Certificate No. :UQ-23069

Rev No. : 01.01

Certificate

OF COMPLIANCE

We hereby declare that the technical file of the product confirms with the requirement of Medical Device Regulations 2002 (SI 2002 n. 618) (UK MDR 2002) as amended by Medical Device Regulation. (EU Exit) Regulation 2020 (MDRs 2020).

Manufacturer: ABDOS LABTECH PRIVATE LIMITED

Address : PLOT NO-01 SHIV GANGA INDUSTRIAL ESTATE LAKESHWARI BHAGWANPUR, 247667(UK) INDIA

Products : Micro Tips(Last Drop & Filter) , Micro Centrifuge Tubes (Last Drop), Centrifuge Tubes (MaxiRCF), Cryo Vials, Storage Vial, Freezing Tubes, Ria Vial/Plastic Tube & Cap, Sample Container, Petridishes, Pipettes (Elegant, Premium, Premium Plus Electronic), Rave, Stepmate Pipette, Equipment's(Swirl, Swirlex, Swirltop, Thermomix, Hotblock, Wavex, Aerovac), Tissue Culture (Flask, Plate & Dishes), Silicone Mats, Magnetic Stirrer Bar Erlenmeyer Flask, Gloves, Pasteur Pipettes, Serological Pipettes, Beaker, Measuring Cylinder,PCR Tubes & Plates, Screw Cap Vial, Bioreaction Tube, Roller Bottles, Media bottles, Volumetric & Conical Flask, Bottles, Carboys, Aspirators, Biohazard & Autoclavable Bags, Racks & Boxes, Loops, Spreader & Scrapers, Scoops, Spatula, Funnel, Bottle Top Filtration System, Storage & Freezer Colour Change Rack, Deepwell Plates,Vaccum Pump & Desiccator, Electrophoresis & Power Supply, Sharp Container, Bottle Top Dispenser and UV Transilluminators

The Certification body has performed an audit of the above product quality system covering design, manufacture and final inspection of the certified product. The Quality system has been assessed, approved and is subject to continuous surveillance according to the Medical Device Regulations 2002 (SI 2002 n. 618) (UK MDR 2002) as amended by Medical Device Regulation. (EU Exit) Regulation 2020 (MDRs 2020).

Technical File Ref:- ALP-TC01

This Certificate is issued under the following Conditions:

1. It applies only to the quality system maintenance 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 UKCA Mark as shown below can be used, under the responsibility of the manufacturer, after completion of an EC Declaration of conformity and compliance Medical Device Regulations 2002 (SI 2002 n. 618) (UK MDR 2002) as amended by Medical Device Regulation. (EU Exit) Regulation 2020 (MDRs 2020),

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.

Date of initial registration : Dec 02, 2023

Certificate expiry : Dec 01, 2025

Date of this certificate : Nov 08, 2024

Re-certification due : Dec 02, 2026

To check the status of the certificate, visit: https://www.eac-council.org/validity.php

(Authorized Signatory)

European Certification Council
101,Colbrand Grove Birmingham, B15 2BS,United Kingdom
www.eac-council.org info@eac-council.org