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Note on research as a “minor non-monetary benefit” according to the MiFID II regulation: This research meets the requirements for being classified as a “minor non-monetary benefit”. For more information, see the disclosure under “I. Research under MiFID II”

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## **GBC AG is initiating coverage on Defence Therapeutics Inc.**

### **06/12/2020 – GBC Management interview with Sébastien Plouffe, CEO of Defence Therapeutic Inc.**

Company: Defence Therapeutics Inc. <sup>\*5a,5b,7,11</sup>

ISIN: CA24463V1013

Analyst: Julien Desrosiers

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*\*catalogue of potential conflicts of interests on page 5*

### **„We can work on any disease and adapt to any emergent pandemic such as the one related to SARS-CoV2“**

In its current letter to shareholders, Defence Therapeutics reports the successful completion of its protein-based COVID vaccine (AccuVAC-PT001) toxicology studies in non-rodent rabbit model. The AccuVAC-PT001 vaccine is an injectable vaccine capable of inducing a powerful and sustainable antibody response in both rodent and non-rodent models. The vaccine proprietary recipe consists of chemically modifying the COVID Spike protein with the Accum™ moiety. Defence Therapeutics is currently working on two different protein-based vaccine candidates. In Parallel, the company continues the development of their ACCUM™ based cancer treatment R&D. GBC analyst Julien Desrosiers spoke with Defence Therapeutics CEO Sébastien Plouffe:

**GBC AG:** For investors that have never heard about Defence Therapeutics, can you explain what is at the center of your company?

**Sébastien Plouffe:** The highlight of Defence is that it has a proprietary technology platform with very wide versatility application. For instance, our Accum™ molecule can be linked to any protein, RNA or DNA molecules and thus, can be used to target any type of cancer or infectious diseases. Although the company's main approach was to focus on ADCs, we quickly applied the Accum™ technology to fight a common and recent pandemic. In other words, we can work on any disease and adapt to any emergent pandemic such as the one related to SARS-CoV2.

**GBC AG:** How did your company become involved in the ACCUM™ story and secured the rights?

**Sébastien Plouffe:** I was looking to acquire a strong technology in the Biotech sector and in 2016, I was introduced to the Accum™ technology. We were immediately impressed by its huge versatility and potential that we have decided to option it. Defence completed the acquisition in May 2020 and since we are actively working to test and develop all kinds of applications and studies against cancer and infectious diseases.

**GBC AG:** With the ACCUM™ technology, you are currently focusing on two main development models: ACCUM™ Technology and ACCUM™ Vaccine. Can you describe in which development stage are the two pathways and in what way they differ?

**Sébastien Plouffe:** The Accum™ technology is based on the fact that a given molecule can be modified with a small moiety (called the Accum™) to enhance its accumulation in target cells. For example, we can attach the Accum™ on an ADC to enhance its accumulation in target cell or attach it to an antigen and deliver it very efficiently to an antigen presenting cell (example dendritic cells) to prime an immune response.

**GBC AG:** ADC (anti body drug conjugate) are usually tailored made to act on one specific target. They are developed with the triptych concept of Protein, Linker and Payload. These three must play their role in perfect harmony to impact the desired target. How can ACCUM™ circumvent this issue?

**Sébastien Plouffe:** In fact, Accum™ does not interfere with the function of any of these components. On the contrary, it works in concert with them by enhancing the function of that given ADC. In other words, Accum™ does not block or mitigate the linker, payload activity or antibody specificity. It just makes sure that the entire package escapes the endosome and makes it to the intracellular target.

**GBC AG:** Can you specifically discuss the latest success you have achieved with ACCUM™?

**Sébastien Plouffe:** We are developing a large pipeline of products. The Accum™ is being used to develop: i) a DC vaccine targeting 4 different indications (melanoma, breast cancer, lymphoma and colon cancer), ii) an ADC against breast cancer, iii) the use of the "naked" Accum™ molecule as an anti-cancer molecule against breast cancer, iv) two different COVID vaccines including one intranasal, and v) two vaccines targeting HPV and cervical cancer.

**GBC AG:** ADC's FDA approval failure have two main causes: Toxicity and complexity of the ADCs manufacturing challenges. What can we expect from ACCUM™ regarding these two main issues?

**Sébastien Plouffe:** The Accum™ will be applied to an ADC that is already available on the market. We have evidence (using Trastuzumab for example) that Accum™ does not increase the toxicity of the ADC. On the contrary! By improving its potency, the potency of the ADCs can be enhanced by lowering the dosing or shortening the regimen as it would need less of the ADC to achieve strong outcomes. As such, the toxicity of the product would be further improved.

**GBC AG:** ADC are very difficult to manufacture and need exclusive equipment, labs and require a very complex supply chain. As you are currently in the process of selecting a manufacturing partner, and how has the COVID crisis changed your expectations and requirements in terms of manufacturing processes as well as securing the supply chain.

**Sébastien Plouffe:** In fact, we are working on two different models. In the first model, we wish to develop the ADC program in partnership with a large pharma as Accum™ can be used to enhance ANY ADC. In parallel, we are working on developing our own ADC with self-cleavable linkers and Accum™ moieties as a cleavable payload on their own. We are aware that the pandemic is delaying all steps of development, but we are actively working to bypass each of these obstacles by planning our objectives and key steps in advance.

**GBC AG: What team of scientists are driving the two pathways and where is the research conducted?**

**Sébastien Plouffe:** We have an established collaboration with the laboratory of Dr. Moutih Rafei, our VP – Research and Development (Université de Montréal) who is an immunologist by training and has extensive experience in the fields of cell therapy, immune-oncology and infectious diseases. In addition, our co-inventor of the technology, Dr. Simon Beaudoin, a biochemist by training with a strong expertise in immuno-conjugation and optimization of ADCs, is actively working on the continued development of the Accum™ technology at our laboratories located in Montreal (CQIB) and at Sherbrooke. We are currently expanding our scientific team.

**GBC AG: As you described ACCUM™ to be scalable and versatile, how will you select which type of disease you will focus on?**

**Sébastien Plouffe:** We have already established our target indications. We will be working towards a treatment for melanoma, breast cancer, COVID and HPV. These indications were selected strategically using three different criteria: i) the widespread of the indication/cancer, ii) absence of a potent treatment/cure, and iii) the possibility of using our products in combination with commercially-available immune-checkpoint blockers.

**GBC AG: Can you explain us how your latest press release is a game changer for Defence Therapeutics and how well you are financed at the moment?**

**Sébastien Plouffe:** The versatility of the Accum™ technology is a game changer by itself as it allows us to develop a wide range of products targeting different indications. Defence's strategy is to initiate a minimum of 2 Phase I trials in 2022 and to co-develop some products in partnership with large pharmas. We are currently well financed, and we may come to the market in Q1 of 2022 to increase our liquidity to achieve more goals before our venue to the NASDAQ planned for Q2 of 2022.

**GBC: Mr. Plouffe, thank you for the interview.**

**Note: GBC initiates Coverage with this management interview and is working on the initial coverage report.**

## ANNEX

### I.

#### **Research under MiFID II**

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The analysts responsible for this analysis are:

**Julien Desrosiers, Finanzanalyst, Financial Analyst**

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