

# EC Declaration of Conformity



**Manufacturer:**

**SCW MEDICATH LTD.**

NO.4, Baolong 6th Road, Baolong Industrial  
Town, Longgang District, Shenzhen, China

We, the manufacturer, herewith declare that the products

**whose single Authorized Representative:**

**OBELIS S.A.**

Bd. Général Wahis 53  
1030 Brussels, Belgium

## Guide Wires

Models: PTFE coated Guide wire, Seldinger Guide wire

meet the provisions of Directive 93/42/EEC which apply to them.

The medical device has been assigned to class IIa according to Annex IX Rule 6 of the Directive 93/42/EEC. It bears the mark

**CE 0197**

The product concerned has been manufactured under a quality management system according to Annex II of Directive 93/42/EEC.

Compliance of the designated product with the Directive 93/42/EEC has been assured via assessment of the quality management system by the Notified Body

**TÜV Rheinland LGA Products GmbH**  
**Tillystraße 2**  
**90431 Nürnberg**  
**Deutschland**

Certificate No.: HD 60139711 0001

Issue date: 05.08.2019

Expiry date: 27.05.2024

following the procedure relating to the EC Declaration of Conformity set out in Annex II of Directive 93/42/EEC.

This Declaration of conformity is valid in connection with the release document for the respective batch of produced devices.

The above mentioned declaration of conformity is exclusively under the responsibility of

**SCW MEDICATH LTD.**  
**NO.4, Baolong 6<sup>th</sup> Road, Baolong Industrial Town,**  
**Longgang District, Shenzhen, China**

Shenzhen, 2019/11/07

Place, date

Miriam Xie, RA

Legally binding signature, Function