

# 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

**04<sup>th</sup> April 2019**

Date of this Certificate

**04<sup>th</sup> April 2022**

Certificate Expiry

**03<sup>rd</sup> April 2023**

Recertification Due (subject to the company maintaining its  
system to the required standard)

**03<sup>rd</sup> April 2025**

*Daniel..*

Authorised Signatory



# Certificate of Compliance



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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