

# Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485:2016

This is to certify that:

Evik Diagnostic Innovations Inc.  
1240 Teron Road, Unit 4  
Kanata  
Ontario  
K2K 2B5  
Canada

Holds Certificate No:

**FM 694551**

and operates a Quality Management System which complies with the requirements of ISO 13485:2016 for the following scope:

Manufacture of reagent beads and assembly of beads into cartridges.



For and on behalf of BSI:

\_\_\_\_\_  
Gary E Slack, Senior Vice President - Medical Devices

Original Registration Date: 2019-02-22

Latest Revision Date: 2020-10-26

Effective Date: 2019-02-22

Expiry Date: 2022-02-21

Page: 1 of 2



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Certificate No: **FM 694551**

Location	Registered Activities
Evik Diagnostic Innovations Inc. 1240 Teron Road, Unit 4 Kanata Ontario K2K 2B5 Canada	Manufacture of reagent beads and assembly of beads into cartridges.
Evik Diagnostic Innovations Inc 105 Schneider Road Kanata Ontario K2K 1Y3 Canada	Manufacture of reagent beads and assembly of beads into cartridges.



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Page: 2 of 2

This certificate remains the property of BSI and shall be returned immediately upon request.  
An electronic certificate can be authenticated [online](#). Printed copies can be validated at [www.bsigroup.com/ClientDirectory](http://www.bsigroup.com/ClientDirectory)  
To be read in conjunction with the scope above or the attached appendix.

Americas Headquarters: BSI Group America Inc., 12950 Worldgate Drive, Suite 800, Herndon, VA 20170-6007 USA  
A Member of the BSI Group of Companies.