CERTIFICATE OF REGISTRATION

Princeton BioMeditech Corporation

4242 U.S. Highway 1
Monmouth Junction, New Jersey 08852 UNITED STATES

D-U-N-S ID No. 362917692

UL Medical Regulatory Services of UL LLC®(UL) issues this certificate to the Firm named above, after auditing the Firm's quality management system and finding it in conformance per the defined scope with respect to:

ISO 13485:2016

with additional regulatory requirements listed on final page of this certificate.

The design, development, manufacture, and service of in vitro diagnostic medical devices and in vitro diagnostic analyzers used in the diagnosis of cancer, disease status, drugs of abuse, cardiac markers, fertility testing and pregnancy testing including home use and near patient in vitro diagnostic devices.

The servicing of in vitro diagnostic analyzers used in the diagnosis of cancer, disease status, drugs of abuse, pregnancy testing and cardiac markers.

MDSAP
MEDICAL DEVICE SINGLE AUDIT PROGRAM

Authorized by

Michael J. Windler, P.E.
Manager of Global Regulatory Service
Distinguished Member of the Technical Staff
UL Life and Health Sciences
UL LLC

Check Certificate Status: here

File Number	A12626	Cycle Start Date	October 26, 2018
Certificate Number	1741.181026	Effective Date	October 26, 2018
Initial Issue Date	October 26, 2018	Expiry Date	October 25, 2021

This quality system registration is included in UL's Directory of Registered Firms and applies to the provision of goods and/or services as specified in the scope of registration from the address(es) shown above. By issuance of this certificate the firm represents that it will maintain its registration in accordance with the applicable requirements. This certificate is not transferable and remains the property of UL Medical and Regulatory Services of UL LLC.
Certificates may be verified by visiting the Online Certifications Directory on UL.com.

UL REGISTERED FIRM

UL Medical and Regulatory Services UL, LLC is an MDSAP Recognized Auditing Organization

UL LLC
333 Pfingsten Road
Northbrook, IL 60062-2096 USA

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Additional Regulatory Requirements

Australia:
- Therapeutic Goods (Medical Devices) Regulations, 2002, Schedule 3 Part 1 (excluding Part 1.6) – Full Quality Assurance Procedure

Canada:
- Medical Devices Regulations – Part 1- SOR 98/282

United States:
- 21 CFR 820
- 21 CFR 803
- 21 CFR 806
- 21 CFR 807 – Subparts A to D
- 21 CFR 821 (where applicable)

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