FORM 7

MONTHLY PROGRESS REPORT

Name of Listed Issuer: Canntab Therapeutics Limited (the "Issuer").

Trading Symbol: PILL

Number of Outstanding Listed Securities: 25,306,601

Date: January 3, 2019

Report on Business

1. Provide a general overview and discussion of the development of the Issuer's business and operations over the previous month. Where the Issuer was inactive disclose this fact.

On December 5, 2018 the Issuer announced the launch of a research partnership with world-renowned orthopaedic surgeon, Dr. Don Garbuz, M.D. (Head of the Division of Lower Limb Reconstruction and Oncology of the Department of Orthopaedics at the University of British Columbia), subject to approval of the University of British Columbia where the proposed clinical trial will take place. The aim of the study is to examine and quantify the efficacy of Canntab products in decreasing the duration and quantity of postoperative use of opioid medications and in the management of preoperative pain. Dr. Garbuz shall be the Lead Investigator for the research study.

The Company is seeking approval from UBC to conduct with Dr. Garbuz as lead investigator, a randomized, double-blind, placebo-controlled clinical trials to determine the efficacy of Canntab's products in helping effectively treat pain in patients. The study will look at the use of Canntab tablets to treat pain after knee replacement surgery. Patient safety and tolerability will also be assessed. Further, Canntab's patent pending filing for addiction treatment therapy for opioids and other painkillers, and its specific proprietary tablet formulation, will be the starting point for the dosing delivery mechanism to be used in the study.

Canntab advises that it will create the tablets for the study in Coburg, Ontario with FSD Pharma, a Licensed Producer under the ACMPR in furtherance of their collaboration and license agreement dated September 17, 2018.

Zeeshan Saeed, President and Founder of FSD Pharma commented, "We are pleased that Canntab will be producing its tablets for this important study at our facility in Cobourg. The aim of this work is clearly in line with FSD Pharma's direction and plans to develop pharmaceutical cannabinoid-based treatments for pain to counter the opioid crisis, including opioid abuse, dependence and overdose."

Dr. Don Garbuz will be overseeing the study as lead investigator pursuant to a consulting agreement to be entered into between Canntab and Dr. Garbuz. Dr. Garbuz is a renowned orthopaedic surgeon and medical researcher, a Professor at the University of British Columbia and is credited with over 120 peer reviewed publications. Dr. Garbuz has been awarded the Hip Society Award for Best Research Paper four times; in 2009, 2011, 2013 and 2017.

"I am excited to be partnering with Canntab to look at conducting a trial that will analyze the efficacy of cannabis as a replacement for opioids, in postoperative care of orthopedic surgical patients undergoing hip/knee replacements at the UBC Hospital. This study will help advance our understanding of the potential effects of cannabis as a treatment for pain and will become the basis for conducting future cannabinoid and pain management clinical trials." said Dr. Don Garbuz, M.D. "This important study which aims to help us better understand the potential impact of cannabis as an alternative for opioids, so that patients might face better outcomes."

There is an opioid epidemic in the United States and Canada. The quality of life and addiction issues that stem from the widespread use of opioids is a major concern for individuals, communities, health care professionals and all levels of government. Most, if not all, patients undergoing total knee and hip arthroplasty use opioid medication to control their pain postoperatively. Their opioid usage is proving to be detrimental to overall health outcomes in some patients, and sometimes fatal. It is not uncommon for patients to misuse these medications after surgery, with nearly 5% of previously opiate-naïve patients becoming chronic abusers. A study from The Centers for Disease Control and Prevention (CDC) found that opiate prescriptions with medication supplies lasting greater than 8 days increased the likelihood of long-term abuse at 1 year (references below).

The financial arrangement calls for an initial upfront fee to create the clinical protocol while securing the participation and, required approvals of the clinical trial by UBC and the independent team comprising the Ethics Committee. Following those approvals, funds will be disbursed partially on certain milestones being achieved. The total cost of the study is estimated at approximately $600,000 with an anticipated time frame to complete of between 12 to 18 months.

2. Provide a general overview and discussion of the activities of management.

   None other than as described above.

3. Describe and provide details of any new products or services developed or offered. For resource companies, provide details of new drilling, exploration or production programs and acquisitions of any new properties and attach any mineral or oil and gas or other reports required under Ontario securities law.

   None other that as described in paragraph 1 above.

4. Describe and provide details of any products or services that were discontinued. For resource companies, provide details of any drilling, exploration or production programs that have been amended or abandoned.

   N/A

5. Describe any new business relationships entered into between the Issuer, the Issuer's affiliates or third parties including contracts to supply products or services, joint venture agreements and licensing agreements etc. State whether the relationship is with a Related Person of the Issuer and provide details of the relationship.

   None other that as described in paragraph 1 above.

6. Describe the expiry or termination of any contracts or agreements between the Issuer, the Issuer's affiliates or third parties or cancellation of any financing arrangements that have been previously announced.

   N/A

7. Describe any acquisitions by the Issuer or dispositions of the Issuer's assets that occurred during the preceding month. Provide details of the nature of the assets acquired or disposed of and provide details of the consideration paid or payable together with a schedule of payments if applicable, and of any valuation. State how the consideration was determined and whether the acquisition was from or the disposition was to a Related Person of the Issuer and provide details of the relationship.

   N/A

8. Describe the acquisition of new customers or loss of customers.

   N/A

9. Describe any new developments or effects on intangible products such as brand names, circulation lists, copyrights, franchises, licenses, patents, software, subscription lists and trade-marks.

   N/A

10. Report on any employee hirings, terminations or lay-offs with details of anticipated length of lay-offs.

    N/A

11. Report on any labour disputes and resolutions of those disputes if applicable.

    N/A

12. Describe and provide details of legal proceedings to which the Issuer became a party, including the name of the court or agency, the date instituted, the principal parties to the proceedings, the nature of the claim, the amount claimed, if any, if the proceedings are being contested, and the present status of the proceedings.

    N/A

13. Provide details of any indebtedness incurred or repaid by the Issuer together with the terms of such indebtedness.

    N/A

14. Provide details of any securities issued and options or warrants granted.

    N/A

15. Provide details of any loans to or by Related Persons.

N/A

16. Provide details of any changes in directors, officers or committee members.

N/A

17. Discuss any trends which are likely to impact the Issuer including trends in the Issuer's market(s) or political/regulatory trends.

N/A

Certificate Of Compliance

The undersigned hereby certifies that:

1. The undersigned is a director and/or senior officer of the Issuer and has been duly authorized by a resolution of the board of directors of the Issuer to sign this Certificate of Compliance.

2. As of the date hereof there were is no material information concerning the Issuer, which has not been publicly disclosed.

3. The undersigned hereby certifies to the Exchange that the Issuer is in compliance with the requirements of applicable securities legislation (as such term is defined in National Instrument 14-101) and all Exchange Requirements (as defined in CNSX Policy 1).

4. All of the information in this Form 7 Monthly Progress Report is true.

Dated, January 3, 2019.

Richard Goldstein
Name of Director or Senior Officer

"signed" Richard Goldstein
Signature
CFO
Official Capacity

Issuer Details Name of Issuer Canntab Therapeutics Limited.	For Month End December, 2018	Date of Report YY/MM/DD 19/01/04
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