



Acutis Diagnostics Leverages SOPHiA GENETICS Technology to Create New Next Generation Sequencing Test

The New Assay Will Screen Patients for Clinical Trials and Help Increase Data Analysis for Clinical Researchers

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BOSTON & LAUSANNE, Switzerland--([BUSINESS WIRE](#))--SOPHiA GENETICS (Nasdaq: SOPH), a cloud-native software company in the healthcare space and a leader in data-driven medicine, today announced that Acutis Diagnostics, a specialized medical laboratory, will use SOPHiA DDM™ to develop a new genomic assay. This new next-generation sequencing (NGS) test will build on Acutis Diagnostics' track record of approved lab-developed tests that support clinical trials, cancer research and treatment.

To create the new assay, Acutis Diagnostics will pair [SOPHiA DDM™ for Hereditary Cancers Solution](#) with their laboratory analysis technologies. The test is in support of genomic screening for patient enrollment in oncology clinical trials and for retrospective analyses to help explain clinical outcomes and drug target discovery.

"The use of SOPHiA DDM™ will help us quickly launch this new assay to aid clinical researchers in speeding up the process for launching select clinical oncology trials and ensuring the most appropriate patients are enrolled," said Dr. Abdel Halim, Chief Scientific Officer and Executive Vice President, Acutis Diagnostics. "Additionally, the test is to provide researchers with a robust set of data and insights to support clinical decision-making and future research."

"Our work at SOPHiA GENETICS is focused on creating technology and solutions that make it easier for everyone to practice and benefit from data-driven medicine," said Ken Freedman, Chief Revenue Officer, SOPHiA GENETICS. "The new assay from Acutis Diagnostics will be designed to enable clinical researchers to work faster, and with a sound set of insights around their data, ultimately benefitting their current and future patients, as well as the industry as a whole."

An estimated 20% of cancer patients have a family history of cancer.¹ Next-generation sequencing (NGS) is the technology of choice in detecting biomarkers for hereditary cancers and identifying mutations that help to inform treatment options. The SOPHiA GENETICS hereditary cancer solutions are NGS-based applications that use artificial intelligence and machine learning with patented technologies to analyze raw NGS data and provide simplified insights. This technology enables clinical researchers to accurately characterize the complex mutational landscape associated with major hereditary cancer disorders.

Through the use of SOPHiA GENETICS' technology, the new assay from Acutis Diagnostics is expected to further clinical research and treatments for those who encounter hereditary cancers.

For more information on SOPHiA GENETICS, visit [SOPHiAGENETICS.COM](https://www.sophiagenetics.com), or connect on [Twitter](#), [LinkedIn](#), [Facebook](#), and [Instagram](#).

About SOPHiA GENETICS

SOPHiA GENETICS (Nasdaq: SOPH) is a cloud-native software company in the healthcare space dedicated to establishing the practice of data-driven medicine as the standard of care and for life sciences research. It is the creator of the SOPHiA DDM™ Platform, a cloud-native platform capable of analyzing data and generating insights from complex multimodal data sets and different diagnostic modalities. The SOPHiA DDM™ Platform and related solutions, products and services are currently used by a broad global network of hospitals, academic centers, laboratories and biopharma institutions. For more information, visit [SOPHiAGENETICS.COM](https://www.sophiagenetics.com), or connect on [Twitter](#), [LinkedIn](#), [Facebook](#), and [Instagram](#). **Where others see data, we see answers.**

SOPHiA GENETICS products are for Research Use Only and not for use in diagnostic procedures, unless specified otherwise. The information in this press release is about products that may or may not be available in different countries and, if applicable, may or may not have received approval or market clearance by a governmental regulatory body for different indications for use. Please contact support@sophiagenetics.com to obtain the appropriate product information for your country of residence.

SOPHiA GENETICS Forward-Looking Statements:

This press release contains statements that constitute forward-looking statements. All statements other than statements of historical facts contained in this press release, including statements regarding our future results of operations and financial position, business strategy, products, and technology, as well as plans and objectives of management for future operations, are forward-looking statements. Forward-looking statements are based on our management's beliefs and assumptions and on information currently available to our management. Such statements are subject to risks and uncertainties, and actual results may differ materially from those expressed or implied in the forward-looking statements due to various factors, including those described in our filings with the U.S. Securities and Exchange Commission. No assurance can be given that such future results will be achieved. Such forward-looking statements contained in this press release speak only as of the date hereof. We expressly disclaim any obligation or undertaking to update these forward-looking statements contained in this press release to reflect any change in our expectations or any change in events, conditions, or circumstances on which such statements are based, unless required to do so by applicable law. No representations or warranties (expressed or implied) are made about the accuracy of any such forward-looking statements.

¹ Al Harthi, F. S., et al. (2020) 'Familial/inherited cancer syndrome: a focus on the highly consanguineous Arab population', npj Genomic Medicine, 5, 3

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