

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 Dental Instruments)
Registration no DCS/13530283**

Manufactured By:

**SARFRAZ AND BROTHERS
PAKKI KOTLI DASKA ROAD NEAR SYEDA AYESHA MADARASA,
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: May 09, 2017

Certificate Expiry Date: May 08, 2018

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

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ATTESTED TO BE TRUE COPY

