UKcert
Certification & Inspection

Certificate of Registration

This is to certify that the
Quality Management System for Medical Device
of

Labdex Ltd.

at

71-75 Shelton Covent Garden, London WC 2H 9JQ United Kingdom

has been independently assessed and
is compliant with the requirements of:

ISO 13485:2016

For the following scope of activities:

Design, Manufacturing & Exports of Lab Equipment including Ovens, Incubators , Sample Concentrators, Shakers, Mixers, Homogenizers, Stirrers, Spectrophotometers, Laboratory Furnaces & Autoclave, Centrifuges, Liquid Handling Products, General Lab, FTIR, Balances, Products & Lab Supplies

Certificate Number : UQ-52543

Date of Initial Registration	04th April 2019
Date of this Certificate	04th April 2022
Certificate Expiry	03rd April 2023
Recertification Due (subject to the company maintaining its system to the required standard)	03rd April 2025

Authorised Signatory

UKAF

UK Certification & Inspection Ltd.

This certificate is the property of UK Certification & Inspection Limited and shall be returned immediately on request.
71-75 Shelton Street, Covent Garden, London, WC2H 9JQ, United Kingdom

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Certificate of Compliance

CE

We hereby declare that the technical file of product complied with the requirement of directives 2014/35/EU Low Voltage Directive & 2014/32/EU Measuring Instruments Directive.

Certificate No. : CE-9617

Manufacturer

Name : **Labdex Ltd.**

Address : **71-75 Shelton Covent Garden, London WC 2H 9JQ UK**

Products : **Manufacturing, Development and Distribution of Laboratory Equipment Including Autoclaves, Balances, Centrifuges, Homogenizers, Laboratory Incubators, Labwares, Laboratory Shakers & 3D Rockers, Laboratory Furnaces, Magnetic Stirrer and Hot Plates, Mixer and Vortexers, FTIR, Microplate Readers and Washers, Oil Bath, Sample Concentrators, Sieve Shakers & Spectrophotometers**

The Certification body has performed an audit of the above product quality system covering the design, manufacture and final inspection of the certified product. The quality system has been assessed, approved and is subject to continuous surveillance according to Directive 2014/35/EU Low Voltage Directive & 2014/32/EU Measuring Instruments Directive.

This certificate is issued under the following conditions:

1. It applies only to the quality system maintained in the manufacture of above referenced models and it does not substitute the design or type-examination procedures, if requested.
2. The certificate remains valid until the manufacturing conditions or the quality systems are changed.
3. The certificate validity is conditioned by positive results or surveillance audits.

The CE mark as shown above can be used, under the responsibility of the manufacturer, after completion of an EC Declaration of conformity and compliance with all relevant EC Directives. The statement is based on a single evaluation of one sample of above mentioned product. It does not imply an assessment of the whole production.

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