LORCAN & FYON

CE

CERTIFICATE OF CE COMPLIANCE

Certificate: PAK/1997039/39-LF

Application of MDR 2017/745 for Class IIa & IIb Medical Devices

This to Certify that the Submitted Products Fall Under:

CLASS IIa & IIb MEDICAL DEVICES

Sterile Single Use Surgical & Dental Instruments,
Electro Surgical Instruments, Implants.

Manufactured by:

PROWA MEDICAL INSTRUMENTS

Address to which this Certificate refers

P.O. Box 3036, 9-KM Daska Road, Sialkot 51310 - Pakistan

These products meet all relevant requirements of the Medical Device Regulation (MDR) 2017/745. The technical file for these devices,including the EC Declaration of Conformity in line with Annex IV, has been thoroughly reviewed, verifying compliance with the applicable MDR standards.

Limitations:

The manufacturer is obligated to notify LORCAN & FYON of any significant modification to the products or manufacturing processes to ensure this certificate's continued validity.

Date of Initial Registration: 13-01-2025

Certificate Renewal: 12-01-2026

Certificate Issued on: 13-01-2025

This Certifie of Registration is granted in accordance with the Board-approved regulations.

AUTHORIZED SIGNATORY

LORCAN & FYON

LORCAN & FYON

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LORCAN & FYON

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