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VolitionRX Announces the Successful Conclusion of the Logistics and Pathway Design Study Conducted in Denmark.

"The logistics and feasibility of a Triage Concept seems very plausible."

ISNES, Belgium, Feb. 5, 2018 /PRNewswire/ -- VolitionRx Limited (NYSE AMERICAN: VNRX) ("Volition") today announced the successful conclusion to the Logistics and Pathway Design Study conducted by Hvidovre Hospital and The Danish Research Group. The study was established to answer the very practical, logistical questions of how a triage blood test could fit into the current Danish screening program for colorectal cancer ("CRC"). The study showed that it was indeed possible to collect, process, gather and ship the blood test from all five participating hospitals to a central laboratory within 24 hours. Furthermore, analysis from the central laboratory was shown to provide the needed data in due time (in accordance with current Danish Legislation) to decide whether or not to perform a colonoscopy.

Cameron Reynolds, Chief Executive Officer, commented "We are delighted that the team in Denmark has completed the Logistics and Pathway Design Study and positively concluded that the logistics and feasibility of using a triage concept, where the triage test is used in addition to a fecal immunochemical test to screen for CRC, seems very plausible and could indeed fit into the current Danish screening pathway. We also further welcome their conclusion that the inclusion of the triage concept could be advantageous. "

Volition has previously reported that many healthcare systems in Europe, including Denmark, are struggling to meet the increased colonoscopy demand that has come from the implementation of solely fecal-based colorectal cancer screening programs. This study demonstrates that a colorectal cancer screening triage blood test may prove beneficial in other national screening programs.

About Volition

Volition is a multi-national life sciences company developing simple, easy to use blood-based cancer tests to accurately diagnose a range of cancers. The tests are based on the science of Nucleosomics®, which is the practice of identifying and measuring nucleosomes in the bloodstream or other bodily fluid - an indication that disease is present.

As cancer screening programs become more widespread, Volition's products aim to help to diagnose a range of cancers quickly, simply, accurately and cost effectively. Early diagnosis has the potential to not only prolong the life of patients, but also to improve their quality of life.

Volition's research and development activities are currently centered in Belgium, with additional offices in London, Texas and Singapore, as the company focuses on bringing its

diagnostic products to market first in Europe, then in the U.S. and ultimately, worldwide.

For more information about Volition, visit Volition's website (http://www.volitionrx.com) or connect with us via:

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Statements in this press release may be "forward-looking statements" within the meaning of Section 27A of the Securities Act of 1933, as amended, and Section 21E of the Securities Exchange Act of 1934, as amended, that concern matters that involve risks and uncertainties that could cause actual results to differ materially from those anticipated or projected in the forward-looking statements. Words such as "expects," "anticipates," "intends," "plans," "aims," "targets," "believes," "seeks," "estimates," "optimizing," "potential," "goal," "suggests," "could," "would," "should," "may," "will" and similar expressions identify forward-looking statements. These forward-looking statements relate to the effectiveness of Volition's bodily-fluid-based diagnostic tests as well as Volition's ability to develop and successfully commercialize such test platforms for early detection of cancer. Volition's actual results may differ materially from those indicated in these forward-looking statements due to numerous risks and uncertainties. For instance, if Volition fails to develop and commercialize diagnostic products, it may be unable to execute its plan of operations. Other risks and uncertainties include Volition's failure to obtain necessary regulatory clearances or approvals to distribute and market future products in the clinical IVD market; a failure by the marketplace to accept the products in Volition's development pipeline or any other diagnostic products Volition might develop; Volition will face fierce competition and Volition's intended products may become obsolete due to the highly competitive nature of the diagnostics market and its rapid technological change; and other risks identified in Volition's most recent Annual Report on Form 10-K and Quarterly Reports on Form 10-Q, as well as other documents that Volition files with the Securities and Exchange Commission. These statements are based on current expectations, estimates and projections about Volition's business based, in part, on assumptions made by management. These statements are not guarantees of future performance and involve risks, uncertainties and assumptions that are difficult to predict. Forward-looking statements are made as of the date of this release, and, except as required by law, Volition does not undertake an obligation to update its forward-looking statements to

reflect future events or circumstances.

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