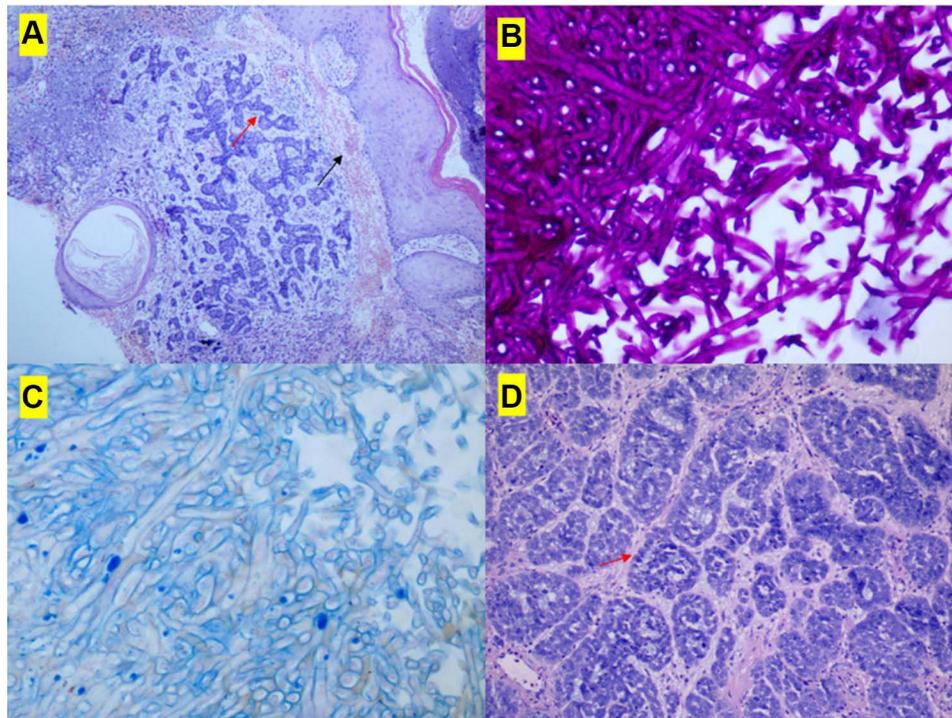


Figure 2 Histological features. (A) Aspergillus clusters can be seen (red arrow), primarily neutrophil (black arrow)-dominated inflammatory necrotic tissue and keratinized epithelium (HE $\times 100$). (B) PAS-positive hyphae of aspergillus (PAS $\times 400$). (C) Fungi are Alcian blue positive (AB $\times 400$). (D) Basal cell-like tumor masses can be seen in the dermis, with extensive infiltration between fibers, fenestrated arrangement of cells around the masses (red arrow) (HE $\times 200$).

the disease. After antifungal treatment, the fungal infection at the ulcer improved, and at this time, no fungal organism was seen on pathological biopsy, which allowed the diagnosis of BCC to be confirmed.

We presented a rare case of basal cell carcinoma complicated by primary cutaneous aspergillosis. Two biopsies were performed successively in combination with history and physical examination, and the diagnosis was finally confirmed. Early detection and diagnosis as well as early treatment are of great benefit to this disease. Many primary dermatologists do not know enough about skin cancer, so there are some difficulties in the diagnosis of such disease. We suggest more medical training to improve the diagnostic ability of skin tumors to better serve patients.

Ethics Statement

Written informed consent for publication of their clinical details and clinical images was obtained from the proxy. No institutional approval was required.

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Disclosure

The authors report no conflicts of interest in this work.

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