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/1247322549-A

Manufactured By:

OREBRO INTERNATIONAL

P.O. Box 2468, Street No. 12, Khan Mahal Road, Tariq Pura,
Sialkot, 51310 – 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

Certificate Issue Date: 24 July, 2024
Date of Initial Registration : 24 July ,2024

Certificate Expiry Date: 23 July, 2025
Re-assessment Date :23 July,2027

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

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