NEWS RELEASE

Quest Diagnostics to Speak at the Jefferies Virtual Healthcare Conference

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SECAUCUS, N.J., May 29, 2020 /PRNewswire/ -- Quest Diagnostics Incorporated (NYSE: DGX), the world's leading provider of diagnostic information services, announced that it is scheduled to speak at the Jefferies Virtual Healthcare Conference. Mark Guinan, Executive Vice President and CFO and Jim Davis, Executive Vice President, General Diagnostics, will discuss the company's vision, goals and two-point strategy to accelerate growth and drive operational excellence, and the company's current perspective on the impact of the COVID-19 pandemic. The presentation is scheduled for Wednesday, June 3, 2020 at 1:30 p.m. Eastern Time.

The presentation will be webcast live during the conference and will be available on the company's investor relations page which can be accessed at **ir.QuestDiagnostics.com**. In addition, the archived webcast will be available within 24 hours after the conclusion of the live event and will remain available until July 3, 2020.

About Quest Diagnostics

Quest Diagnostics empowers people to take action to improve health outcomes. Derived from the world's largest database of clinical lab results, our diagnostic insights reveal new avenues to identify and treat disease, inspire healthy behaviors and improve health care management. Quest Diagnostics annually serves one in three adult Americans and half the physicians and hospitals in the United States, and our 47,000 employees understand that, in the right hands and with the right context, our diagnostic insights can inspire actions that transform lives. **www.QuestDiagnostics.com**.

About COVID-19 testing at Quest Diagnostics

Quest Diagnostics is at the forefront of the response to the COVID-19 pandemic, working to broaden access to laboratory insights to help us all lead healthier lives. We provide both molecular diagnostic and antibody serology tests to aid in the diagnosis of COVID-19 and immune response. Our COVID-19 test services are

based on tests that have received FDA emergency use authorization and which also meet our high standards for quality. We are providing these test services under the Public Readiness and Emergency Preparedness Act. We provide data on COVID-19 testing to various federal and state public health authorities, including the Centers for Disease Control and Prevention, and participate in studies with government and private institutions, aiding COVID-19 public health response and research. Through our team of dedicated phlebotomists, air fleet team, couriers and laboratory professionals, Quest Diagnostics works hard every day to help patients and communities across the United States access quality COVID-19 testing.

The antibody tests and the molecular tests (together "All tests") have not been FDA cleared or approved; All tests have been authorized by FDA under EUAs for use by authorized laboratories; The antibody tests have been authorized only for the detection of IgG antibodies against SARS-CoV-2, not for any other viruses or pathogens; The molecular tests have been authorized only for the detection of nucleic acid from SARSCoV-2, not for any other viruses or pathogens; and, All tests are 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 Act, 21 U.S.C. § 360bbb-3(b)(1), unless the authorization is terminated or revoked sooner.

For more information about the latest developments with our COVID-19 testing, visit: **newsroom.questdiagnostics.com/COVIDTestingUpdates**

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