



Certificate of Compliance

Technical file of the company mentioned below has been observed 93/42/EEC for Medical device directive has been taken as references for the processes.

Product Requirements of the organization has been assessed in accordance with the Directive applicable of product for: **Surgical, General Surgery, Oral Surgeries, Plastic/Reconstructive Surgeries, Micro-Surgery, Liposuction, Orthopedic, Ophthalmic, Endoscopic, ENT, Hospital, Hollow Ware & Veterinary Instruments**

Certificate Number: TCE/PK87/01842904
Company Name: **IMPLANT SURGICAL COMPANY**
Company Address: Fateh Garh, Ghos Pura, Sialkot 51310 - Pakistan

Applicable EC Directives & Annex: the mentioned products has been compliance with the essential requirements specified in the EU Medical Device Directive MDD 93/42/EEC as amended by 2007/47/EC Annex VII.

This certificate permits the above named organization to issue declarations of performance in support of CE Marked product with to the scope of certification as stated within this certificate; thereby confirming to interested parties the above named organization has produced the product to the agreed specification.

Initial Certification Date: 22 April 2022
Issue Date: 10 June 2024
Expiry Date: 09 June 2025
Revision: 00



This Certificate is issued against the product sample conforming to the essential requirements as per the Directive. The product liability rests with the manufacturer or his representative in accordance with Council Directive 85/374/EEC.

After preparation of necessary technical documentation as well as the EC conformity declaration the required CE marking can be affixed on the product. Other relevant directives have to be observed.



To verify the certificate please scan the above QR code.

Approved on Behalf of the Certification Board



Signature By Scheme Manager