

CERTIFICATE



CE

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 ENT Instruments)
Registration no DCS/857262**

Manufactured By:

**[REDACTED] INSTRUMENTS CO.
11-B Model Town Ugoki, 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 VII.

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: November 28, 2014 Certificate Expiry Date: November 27, 2015
This Certificate of Registration is granted subject to the Regulations approved by the Board

www.dynamexcertification.org

