Certification & Inspection

Certificate of Compliance

CE

Gonfirms that the technical life of Product complied with the Requirement of ln Vitro Diagnostic Medical Devices Directive 98/79/EC

Certificate Number: UQ-7228

Manufacturer

COMPANY NAME :- LABY INSTRUMENTS INDUSTRY
FACTORY ADDRESS :- 62, FIRST FLOOR, INDUSTRIAL ESTATE, AMBALA CANTT-133006
PRODUCTS :- BENCH TOP LABORATORY CENTRIFUGE, MICRO-CENTRIFUGE, HEMOCRITE CENTRIFUGE, BENCH TOP REFRIGERATED CENTRIFUGE, PRP CENTRIFUGE, CYTO CENTRIFUGE, BLOOD BAG REFRIGERATED CENTRIFUGE, GEL ELECTROPHORESIS HORIZONTAL AND VERTICAL, PAPER ELECTROPHORESIS HORIZONTAL AND VERTICAL, DIGITAL AND ANALOG POWER SUPPLIES FOR ELECTROPHORESIS, HAEMOGLOBINMETER (SAHLI'S), HAEMOGLOBINMETER METER DIGITAL MICROPROCESSOR BASED MAGNETIC STIRRER WITH CERAMIC TOP, PAPER CHROMATOGRAPHY KIT, PH METER DIGITAL, PHOTO ELECTRIC COLORIMETER DIGITAL, U.V TRANSILLUMINATOR, NEEDLE & SYRINGE DESTROYS, GEL DOCUMENTATION SYSTEM, ANAEROBIC CULTURE JAR, BLOOD BAG TUBE SEALER (PORTABLE ELECTRIC), WESTERN BLOTTING SYSTEM, GEL CASTER, VORTEX MIXTURE, CELL WASH CENTRIFUGE MACHINE, FLOOR MODEL REFRIGERATED CENTRIFUGE MACHINE, NON REFRIGERATED CENTRIFUGE MACHINE, REFRIGERATED CENTRIFUGE MACHINE FOR GENERAL AND RESEARCH PURPOSE, MEDICAL CENTRIFUGE.

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 Vitro Diagnostic Medical Devices Directive 98/79/EC

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 with applicable CE requirements or The quality systems are changed.
3. The certificate validity is conditioned by positive results or surveillance audits.
4. After fulfilling the relevant Standard testing performance, the manufacturer shall affix to each device, of the referenced models
5. 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.

Validity of this certificate can be verified at www.ukcert.uk/verify

Date of Certification	23. July. 2025
1 st Surveillance Audit Due	22. July. 2026
2 nd Surveill ance Audit Due	22. July. 2027
Certificate Expiry (subject to the company maintaining its)	22. July. 2028

system to the required standard)

Authorised Signatory

CE

UK Certification & Inspection Ltd.