

# CE Compliance Certificate



Application of Medical Device Regulation MDR 2017/ 745 for Class I Medical Devices.

This is to certify that the products submitted are:

**MEDICAL DEVICES CLASS I  
(Re-Useable Surgical and Dental Instruments)  
Registration no DCS/126672008-A**

Manufactured By:

**MEDITOOLS INTERNATIONAL  
FAC No. 33, S.I.E., No 1, Gole Canteen, Sialkot – Pakistan.**

Comply with the applicable requirements of the MDR 2017/ 745, The Technical file of the devices have been assessed, including the EC declaration of conformity according to the Annex IV confirming that the medical devices are fulfilling the applicable requirements of MDR 2017/745.

#### Limitations:

The manufacturer must inform DCS of any substantial changes occurred in the Product or process in order to examine whether this certificate remains valid.

CHAIRMAN

SCHEME MANAGER

Certificate Issue Date: February 11, 2026

Certificate Expiry Date: February 10, 2027

This Certificate of Registration is granted subject to the Regulations approved by the Board

[www.dynamexcertification.org](http://www.dynamexcertification.org)

