

**Application of metagenomic next-generation sequencing and targeted metagenomic next-generation sequencing in diagnosing pulmonary infections in immunocompetent and immunocompromised patients**

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### *Characteristics of pulmonary infection and mNGS detection*

In this study, causative pathogens were defined according to the results of mNGS and culture methods, clinical symptoms, other laboratory results, and clinical experience. Patients with pulmonary infection was divided into single infection ( $n = 166$ ) and mixed infection ( $n = 87$ ) groups, and the former accounted for about two thirds of cases. For single infection, bacteria (44.3%, 112/255) were the most common pathogens, followed by fungi (10.7%, 27/255) and viruses (6.7%, 27/255) (Figure S1a). In addition to single infection, our results show that patients with pulmonary infection can also be simultaneously infected by different kinds of pathogens, including bacterial and fungal co-infection, bacterial and viral co-infection, fungal and viral co-infection, and bacterial, fungal and viral co-infection (Figure S1a).

Besides, mNGS detection results are summarized in Figure S1b. We found that the most common bacterial pathogens detected by mNGS were *Klebsiella pneumoniae* ( $n = 69$ ), *Acinetobacter baumannii* ( $n = 47$ ), *Enterococcus faecium* ( $n = 39$ ), and *Staphylococcus epidermidis* ( $n = 38$ ). The most common fungal pathogens were *Candida albicans* ( $n = 116$ ), *Candida parapsilosis* ( $n = 28$ ), *Pneumocystis jirovecii* ( $n = 24$ ), *Candida tropicalis* ( $n = 24$ ), and *Candida glabrata* ( $n = 23$ ). Human gammaherpesvirus 4 (EBV) ( $n = 42$ ), Human alphaherpesvirus 1 ( $n = 33$ ), and Human betaherpesvirus 5 (CMV) ( $n = 30$ ) were the most common viral pathogens.



Figure S1 Characteristics of pulmonary infection and mNGS detection results. a, Different kinds of infections. b, mNGS detection.