# 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/597716-A**

Manufactured By:

## RENIX INTERNATIONAL

**NAWAN PIND ARRAYIAN, IQBAL PURA,**
**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

Certificate Issue Date: January 06, 2022 Certificate Expiry Date: January 05, 2023

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

www.dynamexcertification.org