CERTIFICATE

DCS

Dynamex Certification Services

CE

Registration No. DCS/9636170

Application of Council Directive 93/42/EEC as updated directive 2007/47/EC for Class I Medical Devices

This is certifying that the products submitted are:

MEDICAL DEVICES CLASS I

(Re-Useable, Non-Powered Surgical and Dental Instruments)

Manufactured By:

METRO MEDICAL

Metro Medical 35-B, Industrial Estate, P.O.Box # 943, Sialkot-Pakistan

Comply with the applicable requirements of the Directive 93/42/EEC as updated directive 2007/47/EC for Class I Medical Devices and Manufacturer is registered with Medicines and Health Regulatory Agency (MHRA) vide reference CA016396 dated 21 August, 2017 http://aic.mhra.gov.uk/era/pdr.nsf/Search?SearchView&Query=%5BManName%5DContains(Metro+Medical)&SearchOrder=4

The Technical file of the products have been assessed according to the procedure of Conformity Assessment described in the Annex -I, Annex 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. Conformance to all the regulatory requirements is the sole responsibility of the manufacturer including the appointment of EU Authorized Representative and registration with concerned competent authority

CHAIRMAN

SCHEME MANAGER

Certificate Issue Date: August 21, 2018

Certificate Expiry Date: August 20, 2019

DYNAMEX CERTIFICATION SERVICES

DCS

www.dynamexcertification.org