NEWS RELEASE **REGULATED INFORMATION**

MDxHealth Licensee Exact Sciences Receives FDA Panel Recommendation for Approval of Colorectal Cancer Screening Test

MDxHealth's Epigenetic Biomarker Used in Cologuard Test

IRVINE, CA, and HERSTAL, BELGIUM – March 28, 2014 – MDxHealth SA (NYSE Euronext: MDXH), a leading molecular diagnostic company that develops and commercializes epigenetic tests to improve the diagnosis and treatment of cancer patients, today announced that the Molecular and Clinical Genetics Panel of the United States Food and Drug Administration's (FDA) Medical Devices Advisory Committee strongly endorsed approval of Cologuard™. The Committee determined by a unanimous vote of 10 to zero that Exact Sciences has demonstrated safety, effectiveness and a favorable risk benefit profile of Cologuard, the company's stool-based DNA (sDNA), non-invasive colorectal cancer screening test. Cologuard incorporates one of MDxHealth's epigenetic biomarkers and methylation specific PCR (MSP) technology, which was licensed to Exact Sciences in July 2010.

"Colorectal cancer is the second leading cause of cancer death in the United States yet often viewed as the most preventable," stated Dr. Jan Groen, CEO of MDxHealth. "We are very proud to see that one of our epigenetic biomarkers is a key component of the biomarker panel incorporated in the Cologuard screening test."

Cologuard is designed to detect specific changes in a patient's DNA that appear in the stool, which could indicate the presence of colorectal cancer or pre-cancerous polyps. The test also identifies the presence of blood in the stool, another indicator of possible colorectal cancer.

"We are pleased the Committee strongly supported Cologuard's approval," said Kevin T. Conroy, chairman and chief executive of Exact Sciences. "We look forward to continuing our work with the FDA to complete its review of Cologuard and remain committed to addressing the growing unmet needs in colorectal cancer screening. We thank the FDA and its advisory committee for its careful consideration of Cologuard. We also appreciate the opportunity to participate in the innovative FDA/CMS parallel review program."

Cologuard is an investigational device currently under review by the U.S. Food and Drug Administration (FDA) and is not available for sale in the United States. The Committee's recommendation will be taken into consideration by the FDA in its final review of the Cologuard test. The FDA is not bound by the Committee's guidance but often follows this expert guidance.

About MDxHealth

MDxHealth is a molecular diagnostics company that develops and commercializes advanced epigenetic tests for cancer assessment and the personalized treatment of patients. The company's first commercial product, ConfirmMDx® for Prostate Cancer, has been validated to help distinguish patients who have a true-negative biopsy from those who may have undetected cancer, thereby aiding in the reduction of unnecessary repeat

biopsies. MDxHealth helps to address a large and growing unmet medical need for better cancer diagnosis and treatment information. The company has a proprietary platform and a strong epigenetic product pipeline focused on the development of products for prostate, brain, bladder and lung cancers. For more information visit www.mdxhealth.com and follow us on Twitter at: twitter.com/mdxhealth.

For more information:

Dr. Jan Groen, CEO
MDxHealth
US: +1 949 812 6979
BE: +32 4 364 20 70
info@mdxhealth.com

Mike Sinclair
Halsin Partners
UK: +44 20 7318 2955
Cell: +44 7968 022075
msinclair@halsin.com

Len Hall
Allen & Caron, Inc
US: +1 949 474 4300
len@allencaron.com

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