September 26, 2016

Myriad to Present New myRisk Hereditary Cancer Data Further Demonstrating the Benefits of Gene Panel Testing

Three Studies to Be Featured at the NSGC 35th Annual Education Conference

SALT LAKE CITY, Sept. 26, 2016 (GLOBE NEWSWIRE) -- Myriad Genetics, Inc. (NASDAQ:MYGN), a leader in molecular diagnostics and personalized medicine, today announced it will present three new studies at the National Society of Genetic Counselors Annual Education Conference being held Sept. 28-Oct. 1, 2016 in Seattle, Washington.

"We are excited to be co-sponsoring the 35th National Society of Genetic Counselors Annual Education Conference," said Susan Manley, L.C.G.C., M.B.A., vice president of Medical Services, Myriad Genetic Laboratories. "We're looking forward to presenting important new scientific data, which will improve the practice of genetic counseling. We also are excited to host a patient advocacy event in conjunction with the Conference, which will help raise funds for important patient advocacy groups including Be the Difference Foundation, Hereditary Colon Cancer Foundation and Li-Fraumeni Syndrome Association."

Below is a list of the featured presentations at NSGC (#NSGC16):

Poster Presentations

- **Title:** Average Age of Diagnosis of Ovarian Cancer for Women with Pathogenic Variants in BRIP1, RAD51C and RAD51D.
  Presenter: Susana San Roman.
  Date: Friday, September 30, 2016: 11:30 am — 12:45 pm PT.
  Poster: C-117.

- **Title:** Ancestry-Based Cancer Risks Associated with APC I1307K.
  Presenter: Lavania Sharma.
  Date: Friday, September 30, 2016: 11:30 am — 12:45 pm PT.
  Poster: C-123.

- **Title:** Psychosocial Outcomes of Identifying High or Moderate Risk Mutation Carriers by Hereditary Cancer Panel Testing.
  Presenter: Julie Culver.
  Date: Thursday, September 29, 2016: 5:45 pm — 7:00 pm PT.
  Poster: B-71.

For more information about these presentations, including a complete list of abstracts and presentations, please visit the NSGC website at http://nsgc.org/p/cm/ld/fid=423.

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 relating to the Company presenting three new studies at the National Society of Genetic Counselors Annual Education Conference being held Sept. 28-Oct. 1, 2016 in Seattle, Washington; presenting important new scientific data, which will improve the practice of genetic counseling; hosting a patient advocacy event in conjunction with the Conference, which will help raise funds for important patient advocacy groups including Be the Difference Foundation, Hereditary Colon Cancer Foundation and Li-Fraumeni Syndrome Association; 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 described or implied in the forward-looking statements. These risks and uncertainties include, but are not limited to: the risk that sales and profit margins of our existing molecular diagnostic tests and pharmaceutical and clinical services may decline or will not continue to increase at historical rates; risks related to our ability to transition from our existing product portfolio to our new tests, including unexpected costs and delays; risks related to changes in the 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 tests and any future tests are terminated or cannot be maintained on satisfactory terms; risks related to delays or other problems with operating our laboratory testing facilities; risks related to public concern over our 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, including but not limited to our acquisition of a healthcare clinic in Germany and Sividon Diagnostic and our planned acquisition of Assurex Health; risks related to our projections about our business, results of operations and financial condition; risks related to the potential market opportunity for our products; 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; 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 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.

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