

# CERTIFICATE



**Registration No. DCS/9479903**

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:

**CLASS I MEDICAL DEVICES  
(Re-Useable, Non-Powered Surgical Instruments)**

Manufactured By:

**SURGICON PVT LTD**

**P.O. Box: No. 244, Khadim Ali Road, Sialkot-Pakistan**

Comply with the applicable requirements of the Directive 93/42/EEC as updated directive  
2007/47/EC for Class I Medical Devices

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

Issue Date: June 29, 2018

SCHEME MANAGER

Expiry Date: June 28, 2019

