Certificate of Compliance

We hereby declare that the technical file of product complied with the requirement of **Medical Devices Directive 93/42/EEC.**

Manufacturer:

Name : SEGURO LIFESCIENCES

Address: PLOT 2 (i) A EPIP PHASE 1, JHARMAJARI, BADDI, DIST. SOLON. HIMACHAL PRADESH 173205, INDIA

Product: ANGIOGRAPHY ACCESSORIES, INFLATION DEVICE, Y CONNECTOR, MANIFOLD, INTRODUCER NEEDLE, INTRODUCER SHEATH, HYDROPHILIC,PTFE,PTCA GUIDE WIRE, ANGIO KIT, ANGIOGRAPHY & PERIPHERAL CATHETER, GUIDING & EXTENSION CATHETER, SC,NC,CTO HIGH PRESSURE,SCORING,CUTTING PTCA BALLOON CATHETER, STEARABLE MICROCATHETER, SUTURE, SEGURO SOLIDIFIER, ABSORB+, RADIAL BAND, TUBINGS,HIGH PRESSURE SYRINGES

The certification body has performed an audit of the above product quality system covering the design, manufacture and final Inspection of certified product. The quality system has been assessed, approved and is subject to continuous surveillance according **Medical Devices Directive 93/42/EEC.**

This certificate issued under the following conditions:

1. It applies only to the quality system maintained in the manufacturing of above referenced models and it does not substitute the design or type examination procedures, if requested
2. The certificate remains valid until the manufacturing conditions or the quality system are changed
3. The certificate validity is conditioned by positive results of surveillance audits
4. After fulfilling relevant EEC legislation, the manufacturer shall affix to each device, of the referenced models The CE mark as shown below can be used under the responsibility of the manufacturer after completion of an EEC Declaration of conformity and compliance with all relevant EEC Directives. The statement is based on a single evaluation of one sample of above mentioned product. Not imply an assessment of the whole production.

:: Certificate No :: CE/26M03998

Date of initial registration:	**01 July 2026**
First Surveillance Audit on or before:	**30 June 2027**
Second Surveillance Audit on or before:	**30 June 2028**
Re-certification Due:	**30 June 2029**

This Certificate is property of MQA and remains valid Subject to satisfactory surveillance audits.

Authorized Signatory

MQA CERTIFICATION SERVICES
130 Thessaly Rd, Nine Elms, London
SW8 5EJ, United Kingdom

UKAF-CB-011

To check validity of the certificate please visit at www.mqacertification.com