



**EC Certificate**  
**Directive 93/42/EEC Annex II, excluding Section 4**  
**Full Quality Assurance System**  
**Medical Devices**

**Registration No.:** HD 60144232 0001

**Report No.:** 17047213 010

**Manufacturer:** SCW Medicath Ltd.  
No. 4 Baolong 6th Road  
Baolong Industrial Town  
Longgang District, Shenzhen  
518116 Guangdong  
P.R. China

**Products:** Medical Devices  
  
(see attachment for products included)  
  
Replaces Approval, Registration No.: HD 60139711 0001

**Expiry Date:** 2024-05-26

The Notified Body hereby declares that the requirements of Annex II, excluding section 4 of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex II, section 5 of the aforementioned directive. For placing on the market of class III devices covered by this certificate an EC design-examination certificate according to Annex II, section 4 is required.

**Effective Date:** 2020-05-26

**Date:** 2020-05-26

Notified Body

Fuxiu Sheng



**TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg**  
TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.

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

**Attachment to  
Certificate**

**Registration No.:** HD 60144232 0001  
**Report No.:** 17047213 010

**Manufacturer:** SCW Medicath Ltd.  
No. 4 Baolong 6th Road  
Baolong Industrial Town  
Longgang District, Shenzhen  
518116 Guangdong  
P.R. China

**Products:**

- Disposable Pressure Transducers
- Introducer Sets
- Guide Wires
- Angiographic Syringes
- Hemodialysis Catheterization Kits
- Patient-Controlled Analgesic Infusion Pumps
- Disposable Infusion Pumps
- Tracheostomy Tube Kits
- Percutaneous Nephrostomy Sets
- Ureteral Stent Sets
- Drainage Catheter Sets
- Transradial Introducer Sets
- Introducer Needles
- I.V Cannulas
- Cervical Ripening Balloon
- Postpartum Balloon

**Date:** 2020-05-26

**Notified Body**



**Fuxiu Sheng**



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

**Attachment to  
Certificate**

**Registration No.:** HD 60144232 0001  
**Report No.:** 17047213 010

**Manufacturer:** SCW Mediatech Ltd.  
No. 4 Baolong 6th Road  
Baolong Industrial Town  
Longgang District, Shenzhen  
518116 Guangdong  
P.R. China

**Products:**

- Locking Drainage Catheters
- Percutaneous Access Sets
- ERCP Guidewires
- Manifolds
- Stopcocks
- Manifold Sets
- Connecting Tubings

Aspects of manufacture concerned with securing and maintaining sterile conditions:

- Dose-control Syringes
- Balloon Inflation Devices
- Colored Piston Specialty Syringes
- Infusion Sets with Needleless Adapters
- Pressure Bandages
- Hemostasis Valve Sets
- Injection Caps

**Date:** 2020-05-26

**Notified Body**



**Fuxiu Sheng**



TÜV Rheinland LGA Products GmbH • 51105 Köln

*SCW Medicath Ltd.  
No.4, Baolong 6th Road,  
Baolong Industrial Town,  
Longgang District, Shenzhen,  
518116 Guangdong,  
P.R. China*

Contact

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Date April 02, 2024

### **Notified Body Confirmation Letter**

Reference. : 10924200

To whom it may concern,

### **Confirmation of the status of a formal application, written agreement, and appropriate surveillance in the framework of Regulation EU 2023/607 amending Regulations (EU) 2017/745 and (EU) 2017/746 as regards the transitional provisions for certain medical devices and in vitro diagnostic medical devices.**

This letter confirms that **TÜV Rheinland LGA Products GmbH**, a Notified Body (NB) designated against Regulation (EU) 2017/745 (MDR) and identified by the number **0197** on NANDO, has received a formal application in accordance with Section 4.3, first subparagraph of Annex VII of MDR and has signed a written agreement in accordance with Section 4.3, second subparagraph of Annex VII of MDR with the following manufacturer:

*SCW Medicath Ltd.  
No.4, Baolong 6th Road,  
Baolong Industrial Town,  
Longgang District, Shenzhen,  
518116 Guangdong,  
P.R. China*  
SRN Number (if available): CN-MF-000019140

The devices covered by the formal application and the written agreement mentioned above are identified in the tables below. Table 1 identifies the devices for which an MDR application has been received, written agreement concluded and for which the NB is also responsible for appropriate surveillance under the applicable Directive. Table 2 identifies the devices for which an MDR application has been received and a written agreement concluded, but the NB has not yet taken the responsibility for appropriate surveillance of the corresponding devices under the applicable Directive.

In the case of devices covered by certificates issued under Directive 90/385/EEC (AIMDD) or Directive 93/42/EEC (MDD) that expired after May 26, 2021 but before March 20, 2023 without having been withdrawn, this letter also confirms that the manufacturer either signed the written agreement under MDR by the date of MDD/AIMDD certificate expiry; or provided evidence that a competent authority of a Member State had granted a derogation or exemption from the applicable conformity assessment procedure in accordance with Article 59(1) of MDR or Article 97(1) of the MDR respectively, by March 20, 2023 for the relevant devices.

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LGA Products GmbH

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Board of Management

Dipl.-Ing.  
Thomas Weigand, Spokesman

Dipl.-Kfm.  
Dr. Jörg Schlösser

Nuremberg HRB 26013  
VAT No.: DE 811835490

Chairman of the  
Supervisory Board

Dr.-Ing. Michael Fübi

The transition timelines that apply to the devices covered by this letter, subject to the manufacturer's continued compliance to the other conditions specified in Article 120.3c of MDR (as amended by (EU) 2023/607), are shown below:

- May 26, 2026 for Class III custom-made implantable devices
- December 31, 2027 for Class III devices and Class IIb implantable devices excluding Well-established technologies (WET - sutures, staples, dental fillings, dental braces, tooth crowns, screws, wedges, plates, wires, pins, clips and connectors)
- December 31, 2028 for other Class IIb devices, Class IIa, Class I devices placed on the market in sterile condition or have a measuring function
- December 31, 2028 for devices not requiring the involvement of a notified body under MDD but requiring it under MDR (e.g., class I devices that qualify as re-usable surgical instruments)

On behalf of the Notified Body



Samuel QIN

Certification body

**Table 1: Devices covered by this letter and for which the NB is also responsible for appropriate surveillance of the corresponding devices under the applicable Directive:**

Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified at the pre-application stage)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
<b>Postpartum Balloon</b> <b>Basic UDI-DI:</b> <b>69388205000000000000006027</b>	Class IIa	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Postpartum Balloon</b> <b>Basic UDI-DI:</b> <b>6938820500000000000000362A</b>	Class IIa	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Cervical Ripening Balloon</b> <b>Basic UDI-DI:</b> <b>69388205000000000000006129</b>	Class IIa	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Cervical Ripening Balloon</b> <b>Basic UDI-DI:</b> <b>6938820500000000000000372C</b>	Class IIa	N/A	Certificate # HD 60144232 0001 NB #0197

<b>Device name or Basic UDI-DI (under MDR application)</b>	<b>MDR Device classification (as proposed by the manufacturer and verified at the pre-application stage)</b>	<b>If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device</b>	<b>MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification</b>
<b>Guide Wire</b>  <b>Basic UDI-DI:</b> <b>693882050000000000000013ZT</b>	Class IIa	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Hemostasis Valve Sets</b>  <b>Basic UDI-DI:</b> <b>693882050000000000000012ZR</b>	Class I devices placed on the market in sterile condition	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Stopcock</b>  <b>Basic UDI-DI:</b> <b>693882050000000000000014ZV</b>	Class IIa	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Balloon Inflation Device</b>  <b>Basic UDI-DI:</b> <b>693882050000000000000002ZN</b>	Class I devices placed on the market in sterile condition	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Connecting Tubing</b>  <b>Basic UDI-DI:</b> <b>693882050000000000000004ZS</b>	Class IIa	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Manifold</b>  <b>Basic UDI-DI:</b> <b>69388205000000000000001828</b>	Class IIa	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Pressure Bandage</b>  <b>Basic UDI-DI:</b> <b>693882050000000000000006ZW</b>	Class I devices placed on the market in sterile condition	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Colored piston specialty Syringe</b>  <b>Basic UDI-DI:</b> <b>6938820500000000000000022ZU</b>	Class I devices placed on the market in sterile condition	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Dose-control Syringe</b>  <b>Basic UDI-DI:</b> <b>6938820500000000000000020ZQ</b>	Class I devices placed on the market in sterile condition	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Manifold Set</b>  <b>Basic UDI-DI:</b> <b>693882050000000000000002729</b>	Class IIa	N/A	Certificate # HD 60144232 0001 NB #0197

Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified at the pre-application stage)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
<b>Ureteral Stent Set</b>  <b>Basic UDI-DI:</b> <b>69388205000000000000031ZV</b>	Class IIa	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Tracheostomy Tube Kits</b>  <b>Basic UDI-DI:</b> <b>693882050000000000000292D</b>	Class IIa	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Tracheostomy Tube Kits</b>  <b>Basic UDI-DI:</b> <b>693882050000000000000622B</b>	Class IIa	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Tracheostomy Tube Kits</b>  <b>Basic UDI-DI:</b> <b>693882050000000000000632D</b>	Class IIa	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Percutaneous Nephrostomy Sets</b>  <b>Basic UDI-DI:</b> <b>693882050000000000000282B</b>	Class IIa	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Locking Drainage Catheter</b>  <b>Basic UDI-DI:</b> <b>693882050000000000000392G</b>	Class IIa	N/A	Certificate # HD 60144232 0001 NB #0197
<b>ERCP Guidewire</b>  <b>Basic UDI-DI:</b> <b>69388205000000000000040ZW</b>	Class IIa	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Percutaneous Access Set</b>  <b>Basic UDI-DI:</b> <b>693882050000000000000382E</b>	Class IIa	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Infusion Sets with needleless adapter</b>  <b>Basic UDI-DI:</b> <b>6938820500000000000002525</b>	Class I devices placed on the market in sterile condition	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Drainage Catheter Sets</b>  <b>Basic UDI-DI:</b> <b>69388205000000000000023ZW</b>	Class IIa	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Introducer Needles</b>  <b>Basic UDI-DI:</b> <b>6938820500000000000002627</b>	Class IIa	N/A	Certificate # HD 60144232 0001 NB #0197

<b>Device name or Basic UDI-DI (under MDR application)</b>	<b>MDR Device classification (as proposed by the manufacturer and verified at the pre-application stage)</b>	<b>If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device</b>	<b>MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification</b>
<b>Introducer Sets</b>  <b>Basic UDI-DI:</b> <b>6938820500000000000011ZP</b>	Class IIa	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Transradial Introducer Sets</b>  <b>Basic UDI-DI:</b> <b>69388205000000000000030ZT</b>	Class IIa	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Hemodialysis Catheterization Kit</b>  <b>Basic UDI-DI:</b> <b>69388205000000000000021ZS</b>	Class IIa	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Hemodialysis Catheterization Kit</b>  <b>Basic UDI-DI:</b> <b>693882050000000000000662K</b>	Class IIa	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Hemodialysis Catheterization Kit</b>  <b>Basic UDI-DI:</b> <b>693882050000000000000672M</b>	Class IIa	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Angiographic Syringes</b>  <b>Basic UDI-DI:</b> <b>69388205000000000000001ZL</b>	Class IIa	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Disposable Infusion Pumps</b>  <b>Basic UDI-DI:</b> <b>693882050000000000000652H</b>	Class IIb excluding Class IIb implantable non-WET	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Patient-Controlled Analgesic Infusion Pumps</b>  <b>Basic UDI-DI:</b> <b>693882050000000000000642F</b>	Class IIb excluding Class IIb implantable non-WET	N/A	Certificate # HD 60144232 0001 NB #0197
<b>Disposable Pressure Transducers</b>  <b>Basic UDI-DI:</b> <b>6938820500000000000000192A</b>	Class IIb excluding Class IIb implantable non-WET	N/A	Certificate # HD 60144232 0001 NB #0197



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified at the pre-application stage)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
I.V Cannulas  Basic UDI-DI: 693882050000000000000002423	Class IIa	N/A	Certificate # HD 60144232 0001 NB #0197
Injection Cap  Basic UDI-DI: 69388205000000000000000592N	Class I devices placed on the market in sterile condition	N/A	Certificate # HD 60144232 0001 NB #0197

**Table 2: Devices covered by this letter and for which the NB is NOT responsible for appropriate surveillance of the corresponding devices under the applicable Directive:**

Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified at the pre-application stage)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
N/A	N/A	N/A	N/A

**Confirmation Letter Revision History**

Date	NB internal reference traceable to each version of the letter	Action
2024/04/02	SCWME_CL607_2024-04-02	Initial issue

# EC Declaration of Conformity



**Manufacturer:**

**SCW MEDICATH LTD.**

NO.4, Baolong 6th Road, Baolong Industrial  
Town, Longgang District, Shenzhen, 518116,  
Guangdong, P.R. China

**whose single Authorized Representative:**

**OBELIS S.A.**

Bd. Général Wahis 53  
1030 Brussels, Belgium

We, the manufacturer, herewith declare that the products

## **Angiographic Syringes**

Models: DSA-60-A1, DSA-150-A1, CT-100-X1, CT-200-A1, MRI-115-A1,  
DSA-100-B1, DSA-150-B1, CT-200-B1, MRI-65-A1, CT-200-A2,  
DSA-200-A1, DSA-130-A1, CT/DSA-150-B1, MRI-60-B1, CT-200-NE,  
DSA-90-NE, DSA-120-NE, CT-100-NE, CT-190-A1, CT-190/190-A1,  
CT-200/200-A2, CT-200/200-B1, CT-100/100-NE, CT-100/200-NE,  
CT-200/200-NE, CT-200-EM, CT-200/200-EM, CT-200-MED,  
CT-200/200-MED, MRI-65/115-A1, MRI-65/65-A1, MRI-115/115-A1,  
MRI-60/60-B1, MRI-60/60-NE, MRI-65-MED, MRI-65/200-MED

**GMDN Code: 15286**

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 2 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 60144232 0001

Issue date: 26.05.2020

Expiry date: 26.05.2024

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, 518116, Guangdong, P.R. China**

*Miriam Xie*

Shenzhen, 2022/09/23

*Place, date*

Legally binding signature, Function