



## Certificate of Compliance

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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:

- . 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

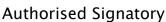
Date of this Certificate 24th March 2025

Certificate Expiry 23rd March 2028

Recertification Due (subject to the company maintaining its 23rd March 2028

system to the required standard)





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
Website:- www.ukcertifications.org.uk, email:- info@ukcertifications.org.uk
Company No. 11847851