October 9, 2015

Myriad to Present New Data on Its myRisk(TM) Hereditary Cancer Test at the CGA-ICC Annual Meeting

Company to Highlight Data for Expanding Indications and Variant Classification for Hereditary Colon Cancer

SALT LAKE CITY, Oct. 9, 2015 (GLOBE NEWSWIRE) -- Myriad Genetics, Inc. (NASDAQ:MYGN), a leader in molecular diagnostics and personalized medicine, today announced that it will present new data on its myRisk™ Hereditary Cancer molecular diagnostic test at the 18th Annual Meeting of the Collaborative Group of the Americas - Inherited Colorectal Cancer (CGA-ICC) being held Oct. 11 to 12, 2015 in Baltimore, Md.

"Myriad is committed to improving the care of patients and families with inherited colorectal cancer syndromes," said Johnathan Lancaster, M.D., Ph.D., chief medical officer, Myriad Genetic Laboratories. "These syndromes are under-diagnosed and our new data using the myRisk Hereditary Cancer gene panel test highlight how next generation sequencing technology can identify more patients at elevated risk of hereditary colon cancer. This information empowers patients and their physicians to take steps that may reduce the risk of a cancer developing. Our recent studies focus on expanding our understanding of the gene mutations associated with colorectal cancer syndromes, and we believe that these data provide further evidence that testing guidelines need to be revised to ensure that patients continue to have access to advances in sequencing technology that may be life-saving."

A list of the Myriad presentations at CGA-ICC (#CGA2015) follows:

Podium Presentations

- Title: Pan-cancer gene panel results for patients with > 5 adenomas.

  Date: Monday, Oct. 12, 2015, 9:00 a.m. ET.

- Title: Ohio Colorectal Cancer Prevention Initiative: Germline mutation spectrum in 250 colorectal cancer patients diagnosed under age 50.

  Date: Monday, Oct. 12, 2015: 9:10 a.m. ET.

Poster Presentation

- Title: Clinical Presentations of Patients and Families Identified with Pathogenic Variants in *CDH1*.

  Date: Monday, Oct. 12, 2015: 7:30 a.m. ET.

For more information about the meeting, please visit the CGA website at: http://www.cgaicc.com/Events/event_landing.cfm.

About Myriad myRisk™ Hereditary Cancer Testing

The Myriad myRisk Hereditary Cancer test uses next-generation sequencing technology to evaluate 25 clinically significant genes associated with eight hereditary cancer sites including: breast, colon, ovarian, endometrial, pancreatic, prostate and gastric cancers and melanoma. For more information visit: https://www.myriad.com/products-services/hereditary-cancers/myrisk-hereditary-cancer/.

About Myriad Genetics

Myriad Genetics Inc., is a leading personalized medicine company dedicated to being a trusted advisor transforming patient lives worldwide with pioneering molecular diagnostics. Myriad discovers and commercializes molecular diagnostic tests that: determine the risk of developing disease, accurately diagnose disease, assess the risk of disease progression, and guide

treatment decisions across six major medical specialties where molecular diagnostics can significantly improve patient care and lower healthcare costs. Myriad is focused on three strategic imperatives: transitioning and expanding its hereditary cancer testing markets, diversifying its product portfolio through the introduction of new products and increasing the revenue contribution from international markets. For more information on how Myriad is making a difference, please visit the Company's website: www.myriad.com.

Safe Harbor Statement

This press release contains "forward-looking statements" within the meaning of the Private Securities Litigation Reform Act of 1995, including statements related to presenting new data at CGA-ICC; advancing hereditary cancer testing or improving patient care; helping patients and their families understand their risk of developing colorectal cancer; and the Company's strategic directives under the caption "About Myriad Genetics." These "forward-looking statements" are based on management's current expectations of future events and are subject to a number of risks and uncertainties that could cause actual results to differ materially and adversely from those set forth in or implied by forward-looking statements. These risks and uncertainties include, but are not limited to: the risk that sales and profit margins of our molecular diagnostic tests and pharmaceutical and clinical services may decline; risks related to our ability to transition from our existing product portfolio to our new tests, including unexpected costs and delays; risks related to decisions or changes in governmental or private insurers' reimbursement levels for our tests or our ability to obtain reimbursement for our new tests at comparable levels to our existing tests; risks related to increased competition and the development of new competing tests and services; the risk that we may be unable to develop or achieve commercial success for additional molecular diagnostic tests and pharmaceutical and clinical services in a timely manner, or at all; the risk that we may not successfully develop new markets for our molecular diagnostic tests and pharmaceutical and clinical services, including our ability to successfully generate revenue outside the United States; the risk that licenses to the technology underlying our molecular diagnostic tests and pharmaceutical and clinical services and any future tests and services are terminated or cannot be maintained on satisfactory terms; risks related to delays or other problems with operating our laboratory testing facilities and our healthcare clinic; risks related to public concern over genetic testing in general or our tests in particular; risks related to regulatory requirements or enforcement in the United States and foreign countries and changes in the structure of the healthcare system or healthcare payment systems; risks related to our ability to obtain new corporate collaborations or licenses and acquire new technologies or businesses on satisfactory terms, if at all; risks related to our ability to successfully integrate and derive benefits from any technologies or businesses that we license or acquire; risks related to our projections about our business, results of operations and financial condition; risks related to the potential market opportunity for our products and services; the risk that we or our licensors may be unable to protect or that third parties will infringe the proprietary technologies underlying our tests; the risk of patent-infringement claims or challenges to the validity of our patents or other intellectual property; risks related to changes in intellectual property laws covering our molecular diagnostic tests and pharmaceutical and clinical services and patents or enforcement in the United States and foreign countries, such as the Supreme Court decision in the lawsuit brought against us by the Association for Molecular Pathology et al; risks of new, changing and competitive technologies and regulations in the United States and internationally; and other factors discussed under the heading "Risk Factors" contained in Item 1A of our most recent Annual Report on Form 10-K for the fiscal year ended June 30, 2015, which has been filed with the Securities and Exchange Commission, as well as any updates to those risk factors filed from time to time in our Quarterly Reports on Form 10-Q or Current Reports on Form 8-K. All information in this press release is as of the date of the release, and Myriad undertakes no duty to update this information unless required by law.

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