

Respiratory Pathogens

Our respiratory pathogen testing utilizes quantitative Real-Time PCR to rapidly analyze your patient's sample in 24 hours. RT-PCR technology precisely detects the correct pathogen(s) and identifies antibiotic drug resistance. This allows providers the ability to prescribe timely and effective treatment.

Rapid and accurate solution eliminates guesswork in diagnosing and treating upper respiratory infections

Acute respiratory infection is a significant cause of morbidity and mortality in young, geriatric and immunocompromised patients. Co-infection is also high within these populations and access to advanced technology is essential to detect multiple pathogens at once. False negative test results can lead to a delayed diagnosis and poor clinical outcomes.

Molecular diagnostic testing quickly identifies pathogens and detects potential antibiotic resistance, so effective treatment can begin sooner.

Accurate diagnosis within 24 hours with real-time PCR for pathogen identification and antibiotic resistance detection

- PCR, a molecular technique, can be used to precisely analyze the genetic material of pathogens
- Provides a more definitive diagnosis than POC antigen assays
- 24-hour turnaround from receipt of specimen
- More accurate than conventional culture¹
- CAP and CLIA accredited

Helps improve clinical confidence, decrease patient risks

- Detects polymicrobial infections
- Unaffected by concurrent antibiotic use
- Identifies potential antibiotic resistance
- Aids in quick clinical decision-making
- Reduces false negative results
- Aids in antibiotic stewardship
- Reduces potential unnecessary drug exposure and adverse events

1. Rhoads, D., Wolcott, R., Sun, Y., Dowd, S. (23 February 2012). Comparison of culture and molecular identification of bacteria in chronic wounds. *Int. J. Mol. Sci.*, 13, 2535-2550. Retrieved from www.mdpi.com/journal/ijms

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Acinetobacter baumannii	Human metapneumovirus
Adenovirus Respiratory	Influenza virus A, B
HAdV-B	Klebsiella pneumoniae
Aspergillus fumigatus, niger	Legionella pneumophila
Bordetella pertussis ,	Moraxella catarrhalis
parapertusis, bronchiseptica	Mycobacterium tuberculosis
Chlamydia trachomatis	and avium/intracellulare,
Chlamydophila pneumoniae	kansasii groups
Citrobacter freundii	Mycoplasma pneumoniae
Coccidioides immitis,	Parainfluenza virus (types
posadasii	1,2,3,4)
Coronavirus - 229E, NL63,	Proteus mirabilis
OC43, and HKU1	Pseudomonas aeruginosa
Cytomegalovirus (CMV,	Respiratory syncytial virus
Human Herpes Virus 5)	Rhinovirus A, B, C
E. coli	Serratia marcescens
Enterovirus A + B probes	Staphylococcus aureus
Enterovirus C + D probes	Streptococcus agalactiae
Haemophilus influenzae	(group B strep. (GBS))
Herpes simplex virus 1 and 2	Streptococcus pneumoniae
(HSV-1, HSV-2)	Streptococcus pyogenes
Human herpes virus 3	
(Varicella zoster virus)	

Antibiotic Resistance

VanA, VanB (Vancomycin Resistance genes)	IMP, NDM, VIM Groups (Class B metallo beta lactamase)
mecA (Methicillin resistance gene)	ACT, MIR, FOX, ACC Groups (AmpC beta lactamase)
ermB, C; mefA (Macrolide Lincosamide Streptogramin Resistance)	OXA-48,-51 (Class D oxacillinase)
qnrA2 (Fluoroquinolone resistance genes)	PER-1/VEB-1/GES-1 Groups (Minor Extended Spectrum beta lactamases)
tet M (Tetracycline resistance genes)	dfr (A1, A5), sul (1, 2) probes (Trimethoprim/Sulfamethoxazole resistance)
SHV, KPC Groups (Class A beta lactamase)	
CTX-M1 (15), M2 (2), M9 (9), M8/25 Groups (Class A beta lactamase)	