Certificate of Compliance

CE

EURO CERT

We hereby declare that the technical file of product complied with the requirement of **Medical Devices Regulation (EU) 2017/745**

Certificate No.: CE-3076

Name	:	**DOLPHIN PHARMACY INSTRUMENTS PVT. LTD.**
Address	:	**J.B.F.HOUSE, GROUND FLOOR, 13, OLD POST OFFICE LANE, KALBADEVI ROAD, MUMBAI – 400002. INDIA**
Product	:	**MANUFACTURER AND EXPORTER AND STOCKIST OF PHARMACY INSTRUMENTS, SCIENTIFIC INSTRUMENTS, GLASSWARE, LABORATORY CHEMICALS, CRUDE DRUGS, OILS**

The quality system file has been assessed, approved and is subject to continuous Surveillance according to the Council Directive on **Medical Devices Regulation (EU) 2017/745**

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. .

Validity of this certificate can be verified at www.eurocertverify.com

Date of Certification	5th August 2023
1st Surveillance Audit Due	4th August 2024
2nd Surveillance Audit Due	4th August 2025
Certificate Expiry (subject to the company maintaining its system to the required standard)	4th August 2026

Authorised Signatory