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/412531-A

Manufactured By:

DIVERSITY CORPORATION (FZE)

Office No. R3-13A, P.O. Box 121038, SAIF Zone Sharjah,
United Arab Emirates.

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: May 27, 2021

Certificate Expiry Date: May 26, 2022

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

www.dynamexcertification.org