

## CERTIFICATE OF COMPLIANCE

This is to certify, that the hereunder described items of Medical Devices Directive 93/42/EEC as amended by Directive 2007/47/EC have proven their conformity to the Safety and Health requirements of the Directive.

Manufactured By: **IIQ INSTRUMENTS**

Address: Wazirabad Road, Noul Chowk Dhinglay  
51310 - Sialkot Pakistan

Products: **SURGICAL INSTRUMENTS , DENTAL INSTRUMENTS  
ORTHOPEDIC INSTRUMENTS  
ENT DIAGNOSTIC INSTRUMENTS  
PODIATRY INSTRUMENTS  
HOLLOWWARE AND SURGICAL / DENTAL SETS**

Classification: **Class I**  
re-usable, non-powered and non-measuring devices  
(according to the Manufacturer's declaration)

The product liability rests with the manufacturer by applicable regulations and standards, after fulfilling the relevant EU legislation requirements and withdrawing an EC declaration of conformity as per Annexure-VII, module A of the product, you have therefore licensed to CE mark the product(s) listed above per Article 17 of the Medical Device Directive. The manufacturer is responsible for the CE Marking process and is not to be exempted from carrying out all necessary compliance activities.

*This certificate shall not be reproduced except in full and remains the property of CNC Services, which must be returned on request. The holder must inform CNC Services of any substantial changes that occurred in the product or process to examine whether this certificate remains valid.*



For CNC Services

**CE**

  
Project Engineer

Date of Issue: 16<sup>th</sup> February 2025  
Date of Expiry: 15<sup>th</sup> February 2026  
Certificate Number: **EC/5377/25**