LABORATORY TEST DIRECTORY

Pneumonia Panel by PCR

Last Literature Review: June 2023 Last Update: July 2023

Pneumonia is an infection of the lungs that may be caused by a variety of different bacterial, fungal, and viral pathogens and can affect patients with normally functioning immune systems as well as those with underlying immune compromise. Pneumonia is a leading cause of morbidity and mortality worldwide in individuals of all ages. In hospitalized patients with severe community-acquired pneumonia, multiplex molecular methods such as polymerase chain reaction (PCR) from lower respiratory tract specimens may be appropriate to determine etiology and inform treatment.¹

This panel detects the presence of 18 bacterial respiratory pathogens, seven antimicrobial resistance markers, and eight respiratory viruses.

² Rapid diagnosis is critical to achieving positive patient outcomes.

Test Interpretation

Sensitivity and Specificity

Sensitivity/specificity versus culture or PCR predicate methods (per FDA clinical trials data):

- · Sensitivity: 96.2%
- · Specificity: 98.3%

Targets Detected

Bacterial targets

- Acinetobacter calcoaceticus / baumannii complex
- Chlamydia pneumoniae^a
- Enterobacter cloacae complex
- Escherichia coli
- Haemophilus influenzae
- Klebsiella aerogenes
- Klebsiella oxytoca
- Klebsiella pneumoniae group
- Legionella pneumophila^a
- Mycoplasma pneumoniae^a
- Moraxella catarrhalis
- Proteus spp.
- Pseudomonas aeruginosa
- · Serratia marcescens
- Staphylococcus aureus
- Streptococcus agalactiae
- Streptococcus pneumoniae
- Streptococcus pyogenes

Antimicrobial resistance markers

- Carbapenemases (conveys resistance against 1 or more carbapenem antibiotics)
 - o IMP
 - o KPC
 - o NDM
 - o Oxa-48-like
 - o VIM

ESBL, extended spectrum beta-lactamase; MREJ, mec right-extremity junction; MRSA, methicillin-resistant Staphylococcus aureus

Featured ARUP Testing

Pneumonia Panel by PCR 3016457

Method: Semi-Quantitative Polymerase Chain Reaction (PCR)/Qualitative Polymerase Chain Reaction (PCR)

Aids in the diagnosis of bacterial and viral pneumonia from lower respiratory tract specimens. For use in individuals with clinically compatible signs and symptoms.

^aTesting for this bacterium is qualitative. Testing for all other targeted bacteria is semiquantitative.

- ESBL (conveys resistance to most penicillins, cephalosporins, and aztreonam)
 - o CTX-M
- MRSA (conveys resistance to methicillin, penicillin, and penicillin-like drugs)
 - mecA/mecC and MREJ

Viral Targets

- Adenovirus
- Coronavirus
- · Human metapneumovirus
- Human rhinovirus/enterovirus
- Influenza A
- Influenza B
- Parainfluenza virus
- Respiratory syncytial virus

ESBL, extended spectrum beta-lactamase; MREJ, mec right-extremity junction; MRSA, methicillin-resistant Staphylococcus aureus

Results

This panel detects the presence of 18 bacterial respiratory pathogens, seven antimicrobial resistance markers, and eight respiratory viruses. Bacterial targets are reported with a relative abundance, akin to conventional culture and/or Gram stain reporting (eg, rare, 1+, 2+, 3+).

Limitations

- This panel is not a replacement for traditional microbiologic testing assays and is instead intended to be a complementary method with a faster turnaround time and higher sensitivity.
- This panel detects only the most common causes of pneumonia. It is not intended as a replacement for respiratory culture, blood cultures, or urine antigen testing for Legionella or Streptococcus pneumoniae. Refer to the Laboratory Test Directory for additional test offerings.
- · When ordering this test, a paired respiratory culture is required.
- · Fungal pathogens are not detected by this panel and should be tested as appropriate using conventional methods.
- Bacterial semiquantitative values by PCR may not directly correlate with culture due to organism viability in culture (eg, as a result of empiric antibiotics prior to specimen collection).
- Bacteria may grow in culture in very low quantities that are not reported by this panel due to predetermined cutoff thresholds. These scenarios should not be considered discrepant results.
- SARS-CoV-2 (COVID-19) is not detected by this panel; if indicated, order a separate COVID-19 test.

References

- 1. Martin-Loeches I, Torres A, Nagavci B, et al. ERS/ESICM/ESCMID/ALAT guidelines for the management of severe community-acquired pneumonia. Intensive Care Med. 2023:1-18.
- 2. Evans SE, Jennerich AL, Azar MM, et al. Nucleic acid-based testing for noninfluenza viral pathogens in adults with suspected community-acquired pneumonia. An official American Thoracic Society Clinical Practice Guideline. Am J Respir Crit Care Med. 2021;203(9):1070-1087.

Related Information

Community-Acquired Pneumonia - CAP Respiratory Viruses

^aTesting for this bacterium is qualitative. Testing for all other targeted bacteria is semiquantitative.