COVID-19 Employee Testing Program

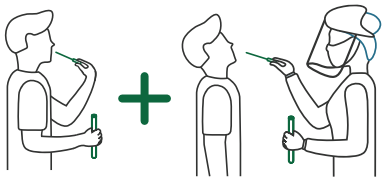
PathGroup
Physician Centered. Patient Focused.

Excellence. Accuracy. Reliability. Superior Service.

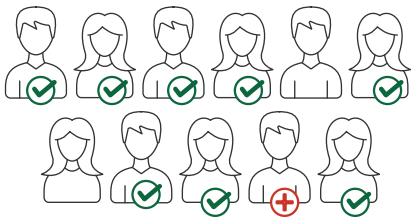
These are the values that PathGroup was founded upon over 50 years ago in Nashville, Tennessee. Consistent and unparalleled reliability in laboratory diagnostics and superior, personalized customer service remain the basis of everything we do today. While working together with our partners to confront the COVID-19 pandemic, we bring these core values to everyone we serve.

The PathGroup Turnkey Employee Testing Program:

As you focus on a safe return to work for your employees, PathGroup's turnkey Employee Testing Program can assist with COVID-19 diagnostic and antibody testing to support your organization. Customizable for your company, with self-collection options and rapid turnaround time for results, PathGroup's Employee Testing Program can help you quickly and **safely get your employees back to work.**



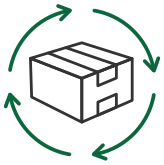
Self-collection* and on-site collection options available



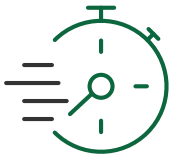
Ongoing on-demand or surveillance testing of employees throughout the year

All testing supplies needed for collection, including nasal swabs and serum tubes

Customizable reporting available to meet your organization's needs



Transportation/logistics via PathGroup couriers to our centralized laboratory in Nashville, TN



Rapid turnaround time for results

SARS-CoV-2, RNA (COVID-19) Molecular (Diagnostic) Testing

- Qualitative detection of nucleic acid from SARS-CoV-2
- SARS-CoV-2 assays utilize Transcription Mediated Amplification (TMA) or real-time RT-PCR technologies
- Nasal, nasopharyngeal (NP) or oropharyngeal (OP) collection
- Swabs and transport media collection kits provided for testing

SARS-CoV-2 (COVID-19) Serology (Antibody) Testing

- Qualitative detection of total antibodies (including IgG, IgA and IgM) to SARS-CoV-2 in serum for individuals with prior COVID-19 infection
- Intended for use as an aid in identifying patients with an adaptive immune response to SARS-CoV-2, indicating recent or prior infection
- Serum collection in standard Serum Separator Tube (SST)

*Self-collection – available by August 1st

To Learn More About COVID-19 Testing at PathGroup, contact your PathGroup representative or call **1.888.410.4618** or email **COVID19Testing@pathgroup.com**

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06.2020

How PathGroup Delivers the COVID-19 Answers You Need:

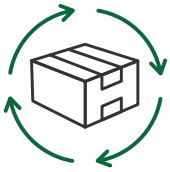
Immediate capacity available for both **molecular (diagnostic) and serology (antibody) COVID-19 testing**

FDA Emergency Use Authorization (EUA)

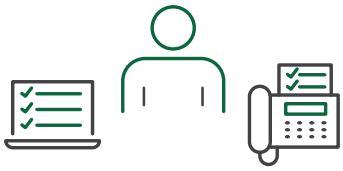
Roche HOLOGIC®

High-throughput testing platforms from **Roche and Hologic***

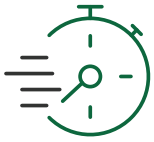
All collection supplies provided (swabs, serum tubes)



Comprehensive logistics solutions for specimen pickup and delivery to PathGroup



Ordering and results via **interface, online portal or manual requisitions/fax**



Rapid turnaround time for results

The PathGroup Difference:

Physician leadership and superior service are the PathGroup Difference

Physician leadership guides every decision we make. For COVID-19, PathGroup has chosen only market-leading FDA EUA testing platforms that share our commitment to quality, accuracy, and reliability. You can trust the results you receive from PathGroup.

Superior service means PathGroup provides individualized service to every client. With the urgency of COVID-19 testing and results, we continue our long-standing commitment to providing dedicated account management and personal client service resources to your organization to answer any questions or needs that arise.

About PathGroup:

Founded in 1965 in Nashville, TN, PathGroup is a premier provider of anatomic, clinical, and molecular pathology laboratory services in the United States. The company continues to be privately held and physician-led, with over 175 pathologists on staff. PathGroup has expanded to meet the needs of hospitals, physicians, and other healthcare providers during its 50+ years of operation and now serves clients in 25 states, including over 100 hospitals and over 15,000 physicians.

FDA EUA disclaimer:
These tests have not been FDA cleared or approved. These tests have been authorized by FDA under an EUA for use by authorized laboratories. These tests have been authorized only for the detection of nucleic acid from SARS-CoV-2, not for any other viruses or pathogens. These tests are only authorized for the duration of the declaration that circumstances exist justifying the authorization of emergency use of in vitro diagnostic tests for detection and/or diagnosis of COVID-19 under Section 564(b)(1) of the Act, 21 U.S.C. §360bbb-3(b)(1), unless the authorization is terminated or revoked sooner.

Antibody cross-reaction disclaimer:
This test has not been reviewed by the FDA. Negative results do not rule out SARS-CoV-2 infection, particularly in those who have been in contact with the virus. Follow-up testing with a molecular diagnostic should be considered to rule out infection in these individuals.
Results from antibody testing should not be used as the sole basis to diagnose or exclude SARS-CoV-2 infection or to inform infection status. Positive results may be due to past or present infection with non–SARS-CoV-2 coronavirus strains, such as HKU1, NL63, OC43, or 229E.

Disclaimer:
* ROCHE is a registered trademark of Hoffman-La Roche Inc.
Hologic is a registered trademark of Hologic, Inc

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