Certificate of Registration of Quality Management System to I.S. EN ISO 13485:2016

The National Standards Authority of Ireland certifies that:

BioTek Instruments, Inc
100 Tigan Street
Highland Park
PO Box 998
Winooski, VT 05404-0998
USA

has been assessed and deemed to comply with the requirements of the above standard in respect of the scope of operations given below:

The Design, Manufacture, Distribution, Service and Installation of In-Vitro Diagnostic Microplate Readers, Automated Microscopes, Washers, Dispensers, Stackers, Incubators, Pipetting Work Stations and Accessories.

Additional sites covered under this multi-site certification are listed on the Annex (File No. MD19.2126)

Approved by:
Geraldine Larkin
Chief Executive Officer

Approved by:
Caroline Dore Geraghty
Director of Medical Devices /
Head of Notified Body

Registration Number: MD19.2126
Certification Granted: August 14, 1996
Effective Date: July 13, 2021
Expiry Date: July 12, 2022

National Standards Authority of Ireland, 1 Swift Square, Northwood, Santry, Dublin 9, Ireland T +353 1 807 3800

Annex to Certificate Number: MD19.2126

Scope of Registration:

The Design, Manufacture, Distribution, Service and Installation of In-Vitro Diagnostic Microplate Readers, Automated Microscopes, Washers, Dispensers, Stackers, Incubators, Pipetting Work Stations and Accessories.

Activity	Location
Headquarters, Design, Manufacturing, Warehouse, Distribution, Service/Repair and Installation	BioTek Instruments, Inc 100 Tigan Street Highland Park Winooski, VT 05404-0998 USA File No: MD19.2126
Administration, Distribution, Warehouse, Service/Repair and Installation	BioTek Instruments GmbH Kocherwaldstrasse 34 D-74177 Bad Friedrichshall Germany File No: MD19.2126/A

Verified by:
Operations Manager