



Ventripoint Receives European Union Medical Device Regulation Certification

Toronto, Ontario – The Newswire – May 30, 2023 - Ventripoint Diagnostics Ltd. ("Ventripoint" or the "Company"), (TSXV:VPT; OTC:VPTDF) is pleased to announce that it has successfully obtained European Union Medical Device Regulation (EU MDR) certification for its cardiac diagnostic system. This significant milestone further underscores Ventripoint's dedication to delivering state-of-the-art diagnostic tools to healthcare professionals and improving patient outcomes.

The EU MDR certification came into effect in May 2021. All medical devices certified under the previous Medical Device Directive (MDD) in the EU must certify to the new requirements (MDR 2017/745) to be sold in the European Market.

By receiving its EU MDR certification, Ventripoint Diagnostics demonstrates its ability to meet the evolving regulatory landscape and provide a safe and effective cardiac diagnostic tool for hospitals and cardiac clinics. Ventripoint Diagnostics is poised to expand its presence in the European market and further its mission to transform the way cardiac diseases are diagnosed and managed.

About Ventripoint Diagnostics Ltd.

Ventripoint has become an industry leader in the application of AI (Artificial Intelligence) to echocardiography. Ventripoint's VMS products are powered by its proprietary KBR technology, which is the result of a decade of development and provides accurate volumetric cardiac measurements equivalent to MRI. This affordable, gold-standard alternative allows cardiologists greater confidence in the management of their patients. Providing better care to patients serves as a springboard and basic standard for all of Ventripoint's products that guide our future developments. In addition, VMS+ is versatile and can be used with all ultrasound systems from any vendor supported by regulatory market approvals in the U.S., Europe and Canada.

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Forward Looking Statements

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Since forward-looking statements and information address future events and conditions, by their very nature they involve inherent risks and uncertainties. Actual results could differ materially from those currently anticipated due to a number of factors and risks. Factors which could materially affect such forward-looking information are described in the risk factors in the Company's most recent annual management's discussion and analysis that is available on the Company's profile on SEDAR at www.sedar.com. Readers are cautioned that the foregoing list of factors is not exhaustive. The forward-looking statements included in this news release are expressly qualified by this cautionary statement. The forward-looking statements and information contained in this news release are made as of the date hereof and the Company undertakes no obligation to update publicly or revise any forward-looking statements or information, whether as a result of new information, future events or otherwise, unless so required by applicable securities laws.