

Letters to the Editor**Isavuconazole Successfully Treated Pulmonary Mucormycosis in Acute Myeloid Leukemia: A Case Report****Keywords:** Isavuconazole; Pulmonary mucormycosis; Acute myeloid leukemia; mNGS.**Published:** May 01, 2025**Received:** February 12, 2025**Accepted:** April 12, 2025**Citation:** Chen H., Chen K., Kong W. Isavuconazole successfully treated pulmonary mucormycosis in acute myeloid leukemia: a case report. *Mediterr J Hematol Infect Dis* 2025, 17(1): e2025031, DOI: <http://dx.doi.org/10.4084/MJHID.2025.031>

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To the editor.

Mucormycosis is an infrequent but deadly fungal infection with high mortality rates (50%), particularly in pulmonary cases.¹ Among patients with hematological malignancies, mortality is even higher at 76%, as reported in an Eastern China study.² Early diagnosis remains challenging due to the low sensitivity and specificity of diagnostic tests, often delaying treatment.³ Recent studies highlight the potential of metagenomic next-generation sequencing (mNGS) for timely diagnosis. Isavuconazole has also shown efficacy comparable to amphotericin B and is well-tolerated.^{4,5} Here, we report a case of successful treatment of *Cunninghamella bertholletiae* with isavuconazole in a patient with hematological malignancy.

A 31-year-old Chinese male was admitted to the hematology department on September 30, 2024, with a continuous fever lasting over five days and bilateral axillary lymph node enlargement. Piperacillin-tazobactam was empirically initiated for suspected bloodstream infection. On October 1, blood tests revealed hemoglobin at 6.1 g/dL, red blood cell count at $1.71 \times 10^{12}/L$, white blood cell count at $116.27 \times 10^9/L$, absolute monocyte count at $77.32 \times 10^9/L$, and platelets at $36.0 \times 10^9/L$ (**Figure 1A**). The patient's maximum temperature was 39.7°C (**Figure 1B**), and interleukin-6 (IL-6) levels were 24.77 pg/mL. Bone marrow analysis confirmed acute myeloid leukemia (AML) M4 subtype with 58% myeloid blasts and mutations in FLT3, KRAS, PHF6, RUNX1, and WT1. Flow cytometry showed positivity for CD38, HLA-DR, CD13, CD33, CD11b, CD14, CD64, CD4 (partial), and cMPO (partial), while CD34, CD117, and lymphoid markers were negative. The patient was started on the DAV regimen (daunorubicin, cytarabine, venetoclax) on October 5.

Despite six days of piperacillin-tazobactam, the patient had persistent fevers, and a chest CT showed no signs of infection. The anti-infective regimen was adjusted to piperacillin-tazobactam combined with moxifloxacin on October 6 and further switched to

imipenem-cilastatin with voriconazole on October 10 due to persistent fever. On October 23, chest CT revealed a patchy consolidation in the lower lobe of the right lung (**Figure 1C**). Peripheral blood mNGS identified *Cunninghamella bertholletiae* (Reads: 23; Relative abundance: 3.24%). Blood cultures and G and GM tests were repeatedly negative during treatment. A histopathologic specimen could not be obtained due to severe thrombocytopenia. Based on the 2019 EORTC/MSG criteria⁶ and the mNGS findings, possible pulmonary mucormycosis was diagnosed, and the treatment regimen was changed to imipenem-cilastatin with isavuconazole. Following isavuconazole administration, the patient's fever resolved. A chest CT on November 5 showed significant improvement in the pulmonary consolidation, and the patient was discharged on oral isavuconazole. By December 4, follow-up CT scans revealed further resolution of the lung lesion (**Figure 1D**). The patient continues oral isavuconazole without adverse events and is awaiting subsequent chemotherapy.

Although the patient was successfully treated, this study has limitations. Due to the patient's extremely low platelet count, a pathological examination could not be performed, preventing definitive confirmation of the fungal infection and allowing only a probable diagnosis. Additionally, the minimum inhibitory concentration of isavuconazole against *Cunninghamella bertholletiae* was not obtained.

In conclusion, this case highlights the diagnostic value of mNGS for pulmonary mucormycosis in hematological malignancies and supports the efficacy and tolerability of isavuconazole in treating this condition.

Ethics approval and consent to participate. The study protocol was approved by the Medical Ethics Committee of the First People's Hospital of Zigong. The patient gave informed consent for the publication of this case report.

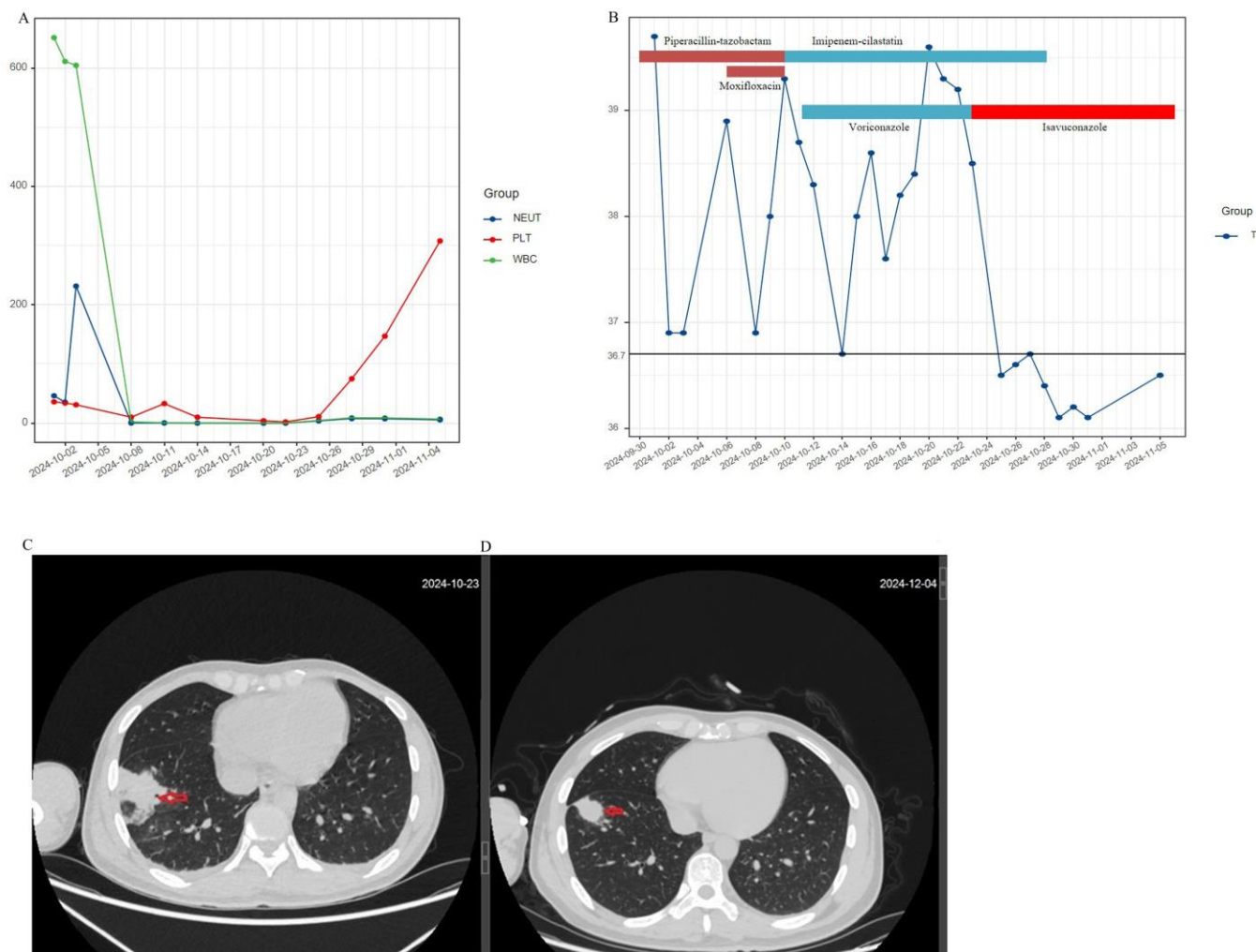


Figure 1A, B, C and D.

Hong Chen¹, Kun Chen² and Wenqiang Kong³.

¹ Zigong Gongjing Hospital of Traditional Chinese Medicine, Zigong, 643000, Sichuan, China.

² Department of Hematology, Zigong First People's Hospital, Zigong, 643000, Sichuan, China.

³ Department of Pharmacy, Zigong First People's Hospital, Zigong, 643000, Sichuan, China.

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Correspondence to: Wenqiang Kong. Department of Pharmacy, Zigong First People's Hospital, No 42, Shangyihao Branch Road, Zigong, 643000, Sichuan, China. E-mail: wqkongpharmacist@outlook.com

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