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Transmission Electron Microscopy Improves the Diagnostic Sensitivity in Nonbacterial Etiology of Severe Pneumonia: A Retrospective Study

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ABSTRACT

Background: Severe pneumonia is responsible for great mortality and morbidity worldwide, and early-applied effective anti-infective therapy can improve the prognosis of patients. However, identification of infectious agents in severe pneumonia remains a major challenge so far. In this study, the potential utility of transmission electron microscopy (TEM) in detecting nonbacterial pathogens in patients with severe pneumonia was retrospectively evaluated.

Materials and Methods: A total of 106 patients diagnosed with severe pneumonia at our hospital from September 2015 to December 2017 were included, and their baseline clinical characteristics were collected. Nonbacterial infectious agents detected by TEM in bronchoalveolar lavage fluid (BALF) and serological tests were summarized. The detection rates were further compared between TEM and serological tests.

Results: BALF examination under the transmission electron microscope revealed 24 viruses, 16 mycoplasmas, 18 chlamydia, 2 fungi and 74 bacteria in 99 samples, among which 61 samples were mixed infections. The combined use of serological tests and TEM significantly improved the detection rate of nonbacterial infectious agents in patients with severe pneumonia.

Conclusions: Our data support that implementation of TEM could improve the sensitivity for detecting viruses, atypical pathogens and mixed infections in BALF from patient of severe pneumonia. Therefore, TEM may be used as an auxiliary diagnostic method of other microbiological tests in severe pneumonia.

Key Indexing Terms: Severe pneumonia; Transmission electron microscopy; Bronchoalveolar lavage fluid; Pathogens; Etiologic diagnosis.

INTRODUCTION

Pneumonia affects approximately 450 million people globally and causes nearly 4 million deaths each year.¹ Severe pneumonia is a common reason for intensive care unit admission and represents a major concern for physicians due to its high mortality rate.²³ Without appropriate treatment, it can rapidly progress to respiratory failure, septic shock and even cause death within several days. Early etiologic diagnosis, which would facilitate the prompt initiation of anti-infective therapy, is critical for improving the clinical outcomes of patients with severe pneumonia.

Severe pneumonia may have a bacterial, viral, fungal or other atypical etiology, including *Mycoplasma pneumoniae*, *Chlamydia pneumoniae* and *Legionella pneumophila*. Identification of causative agents in severe nonbacterial pneumonia is very challenging. Microbiological culture of lower respiratory tract specimens, blood and lung tissue is considered as the gold standard for etiologic diagnosis of pneumonia. However, for viruses and atypical pathogens, culture analysis is rarely used due to delay in the results and complicated procedures.⁴⁻⁷ While serological tests and polymerase chain reaction (PCR) targeting specific pathogens are routinely deployed, their clinical usefulness is limited in identifying a novel or rare pathogen and mixed infections.⁸⁻⁹ It has been reported that causative agents cannot be confirmed in nearly half of pneumonia episodes despite careful testing.¹⁰ Thus, anti-infective therapy is empirically applied based on clinical features and thoracic images in the early stage of pneumonia, which may increase the rate of treatment failure and cause drug-related toxicity. Therefore, there is an urgent need to develop a comprehensive assay system for prompt etiologic diagnosis of severe pneumonia.
Transmission electron microscopy (TEM) has been used in diagnostic microbiology since the 1960s, and had a profound impact on our knowledge and understanding of microorganisms. While TEM is well-recognized as a useful tool for rapid morphologic identification of infectious agents in emergent situations and infectious diseases of unknown cause, its potential diagnostic value in nonbacterial etiology of severe pneumonia remains unexplored yet.

Bronchoalveolar lavage is a common method to obtain valid samples from the lower respiratory tract. Bronchoalveolar lavage fluid (BALF) has the advantages of improving the detection of etiologic agents in community-acquired pneumonia and being suitable for multiple detection methods, including microscope. Therefore, in this study, we retrospectively analyzed the potential diagnostic value of TEM examination of BALF samples in severe pneumonia.

METHODS

Participants
A retrospective review of the medical records was made of all patients who were diagnosed with severe pneumonia and treated at Xiangya Hospital, Central South University (Changsha, Hunan, China) between September 2015 and December 2017. The diagnosis of severe pneumonia was made according to the Infectious Diseases Society of America and the American Society for Microbiology criteria. Major criteria include the requirement for mechanical ventilation or need for vasopressors. Minor criteria are listed as follows: (1) BUN >20 mg/dL; (2) confusion or disorientation; (3) hypotension requiring aggressive fluid resuscitation; (4) hypothermia with core temperature <96.8 F; (5) leukopenia with white blood cell count <4000 per mm³; (6) multilobar infiltrates; (7) PaO₂ to FiO₂ ratio <250; (8) respiratory rate >30 breaths/minute; (9) thrombocytopenia with platelet count <100,000 per mm³. Patients who met at least 1 major criterion and ≥3 minor criteria were diagnosed as severe pneumonia. BALF samples from patients who had failed initial anti-infective treatments were examined by TEM. Patients were excluded if they were newborns who never left the hospital or if TEM examination and serological tests were not performed. A total of 106 patients were included in this study.

Bronchoalveolar Lavage
Bronchoalveolar lavage was performed within 48 hours after patient admission by flexible bronchoscopy under local anesthesia with lidocaine in a single-center (Xiangya Hospital) setting using standard methods. Briefly, the sampling area was selected based on the infiltration location on a chest radiograph. Three 20 mL fractions of sterile saline were instilled into the relevant lobe and segment of the lung. BALF was retrieved by gentle syringe suction, put into sterile containers, and immediately submitted to the Department of Pathology for TEM examination.

TEM Examination of BALF
TEM examination of BALF was performed by the Department of Pathology, Xiangya Hospital. BALF was centrifuged at 3000 rpm for 10 minutes and the supernatant was removed. Subsequently, the pellet remaining on the bottom of the tube after centrifuge was fixed in 2.5% glutaraldehyde and 1% osmium tetroxide, dehydrated through a graded series of acetone (50%, 70%, 90% and 100%) and embedded in a mixture of epon substitute and araldite. A total of 6 thin sections (50-100 nm) from each sample were stained with 3% uranyl acetate and Reynolds’s lead citrate. Imaging was performed at 200 kV using a Hitachi H7700 Transmission Electron Microscope (Hitachi, Tokyo, Japan). Digital images of the specimens were acquired using an AMT Advantage XR 12 CCD camera (AMT, Danvers, MA) and analyzed by 2 experienced electron microscopists. In total, 48 hours are needed for TEM examination of each batch of BALF samples.

Serological Tests
Serological tests were performed by the Department of Clinical Laboratory, Xiangya Hospital. Serum samples were collected from patients before antibiotic treatment (within 24 hours after hospital admission) if feasible, depending on the patients’ situations. Serum immunoglobulin (Ig)M antibodies specific to common pathogens in the respiratory tract (including Adenovirus, Respiratory syncytial virus, Influenza B virus, Influenza A virus, Human parainfluenza viruses, M pneumoniae, C pneumoniae, L pneumophila and Coxiella burnetii) were detected using indirect immunofluorescence assay.

Statistical Analysis
The McNemar test was performed to compare the detection rates of different diagnostic methods. All statistical analyses were carried out using SPSS Statistics version 24.0 (IBM Corporation, Armonk, NY). A P value < 0.05 was considered to be statistically significant.

To ensure the quality of any reporting of the results from the present study, the recommended guidelines based on the criteria published by the Standards for Reporting of Diagnostic Accuracy initiative for the accurate reporting of investigations of diagnostic studies were followed.

RESULTS
Baseline Characteristics of Patients with Severe Pneumonia
Baseline characteristics of 106 patients included in this study were summarized in Table 1. Eighty-one of
106 (76.4%) of these patients were males, and 86 of 106 (81.1%) were older than 40 years of age. Forty-eight out of 106 patients (45.3%) did not have a smoking history. Forty-eight out of 106 (76.4%) of these patients were males, and 86 of 106 (81.1%) patients got significantly relieved and discharged from our hospital, while comprehensive therapies did not work on the remaining 18.9% of patients.

**Pathogens Detected by TEM in BALF**

Representative transmission electron micrographs images of pathogens in BALF were shown in Figure 1 and Figure S1, including virus, *chlamydia, mycoplasma*, fungus, *coccus* and *bacillus*. Generally, categories of pathogens were determined according to their sizes, shapes and characteristic structures. For example, most viruses have a diameter between 20 and 300 nm. There are 4 main morphological virus types, including helical, icosahedral, enveloped and complex symmetric viruses. *Mycoplasma* is a *mollicute* genus of bacteria that lack a cell wall around their cell membranes and have a diameter of 100-300 nm, while *chlamydia* is a genus of pathogenic bacteria that are obligate intracellular parasites, which may be found in the form of an elementary body or a reticulate body. Besides, fungi usually grow as hyphae, which are cylindrical, thread-like structures 2-10 μm in diameter and up to several centimeters in length. Fungi also disperse spores or spore-containing propagules for reproduction purpose.

In total, TEM examination of BALF revealed 24 virus-positive samples, 16 mycoplasma-positive samples, 18 *chlamydia*-positive samples, 2 fungus-positive samples and 74 bacterium-positive samples, while no pathogens in the remaining 7 samples were detected. A total of 61 samples were found to be mixed infections by TEM (Table 2). A total of 42 bacteria in 33 samples were identified by bacterial culture analyses of lower respiratory tract specimens, among which 9 samples were mixed infections (Table S1). Even though TEM showed a higher detection rate of bacteria than cultures, it could not differentiate bacterial strains and provide any information of antibiotic sensitivity, thus it had little diagnostic value in detecting bacteria in BALF.

**Comparison of Pathogen-positive Samples Detected by TEM and Serological Tests**

Nonbacterial pathogens such as viruses, *M pneumo- niae, C pneumoniae and L pneumophila* cannot be identified by routine culture method. Specific isolation and culture tests, which are labor-intensive and time-consuming, are rarely used in the hospital for etiologic diagnosis. Instead, pathogen-specific IgM antibody in serum is considered as significantly indicative and routinely measured when there is a clinical suspicion of infection with these pathogens. As shown in Table 3, TEM examination of BALF successfully confirmed most pathogen-positive samples indicated by serological tests, including 14 of 16 (87.5%) virus, 4 of 6 (66.7%) mycoplasma and 2 of 2 (100%) *chlamydia*. Strikingly, TEM also detected infectious agents in a number of samples from patients reported as negative results in serological tests (Table 3).

**Table 1. Baseline clinical characteristics of patients.**

| Clinical characteristics                  | Patients number (n = 106) |
|-------------------------------------------|--------------------------|
| Gender (%)                                |                          |
| Male                                      | 81 (76.4)                |
| Female                                    | 25 (23.6)                |
| Age (years) (%)                           |                          |
| ≤18                                       | 8 (7.5)                  |
| 19-40                                     | 12 (11.3)                |
| 41-60                                     | 44 (41.5)                |
| >60                                       | 42 (39.6)                |
| Smoking history (%)                       |                          |
| Nonsmoker                                 | 48 (45.3)                |
| Present smoker                            | 34 (32.1)                |
| Former smoker                             | 24 (22.6)                |
| Clinical manifestations (%)               |                          |
| Fever                                     | 83 (78.3)                |
| Cough                                     | 78 (73.6)                |
| Expectoration                             | 36 (34.0)                |
| Dyspnea                                   | 72 (67.9)                |
| CT findings (%)                           |                          |
| Consolidation                             | 56 (52.8)                |
| GGO                                       | 38 (35.8)                |
| Multiple nodules/masses                   | 8 (7.5)                  |
| Pleural effusion                          | 22 (20.8)                |
| ARDS (%)                                  | 18 (17.0)                |
| Comorbidities (%)                         |                          |
| Hypertension                              | 29 (27.4)                |
| Heart disease                             | 34 (32.1)                |
| Chronic renal failure                     | 36 (34.0)                |
| Diabetes mellitus                         | 14 (13.2)                |
| COPD                                      | 16 (15.1)                |
| Cerebrovascular events                    | 14 (13.2)                |
| Malignancy                                | 10 (9.4)                 |
| Hypothyroidism                            | 9 (8.5)                  |
| Immunosuppression                         | 2 (1.9)                  |
| Pre-admission antibiotics (%)             | 100 (94.3)               |
| Average hospital length of stay (days)    | 12.9                     |
| Hospital outcome (%)                      |                          |
| Treatment failure                         | 20 (18.9)                |
| Relieved and discharged                   | 86 (81.1)                |

Abbreviations: ARDS, acute respiratory distress syndrome; COPD, chronic obstructive pulmonary disease; CT, computer tomography; GGO, ground glass opacity.
FIGURE 1. Representative transmission electron micrographs of nonbacterial pathogens detected in BALF from patients with severe pneumonia. A, Coronavirus (red arrows), showing a diameter of about 100 nm and crown-like projections on the surfaces; B, mycoplasma, showing non-helical spherical cell morphology with a diameter of 200-300 nm and the absence of cell wall; C, chlamydia, showing intracellular irregular spherical cell morphology with a diameter of 0.3-0.8 μm. Both forms of chlamydia were seen, including the elementary body (smaller, orange arrow) and the reticulate body (larger, red arrow); D, fungus, showing spheroid fungal cells with a diameter of 2-5 μm, and some were dispersing spores (red arrow). Scale bars were marked in each panel.
DISCUSSION

Identification of infectious agents is challenging in patients with severe pneumonia. In the present study, we found that TEM had a good detectability of nonbacterial pathogens in BALF obtained from patients with severe pneumonia, and the implementation of TEM could significantly improve the diagnostic sensitivity of nonbacterial etiology. To the best of our knowledge, this is the first retrospective study reporting the usefulness of TEM in the etiologic diagnosis of severe pneumonia.

Severe pneumonia is life-threatening. Without any clear indications of causative agents, antibiotic treatment is usually determined empirically, which may cause severe drug resistance, higher cost and increased mortality risk. Therefore, microbiological tests are strongly recommended for patients with severe pneumonia, in whom the probability of changing the empirical treatment is high, to reduce treatment failure and prevent antibiotic overuse.10,17,18 As the gold standard of etiologic diagnosis, microbiological culture is not efficient in severe pneumonia due to its time-consuming procedures and high incidence of false negative results.19 More importantly, culture analysis is rarely used to detect viruses and atypical pathogens due to delay in the results and complicated procedures. PCR and serological tests are popular due to the high-speed and capacity of processing large quantities of samples at the same time; however, they are not suitable for detecting novel/rare pathogens or mixed infections. Therefore, additional techniques that could facilitate etiologic diagnosis of severe pneumonia are urgently needed, especially when initial empirical anti-infective therapy has failed. In the present study, our data demonstrated that TEM had a good sensitivity in detecting nonbacterial agents due to its time-consuming procedures and high incidence. In severe pneumonia, TEM examination of BALF samples only takes 48 hours, which means that clinical physicians could get fast information of what kind of infectious agents are existing in the lungs of patients. Unlike serological test and PCR, TEM does not require any organism-specific

Improved Detection Rate of Infectious Agents by Combining TEM with Serological Tests

To investigate whether the application of TEM could facilitate etiologic diagnosis of severe pneumonia, we further compared the pathogen detection rate of serological tests and TEM. It revealed that the detection rate of TEM was significantly higher than serological tests (Table 4). Furthermore, the number of samples confirmed as pathogen-positive markedly increased when combining serological tests and TEM (Table 4).

TABLE 2. Pathogens identified by TEM examination of BALF.

| Pathogens | Number of positive samples |
|-----------|---------------------------|
| Viruses   | 24                        |
| Mycoplasma| 16                        |
| Chlamydia | 12                        |
| Fungi     | 2                         |
| Bacteria  | 74                        |
| Coccili   | 45                        |
| Bacilli   | 39                        |
| Mixed infections | 61                      |

Abbreviations: BALF, bronchoalveolar lavage fluid; TEM, transmission electron microscopy.

TABLE 3. Comparison of viruses, mycoplasma or chlamydia positive samples detected by serological tests and TEM.

| Pathogens | TEM | Serum IgM Positive | Negative | Total |
|-----------|-----|--------------------|----------|-------|
| Viruses   | Positive | 14             | 2        | 16    |
|           | Negative  | 10             | 80       | 90    |
|           | Total     | 24             | 82       | 106   |
| Mycoplasma| Positive | 4              | 2        | 6     |
|           | Negative  | 12             | 88       | 100    |
|           | Total     | 16             | 90       | 106   |
| Chlamydia | Positive | 2              | 0        | 2     |
|           | Negative  | 16             | 88       | 104    |
|           | Total     | 18             | 88       | 106   |

Abbreviations: IgM, immunoglobulin M; TEM, transmission electron microscopy.

TABLE 4. Detection rates of infectious agents by combining TEM with serological test.

| Methods               | Viruses | Mycoplasma | Chlamydia |
|-----------------------|---------|------------|-----------|
| IgM assay             | 16/106  | 6/106      | 2/106     |
| TEM                   | 24/106²  | 16/106⁵  | 18/106²  |
| IgM assay + TEM       | 30/106²  | 18/106⁵  | 18/106²  |

Abbreviations: IgM, immunoglobulin M; TEM, transmission electron microscopy

² P < 0.05 vs. culture (or IgM).
reagents for detecting pathogens. However, there are several drawbacks to the TEM technique. First of all, samples must be processed individually for TEM examination, leading to a low throughput. The field of view under TEM is also relatively small, increasing the possibility that the region analyzed may not be representative of the whole sample, which could cause false negative results. In addition, the accuracy of TEM examination is dependent on the skill and experience of microscopists. Thus, in this study, 2 experienced electron microscopists were involved in imaging analysis of each sample to avoid misdiagnosis. Furthermore, the diagnostic value of TEM in bacterial pneumonia is limited, because routine TEM cannot differentiate bacterial strains and provide any information of antibiotic sensitivity, thus, had few benefits to subsequent antibiotic therapies. So far, advanced electron microscopy (EM) techniques have been applied to microbiology in basic and clinical studies, including cryogenic electron tomography, immuno-EM, 3-dimensional EM, and correlated light microscopy and TEM. When TEM is integrated with immunohistochemistry or other molecular labeling techniques, it would become practical for electron microscopists and clinical physicians to distinguish various pathogens on the strain level under TEM. Recently, rapid methods of sample preparation and transportable EM equipment, which makes EM an easier and faster tool to use, have also been reported to improve the identification of infectious agents. These studies and our data collectively support that in future, TEM may be used along with other diagnostic tools, as one part of a comprehensive assay system for microbiological diagnosis of severe pneumonia.

Several limitations should be noted in the present study. First of all, no BALF samples from healthy volunteer subjects were obtained, which made the diagnostic specificity of TEM unevaluable and the interpretation of TEM examination difficult, which we plan to address in future studies. In addition, the major disadvantage of TEM used in this study was that it did not identify a pathogen on the strain level. Hopefully, the application of more advanced techniques and combined use with other microbial tests would enhance the diagnostic accuracy of TEM, thus further improving its diagnostic value. Moreover, even though TEM examination of BALF appears to have a potential utility in improving diagnostic sensitivity in severe pneumonia caused by nonbacterial pathogens, its cost-effectiveness is still unknown. Therefore, clinical trials with large patient numbers must be conducted to determine whether the implementation of TEM could benefit patients with severe pneumonia through promoting the prompt initiation of anti-infective treatments.

CONCLUSIONS

In summary, our data supported that TEM had a good sensitivity in detecting nonbacterial pathogens in BALF, including viruses, atypical pathogens and mixed infections. The combined use of serological tests and TEM could improve the detection rate of nonbacterial infectious agents in patients with severe pneumonia. Considering the severity and rapid progression of severe pneumonia, a single diagnostic method is not recommended in such a case. Implementation of TEM would be a judicious strategy for the etiologic diagnosis of severe pneumonia caused by nonbacterial pathogens in the future.

SUPPLEMENTARY MATERIALS

Supplementary material associated with this article can be found, in the online version, at https://doi.org/10.1016/j.amjms.2018.11.012.

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