CE Compliance Certificate



Application of Council Directive 93/42/EEC of 14 June 1993 as updated directive 2007/47/EEC for Class I Medical Devices.

This is certify that the products submitted are:

MEDICAL DEVICES CLASS I

(Re-Useable Surgical and Dental Instruments)
Registration no DCS/12120625137-A

Manufactured By:

WITTEX INTERNATIONAL

Wazirabad Road, Harar, 51310, Sialkot-Pakistan.

Comply with the applicable requirements of the Directive 93/42/EEC as updated directive 2007/47/EEC, The Technical file of the devices have been assessed according to the procedure of conformity Assessment described in the Module A, Annexure V.

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

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Certificate Issue Date: May 04, 2023 Certificate Expiry Date: May 03, 2024
This Certificate of Registration is granted subject to the Regulations approved by the Board

www.dynamexcertification.org

