

Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485:2016

This is to certify that:

CLEIO Inc.
1505 Royale
Trois-Rivieres
Québec
G9A 4J9
Canada

Holds Certificate Number:

FS 703022

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

Design and Development services for medical device manufacturers, including industrial and user experience design, mechanical, electronics, software and firmware engineering and design quality management.

For and on behalf of BSI:

Graeme Tunbridge, Senior Vice President Medical Devices

Original Registration Date: 2020-05-06

Latest Revision Date: 2023-03-31

Effective Date: 2023-05-06

Expiry Date: 2026-05-05

Page: 1 of 2



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Certificate No: FS 703022

Location	Registered Activities
CLEIO Inc. 4080, boulevard Le Corbusier bureau 201 Laval Québec H7L 5R2 Canada	Design and development of medical devices for various applications.
CLEIO Inc. 1505 Royale Trois-Rivieres Québec G9A 4J9 Canada	Design and Development services for medical device manufacturers, including industrial and user experience design, mechanical, electronics, software and firmware engineering and quality management.



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Page: 2 of 2

This certificate was issued electronically and remains the property of BSI and is bound by the conditions of contract.

An electronic certificate can be authenticated [online](#).

Printed copies can be validated at www.bsigroup.com/ClientDirectory

Issuing Body: BSI Group The Netherlands B.V., John M. Keynesplein 9, 1066 EP Amsterdam, The Netherlands

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Contact Office: 12950 Worldgate Drive, Suite 800, Herndon, VA 20170-6007 USA.