Hemostemix Announces It Is Trading on the Frankfurt Stock Exchange and It Has Retained HE Capital Markets Ltd.

written by Raj Shah | April 8, 2021

April 8, 2021 (Source) – Hemostemix Inc. (TSXV: HEM) (OTC: HMTXF) (FSE: 2VF0.F) ("Hemostemix" or the "Company") is pleased to announce it is trading on the Frankfurt Stock Exchange under symbol 2VF0.F and it has engaged HE Capital Markets Ltd. (HE Capital) to design and implement a North American and European multimedia digital advertising campaign on certain investor-focused and financial market websites. HE Capital will also provide other media communications services to raise the company's overall corporate profile in North America and Europe.

Thomas Smeenk, CEO, said: "With the listing on Frankfurt and the engagement of HE Capital, we are deepening our liquidity pools, introducing Hemostemix to a broader investment community. In the USA we are working to up-list the company, and we are exploring how best to reach our audience in Asia."

The cash consideration to be paid by the Company for these services consists of USD $18,000 for an initial three-month campaign. HE Capital acts at arm's length to Hemostemix and does not currently have any interest, directly or indirectly, in the company or its securities. HE Capital may choose to acquire securities of Hemostemix in the future.

HE Capital is a London-based investor relation and capital markets advisory firm, providing comprehensive communications solutions to public company clients in North America and Europe

through a combination of investor relations, public relations, and digital and social media services. HE Capital's office address is 6 Hays Lane, London Bridge, London, SE1 2HB, United Kingdom.

About Hemostemix

Hemostemix is a publicly traded autologous stem cell therapy company. A winner of the World Economic Forum Technology Pioneer Award, the Company developed and is commercializing its lead product ACP-01 for the treatment of CLI, PAD, Angina, Ischemic Cardiomyopathy, Dilated Cardiomyopathy and other conditions of ischemia. ACP-01 has been used to treat over 300 patients, and it is the subject of a randomized, placebo-controlled, double blind trial of its safety and efficacy in patients with advanced critical limb ischemia who have exhausted all other options to save their limb from amputation.

On October 21, 2019, the Company announced the results from its Phase II multi-center randomized double-blind placebo-controlled clinical trial of ACP-01 in critical limb ischemia (CLI) abstract entitled "Autologous Stem Cell Treatment for CLI Patients with No Revascularization Options: An Update of the Hemostemix ACP-01 Trial With 4.5 Year Followup" which noted healing of ulcers and resolution of ischemic rest pain occurred in 83% of patients, with outcomes maintained for up to 4.5 years.

The Company owns 91 patents across five patent families titled: Regulating Stem Cells, In Vitro Techniques for use with Stem Cells, Production from Blood of Cells of Neural Lineage, and Automated Cell Therapy. For more information, please visit www.hemostemix.com.

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Forward-Looking Information*: This news release contains "forward-looking information" within the meaning of applicable Canadian securities legislation. All statements, other than statements of historical fact, included herein are forward-looking information. In particular, this news release contains forward-looking information regarding the financing of Hemostemix, settlement of trade accounts payable with shares. There can be no assurance that such forward-looking information will prove to be accurate. Actual results and future events could differ materially from those anticipated in such forward-looking information. This forward-looking information reflects Hemostemix's current beliefs and is based on information currently available to Hemostemix and on assumptions Hemostemix believes are reasonable. These assumptions include, but are not limited to: the underlying value of Hemostemix and its common shares; the successful resolution of the litigation that Hemostemix is pursuing or defending (the "**Litigation**"); the timing of receipt of its full and complete clinical trials data; the results of ACP-01 research, trials and studies being equivalent to or better than previous research, trials or studies as well as management's expectations of anticipated results; Hemostemix's general and administrative costs remaining constant; the receipt of all required regulatory approvals for research, trials or studies as well as the disclosed or any required or desired financings of Hemostemix, including TSX Venture Exchange acceptance and any third party consents; the*

level of activity, market acceptance and market trends in the healthcare sector; the economy generally; consumer interest in Hemostemix's services and products; competition and Hemostemix's competitive advantages; and,a obtaining satisfactory financing to fund Hemostemix's operations including any research, trials or studies, and the Litigation. Forward-looking information is subject to known and unknown risks, uncertainties and other factors that may cause the actual results, level of activity, performance or achievements of Hemostemix to be materially different from those expressed or implied by such forward-looking information. Such risks and other factors may include, but are not limited to: the ability of Hemostemix to complete its current CLI clinical trial, complete a satisfactory futility analysis and the results of such and future clinical trials; litigation and potential litigation that Hemostemix may face; general business, economic, competitive, political and social uncertainties; general capital market conditions and market prices for securities; delay or failure to receive board or regulatory approvals; the actual results of future operations including the actual results of future research, trials or studies; competition; changes in legislation affecting Hemostemix; the timing and availability of external financing on acceptable terms; long-term capital requirements and future developments in Hemostemix's markets and the markets in which it expects to compete; lack of qualified, skilled labour or loss of key individuals; and risks related to the COVID-19 pandemic including various recommendations, orders and measures of governmental authorities to try to limit the pandemic, including travel restrictions, border closures, non-essential business closures, service disruptions, quarantines, self-isolations, shelters-in-place and social distancing, disruptions to markets, disruptions to economic activity and financings, disruptions to supply chains and sales channels, and a deterioration of general economic conditions including a

possible national or global recession or depression; the potential impact that the COVID-19 pandemic may have on Hemostemix may include a decreased demand for the services that Hemostemix offers; and a deterioration of financial markets that could limit Hemostemix's ability to obtain external financing. A description of additional risk factors that may cause actual results to differ materially from forward-looking information can be found in Hemostemix's disclosure documents on the SEDAR website at www.sedar.com. Although Hemostemix has attempted to identify important factors that could cause actual results to differ materially from those contained in forward-looking information, there may be other factors that cause results not to be as anticipated, estimated or intended. Readers are cautioned that the foregoing list of factors is not exhaustive. Readers are further cautioned not to place undue reliance on forward-looking information as there can be no assurance that the plans, intentions or expectations upon which they are placed will occur. Forward-looking information contained in this news release is expressly qualified by this cautionary statement. The forward-looking information contained in this news release represents the expectations of Hemostemix as of the date of this news release and, accordingly, it is subject to change after such date. However, Hemostemix expressly disclaims any intention or obligation to update or revise any forward-looking information, whether as a result of new information, future events or otherwise, except as expressly required by applicable securities law.