Exploring the Clinical Utility of Metagenomic Next-Generation Sequencing in the Diagnosis of Pulmonary Infection

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ABSTRACT

Introduction: We aimed to explore the real-world clinical application value and challenges of metagenomic next-generation sequencing (mNGS) for pulmonary infection diagnosis.

Methods: We retrospectively reviewed the results of mNGS and conventional tests from 140 hospitalized patients with suspected pulmonary infections from January 2019 to December 2020. The sample types included bronchoalveolar lavage fluid, lung tissue by transbronchial lung biopsy, pleural effusion, blood, and bronchial sputum. Apart from the mNGS reports that our patients received, an extra comprehensive and thorough literature search was conducted.

Results: Significant differences were noticed in the positive detection rates of pathogens between mNGS and conventional diagnostic testing (115/140, 82.14% vs 50/140, 35.71%, $P < 0.05$). The percentage of mNGS-positive patients was significantly higher than that of conventional testing-positive patients with regard to bacterial detection ($P < 0.01$), but no significant differences were found with regard to fungal detection ($P = 0.67$). Significant statistical differences were found between mixed infection cases (15, 22.70%) and single infection cases (4, 7.84%) in terms of diabetes ($P = 0.03$). The most frequent pattern of mixed infection was bacteria and fungi mixed infection (40, 40/89 = 44.94%), followed by bacteria mixed infection (29, 29/89 = 32.58%). The sensitivity of mNGS in pulmonary infection diagnosis was much higher than that of conventional test (89.17% vs 50.00%; $P < 0.01$), but the specificity was the opposite (75.00% vs 81.82%; $P > 0.05$).

Conclusion: mNGS is a valuable tool for the detection of pulmonary infections, especially mixed pulmonary infections. The most common combinations we found were bacterial–fungal coinfection and bacterial–bacterial coinfection. Still, there are many challenges in the clinical application of mNGS in the diagnosis of pulmonary infections. There is still a lot of work to be done in interpreting the mNGS
reports, because both clinical judgment and literature analysis strategy need to be refined.

**Keywords:** Application; Bronchoalveolar lavage fluid (BALF); Bronchoscopy; Clinical diagnosis; Etiology; Metagenomic next-generation sequencing (mNGS); Mixed infection; Pulmonary infection; Transbronchial lung biopsy (TBLB)

### Key Summary Points

**Why carry out this study?**

Pulmonary infections remain important causes of morbidity and mortality in the world. Improved diagnostic methods with better sensitivity, speed, and spectrum for pathogen detection are urgently needed.

We aimed to explore the real-world clinical application value and challenges of metagenomic next-generation sequencing (mNGS) for pulmonary infection diagnosis.

**What was learned from the study?**

Significant differences were noticed in the positive detection rates of pathogens between mNGS and conventional diagnostic testing (115/140, 82.14% vs 50/140, 35.71%, \( P < 0.05 \)) and the sensitivity of mNGS in pulmonary infection diagnosis was much higher than that of conventional test (89.17% vs 50.00%; \( P < 0.01 \)).

mNGS is a valuable tool for the detection of pulmonary infections, especially mixed pulmonary infections. There is still a lot of work to be done in interpreting the mNGS reports, because both clinical judgment and literature analysis strategy need to be refined.

### DIGITAL FEATURES

This article is published with digital features, including a summary slide, to facilitate understanding of the article. To view digital features for this article go to [https://doi.org/10.6084/m9.figshare.14696046](https://doi.org/10.6084/m9.figshare.14696046).

### INTRODUCTION

Pulmonary infections remain important causes of morbidity and mortality in the world [1, 2]. Although various assays exist, rapid and accurate diagnosis of causative pathogens is still difficult. Traditional culture methods, only used for fungal and bacterial detection, are time-consuming and exhibit low rates of positive detection, and therefore cannot meet the clinical needs. Other detection methods like polymerase chain reaction (PCR) and immunology techniques can only identify several specific pathogens. Moreover, infections with multiple pathogens and the appearance of multidrug-resistant pathogens make the identification of the causative agent more difficult. Therefore, improved diagnostic methods with better sensitivity, speed, and spectrum for pathogen detection are urgently needed.

Metagenomic next-generation sequencing (mNGS) is a novel and promising approach which combines high-throughput sequencing with bioinformatics analysis. mNGS requires shorter analysis time, has a wide range of detectable pathogens (all pathogens including bacteria, fungi, viruses, and parasites can be simultaneously identified by a single assay), simple sample-processing, is less affected by prior antibiotic use, and has semiquantitative value in follow-up [3].

Our study summarized the mNGS results for 140 adult patients with suspected pulmonary infection and aimed to evaluate the clinical performance and effect of mNGS in pathogen diagnosis.
METHODS

Ethics Statement

The study was conducted in accordance with the 1964 Helsinki declaration and its later amendments or comparable ethical standards. This study was approved by the ethics committee of The Affiliated Wuxi Second People’s Hospital of Nanjing Medical University. Since the data were completely anonymous, informed consent was waived.

Patients and Sample Collection

We retrospectively reviewed 140 hospitalized patients with suspected pulmonary infections in the Department of Respiratory and Critical Care, The Affiliated Wuxi Second People’s Hospital of Nanjing Medical University from January 2019 to December 2020. The attending physician judged whether the patient had suspected pulmonary infection on the basis of clinical manifestations and imaging examination results.

The inclusion criteria were as follows: (1) patients who were suspected of having suspected pulmonary infections; (2) patients who agreed to undergo the mNGS examination; (3) bronchoalveolar lavage fluid (BALF) samples and detection process passed quality control for mNGS; and (4) patients whose medical data were recorded completely.

The exclusion criteria were as follows: (1) patients who refused to undergo the mNGS examination; (2) BALF samples or detection process failed to pass quality control for mNGS; (3) patients with incomplete clinical and laboratory data;

Bronchoscopy was performed by experienced bronchoscopists to obtain specimens including BALF and lung tissue. Transbronchial lung biopsy (TBLB) was performed and BALF was collected by BF-P-260F ultrathin bronchoscope (Olympus, Tokyo, Japan). DirectPath V1.0 virtual bronchoscopic navigation (Olympus, Tokyo, Japan) was used for assistance in navigation.

Body fluid samples, lung biopsy tissues, and plasma samples were sent to BGI Co., Ltd (Shenzhen, China) for sequencing as described in detail below. The remaining specimens were sent to our microbiological laboratory. The conventional tests were performed by culture of the bacteria or fungi in blood agar or in Sabouraud agar at 35 °C for a maximum period of 5 days. Bacteria or fungi culturing were performed by using the VITEK-II Compact automated microbiological system (bioMérieux, France) [4]. Other conventional diagnostic testing included Chlamydia pneumoniae, Mycoplasma pneumoniae, Legionella pneumophila, cytomegalovirus (CMV), Epstein–Barr virus (EBV), adenovirus, and herpes simplex virus (HSV) serological antibody detection. Xpert testing for Mycobacterium tuberculosis and so on.

Metagenomic Next-Generation Sequencing and Analysis

Body Fluid Sample: Sample Processing and DNA Extraction

A body fluid sample (1.5–3 mL of BALF, pleural effusion, or sputum) was collected from each patient according to standard procedures [5]. A 1.5-mL microcentrifuge tube containing 0.6 mL sample, enzyme, and 1 g of 0.5-mm glass beads was agitated by a vortex mixer, and a 0.3-mL sample was separated into a 1.5-mL microcentrifuge tube and DNA was extracted using the TIANamp Micro DNA Kit (DP316, Tiangen Biotech) following the manufacturer’s operational manual. The extracted DNA samples were used for the construction of DNA libraries [6].

Lung Biopsy Tissue: Sample Processing and DNA Extraction

Lung biopsy tissue was minced into small pieces and ground with a tissue homogenate machine. A 10-mg sample was separated into a 1.5-mL microcentrifuge tube and DNA was extracted using the TIANamp Micro DNA Kit (DP316, Tiangen Biotech) following the manufacturer’s operational manual.
**Plasma Sample: Sample Processing and DNA Extraction**

Blood (3–5 mL) was drawn from patients and centrifuged at 4000 rpm for 10 min at 4 °C. DNA was extracted from 300 μL of plasma using the TIANamp Micro DNA Kit (DP316, Tiangen Biotech, Beijing, China) according to the manufacturer’s operational manual.

**Construction of DNA Libraries**

DNA libraries were constructed through DNA fragmentation (about 150 bp), end repair, adapter ligation, and PCR amplification. An Agilent 2100 Bioanalyzer (Agilent Technologies, Santa Clara, CA, USA) was used for quality control of the DNA libraries (200–300 bp). Then, libraries with confirmed quality were sequenced by the BGISEQ-50 platform (BGI Co., Ltd, Shenzhen, China) [7].

**Sequencing and Bioinformatic Analysis**

High-quality sequencing data were generated by removing low-quality reads, followed by computational subtraction of human host sequences mapped to the human reference genome (hg19) using Burrows–Wheeler alignment [8]. The remaining data by removal of low-complexity reads were classified by simultaneously aligning to four microbial genome databases (bacteria, fungi, viruses, and parasites) which were downloaded from the National Center for Biotechnology Information (NCBI) (ftp://ftp.ncbi.nlm.nih.gov/genomes/). RefSeq contains 4945 whole genome sequence of viral taxa, 6350 bacterial genomes or scaffolds, 1064 fungi related to human infection, and 234 parasites associated with human diseases.

**Threshold Criteria for Interpretation of mNGS Result**

SMRN: number of sequences strictly mapped at species level.

- SMRN of microorganisms in sample is higher than in the negative control group.

For different types of microorganisms, the thresholds were set as follows:

- **Bacterial/virus/fungus/mycoplasma/chlamydia:** SMRN ≥ 3.
- **Parasite:** SMRN ≥ 100.
- **Mycobacterium tuberculosis:** SMRN ≥ 1.

Definite pathogens of pulmonary infection; excluding the normal flora of oral cavity, respiratory tract or the skin (via a large number of literature search).

Meets clinical judgment or targeted treatment response by three experienced senior clinicians.

Microorganisms detected by mNGS which meet all the criteria were identified as suspected pathogens.

**Statistical Analyses**

Continuous variables are expressed as mean ± standard deviation (SD) and categorical variables are expressed as numbers (percentage). We tested for differences in continuous variables using t test and categorical variables with chi-square test as appropriate. The 2 × 2 contingency tables were established to determine sensitivity, specificity, positive predictive value (PPV), and negative predictive value (NPV).

The results are presented with 95% confidence intervals (CIs).

Data analyses were performed using SPSS 22.0 software and a two-tailed value of P of 0.05 was considered to represent a significant difference.

**RESULTS**

**Baseline Characteristics of Patients**

A total of 140 patients (62 men, 44.28%; 78 women, 55.72%) with suspected pulmonary infection were included in this study (Table 1).

The mean age was 59.16 ± 14.89 years and the majority of patients were above 40 but at most 70 years old (61.43%).

Among them, 75 (75/140 = 53.57%) patients had underlying diseases, including 23 cases of type 2 diabetes (16.43%), 16 (11.43%) cases of connective tissue disease, 14 (10.00%) cases of bronchiectasis, 10 (7.14%) cases of...
extrapulmonary malignancies, 7 (5.00%) cases of previous history of tuberculosis, 7 (5.00%) cases of previous history of lung cancer, 5 (3.57%) cases of chronic obstructive pulmonary disease, and 1 case of bronchial asthma.

A total of 128 patients underwent bronchoscopy and 9 patients underwent TBLB. The main sample types of mNGS in our patients were BALF (119, 85.00%), while other sample types included lung tissue (9), pleural effusion (6), blood (5), bronchial sputum (1), and so on. The mean time from admission to sampling was 4.3 days.
Concordance Between mNGS and Conventional Diagnostic Testing

In the present study, mNGS and conventional diagnostic testing (including culture) were both positive for pathogens detection in 48 (48/140 = 34.29%) cases. A total of 67 (67/140 = 48.10%) cases were positive for pathogens detection by mNGS only. There were only 2 cases positive for pathogens detection by conventional test only. mNGS and conventional test were both negative for pathogens detection in 23 (23/140 = 16.43%) cases (Fig. 1).

For the 48 both-positive cases, results of mNGS and conventional tests were totally matched in 2 (2/48 = 4.17%) cases and were completely mismatched in 26 (26/48 = 54.17%) cases. A total of 20 (20/48 = 41.67%) cases were found to be partly matched which means at least one detected pathogen was overlapped between conventional diagnostic testing and mNGS (Fig. 1).

Comparison of Pathogenic Detection Between mNGS and Conventional Diagnostic Testing

Significant differences were noticed in the positive detection rates of pathogens between mNGS and conventional diagnostic testing (115/140, 82.14% vs 50/140, 35.71%, \( P < 0.05 \)).

Among pathogens detected by mNGS (Fig. 2a) and conventional test, \textit{Parvimonas micra} (\( n = 8 \)) was the most frequently detected Gram-positive bacteria, followed by \textit{Streptococcus pneumoniae} (\( n = 7 \)), \\textit{Abiotrophia defectiva} (\( n = 7 \)), \textit{Enterococcus faecium} (\( n = 6 \)), \textit{Streptococcus constellatus} (\( n = 5 \)), and \textit{Actinomyces israelii} (\( n = 5 \)).

According to the results of mNGS (Fig. 2b) and conventional test, \textit{Haemophilus parainfluenzae} (\( n = 19 \)) was the most frequently detected Gram-negative bacteria, followed by \textit{Pseudomonas aeruginosa} (\( n = 13 \)), \textit{Acinetobacter baumannii} (\( n = 10 \)), \textit{Klebsiella pneumoniae} (\( n = 9 \)), \textit{Prevotella melaninogenica} (\( n = 8 \)), and \textit{Pyramidobacter piscolens} (\( n = 7 \)).

The most commonly fungi detected by mNGS (Fig. 2c) and conventional test were...
Candida albicans (n = 23), Pneumocystis jirovecii (n = 4), and Aspergillus fumigatus (n = 3).

The results of mNGS showed that a total of 21 patients were virus-positive (21/140 = 15%).
and the most frequently detected viruses were Human gammaherpesvirus 4 (EBV) (n = 16), Human betaherpesvirus 7 (HHV-7) (n = 5), and Human betaherpesvirus 5 (CMV) (n = 4) (Fig. 2d).

In addition, six cases of Mycoplasma, one case of Chlamydia psittaci, and seven cases of M. tuberculosis were also detected by mNGS or conventional test (Fig. 2e).

The percentage of mNGS-positive patients was significantly higher than that of conventional testing-positive patients with regard to bacterial detection (P < 0.01), but no significant
differences were found with regard to fungal detection ($P = 0.67$) (Fig. 3).

*Streptococcus infantis*, *Oribacterium parvum*, *Prevotella pallens*, *Kingella oralis*, *Mogibacterium timidum*, *Treponema maltophilum*, *Dialister invisus*, *Streptococcus parasanguinis*, *Neisseria subflava*, *Rothia dentocariosa*, *Campylobacter showae*, *Actinomyces johnsonii*, *Johnsonella ignava*, *Cardiobacterium hominis*, *Scardovia wiggsiae*, *Cardiobacterium valvarum*, *Desulfovibrio orale*, *Fretibacterium fastidiosum*, *Actinomyces gerencseriae*, etc. were not interpreted as pathogens, because they were known as normal flora of the oral cavity, respiratory tract, or the skin and were not considered to cause pulmonary infection so far (via literature searches).

**Comparison of Clinical Characteristics Between Mixed and Single Pulmonary Infection**

Mixed pulmonary infection was defined when two or more infectious pathogens were identified by mNGS or conventional test. As shown in Table 2, there were 66 cases of mixed pulmonary infection and 51 cases of single pulmonary infection.

No difference in age between mixed infection cases ($60.15 \pm 15.90$) and single infection cases ($58.45 \pm 13.55$) was observed ($P = 0.54$).

There were no significant differences in gender or inflammatory indices between the two groups.

The most common symptoms in both groups were cough, fever, and thoracalgia.

Additionally, there were no significant differences in underlying pulmonary disease, including bronchiectasis, chronic obstructive pulmonary disease (COPD), previous history of tuberculosis, bronchial asthma, and lung cancer, between these two groups.

Significant statistical differences were found between mixed infection cases (15, 22.70%) and single infection cases (4, 7.84%) in terms of diabetes ($P = 0.03$). However, there were no differences between the two groups in terms of conventional testing was not; conventional testing+, only the conventional testing result was positive, mNGS was not

![Fig. 3](image_url)
other underlying extrapulmonary diseases like connective tissue disease, and malignant tumor (11, 16.7% vs 5, 9.80%; 4, 6.10% vs 4, 7.84%, respectively).
No difference in time from admission to test between the two groups (3.97 ± 3.00 vs 4.69 ± 3.83) was observed ($P = 0.26$).

**Diagnostic Performance of the Combination of mNGS and Conventional Diagnostic Testing in Mixed Infection**

A total of 69 patients (69/140 = 49.29%) were diagnosed with a mixed infection using only the mNGS method. When combined with conventional test results, the diagnostic ratio of the mixed infection was increased to 63.57% (89/140) (Fig. 4).

The most frequent pattern of mixed infection was bacteria and fungi mixed infection (40 patients, 40/89 = 44.94%), followed by bacteria mixed infection (29 patients, 29/89 = 32.58%) (Fig. 4).

Eight patients were diagnosed as bacteria, fungi, viruses, and mycoplasma mixed infection (8/89 = 8.99%); eight patients were diagnosed as bacteria, fungi and viruses mixed infection (8/89 = 8.99%) (Fig. 4).

**Comparison of mNGS and Conventional Test in the Diagnosis of Pulmonary Infection**

The performance of mNGS and conventional test in diagnosis of pulmonary infection is shown in Table 3.

The sensitivity and specificity of diagnosing pulmonary infection by mNGS were 89.17% (95% CI 83.61–94.73) and 75.00% (95% CI 63.48–86.52), respectively, with NPV and PPV being 95.54% and 53.57%.

As for conventional test, the sensitivity and specificity of diagnosing pulmonary infection were 50.00% (95% CI 40.98–59.02) and 81.82% (95% CI 65.70–97.94), respectively, with NPV and PPV being 93.65% and 23.38%.

The sensitivity of mNGS in pulmonary infection diagnosis was much higher than that of conventional test (89.17% vs 50.00%; $P < 0.01$), but the specificity was the opposite, with no statistically significant difference (75.00% vs 81.82%; $P > 0.05$).

**DISCUSSION**

In our study, mNGS exhibited better performance than conventional test for detecting bacteria ($P < 0.01$), whereas it was not superior to conventional test with regard to fungal detection ($P = 0.67$). This was inconsistent with
previous findings. According to Miao et al., the sensitivity of mNGS is not better than that of culture for recognizing bacteria, but mNGS has superior feasibility in detecting fungi [9]. Another study found that compared with sequencing, culture is able to identify most bacterium-associated pneumonia [10]. Unlike the aforementioned studies, Xie et al. reported that there was no significant difference between conventional microbiologic test and mNGS with regard to fungal detection; however, mNGS displayed a significantly higher positive rate for detecting pathogenic bacteria [11]. Possible explanations for this divergence are due to different diseases, different sample types, and different test conditions of mNGS and conventional test. Moreover, the positive rate of pathogenic fungi cultivation in our study is relatively higher.

Our results showed that the sensitivity of mNGS in pulmonary infection pathogen detection was much higher than that of the conventional test (89.17% vs 50.00%; \( P < 0.01 \)), while the specificity of the conventional test was better than mNGS (75.00% vs 81.82%; \( P > 0.05 \)). This outcome is inconsistent with the study by Cai et al., who found that the sensitivity and specificity of mNGS were both superior to those of culture [12]. However, another study showed that the specificity of mNGS and conventional test are nearly the same (95.2% and 95.2%, respectively; \( P = 1.0 \)) [13]. Similarly, it has also been reported that the specificity of mNGS was comparable to that of conventional test [9]. On the other hand, other reports have reached similar conclusions to ours [14, 15]. Several explanations might account for these differences.

In general, the overall positive rate of mNGS (115/140, 82.14%) for pathogens in pulmonary infections was significantly higher than that of conventional test (35.71%, \( P < 0.05 \)). This result is consistent with previous studies. In a multicenter, prospective study, Parize et al. showed that clinically relevant pathogens were identified in a significantly higher proportion of patients with mNGS than conventional microbiological methods (36/101 (36%) vs 11/101 (11%), respectively, \( P < 0.001 \)) [16]. Langelier et al. reported that mNGS has a greater capacity for detecting microorganisms compared with current microbial diagnostics [17]. These findings, along with our result, confirm that mNGS can raise the detection rate of respiratory pathogens.

In terms of mixed infection diagnosis, the proportion of mixed pulmonary infections in our patients was much higher than we originally thought. We found that a total of 69 (69/140 = 49.29%) cases were positive for mixed infection by mNGS only. However, when combined with conventional test results, the positive ratio of the mixed infection increased to 63.57% (89/140). These results are in agreement with Fang et al.’s findings which showed that, compared with using solely the mNGS method, the diagnostic ratio of the mixed infection was increased from 55.6% to 58.3% (42/72) when combining mNGS and conventional test [4]. Their results and ours both indicate that the combination of mNGS and conventional test contributes to the diagnostic ratio of the mixed pulmonary infection. Prior studies have shown that mNGS was significantly better at the identification of coinfections than conventional tests like serological antibody, PCR, or culture [16, 18, 19].

This study found that the most common combinations were bacterial–fungal coinfection and bacterial–bacterial coinfection. Unlike our

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result, Wang et al. showed that the most common mixed pulmonary infection was the combination of viruses and fungi [14]. In another study, Legoff et al. observed that bacterial–viral coinfection was the most frequent combinations [20]. In 2020, Wu et al. reported that the most common combinations were viral–bacterial coinfection and bacterial–bacterial coinfection in patients with normal immune function, while bacterial–viral coinfection and bacterial–fungal coinfection were the most frequent combinations in patients with compromised immune function [21]. The inconsistency between our study and other studies might result from the low proportion of immunocompromised patients in this study. Another possible explanation is that our hospital is located in the lower reaches of the Yangtze River, a high humidity area in which fungal infections are common. *C. albicans* was the major fungal pathogen in this region [22]. Furthermore, we observed triple mixed pulmonary infection (bacteria, fungi, and viruses) and fourfold mixed pulmonary infection (bacteria, fungi, viruses, and mycoplasma) in some patients, the majority of whom were diabetic. Similar results have been reported before [21]. Owing to the advantages mNGS, so many patterns of mixed pulmonary infection can be detected. On the other hand, mixed infections can hardly be identified by traditional culture methods, because various microorganisms interact with, and inhibit, each other. It is known that severe pneumonia is usually not caused by a single pathogen but by coinfections. Thus, mNGS is a more favorable method that may improve clinical guidance in the diagnosis and treatment of patients with pneumonia.

Few studies have discussed the value of mNGS in etiological detection for pulmonary infection in patients with diabetes. In terms of diabetes mellitus, in our study, significant statistical differences were observed between mixed pulmonary infection cases (15, 22.70%) and single infection cases (4, 7.84%) (P = 0.03). This outcome is contrary to that of Wu et al. who found no differences between the patients with and without mixed infections in terms of diabetes [21]. Diabetes is a risk factor for suffering pulmonary infection like community-acquired pneumonia, CAP [23]. People with diabetes are at two times greater risk of developing a pulmonary infection than healthy people [24]. In the present study, mNGS was performed on 24 patients with diabetes and lower respiratory tract infection for pathogen identification. Our data show that patients with diabetes are more susceptible to mixed respiratory tract infections, especially bacterial and fungal infections. Patients with diabetes mellitus were prone to invasive fungal infections [25]. The most common pathogens are *A. fumigatus* and *C. albicans* [26, 27]. This is consistent with the results of our current study. In addition, diabetes mellitus increases active pulmonary tuberculosis risk and worsens its outcomes [28]. Two patients with diabetes were diagnosed with active pulmonary tuberculosis by mNGS in this study. These results indicate that for the diabetic population, mNGS are well suited for etiological detection of pulmonary infection.

mNGS is a bias-free assay that can detect all pathogens in the environment; however, this test method itself cannot distinguish pathogenic microorganisms from colonizing microorganisms, background microorganisms, and contaminated microorganisms [29]. For now, no authoritative guide is available to the interpretation of the mNGS report and therefore this remains a great challenge for clinicians [29, 30].

In order to interpret the results of mNGS more accurately and objectively, in our study, we conducted an extra comprehensive and thorough literature search in PubMed in an attempt to identify published cases of pulmonary infection due to pathogens that we detected, without time limits. We found out in mNGS reports that our patients received that some microorganisms which do not cause human pulmonary infections (e.g., *Torque teno virus 15*, *Shigella dysenteriae*, *Cardiobacterium hominis*), some microorganisms which are not pathogenic in humans, or some microorganisms whose pathogenicity is as yet unclear (e.g., *Bacteroides vulgatus*, *Bifidobacterium longum*, *Parabacteroides merdae*, *Pyramidobacter piscolens*, and *Megasphaera micronuciformis*) were all
included in the pathogenic microorganism list. This problem is relatively easy to solve by reading related literature. The most troublesome issue is opportunistic pathogens. Some microorganisms, for instance, *S. constellatus*, are part of the normal flora of the oral or respiratory tracts of humans, but pulmonary infections due to them have been widely reported [31, 32]; on the contrary, although reports have shown that some microorganisms, such as *A. israelii*, can cause respiratory infection [33], the number of articles was quite limited (sometimes only one to two). In fact, the types of pathogens that cause pulmonary infections are constantly evolving. For example, the isolation rate of *H. parainfluenzae* in patients with pneumonia is increasing year by year; although it remains controversial, this bacterium has been increasingly regarded as an important pathogen of respiratory infections [34]. All of the aforementioned problems pose obvious difficulties and challenges to clinicians when performing a literature search. Therefore, we think that the aforementioned two lists in mNGS reports to some extent can facilitate the interpretation, but it is somewhat arbitrary and can only provide limited reference values to clinicians. An extra comprehensive and elaborate literature search should be conducted.

Moreover, these findings could not be generalized. Various clinical factors including age, symptoms and signs, laboratory tests, imaging performance, immune status, care status, the risk for aspiration pneumonia, ventilator use, tracheal intubation, antibiotic treatment history, treatment response, and so on should all be taken into account [4, 16, 21, 35]. In different patients, even for the same bacteria, we may need to change our judgment.

Certainly, one should also consider the particularity and complexity of mNGS in diagnosing pulmonary infections. In the context of the misuse and unreasonable administration of antibiotics, numerous opportunistic pulmonary infections have become a tough clinical problem in immunocompromised patients. In cerebrospinal fluid and brain biopsy samples, which are considered sterile, the result of mNGS is easy to interpret [36–38]. In contrast, it is much more difficult to evaluate mNGS reports in BALF and sputum [39, 40]. Recently, it has been reported that in patients with severe pneumonia, the utility of mNGS for the simultaneous detection of pathogens from both BALF and blood samples is strongly recommended [41].

The present study has certain limitations. Neither RNA sequencing nor drug resistance testing was performed. Nearly all patients were given empirical antimicrobial treatment before pathogen-detecting test sampling, which might decrease the rate of positive culture. The number of some sample types like blood and bronchial sputum is low. Although a comprehensive and thorough literature search was conducted, how to reasonably interpret the results of the literature search is still worth discussing. Similarly, although clinical judgment was made by three experienced senior clinicians, as a result of the lack of unified standards, subjective bias is unavoidable.

In future, we think that assembling an interdisciplinary team (consisting of medical microbiologists, computational biologists, epidemiologists, infectious disease specialists, and other clinicians) and targeted training are critical to better use of mNGS [28]. With decreasing sequencing costs, drug-resistance gene detection by mNGS may become more feasible.

**CONCLUSIONS**

mNGS exhibited better performance than conventional tests for detecting bacteria, whereas it was not superior to conventional tests with regard to fungal detection. The sensitivity of mNGS in pulmonary infection pathogen detection was much higher than that of conventional tests, while the specificity of conventional tests was better than mNGS. The overall positive rate of mNGS for pathogens in pulmonary infections was significantly higher than that of conventional tests. mNGS is a valuable tool for the detection of mixed pulmonary infections and the most common combinations we found were bacterial–fungal coinfection and bacterial–bacterial coinfection. Significant statistical differences were found between mixed infection cases and single infection cases in terms of diabetes. However,
there are still many challenges in the clinical application of mNGS in the diagnosis of pulmonary infections. There is still a lot of work to be done in interpreting the mNGS reports, because both clinical judgment and literature analysis strategy need to be refined.

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Authorship. All named authors meet the International Committee of Medical Journal Editors (ICMJE) criteria for authorship for this article, take responsibility for the integrity of the work as a whole, and have given their approval for this version to be published.

Authors’ Contributions. All authors contributed to the study conception and design. Material preparation was performed by GX, BZ, XR, JJ and TH. Data collection was performed by GX, BZ, XW, LB, YX and HZ. Analysis was performed by GX, BZ, XW and HZ. The first draft of the manuscript was written by GX and BZ. All authors read and approved the final manuscript. GX and BZ are the co-first authors and contributed equally to this study.

Disclosures. Guijuan Xie, Bo Zhao, Xun Wang, Liang Bao, Yiming Xu, Xian Ren, Jiali Ji, Ting He and Hongqing Zhao declare that they have no conflicts of interest to this work.

Compliance with Ethics Guidelines. The study was conducted in accordance with the 1964 Helsinki declaration and its later amendments or comparable ethical standards. This study was approved by the ethics committee of The Affiliated Wuxi Second People’s Hospital of Nanjing Medical University. Since the data were completely anonymous, informed consent was waived.

Data Availability. The datasets generated during and/or analyzed during the current study are available from the corresponding author on reasonable request.

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