



Myriad Genetics to Present Prostate Cancer Data at the ASCO-NCI-EORTC Annual Meeting on Molecular Markers in Cancer

SALT LAKE CITY, Oct 14, 2010 (GlobeNewswire via COMTEX News Network) -- Myriad Genetics, Inc. (Nasdaq:MYGN) announced today that Steve Stone, Ph.D., Vice President of Cancer Genomics, is scheduled to present data on PROLARIS (TM), Myriad's prognostic medicine product for prostate cancer, at the ASCO-NCI-EORTC Annual Meeting on Molecular Markers in Cancer to be held from October 18-20, 2010, at the Westin Diplomat Hotel in Hollywood, Florida.

The presentation entitled: "Use of cell cycle progression genes to differentiate indolent from aggressive prostate cancer" will be given on October 19 from 7:30 - 10:45 am Eastern at the Main Meeting General Session. The abstract will be available and accessible on ASCO's website, www.ASCO.org, at 6:00 pm Eastern on October 14, 2010.

About Myriad Genetics

Myriad Genetics, Inc. is a leading molecular diagnostic company focused on developing and marketing novel predictive medicine, personalized medicine and prognostic medicine products. Myriad's news and other information are available on the Company's Web site at www.myriad.com.

The Myriad Genetics, Inc. logo is available at <http://www.globenewswire.com/newsroom/prs/?pkgid=6336>

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This press release contains "forward-looking statements" within the meaning of the Private Securities Litigation Reform Act of 1995, including statements relating to the scheduled presentation of PROLARIS data at the ASCO-NCI-EORTC Annual Meeting on Molecular Markers in Cancer on October 19, 2010. 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 existing molecular diagnostic products may decline or will not continue to increase at historical rates; the risk that we may be unable to develop or achieve commercial success for additional molecular diagnostic products in a timely manner, or at all; the risk that licenses to the technology underlying our molecular diagnostic products and any future products are terminated or cannot be maintained on satisfactory terms; risks related to delays or other problems with manufacturing our products or operating our laboratory testing facilities; risks related to public concern over our products; risks related to regulatory requirements or enforcement in the United States and foreign countries and changes in the structure of healthcare payment systems; uncertainties about our ability to obtain new corporate collaborations and acquire new technologies on satisfactory terms, if at all; the development of competing products and services; the risk that we or our licensors may be unable to protect the proprietary technologies underlying our products; the risk of patent-infringement claims or challenges of our patents; 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 in our Annual Report on Form 10-K for the year ended June 30, 2010, 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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