



LORCAN & FYON



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 modifications to the products or manufacturing processes to ensure this certificate's continued validity.

Date of Initial Registration: 13-01-2025

Certificate Renewal: 08-01-2027

Certificate Issued on: 09-01-2026

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

AUTHORIZED SIGNATORY



LORCAN & FYON B.V.



AUTHORIZED SIGNATORY



LORCAN & FYON B.V.

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