QIAstat-Dx Respiratory SARS-CoV-2 Panel

For quick and accurate detection of respiratory infections

Influenza-like illnesses aren't just caused by influenza viruses. There are many possible bacterial and viral causes that result in similar, overlapping symptoms. As a result, diagnosing respiratory infections using traditional methods can be slow and challenging. Imagine getting 21 results in about an hour. Powered by our easy workflow and reliable multiplex PCR technology, the QIAstat-Dx Respiratory SARS-CoV-2 Panel makes this possible.

- **Get comprehensive results fast**
  - Detect and differentiate 21 pathogen targets in about an hour
  - Support timely clinical decision-making and rapid therapeutic intervention

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- **Set up your tests quickly and easily**
  - All-in-one assay cartridges require minimal hands-on time
  - Just 300 µL liquid transport medium with no precision pipetting required

- **Receive supportive information**
  - Easily access Ct values and amplification curves for all detected pathogens

Comprehensive pathogen coverage with QIAstat-Dx

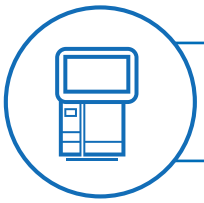
Viral

- SARS-CoV-2
- Influenza A
- Influenza A subtype H1N1/2009
- Influenza A subtype H1
- Influenza A subtype H3
- Influenza B
- Coronavirus 229E
- Coronavirus HKU1
- Coronavirus NL63
- Coronavirus OC43
- Parainfluenza virus 1
- Parainfluenza virus 2
- Parainfluenza virus 3
- Parainfluenza virus 4
- Adenovirus
- Respiratory syncytial virus A/B
- Human metapneumovirus A/B
- Rhinovirus/Enterovirus*

Bacterial

- *Mycoplasma pneumoniae*
- *Bordetella pertussis*
- *Chlamydophila pneumoniae*

*Enterovirus and Rhinovirus are both detected, but not differentiated, with the QIAstat-Dx Respiratory SARS-CoV-2 Panel.



QIAstat-Dx Respiratory SARS-CoV-2 Panel runs on our modular **QIAstat-Dx Analyzer**.

Ordering Information - Assay

Product	Contents	Cat. no.
QIAstat-Dx Respiratory SARS-CoV-2 Panel	Six individually packaged cartridges containing all reagents needed for sample preparation and multiplex RT-real time PCR plus internal control, including six transfer pipettes	691223

The QIAstat-Dx Respiratory SARS-CoV-2 Panel is intended for in vitro diagnostic use.

- This product has not been FDA cleared or approved, but has been authorized for emergency use by FDA under an EUA for use by authorized laboratories;
- This product has been authorized only for the detection and differentiation of nucleic acid of SARS-CoV-2 from multiple respiratory viral and bacterial organisms; and
- This emergency use of this product is only authorized for the duration of the declaration that circumstances exist justifying the authorization of emergency use of in vitro diagnostics for detection and/or diagnosis of COVID-19 under Section 564(b)(1) of the Federal Food, Drug, and Cosmetic Act, 21 U.S.C. § 360bbb3(b)(1), unless the declaration is terminated or authorization is revoked sooner.

Confidently detect respiratory infections with QIAstat-Dx – learn more at **QIAGEN.com/QIAstat-Dx**

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