Certificate of Compliance

CE

We hereby declare that the technical file of product complied with the requirement of EU Council Directive 98/79/EC of 27 October 1998 on (Regulation (EU) 2017/746 on in vitro diagnostic medical devices).

Certificate No.: CE-6562

Manufacturer

Name : **ABDOS LABTECH PRIVATE LIMITED**

Address : **Plot No-1, Shivganga Industrial Area, Lakheswari Bhagawanpur, Roorkee, Uttarakhand-247667, 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/Fixed, Premium Plus Electronic, MultiChannels), 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 (Dermagaurd/Dermagaurd Lite), Pasteur Pipettes, Serological Pipettes, Beaker, Measuring Cylinder, PCR Tubes/Strips & Plates, Screw Cap Vial (Hing/Loop), Bioreaction Tube, Roller Bottles, Media Bottles, Biofill, 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, Vacuum Pump & Desiccators, Electrophoresis & Power Supply, Sharp Container, Bottle Top Dispenser, UV Transilluminators, Safe Bottle Carrier, CT Bottles, Rodac/Contact Plates, Ready to use Plates, Mask, Empty Tip Boxes, Reservoirs & All Products as per ABDOS Catalogue**

Complies with the requirements applicable to it

The quality system file has been assessed, approved and is subject to continuous surveillance according to the EU Council Directive 98/79/EC of 27 October 1998 on (Regulation (EU) 2017/746 on in vitro diagnostic medical devices)

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.
4. After fulfilling the relevant EU legislation, 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.ukcertifications.org.uk/verify

Date of initial Registration	24th March 2025
Date of this Certificate	24th March 2025
Certificate Expiry	23rd March 2028
Recertification Due (subject to the company maintaining its system to the required standard)	23rd March 2028

Authorised Signatory

UK Certification & Inspection Ltd.

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Company No. 11847851