LORCAN & FYON

CE

CERTIFICATE OF CE COMPLIANCE

Certificate: PAK/1997035/35-LF

Application of MDR 2017/745 for Class I Medical Devices

This to Certify that the Submitted Products Fall Under:

CLASS I MEDICAL DEVICES

(Re-useable Surgical and Dental Instruments)

Manufactured by:

PROWA MEDICAL INSTRUMENTS

Address to which this Certicate 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. T he 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: 09-01-2025

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

EU REP CERTIFICATE

AUTHORIZED SIGNATORY

LORCAN & FYON B.V .

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