

Hemostemix Inc.

MANAGEMENT'S DISCUSSION AND ANALYSIS OF THE RESULTS OF OPERATIONS AND FINANCIAL CONDITION

For the three and six months ended June 30, 2025 and 2024 as at August 29, 2025

BASIS OF PRESENTATION

The following interim Management's Discussion and Analysis ("MD&A") covers the operations, financial position and operating results of Hemostemix Inc. (the "Company", "Hemostemix", "we", "us" or "our") for the three and six months ended June 30, 2025 and 2024. It is intended to help readers better understand the operations and key financial results, as they are, in our opinion, at the date of this report and should be read in conjunction with the unaudited condensed consolidated interim financial statements of the Company for the three and six months ended June 30, 2025 and 2024 and the accompanying notes which have been prepared under IFRS® Accounting Standards ("IFRS") as issued by the International Accounting Standards Board ("IASB"). The unaudited condensed consolidated interim financial statements have been reviewed by the Audit Committee of the Company and have been approved by its Board of Directors on August 29, 2025. Additional information relating to the Company is available on SEDAR+ at www.sedarplus.ca as well as the Company's website at www.hemostemix.com.

CAUTIONARY STATEMENT REGARDING FORWARD LOOKING INFORMATION

This MD&A contains certain forward-looking information and forward-looking statements, as defined in applicable securities laws (collectively referred to herein as "forward-looking statements"). These statements relate to future events or the Company's future performance. All statements other than statements of historical fact are forward-looking statements. Often, but not always, forward-looking statements can be identified by the use of words such as "plans", "expects", "is expected", "budget", "scheduled", "estimates", "continues", "forecasts", "projects", "predicts", "intends", "anticipates" or "believes", or variations of, or the negatives of, such words and phrases, or state that certain actions, events or results "may", "could", "would", "should", "might" or "will" be taken, occur or be achieved. Forward-looking statements involve known and unknown risks, uncertainties and other factors that may cause actual results to differ materially from those anticipated in such forward-looking statements. The forward-looking statements in this MD&A speak only as of the date of this MD&A or as of the date specified in such statement. Specifically, this MD&A includes, but is not limited to, forward-looking statements regarding:

- belief that the Company will be successful in raising additional capital to continue as a going concern;
- belief that its products and research and development efforts are targeting diseases and conditions with significant unmet medical treatment needs;
- the Company's goal of creating shareholder value;
- its ability to meet its operating costs for the twelve months ended June 30, 2025;
- belief that 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 positive anticipated results regarding future clinical trials for ACP-01 for other indications;
- the Company's belief that the ACP-01 technology process can be commercialized as effectively or more effectively than other technologies;
- our expectations regarding our ability to arrange for and scale up manufacturing of our products and technologies;
- the plans, costs, and timing for future research and development of the Company's stem cell technologies, including the costs and potential impact of complying with existing and proposed laws and regulations and clinical trials;

- belief that the Company's prior ACP-01 trial data will be sufficient to support regulatory submissions and approvals for additional indications such as ischemic and dilated cardiomyopathy, and chronic limb threatening ischemia;
- management's outlook regarding future trends;
- expectations regarding the performance of critical suppliers and service providers, including its contract manufacturer organization (CMO);
- expectations for additional commercialization partners;
- expectations for our ability to secure commercialization partners to develop our other technologies (NCP-01);
- expectations with respect to existing and future corporate alliances and licensing transactions with third parties, and the receipt and timing of any payments to be made by us or to us pursuant to such arrangements;
- expectations regarding the outcome of litigation;
- plans and objectives of management for future operations;
- our strategy with respect to the protection of our intellectual property;
- final financial performance; and
- general business and economic conditions and outlook.

Various assumptions or factors are typically applied in drawing conclusions or making the forecasts or projections set out in forward-looking information. Those assumptions and factors are based on information currently available to the Company, including information obtained from third-party industry analysts and other third-party sources. In some instances, material assumptions and factors are presented or discussed elsewhere in this MD&A in connection with the statements or disclosure containing the forward-looking information. You are cautioned that the following list of material factors and assumptions is not exhaustive. The factors and assumptions include, but are not limited to, assumptions that there may be no:

- unforeseen changes in the legislative and operating framework for the business of the Company;
- unstable competitive environment; and
- significant events occurring outside the ordinary course of business such as a natural disaster or other calamity.

These statements are only predictions and involve known and unknown risks, uncertainties and other factors including the risks set out in the section entitled "Risks and Uncertainties" below, which may cause the Company's or its industry's actual results, levels of activity, performance and achievements to be materially different from any future results, levels of activity or performance expressed or implied by these forward-looking statements. These risks and uncertainties include, but are not limited to the following risks:

- the successful and timely completion of research and development initiatives;
- negative results from the Company's animal studies and clinical trials;
- negative results of current litigation and potential litigation that the Company may face;
- risks associated with 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;
- risks associated with future developments in the Company's markets and the markets in which it expects to compete;
- lack of qualified, skilled labour or loss of key individuals;
- the viability and marketability of the Company's technologies;
- the effects of government regulation on the Company's business;
- the development of superior technology by the Company's competitors;

- the failure of consumers and the medical community to accept the Company's technology as safe and effective;
- risks associated with the performance of commercial partners and critical suppliers and service providers;
- risks associated with the Company's ability to obtain and protect rights to its intellectual property;
- risks associated with the Company's ability to raise additional capital to support operations;
- reliance on third parties to plan, conduct and monitor our clinical trials;
- other factors beyond the Company's control.

Although the Company believes that the expectations reflected in the forward-looking statements are reasonable, the Company cannot guarantee future results, levels of activity or performance. Further, any forward-looking statement speaks only as of the date on which such statement is made, and except as required by applicable law, the Company undertakes no obligation to update any forward-looking statement to reflect events or circumstances after the date on which such statement is made or to reflect the occurrence of unanticipated events. New factors emerge from time to time, and it is not possible for management to predict all of such factors and to assess in advance the impact of such factors on the Company's business or the extent to which any factor, or combination of factors, may cause actual results to differ materially from those contained in any forward-looking statement.

THE COMPANY

Hemostemix is a biotechnology Company whose principal business is to develop, manufacture and commercialize blood-derived stem cell therapies for medical conditions not adequately addressed by current treatments. Hemostemix, an entity under the Business Corporations Act (Alberta) was formed in November 2014. On November 27, 2014, shares of the Company began trading on the TSX Venture Exchange (the "Exchange") under the symbol "HEM". In October 2018, the Company was approved for listing its common shares for trading on the OTCQB Venture Market, a US trading platform that is operated by the OTC Markets Group in New York. Our shares now trade on the OTCQB under the symbol "HMTXF". In April 2021, the Company was approved for listing its common shares for trade on the Frankfurt Stock Exchange under the symbol "2VFO". The Company's head office is located at suite 1150, 707-7th Avenue SW, Calgary, AB T2P 3H6.

The unaudited condensed consolidated interim financial statements of the Company comprise the accounts of Hemostemix, Hemostemix Quebec Inc., Hemostemix PR Inc., PreCerv Inc., Hemostemix Ltd., and Kwalata Trading Limited, the Company's wholly-owned subsidiaries. Kwalata Trading Limited ("Kwalata"), incorporated under the laws of Cyprus, was established to own our intellectual property ("IP"). Hemostemix Ltd., another wholly-owned subsidiary, was incorporated under the laws of Israel to conduct manufacturing and perform research and development. Effective October 1, 2017, Hemostemix Ltd. ceased operations (see "Wholly-Owned Subsidiary"). On June 14, 2022, the Company incorporated PreCerv as a wholly-owned subsidiary. PreCerv obtained a global field of use license to NCP-01 and ACP-01 to treat conditions of the central and peripheral nervous system, including but not limited to Neuropathic pain syndromes, Traumatic spinal cord injury, chronic brainstem injury, traumatic brain injury, peripheral nerve injury. On August 15, 2023, the Company incorporated Hemostemix Quebec Inc. as a wholly-owned subsidiary. On August 22, 2024, the Company incorporated Hemostemix PR Inc. as a wholly-owned subsidiary.

BUSINESS OVERVIEW

We are a clinical stage biotechnology Company with a patented stem cell technology platform whose principal business is to develop, manufacture and commercialize blood-derived stem cell therapies to treat various diseases not adequately addressed by current therapeutics. The intellectual property of the Company broadly covers synergetic cell populations that can be differentiated into angiogenic cell precursors (“ACPs”, including the lead cell product ACP-01), neural cell precursors (“NCPs”) and cardiomyocyte cell precursors.

CORPORATE, PRODUCT, CLINICAL TRIAL AND FINANCING UPDATE

The following items highlight the Company’s activities during the six months ended June 30, 2025 and any subsequent development up until the date hereof.

Corporate Update

On January 7, 2025, the Company issued 4,085,461 shares at \$0.10 per share to satisfy the interest owing from January 1, 2023 to September 30, 2024 of \$408,546 on the outstanding \$2,750,000 Convertible Debenture.

On January 28, 2025, 50,000 warrants at a price of \$0.25 were exercised.

On January 30, 2025, 180,000 options at a price of \$0.07 were exercised by a director of the Company.

On February 10, 2025, 57,140 warrants at a price of \$0.175 were exercised.

On February 11, 2025, 57,140 debenture warrants at a price of \$0.20 were exercised.

On February 19, 2025, the Company announced an arm's length perpetual royalty-free global license to CytolImmune’s Bioreactor stem cell technologies. Subject to the TSXV Exchange acceptance, Hemostemix will pay CytolImmune \$5,000,000 (twenty million shares) for the perpetual global royalty free license.

On March 5, 2025, 16,000 warrants at a price of \$0.05 were exercised.

On March 7, 2025, 200,000 warrants at a price of \$0.12 were exercised.

On May 11, 2025, the Company announced that it had closed a non-brokered private placement of \$336,500 USD, issuing 1,634,466 common shares at \$0.295CAD each to CytolImmune Therapeutics LLC, following the approval of the TSXV Exchange.

On June 27, 2025, 20,980 warrants at a price of \$0.12 were exercised.

On June 27, 2025, 38,400 warrants at a price of \$0.12 were exercised.

On June 27, 2025, 147,200 warrants at a price of \$0.05 were exercised.

On June 27, 2025, 294,400 warrants at a price of \$0.05 were exercised.

On June 27, 2025, 73,600 warrants at a price of \$0.05 were exercised.

Product Update

Angiogenic Cellular Precursor (ACP-01)

Our main product, ACP-01, is created from a process we discovered, developed and patented. From the patient's blood a synergetic cell population is isolated, cultured, differentiated into our products, then injected into the patient's ischemic tissue or organ(s). Our process for harvesting stem cells is less invasive, as the stem cells are taken from the patient's blood, which is a simplified process as compared to taking stem cells from fatty tissue or bone marrow. Hemostemix's proprietary technology is a personalized regenerative therapy that is administered to a patient within 7 days of the initial blood draw.

Based on four open label studies, [including] a randomized phase I study, a randomized double blind phase II Clinical trial and the retrospective study of 53 consecutive patients treated for ischemic and dilated cardiomyopathy, we believe that ACP-01 has applications in the treatment of other vascular diseases such as peripheral arterial disease ("PAD"), angina pectoris, and other diseases of ischemia.

Regulatory Update for ACP-01

In the first quarter of 2019, the Company submitted an application to the US Food and Drug Administration ("FDA") for Orphan Drug Designation ("ODD") for ACP-01 for the treatment of patients with CLI. The Orphan Drug Act provides for granting special status to a drug or biological product to treat a rare disease or condition upon request of a sponsor. The FDA defines rare diseases as those affecting fewer than 200,000 people in the United States at any given time. Our application sought ODD for the treatment of end-stage CLI patients. The FDA responded to the Company's application stating that based on the information and data they reviewed ACP-01 had the potential to treat all patients suffering from CLI, not just those with end-stage CLI. Based on the potential to treat such a large patient population, ACP-01 did not qualify for Orphan Drug Status as a treatment of end-stage CLI.

Neural Cellular Precursor (NCP-01)

On October 19, 2022 the Company and its wholly-owned subsidiary, PreCerv Inc., announced the preclinical commencement of PreCerv Inc.'s study of NCP-01 at Clemson University, South Carolina. The objectives of the study, a spinal cord repair model, are to evaluate the fate of NCP-01 cells following injection into the spinal fluid pathway, and evaluate the dosing effect on neuropathic pain and motor function recovery.

Intellectual Property

Our proprietary technologies are based on more than 16 years of clinical data, four open label studies, a randomized phase I study, a randomized double blind phase II Clinical trial and the retrospective study of 54 consecutive patients treated for ischemic and dilated cardiomyopathy, and a truncated randomized, placebo-controlled, double blind Phase II clinical trial of Critical Limb Ischemia.

The Company continuously monitors its patent portfolio and vigorously defends its intellectual property rights. The Company has 91 patents, organized into five patent families, issued in more than 25 jurisdictions.

The five patent families are:

Family Patent	Status	Title
1	Granted in several countries including in the US Pending in Canada and Thailand	In-vitro techniques for use with stem cells
2	Granted in several countries including Canada To be filed in US	Production from blood of cells of neural lineage

3	Granted in Singapore Pending in Canada, Europe and US	Regulating stem cells
4	Granted in several counties including the US and Canada Pending in Europe	Regulating stem cells
5	Granted Mexico, Singapore	Automated cell therapy

Clinical Trial Updates

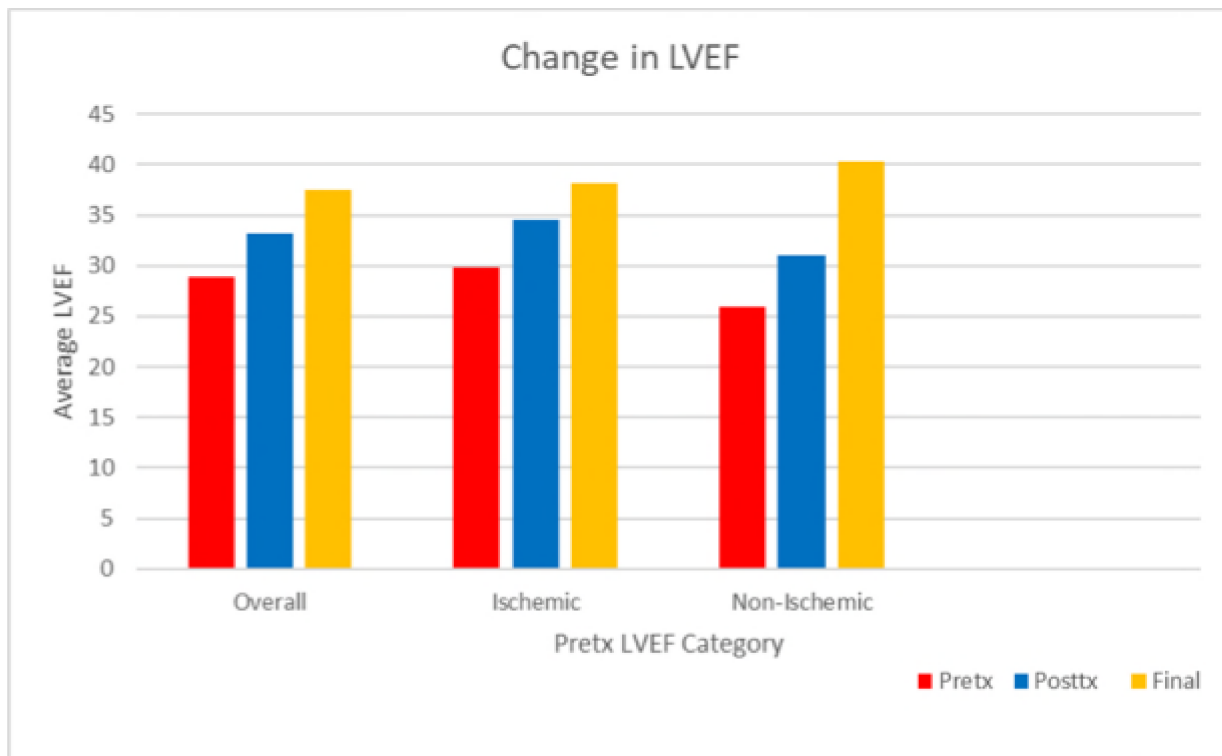
Retrospective Cardiomyopathy Study

On August 30, 2022, the Company announced the results of a new retrospective study of 53 consecutive patients who underwent a trans-catheter, intramyocardial injection of angiogenic cell precursors (ACP-01) as a treatment for heart failure (ischemic and non-ischemic dilated cardiomyopathy).

The 2021 American Heart Association estimated the prevalence of heart failure (HF) in the United States to be 6 million; within 5-years of hospitalization, the death rate amongst this population is approximately 50%. ACP-01 in its capacity to replace damaged cells, secrete growth factors, stimulate angiogenesis and exert an anti-inflammatory effect to minimize scarring, have emerged as a therapeutic option. Hemostemix completed an IRB approved, retrospective, outcomes study to analyse the effect of ACP-01 implants on cardiac function in patients with severe heart failure (New York Heart association Grades 3 and 4). Cardiac function was measured in terms of ejection fraction of the left ventricle (LVEF %).

At first follow-up (average 4 months) after ACP-01 cell implantation, for all types of heart failure, the LVEF was increased by 4.6% (from 28.6% to 33.2%), representing a statistically significant improvement ($p < 0.0011$). On final follow-up (average 12 months after cell implantation) for all patients, the LVEF% had improved by 7.69%, which was statistically significant ($p < 0.003$).

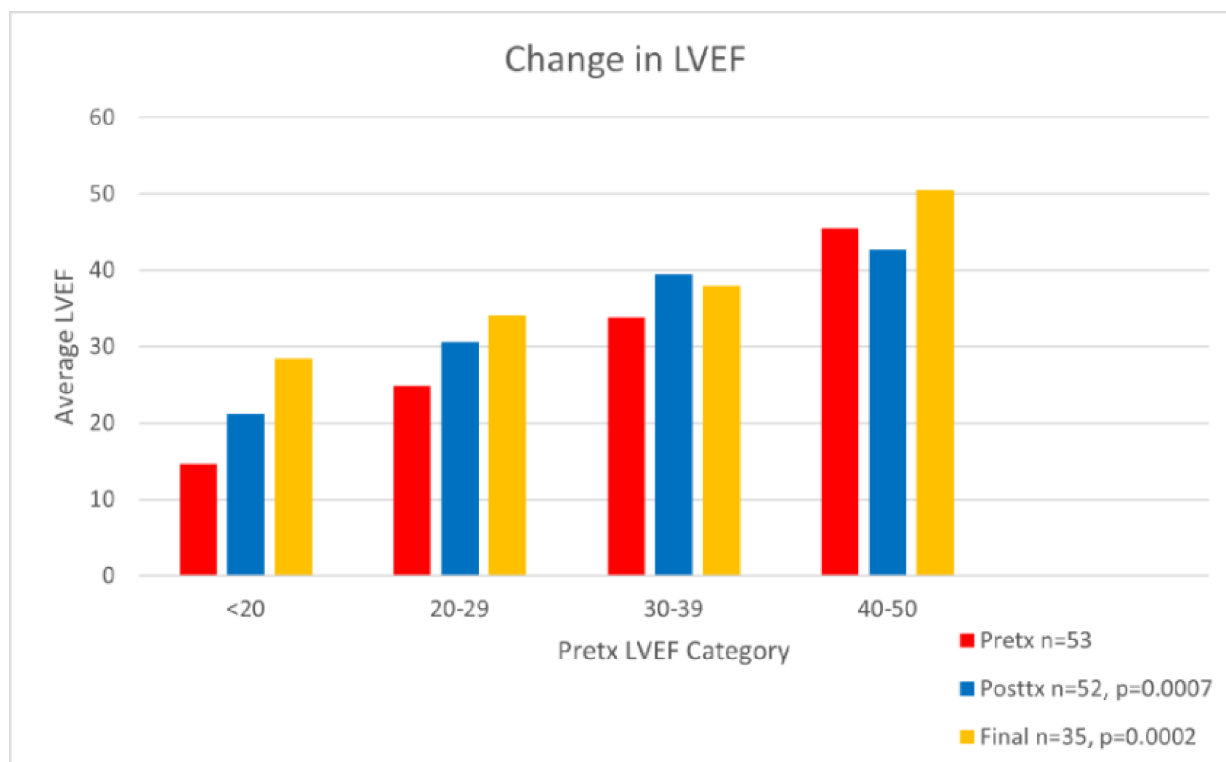
When analyzing ischemic heart failure alone ($n=41$), LVEF increased from 29.9% before implantation, to 34.5%, and to 38.2% at final follow-up, for an overall improvement of 8.37%, which was statistically significant ($p < 0.003$). There was greater improvement in the non-ischemic dilated cardiomyopathy patients ($n=8$), who improved from 25.94% before treatment to 40.29% at final follow-up, for an overall improvement of 14.35% ($p < 0.002$).



To determine whether the observed improvements after ACP-01 implantation occurred at different levels of cardiomyopathy severity, the patients were divided into quartiles of pre-procedural LVEF%:

- i. those with extremely severe cardiomyopathy, pre-procedural LVEF% <20%;
- ii. those with severe pre-procedural LVEF% 20-29%;
- iii. those with moderately severe cardiomyopathy pre-procedural LVEF% 30-39%
- iv. those termed “heart failure with mid-range EF”, with pre-procedural LVEF% 40-50% (per American College of Cardiology/American Heart Association/European Society of Cardiology)

Overall, upward mobility from one quartile to the next quartile was significant ($p < 0.0007$) for first follow up, and more significant ($p < 0.0002$) for final follow up. Three of the four groups moved up into the next category of LVEF function. The greatest improvement of LVEF% was generated in patients with extremely severe cardiomyopathy (LVEF% <20%).



Quality of life statements were used also to assess response to treatment. Patients receiving ACP-01 implantation reported "improvement of quality of life" in 66% of cases, no overall change in 28% of cases, and worsened quality of life in 6% of cases. Although not statistically significant, 94% of cases were the same or improved following ACP-01 treatment, a contradistinction to the deterioration of quality of life exhibited by the general population of patients with heart failure.

The study's conclusion was that trans-catheter intramyocardial injection of angiogenic cell precursor (ACP-01) as a treatment for heart failure is feasible and safe. The results are statistically significant and merit randomized, double-blind, placebo-controlled trials to confirm the benefit of this type of cell transplantation.

The calculated LVEF%, derived from MUGA scan, echocardiogram or SPECT was normally distributed, fulfilling the assumption for parametric testing, and treated as continuous variables, expressed as probability density functions. The paired t test compared mean preoperative and postoperative LVEF%. A p value of < 0.05 was considered statistically significant.

Phase II Clinical Trial for Patients with Critical Limb Ischemia

CLI is a severe blockage in the arteries of the lower extremities, which markedly reduces blood-flow. It is a serious form of peripheral arterial disease ("PAD"). PAD is caused by atherosclerosis, the hardening and narrowing of the arteries over time due to the build-up of fatty deposits called plaque. CLI is a chronic condition that results in severe pain in the extremities due to nerve and tissue damage. Complications of poor circulation can include sores and ulcerating wounds that will not heal in the legs and feet. Left untreated, the complications of CLI may result in the amputation of the affected limb.

Most patients with CLI are treated surgically and depending on the severity, the surgery can be minimally invasive (angioplasty or stents) or very invasive (bypass surgery, grafts, or amputation). ACP-01 may be an alternative to surgery, which, based on our prior clinical trials, we believe is safer and more cost effective, as no lengthy hospital stay or recovery time is needed. The prevalence of CLI is increasing, as CLI predominately affects the

growing baby boomer population aged 50 and older. According to The Sage Group LLC, in the United States alone, approximately 20 million people are affected by PAD, and it is estimated that approximately 7-8 million people in the United States and Europe suffer from CLI. The Sage Group LLC estimates that in the United States, medical costs attributable to CLI amount to US\$25 billion annually.

The HS 12-01 clinical trial was a randomized, placebo-controlled, double blind Phase II clinical trial of ACP-01 as a potential treatment of CLI.

On October 21, 2019, the Company was provided a summary of the presentation entitled “Autologous Stem Cell Treatment for CLI Patients with No Revascularization Options: An Update of the Hemostemix ACP-01 Trial with 4.5 Year Follow up” by the lead investigator, Dr. York Hsiang, who gave this update at the 41st Annual Canadian Society for Vascular Surgery Meeting, September 14, 2019. Dr. Hsiang reported on the blinded results from the long-term follow-up of the first cohort of patients enrolled at two trial sites, Vancouver Coastal Health Research Institute (“VCHRI”) and the University Health Network, Peter Munk Cardiac Centre located in Toronto, Ontario.

The following is a summary of Dr. Hsiang's the results and conclusion:

- Twelve patients with CLI with no interventional options were enrolled at two treatment centers (10 male, 2 female, mean age 76)
- Prior to treatment, three patients had ischemic rest pain, eight patients had ulceration, and one patient had gangrene
- Study subjects were randomized 2:1 to receive injection of ACP-01 or placebo into their most affected lower extremity and followed for at least 1 year
- Healing of ulcers and resolution of ischemic rest pain occurred in 10 of the 12 patients (83%)
- There were no clinically significant safety issues
- Outcomes were maintained for up to 4.5 years. (3.5 years for two patients, 3 years for one patient, and one patient who died after ulcer healing secondary to congestive heart failure)
- These blinded preliminary results in the study are promising, and show an acceptable safety profile for ACP-01

On August 31, 2022, the Company reported the unblinded results of the HS 12-01 CLI trial as follows:

Category	Statistic	ACP-01	Placebo	Total
Subjects randomized	n	47	21	68
Subjects randomized but were not treated	n (%)	1 (2.1)	0	1 (1.5)
Subjects treated	n (%)	46 (97.9)	21 (100)	67 (98.5)
Subjects who completed the study	n (%)	27 (57.4)	19 (90.5)	46 (67.6)
Withdrawal from the study	n (%)	20 (42.6)	2 (9.5)	22 (32.4)
Amputation	n (%)	3 (11.1)	2 (10.5)	5 (10.8)

- As originally designed, the clinical trial power analysis required 95 subjects to achieve a statistically significant result. The previous management team, however, truncated the trial to 68 subjects, reducing the power of the analysis of the study's primary endpoint to 25%. The trial demonstrated that ACP-01 was safe and treated patients, as compared to placebo treated patients:
- trended toward improvements in ulcer healing at 3 months
- trended toward ulcer healing at 6 months
- trended toward ulcer healing at the end of study at 12 months
- trended toward a reduction in pain associated with critical limb ischemia at 12 months

Hemostemix published the following Phase II Results in the Journal of Biomedical Research and Environmental Science, February 2024.

Introduction: Critical limb ischemia has a prevalence in the US of 1.33%, with mortality 15-20% and major amputation 10-40% per year. Stem cell treatment has emerged as a treatment option for the 45% of patients for whom revascularization procedures are not possible.

Objective: This study re-examines the data of the Phase II clinical treatment of no option Critical limb ischemia with Hemostemix' angiogenic cell precursors, focusing upon ulcer wound healing, amputation and death rate of this cohort.

Methods: Primary endpoints were changes in ulcer size and major amputation or death within one year of treatment. The secondary endpoint was change in pain level.

Results: From 2015 to 2021, 67 patients with no option Critical limb ischemia were allocated to treatment with ACP-01 (46/67) or placebo (21/67). From this data, only patients who presented with wound ulcers before administration of ACP-01 were reviewed (21 treatment, 8 placebo). Ulcer size in the treated group decreased from a mean of 1.46 cm² to 0.48 mm² (p = 0.01) by 3 months. There was no significant decrease in the size of the ulcers of the placebo group (p < 0.54). At one year there were no complications related to treatment. The treatment group had one amputation (4.8%) and one death (4.8%); the placebo group had 2 amputations (25%) and 1 death (12.5%). Change in pain was not significant in either group at 3 months, but at 1 year was improved in the placebo group (p = 0.01).

Conclusion: The administration of ACP-01 within a program of careful patient follow up is safe and associated with reduced ulcer size and decreased rate of amputation and death. Consideration should be given to re-administration of stem cell treatments every 3-6 months to optimize improvement of Critical limb ischemia. Further studies, more appropriately powered, are warranted.

Neural Cellular Precursor (NCP-01).

On January 7, 2020, the Company announced that it was issued its 87th patent for the generation of NCP-01 from peripheral blood. The patent, Production from Blood of Cells of Neural Lineage, was issued by Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Netherlands, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Monaco, Poland, Portugal, Romania, Slovakia, Slovenia, Spain, Sweden, Switzerland, Turkey and the United Kingdom.

Stock Options and Warrants

On January 22, 2025, the Company granted 2,085,000 stock options to various officers, directors and consultants of the Company. The stock options granted have an exercise price of \$0.28 and an expiry date of January 22, 2030. 1,815,000 of these stock options will vest immediately. The remaining 270,000 stock options will vest 50% immediately with the remaining 50% fully vested on January 31, 2026. The fair value of the stock options were \$542,074 and was estimated on the date of grant using the Black-Scholes model with the following assumptions: expected volatility of 237.81%, risk-free interest rate of 3.02% and an average expected life of 5 years.

During the three months ended March 31, 2025, 275,000 warrants with a Black Scholes value of \$8,106, were exercised into 275,000 common shares for proceeds of \$42,750

During the six months ended June 30, 2025, 704,860 broker warrants were exercised into 704,860 common shares for proceeds of \$48,936.

During the six months ended June 30, 2025, 5,468,417 warrants expired unexercised.

OUTLOOK

The Company continues to strongly believe in the technology based on the safety profile and efficacy of ACP-01 as reported in four heart studies and three critical limb ischemia trials, and its extensive work to optimize and scale its manufacturing processes.

Our ability to accomplish all our future strategic plans is dependent upon obtaining additional financing or executing other strategic options and there is no assurance that we will achieve these objectives. Management will continue to pursue various options to raise additional funding, some which could be dilutive to existing shareholders. Alternatives for raising further capital could include the issuance of additional equity, debt, convertible debentures, government or partnership funding. We intend to seek commercialization partners for our therapy and development partners for accelerating clinical development of novel therapies for significant and unmet medical needs.

CONSOLIDATION AND PRESENTATION

Wholly-Owned Subsidiaries

Hemostemix has five wholly-owned subsidiaries, Kwalata Trading Limited ("Kwalata"), Hemostemix Ltd. ("HEM Israel"), PreCerv Inc. ("PreCerv"), Hemostemix Quebec Inc. and Hemostemix PR Inc. All of the Company's patents and trademarks are registered in the name of Kwalata. HEM Israel was the original manufacturing company.

On June 14, 2022, the Company incorporated PreCerv to enable PreCerv to obtain from Hemostemix a global field of use license to NCP-01 and ACP-01 to treat conditions of the central and peripheral nervous system, including but not limited to the following:

1. Neuropathic pain syndromes.
2. Traumatic spinal cord injury, chronic brainstem injury, traumatic brain injury, peripheral nerve injury. Rare diseases including: syringomyelia, Charcot-Marie tooth disease, and Guillain-Barre syndrome, Amyotrophic lateral sclerosis (ALS), age-related macular degeneration (ARMD), corneal or eye diseases and retinopathies of any cause.
3. Cerebral stroke.

On August 29, 2023, the Company incorporated Hemostemix Quebec Inc. as a wholly-owned subsidiary. Hemostemix Quebec Inc.

On August 22, 2024, the Company incorporated Hemostemix PR Inc. as a wholly-owned subsidiary. Hemostemix PR Inc. to facilitate our planned manufacturing build out in Puerto Rico commencing with our Puerto Rican contract manufacturer.

Functional and Presentation Currency

The unaudited condensed consolidated interim financial statements are presented in Canadian dollars, which is the Company's functional and presentation currency. Each subsidiary determines its own functional currency and items included in the unaudited condensed consolidated interim financial statements of each entity are measured using that functional currency. The functional currency of the subsidiaries is Canadian dollars. Transactions denominated in foreign currency (other than the functional currency) are recorded on initial recognition at the exchange rate at the date of the transaction. After initial recognition, monetary assets and liabilities denominated in foreign currency are translated at the end of each reporting period into the functional currency at the exchange rate at that date. Exchange differences, other than those capitalized to qualifying assets or recorded in equity in hedging transactions, are recognized in profit or loss. Non-monetary assets and liabilities measured at cost in a foreign currency are translated at the exchange rate at the date of the transaction. Non-monetary assets and liabilities denominated in foreign currency and measured at fair value are translated into the

functional currency using the exchange rate prevailing at the date when the fair value was determined.

SELECTED FINANCIAL INFORMATION FOR THE PERIODS

The following table provides selected consolidated financial information for the Company as at and for the three and six months ended June 30, 2025 and 2024.

	Six months ended June 30,	
	2025	2024
Total Assets	926,667	991,863
Total Liabilities	8,090,547	8,196,014
Net loss and comprehensive loss before taxes	(1,513,314)	(994,878)
Basic and diluted loss per share	(0.010)	(0.011)
Weighted average number of shares outstanding	145,315,712	87,122,318

Total Assets decreased primarily as a result of new investments for the current period.

Total Liabilities decreased by \$105,467, related to operations, accounts payable and interest and accretion on the debentures during the year.

Net loss and comprehensive loss before taxes increased to \$1,513,314 for the six months ended June 30, 2025, primarily as a result of increased legal fees, consulting and share based compensation.

RESULTS OF OPERATIONS

Comparison of Expenses

	Three months ended June 30,		Increase (Decrease)	Increase (Decrease)
	2025	2024	\$	%
	\$	\$	\$	%
Research and development	2,747	-	2,747	DIV/0
Consultants	151,269	245,207	(93,938)	(38)
Stock-based compensation	14,858	4,143	10,715	259
Marketing and office expenses	22,050	12,006	10,044	84
Professional fees	259,670	88,779	170,891	192
Travel	31,256	13,928	17,328	100
Foreign exchange (gain) loss	(40,497)	4,723	(45,220)	(957)
Finance expense	173,124	150,533	22,591	15
Gain on write down of payables	(306,837)	-	(306,837)	DIV/0
Depreciation and amortization	12	25	(13)	(52)
Net loss from operations	307,652	519,344	(211,692)	(41)

	Six months ended			
	June 30, 2025 \$	June 30, 2024 \$	Increase (Decrease) \$	Increase (Decrease) %
Research and development	2,747	2,972	(225)	(8)
Consultants	258,781	423,087	(164,306)	(39)
Stock-based compensation	562,408	8,286	554,122	6,687
Marketing and office expenses	70,943	92,662	(21,719)	(23)
Professional fees	480,322	142,274	338,048	238
Travel	63,126	20,500	42,626	208
Foreign exchange loss	42,500	8,958	33,542	374
Finance expense	339,301	296,089	43,212	15
Gain on extinguishment	(306,837)	-	(306,837)	DIV/0
Depreciation and amortization	23	50	(27)	(54)
Net loss from operations	1,513,314	994,878	518,436	52

Analysis of expenses

Research and development (“R&D”)

R&D expense is the cost for the third party manufacturing laboratory which produces ACP-01 that is used in the clinical trials and provides continued research and development work in their laboratory. Unrelated to third party manufacturing, R&D costs for the three and six months ended June 30, 2025 were \$2,747 and \$2,747, respectively compared to \$nil and \$2,972, respectively for the three and six months ended June 30, 2024 representing a decrease of \$(2,747) and \$225, respectively. The majority of the decrease is related to the clinical trials being finished for ACP-01.

Consultants

Consulting fees and salaries changed to \$151,269 and \$258,781, respectively for the three and six months ended June 30, 2025, as compared to \$245,207 and \$423,087, respectively in the corresponding period of the prior period. The change was due to a change in management consultant fees in the current period compared to the previous period.

Stock-based compensation expense (“SBC”)

SBC increased by \$10,715 and \$554,122, respectively in the three and six months ended June 30, 2025, compared to the corresponding periods of 2024. The increase is primarily due to the vesting of the stock options granted on January 22, 2025, and the impact of the change in vesting terms for certain stock options issued. Stock options are granted to certain officers, directors, employees and consultants, with the number, term and vesting period of the options granted being determined at the discretion of the Company’s board of directors and in conjunction with the terms of the Company’s stock option plans, to a maximum of 10% of the outstanding Common Shares.

Marketing and office expenses

For the three and six months ended June 30, 2025, marketing and office expenses changed to \$22,050 and \$70,943, respectively compared to \$12,006 and \$92,662, respectively in the same period of the prior year. Marketing and office expenses includes office administration costs including courier, and utilities as well as marketing and communications costs.

Professional fees

	Three months ended June 30,			Six months ended June 30,		
	2025	2024	% change	2025	2024	% change
Patent costs	95,542	74,502	28	111,165	81,574	36
Accounting fees	45,762	(13,702)	(434)	105,151	26,364	299
Legal - litigation	-	2,856	(100)	7,505	2,856	163
Legal - Other	83,079	14,495	473	106,821	16,742	538
Other Professional fees	(13,779)	7,375	(287)	6,160	8,071	(24)
Investor relations	49,066	3,253	100	143,520	6,667	2,053
Total	259,670	88,779	192	480,322	142,274	238

Professional fees increased to \$259,670 and \$480,322, respectively, for the three and six months ended June 30, 2025, as compared to \$88,779 and \$142,274, respectively in the prior period, primarily as a result of increased investor relations, accounting fees and legal costs relating to the private placements which were incurred in the current period.

Depreciation expense for the three months and three and six months ended June 30, 2025, were \$12 and \$23, respectively compared to \$25 and \$50, respectively in the corresponding periods of the prior period.

Finance expense, for the three months and three and six months ended June 30, 2025 were \$173,124 and \$339,301, respectively compared to \$150,533 and \$296,089, respectively in the corresponding periods of the prior year. Interest expense relates to the interest on the convertible debentures issued in 2021 and 2022.

Foreign exchange for the three months and three and six months ended June 30, 2025 were a loss of \$(40,497) and \$42,500, respectively compared to \$4,723 and \$8,958, respectively in the corresponding prior year period. The change from the current and prior year periods relates to an unrealized foreign exchange gain due to a change in rates, as well as a substantial payables balance in US currency.

LIQUIDITY AND CAPITAL RESOURCES

Hemostemix is a development stage Company that to date has had no revenue and negative operating cash flows, which are expected to continue in the foreseeable future. As a development stage Company, we require significant additional investment for research and development, manufacturing, clinical testing and regulatory submissions prior to commercialization. Since inception, we have financed our cash requirements primarily through issuances of equity and debt securities. Our ability to continue as a going concern is dependent upon obtaining additional investment capital and grant monies.

Based on the foregoing, we will continue to pursue various funding options and opportunities; however, no assurances can be made that we will be successful in raising additional investment capital, to continue as a going concern. If we are not able to raise capital, we will have to reduce our cash requirements by eliminating or deferring spending on research, development and corporate activities.

For the six months ended June 30, 2025, there was a net cash outflow from operating activities of \$463,746 compared to a net cash outflow of \$51,664 for the six months ended June 30, 2024, an increase in outflow of \$412,082 compared to the prior comparative period.

Expressed in tabular form, the decrease from the net cash used for operations is as follows:

Net loss from operations for the period	\$(518,436)
Increase in stock compensation expense	\$554,122
Increase in finance expense	\$42,420
Decrease in depreciation and amortization	\$(27)
Loss on settlement of debt through shares	
Change in subscription receivables	\$184,240
Change in other receivables and prepaid expenses	\$(19,204)
Change in HST/GST receivable	\$748
Change in accounts payable and accrued liabilities	\$(655,945)
Decrease in the net cash used for operations	<u>\$412,082</u>

As at June 30, 2025, the Company had a working capital deficit of \$1,811,031 compared to \$1,780,598 at December 31, 2024, resulting in an increase in working capital deficit of \$30,433.

Outstanding Share Data

As at June 30, 2025, the number of issued and outstanding common shares was 147,521,740 (December 31, 2024 – 140,641,953). As at August 29, 2025, the number of common shares issued and outstanding is 181,052,741.

As at June 30, 2025, the Company had 14,291,694 share purchase options outstanding (December 31, 2024 – 12,386,694). As at August 29, 2025, the number of outstanding share purchase options outstanding is 14,291,694.

As at June 30, 2025, the Company had 72,041,373 share purchase warrants outstanding (December 31, 2024 – 78,489,650). As at August 29, 2025, the number of outstanding warrants was 104,272,370.

MATERIAL ACCOUNTING POLICIES

Refer to Note 2 in the 2024 audited annual consolidated financial statements for a detailed description of our material accounting policies. We have consistently applied the same accounting policies for all periods presented in these consolidated financial statements for the three and six months ended June 30, 2025.

COMMITMENTS & CONTINGENCIES

Commitments

Clinical Trial Costs

In 2024, and continuing into 2025, these costs will primarily relate to analytical and trial planning and initiation activities.

Contingencies

In the ordinary course of operating, the Company may from time to time be subject to various claims or possible claims. Management believes that there are no claims or possible claims that if resolved would either individually or collectively result in a material adverse impact on the Company's financial position, results of operations, or cash flows. These matters are inherently uncertain, and management's view of these matters may change in the future.

Zenith Appraisal & Land Consulting Ltd. Lawsuit

Zenith Appraisal and Land Consulting Ltd's made a claim for compensation, notwithstanding that its principal, David Wood, a former director of the Company, was paid in full and signed a release following his resignation from the Board during April 2020. Zenith's claim is without merit and the Company will defend itself.

RELATED PARTY BALANCES AND TRANSACTIONS

Related party transactions are conducted on the terms and conditions agreed to by the related parties. It is the Company's policy to conduct all transactions and settle all balances with related parties on market terms and conditions.

The following includes all compensation to a consultant and key management personnel:

The Company recorded share-based compensation expense for the three and six months ended June 30, 2025 of \$6,858 and \$534,073, respectively (three and six months ended June 30, 2024 - \$1,308 and \$2,616, respectively) to the current management and directors of the Company.

For the three and six months ended June 30, 2025, the Company incurred \$49,500 and \$99,000, respectively (three and six months ended June 30, 2024 - \$49,500 and \$99,000, respectively) to Mr. Thomas Smeenck for consulting services. As at June 30, 2025, Mr. Smeenck was owed \$237,552 (December 31, 2024 - \$292,999) and this amount was included in accounts payable and accrued liabilities.

For the three and six months ended June 30, 2025, the Company incurred \$50,000 and \$100,000, respectively (three and six months ended June 30, 2024 - \$37,500 and \$75,000, respectively) to a consultant of the Company. As at June 30, 2025, they were owed \$175,672 (December 31, 2024 - \$273,391) and this amount was included in accounts payable and accrued liabilities.

For the three and six months ended June 30, 2025, the Company incurred \$16,243 and \$49,491, respectively (three and six months ended June 30, 2024 - \$1,635 and \$10,480, respectively) to Marrelli Support Services Inc., a company which the CFO is related to. As at June 30, 2025, the Company owed \$7,111 (December 31, 2024 - \$nil) to Marrelli Support Services Inc. for accounting fees, and this amount was included in accounts payable and accrued liabilities.

FINANCIAL INSTRUMENTS & CAPITAL RISK MANAGEMENT

Our financial instruments consist of cash and cash equivalents, subscriptions receivables and accounts payable, debentures and accrued liabilities. As at June 30, 2025, there are no significant differences between the carrying values of these amounts and their estimated market values.

Financial risk management

The Company's financial risk management policies are established to identify and analyze the risks faced by the Company, to set acceptable risk tolerance limits and controls, and to monitor risks and adherence to limits. The financial risk management policies and systems are reviewed regularly to ensure they remain consistent with the objectives and risk tolerance acceptable to the Company and current market trends and conditions. The Company, through its training and management standards and procedures, aims to uphold a disciplined and constructive control environment in which all employees understand their roles and obligations.

Interest rate risk

Interest rate risk is the risk that future cash flows of a financial instrument will fluctuate because of changes in market interest rates. We are exposed to interest rate risk through our cash and cash equivalents. The Company mitigate this risk by investment of excess cash resources in investment grade vehicles while matching maturities with our operational requirements.

Fluctuations in market rates of interest do not have a significant impact on our results of operations due to the short term to maturity of the investments held.

The Company mitigate our exposure to interest rate risk on loans by utilizing fixed rates.

Currency risk

Currency risk is the risk that future cash flows of a financial instrument will fluctuate because of changes in foreign exchange rates. In the normal course of our operations. The Company are exposed to currency risk from the purchase of goods and services in the United States. In addition, the Company are exposed to currency risk to the extent cash is held in foreign currencies. The impact of a 10% increase in the value of the U.S. dollar against the Canadian dollar would have increased our net loss for the June 30, 2025 by approximately \$34,202 (June 30, 2024- \$126,298).

We mitigate our foreign exchange risk by maintaining sufficient foreign currencies, through the purchase of foreign currencies, when cash allows, to settle our foreign accounts payable and future commitments.

Balances in foreign currencies at June 30, 2025 are as follows:

	<u>US Dollars (\$)</u>
Cash and cash equivalents	605,980
Accounts payable and accrued expenses	<u>(948,000)</u>
	(342,020)

Liquidity risk

Liquidity risk is the risk that we will encounter difficulty in meeting obligations associated with financial liabilities. We manage liquidity risk through the management of our capital structure. Accounts payable are all due within the current operating period.

As at June 30, 2025, the Company has a working capital deficit of \$1,811,031 (December 31, 2024 – \$1,780,598). As at June 30, 2025, the Company has an accumulated deficit of \$67,147,390 (December 31, 2024 - \$65,881,711) and is not yet generating operating cash flows. As such, there is material uncertainty about the ability of the Company to continue as a going concern. In order to continue as a going concern, the Company requires additional capital to fund ongoing operations and intends on continuing to raise additional funds through the issuance of equity and/or debt.

Capital risk management

The Company's objectives when managing capital are:

- ensuring sufficient liquidity to support its financial obligations and execute its operating and strategic plans;
- maintaining healthy liquidity reserves and access to capital; and
- minimizing the after-tax cost of capital while taking into consideration current and future industry, market and economic risks and conditions.

To assess its effectiveness in managing capital, management monitors certain key ratios to ensure they are within targeted ranges.

The Company defines its capital as its equity. Its capital management objectives and approach were unchanged during the quarter.

SUBSEQUENT EVENTS

On July 23, 2025, the Company announced it had closed the sale of 15 ACP-01 Therapy Convertible Debentures for proceeds of USD\$517,230, subject to the approval of TSXV.

On July 24, 2025, the Company announced it had closed a non-brokered private placement for gross proceeds of \$2,969,600 (the "Offering"). The Offering consisted of the issuance of an aggregate of 29,696,000 Units at a price of \$0.10 per Unit. Each Unit consists of one common share ("Common Share") in the capital of the Company and one common share purchase warrant ("Warrant"), with each full Warrant entitling the holder to acquire one Common Share at a price of \$0.15 per Common Share for a period of 24 months from the closing of the offering. In connection with the Offering, the Company paid eligible finders aggregate cash finder fees of approximately \$97,600 and issued 976,000 finder's options to purchase Common Shares of the Company at an exercise price of \$0.15 per Common Share within 24 months from the closing date of the Offering.

On July 30, 2025, the Company announced that its previously closed non-brokered private placement had been increased from \$2,969,600 to \$3,000,000. In connection with the increased Offering, the Company paid aggregate cash finder's fees of \$100,032 and issued 1,000,320 finder's options. Each finder's option entitles the holder to purchase one Common Shares at an exercise price of \$0.15, exercisable for a period of 24 months from the closing date of the Offering.

The use of proceeds will be allocated to repayment of CD#1 in full at a 50% discount to face value (\$1,250,000) and general working capital in support of the Company's continuing operational expenses, including marketing and sales of VesCell.

DISCLOSURE CONTROLS, PROCEDURES AND INTERNAL CONTROLS OVER FINANCIAL REPORTING

Management has established and continues to complement a system of disclosure controls and procedures and internal controls over financial reporting. This system is designed to provide reasonable assurance that material information relating to the issuer and its subsidiaries are available and reported to senior management and permits timely decisions regarding public disclosure. As of June 30, 2025, the Company's Chief Executive Officer and Chief Financial Officer have evaluated the effectiveness of the design and operation of the Company's disclosure controls and procedures. Based on this evaluation, the Chief Executive Officer and the Chief Financial Officer have concluded that the Company's disclosure controls and procedures, as defined in Multilateral Instrument 52-109 – Certification of Disclosure in Issuer's Annual and Interim Filings are effective, except as noted below, to ensure that the information required to be disclosed in reports that are filed or submitted under Canadian Securities legislation are recorded, processed, summarized and reported within the time period specified in those rules.

The Company's disclosure controls and procedures are indicative of many small and growing companies. Consequently, management has identified certain weaknesses that currently exist in the disclosure controls and procedures including, but not limited to, the segregation of duties and expertise in specific areas of public disclosure. The existence of these weaknesses is partially compensated for by senior management monitoring these issues, and in the case of complex or extraordinary transactions, consulting with external experts to advise management in their analysis and conclusions.

Throughout the year management continued to address, as required, steps to improve disclosure controls and procedures and internal controls over financial reporting. However, no specific changes to disclosure controls and procedures were made during the period. The Company recognizes this is an ongoing and dynamic process and continues to focus on internal controls related to financial reporting and disclosure controls and procedures and is committed to further improvements in the future.

RISKS AND UNCERTAINTIES

Lack of Product Revenues and History of Losses

To date, Hemostemix has not recorded any revenues from the sale of biopharmaceutical products or earning any licensing revenues, and, as a result, it faces a high risk of business failure. Hemostemix expects to incur additional losses during the periods of research and development, clinical testing, and application for regulatory approval of its product candidates. Hemostemix expects to incur losses unless and until such time as payments from corporate collaborations, product sales and/or royalty or license payments generate sufficient revenues to fund its continuing operations.

Ability to Continue as a Going Concern

The Company's auditors' opinion on its June 30, 2025 financial statements includes an explanatory paragraph in respect of there being doubt about the Company's ability to continue as a going concern.

Biotech Public Market Risks

Prospects for companies in the biotechnology industry generally may be regarded as uncertain given the nature of the industry and, accordingly, investments in biotechnology companies should be regarded as speculative. Biotechnology research and development involves a significant degree of risk. An investor should carefully consider the risks and uncertainties described below. The risks and uncertainties described below are not an exhaustive list. Additional risks and uncertainties not presently known to Hemostemix or that Hemostemix believes to be immaterial may also adversely affect Hemostemix's business. If any one or more of the following risks occur, Hemostemix business, financial condition and results of operations could be seriously harmed. Further, if Hemostemix fails to meet the expectations of the public market in any given period, the market price of Hemostemix shares could decline.

Early Stage Development and Scientific Uncertainty

Hemostemix's products are at an early stage of development. Significant additional investment in research and development, product validation, manufacturing, production scale-up, clinical testing, and regulatory submissions of such product candidates is required prior to commercialization. There can be no assurance that any such products will actually be developed. The development and regulatory processes may require access to raw materials and inputs which may not be available to Hemostemix in sufficient amounts or in a timely fashion to allow Hemostemix to complete the development or receive regulatory approval of any product or process. A commitment of substantial time and resources is required to conduct research and clinical trials if Hemostemix is to complete the development of any product. It is not known whether any of these product or process candidates will meet applicable health regulatory standards and obtain required regulatory approvals, or whether such products can be produced in commercial quantities at reasonable costs and be successfully marketed, or if Hemostemix's investment in any such products will be recovered through sales or royalties. The Company's technology will require significant research and development and preclinical and clinical testing prior to regulatory approval, if required, being obtained in the United States or other countries. The Company may not be able to obtain regulatory approvals, if required, to complete necessary clinical trials for its cell technology, or to commercialize it. The Company's technology may prove to have undesirable and unintended side effects, or other characteristics adversely affecting its safety, efficacy or cost-effectiveness could prevent or limit its use. The Company's technology may fail to provide its intended benefit or achieve benefits equal to or better than its competitor's products at the time of testing or production and, if so, its business may fail.

Clinical Trial Risks

The Company's clinical trials may fail to produce successful results or could be suspended due to unacceptable safety risks, which could cause its business to fail. Clinical trials are subject to extensive regulatory requirements, and are very expensive, time-consuming and difficult to design and implement, in part because they may be subject to rigorous regulatory requirements. The Company's products may fail to achieve necessary safety and

efficacy endpoints during clinical trials. The Company believes that its clinical trials will take a substantial period of time to complete. Furthermore, failure can occur at any stage of the trials, and the Company could encounter problems that cause us to abandon or repeat clinical trials. The commencement and completion of clinical trials may be delayed by several factors, including: unforeseen safety issues; lack of effectiveness during clinical trials; slower than expected rates of patient recruitment; and inability to monitor patients adequately during or after treatment. In addition, the Company or regulatory officials may suspend the Company's clinical trials at any time if it appears that the Company is exposing participants to unacceptable health risks. If the Company's clinical trials fail to produce successful results, or are suspended due to unacceptable safety risks, the Company's business may fail.

Additional Financing Requirements and Access to Capital

Hemostemix will require substantial additional funds for further research and development, planned clinical testing, regulatory approvals, establishment of manufacturing capabilities and, if necessary, the marketing and sale of its products. Hemostemix may attempt to raise additional funds for these purposes through public or private equity or debt financing, collaborations with other biopharmaceutical companies and/or from other sources. There can be no assurance that additional funding or partnership will be available on terms acceptable to Hemostemix and which would foster successful commercialization of Hemostemix products.

Government Regulations

Biotechnology and pharmaceutical companies operate in a high-risk regulatory environment. The manufacture and sale of human diagnostic and therapeutic products is governed by numerous statutes and regulations in the United States, Canada, and other countries where Hemostemix intends to market its products. The subject matter of such legislation includes approval of manufacturing facilities, controlled research and testing procedures, review and approval of manufacturing, preclinical and clinical data prior to marketing approval, as well as regulation of marketing activities, notably advertising and labelling.

The process of completing clinical trials and obtaining required approvals is likely to take several years and require the expenditure of substantial resources. Furthermore, there can be no assurance that the regulators will not require modification to any submissions which may result in delays or failure to obtain regulatory approvals. Any delay or failure to obtain regulatory approvals could adversely affect the ability of Hemostemix to utilize its technology, thereby adversely affecting operations. Further, there can be no assurance that Hemostemix's diagnostic product candidates will achieve levels of sensitivity and specificity sufficient for regulatory approval or market acceptance, or that its therapeutic product candidates prove to be safe and effective in clinical trials or receive the requisite regulatory approval. There is no assurance that Hemostemix will be able to timely and profitably produce its products while complying with all the applicable regulatory requirements. Foreign markets, other than the United States and Canada, generally impose similar restrictions.

Hazardous Materials and Environmental Matters

Certain of Hemostemix's research and development processes may involve the controlled use of hazardous materials. Hemostemix is subject to federal, provincial, and local laws and regulations governing the use, manufacture, storage, handling and disposal of such materials and certain waste products. Although management of Hemostemix believes that its procedures for handling and disposing of such materials comply with the standards prescribed, the risk of accidental contamination or injury from these materials cannot be completely eliminated. In the event of such an accident, Hemostemix could be held liable for damages and such liability could exceed the resources of Hemostemix. Hemostemix is not specifically insured with respect to this liability. Although management of Hemostemix believes that it currently complies in all material respects with applicable environmental laws and regulations, Hemostemix may be required to incur significant costs to comply with environmental laws and regulations in the future. Furthermore, there can be no assurance that the operations, business, or assets of Hemostemix will not be materially adversely affected by current or future environmental laws or regulations.

Patents and Proprietary Technology

Hemostemix's success will depend in part on its ability to obtain, maintain, and enforce patent rights, maintain trade secret protection, and operate without infringing the proprietary rights of third parties. There can be no assurance that pending patent applications will be allowed, that Hemostemix will develop additional proprietary products that are patentable, that issued patents will provide Hemostemix with any competitive advantage or will not be challenged by any third parties, or that patents of others will not have an adverse effect on the ability of Hemostemix to do business.

Furthermore, there can be no assurance that others will not independently develop similar products, duplicate any of the Hemostemix products, or design around the products patented by Hemostemix. In addition, Hemostemix may be required to obtain licenses under patents or other proprietary rights of third parties. No assurance can be given that any licenses required under such patents or proprietary rights will be available on terms acceptable to Hemostemix. If Hemostemix does not obtain such licenses it could encounter delays in introducing one or more of its products to the market, while it attempts to design around such patents, or could find that the development, manufacturing or sale of products requiring such licenses could be foreclosed. In addition, Hemostemix could incur substantial costs in defending itself in suits brought against it on such patents or in suits where it attempts to enforce its own patents against other parties.

Until such time, if ever, that patent applications are filed, the ability of Hemostemix to maintain the confidentiality of its technology may be crucial to its ultimate possible commercial success. While Hemostemix has adopted procedures designed to protect the confidentiality of its technology, no assurance can be given that such arrangements will be effective, that third parties will not gain access to Hemostemix trade secrets or disclose the technology, or that Hemostemix can meaningfully protect its rights to its trade secrets.

Dependence on Collaborative Partners, Licensors and Others

Hemostemix activities will require it to enter into various arrangements with corporate and academic collaborators, licensors, licensees and others for the research, development, clinical testing, manufacturing, marketing, and commercialization of its products. Hemostemix intends to attract corporate partners and enter into additional research collaborations. There can be no assurance, however, that Hemostemix will be able to establish such additional collaborations on favourable terms, if at all, or that its current or future collaborations will be successful. Failure to attract commercial partners for its products may result in Hemostemix incurring substantial clinical testing, manufacturing, and commercialization costs prior to realizing any revenue from product sales or result in delays or program discontinuance if funds are not available in sufficient quantities. If any collaborative partner fails to develop, manufacture, or commercialize successfully any product to which it has rights, or any partner's product to which Hemostemix will have rights, Hemostemix's business may be adversely affected. Failure of a collaborative partner to continue to participate in any particular program could delay or halt the development or commercialization of products generated from such program. In addition, there can be no assurance that the collaborative partners will not pursue other technologies or develop alternative products either alone or in collaboration with others, including Hemostemix's competitors, as a means for developing treatments for the diseases targeted by Hemostemix programs.

Furthermore, Hemostemix may hold licenses for certain technologies and there can be no assurance that these licenses will not be terminated, or that they will be renewed on conditions acceptable to Hemostemix. Hemostemix may negotiate additional licenses in respect of technologies developed by other companies and academic institutions. Terms of license agreements to be negotiated may include, inter alia, a requirement to make milestone payments, which may be substantial. Hemostemix will also be obligated to make royalty payments on the sales, if any, of products resulting from licensed technology and, in some instances, may be responsible for the costs of filing and prosecuting patent applications. Should any of Hemostemix licensees breach their regulatory, clinical, operational or legal requirements this may impact Hemostemix reputation

and/or ability to conduct its business or make progress as anticipated.

Rapid Technological Change

The biotechnology and pharmaceutical industries are characterized by rapid and substantial technological change. There can be no assurance that developments by others will not render Hemostemix proposed products or technologies noncompetitive, or that Hemostemix will keep pace with technological developments. Competitors have developed or are developing technologies that could be the basis for competitive products. Some of these products have an entirely different approach or means of accomplishing the desired diagnostic or therapeutic effect as compared with products to be developed by Hemostemix and could be more effective and less costly than the products to be developed by Hemostemix. In addition, alternative forms of medical treatment may be competitive with Hemostemix products.

Competition

Technological competition from pharmaceutical companies, biopharmaceutical companies and universities are intense and is expected to increase. Potential competitors of Hemostemix have or may develop product development capabilities or financial, scientific, marketing, and human resources exceeding those of Hemostemix. Competitors may develop products before Hemostemix develops its own products, obtain regulatory approval for such products more rapidly than Hemostemix, or develop products which are more effective than those which Hemostemix intends to develop. Research and development by others may render Hemostemix's proposed technology or products obsolete or non-competitive or produce treatments or cures superior to any therapy developed or to be developed by Hemostemix, or otherwise preferred to any therapy developed by Hemostemix.

Status of Healthcare Reimbursement

Hemostemix's ability to successfully market certain diagnostic or therapeutic products may depend in part on the extent to which reimbursement for the cost of such products and related treatments will be available from government health administration authorities, private health insurers and other organizations. Significant uncertainty exists as to whether newly approved healthcare products will qualify for reimbursement. Furthermore, challenges to the price of medical products and services are becoming more frequent. There can be no assurance that adequate third-party coverage will be available to establish price levels, which would allow Hemostemix to realize an acceptable return on its investment in product development.

Acceptance of Technology

The Company's success depends on the acceptance of its stem cell technology by the medical community and consumers as a safe and effective solution. The success of its technology will depend on its acceptance by potential consumers and the medical community. Because its technology is new in the treatment of CLI, the long term effects of using its new technology are unknown. The results of short-term clinical trials do not necessarily predict long-term clinical benefit or reveal adverse effects. If results obtained from future commercial experience indicate that its technology is not as safe or effective as other treatments, adoption of this technology by consumers and the medical community may suffer and its business will be harmed.

Potential Product Liability

Pharmaceutical products involve an inherent risk of product liability claims and associated adverse publicity. Product liability insurance is costly, and availability is limited and may not be available on terms which would be acceptable to Hemostemix, if at all. An inability to maintain sufficient insurance coverage on reasonable terms or to otherwise protect against potential product liability claims could prevent or inhibit the commercialization of Hemostemix's products. A product liability claim brought against Hemostemix, or withdrawal of a product from the market, could have a material adverse effect upon Hemostemix and its financial condition.

Manufacturing

Hemostemix product manufacturing was done at a single facility without secondary backup. Hemostemix's ability to manufacture and conduct its clinical trial may depend on its ability to manufacture and ship product in and out of a third-party manufacturing facility.

Reliance on Key Personnel

Hemostemix is dependent on certain members of its management and scientific staff as well as consultants and contractors, the loss of services of one or more of whom could adversely affect Hemostemix. In addition, Hemostemix's ability to manage growth effectively will require it to continue to implement and improve its management systems and to recruit and train new employees. There can be no assurance that Hemostemix will be able to successfully attract and retain skilled and experienced personnel.

Volatility of Share Price, Absence of Dividends and Fluctuation of Operating Results

Market prices for the securities of biotechnology companies, including Hemostemix, have historically been highly volatile. Factors such as fluctuation of Hemostemix operating results, announcements of technological innovations, patents or new commercial products by Hemostemix or competitors, results of clinical testing, regulatory actions, or public concern over the safety of biopharmaceutical products and other factors could have a significant effect on the share price or trading volumes for the common shares. Hemostemix's shares, may be subject to significant price and volume fluctuations and may continue to be subject to significant price and volume fluctuations in the future. Hemostemix has not paid dividends to date and does not expect to pay dividends in the foreseeable future.

Conflict of Interest

Certain of the directors and senior officers of Hemostemix may, from time to time, be employed by or affiliated with organizations which have entered into agreements with Hemostemix. As disputes may arise between these organizations and Hemostemix, or certain of these organizations may undertake or have undertaken research with competitors of Hemostemix, there exists the possibility for such persons to be in a position of conflict. Any decision or recommendation made by these persons involving Hemostemix will be made in accordance with his or her duties and obligations to deal fairly and in good faith with Hemostemix and such other organizations. In addition, as applicable, such directors and officers will refrain from voting on any matter in which they have a conflict of interest.

No Key Management Insurance

The Company does not currently have key management insurance in place in respect of any of its senior officers or personnel.

No Anticipated Dividends

The Company does not intend to pay dividends on any investment in the shares of stock of the Company. The Company has never paid any cash dividends and currently do not intend to pay any dividends for the foreseeable future. To the extent that the Company requires additional funding currently not provided for in its financing plan, its funding sources may prohibit the payment of a dividend. Because the Company does not intend to declare dividends, any gain on an investment in the Company will need to come through an increase in the stock's price. This may never happen, and investors may lose all of their investment in the Company.

Market Disruption Risks

Geopolitical risks such as war and occupation, terrorism, tariffs and trade wars may in the future lead to increased short-term market volatility and may have adverse long-term effects on world economies and markets generally. Those events could also have an acute effect on individual company's or related groups of companies. These risks could also adversely affect securities markets, inflation and other factors from time to time. Such events could, directly or indirectly, have a material effect on the prospects of the Company.

ADDITIONAL DISCLOSURE FOR VENTURE ISSUERS WITHOUT SIGNIFICANT REVENUE

The Company's main focus is to develop, blood-derived cell therapies primarily for the treatment of severe medical conditions not adequately addressed by current treatments.

To achieve commercialization of its products, the Company must obtain regulatory approval in each respective jurisdiction it intends to market its products. Management of Hemostemix believes it may be possible to achieve this in certain jurisdictions on the basis of positive Phase 2 clinical trial data, but in most jurisdictions additional clinical data from larger clinical trials will be required to obtain such approval.

Hemostemix is selling forward its Therapy Convertible Debentures (TCD) to patients who agree to be treated in a jurisdiction that allows autologous stem cell therapy under exemption. To date 23 TCDs are sold. Future revenues will be generated from the conversion of the TCD to an ACP-01 therapy treatment.