FORM 7

MONTHLY PROGRESS REPORT

Name of Listed Issuer: Luminor Medical Technologies Inc. (the "Issuer").

Trading Symbol: LMT

Number of Outstanding Listed Securities: 34,385,060

Date: December 31, 2017

This Monthly Progress Report must be posted before the opening of trading on the fifth trading day of each month. This report is not intended to replace the Issuer's obligation to separately report material information forthwith upon the information becoming known to management or to post the forms required by Exchange Policies. If material information became known and was reported during the preceding month to which this report relates, this report should refer to the material information, the news release date and the posting date on the Exchange website.

This report is intended to keep investors and the market informed of the Issuer's ongoing business and management activities that occurred during the preceding month. Do not discuss goals or future plans unless they have crystallized to the point that they are "material information" as defined in the Policies. The discussion in this report must be factual, balanced and non-promotional.

General Instructions

(a) Prepare this Monthly Progress Report using the format set out below. The sequence of questions must not be altered nor should questions be omitted or left unanswered. The answers to the items must be in narrative form. State when the answer to any item is negative or not applicable to the Issuer. The title to each item must precede the answer.

(b) The term "Issuer" includes the Issuer and any of its subsidiaries.

(c) Terms used and not defined in this form are defined or interpreted in Policy 1 – Interpretation and General Provisions.

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.

   The operates in two segments through two subsidiaries: Jamaica-Blu Ltd. ("JBlu") and Scout Assessment Corp. ("SACorp").

JBlu holds the exclusive Canadian licence of all current and future cannabis commercial products developed by Rise Research Inc. ("RISE"). RISE's cannabis commercial products are based on a patent pending formula, currently filed with the U.S. Patent and Trademark Office, to create precise cannabis-based formulations that may produce specifically targeted effects for various ailments including diabetes. Currently, RISE's portfolio consists of cannabis-based formulations which support patients with low libido called Jamaica blū and Jamaica pnk. The Company plans to launch its JBlu operations for medical use in Canada in Q1 2018 with the anticipation of the recreational use of cannabis in Canada in Q3 2018.

SACorp owns the company's Scout DS® technology. The Scout DS®, which in 90 seconds rapidly tests individuals without blood draw or the need to fast, provides immediate results and is indicated for use for the non-invasive screening of individuals 18 years or older that are at risk for pre diabetes and/or type 2 diabetes. The Scout DS® has received clearance from Health Canada for commercial distribution, has been CE Marked in the European Union, and is also cleared for sale in other markets. Commercial piloting of the technology has taken place in Canada, and preliminary distribution channels have also been established or advanced prior to the Company's involvement with the technology including in countries that may recognize Health Canada and CE regulatory clearances such as Saudi Arabia, Qatar, the United Arab Emirates, India, Jordan, Brazil, Turkey and Kuwait.

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

JBlu's objectives are to be recognized as a leading and trusted brand in the industry in Canada. The Company plans to commence the selling of products, through a licenced producer, into the ACMPR (medical) in Canada by April 1, 2018. The Company plans to commence the selling of products, through a licenced producer, into the adult use market in Canada by program launch day which is anticipated to be July 1, 2018. The Company plans to have 10,000 participants in clinical study by September 30, 2018.

SACorp's business plan includes validating qualified licensing partners who will be responsible to invest in developing geographically defined markets under the guidance and direction of the Company. Since the Company acquired the Scout DS®, several pilots have been conducted by the Company in the pharmacy and employee workplace screening market segments. The Company will use the data, experience and knowledge it has gained about these market segments in order to provide general direction and guidance to newly established licensing partners. The new licensees would be responsible for all market development costs for their

given territory and all costs related to regulatory approval in any given market (if required) under the direction of the Company.

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.

   N/A

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.

   N/A

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.

Security	Number Issued	Details of Issuance	Use of Proceeds[(1)]
Common shares	48,611	Warrant exercise	$14,583 – working capital
Common shares	500,000	Conversion of debenture	N/A

(1) State aggregate proceeds and intended allocation of proceeds.

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 11, 2018.

Chris Carmichael
Name of Director or Senior Officer

Signature
CFO
Official Capacity

Issuer Details Name of Issuer Luminor Medical Technologies Inc.	For Month End December 2017	Date of Report YY/MM/D 18/01/11
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Contact Name Chris Carmichael	Contact Position CFO	Contact Telephone No. (647) 225-4337
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