EC Certificate

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

Registration No.: HD 60144935 0001

Report No.: 17034752 009

Manufacturer:
EXCELLENTCARE MEDICAL
(HUIZHOU) LTD.
Shatou Industrial Zone
Yuanzhou Town, Boluo County
Huizhou
516123 Guangdong
P.R. China

Products:
Medical Devices

(see attachment for products included)

Replaces Approval, Registration No.: HD 60135433 0001

Expiry Date: 2023-12-24

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: 2020-06-05

Date: 2020-06-05

Notified Body

Wenxiang Zhang

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.

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Doc. 1/1, Rev. 0

TÜV Rheinland
LGA Products GmbH
Tillystraße 2, 90431 Nürnberg

Attachment to Certificate
Registration No.: HD 60144935 0001
Report No.: 17034752 009

Manufacturer:
EXCELLENTCARE MEDICAL (HUIZHOU) LTD.
Shatou Industrial Zone
Yuanzhou Town, Boluo County
Huizhou
516123 Guangdong
P.R. China

Products:

- Laryngeal Mask Airways
- Anaesthetic Face Masks
- Breathing System Filters
- Breathing Circuits
- Suction Tubes
- Oxygen Masks
- Nasal Cannulas
- Nebulizers

Notified Body

TÜV Rheinland LGA Products GmbH
Zertifizierungsstelle

Date: 2020-06-05

Wenxiang Zhang

Business Stream Products
Certification Department

TÜVRheinland®
LGA
Precisely Right.

TÜV Rheinland LGA Products GmbH • 51105 Köln

EXCELLENTCARE MEDICAL (HUIZHOU) LTD.
Shatou Industrial Zone,
Yuanzhou Town,
Boluo County, Huizhou,
516123, Guangdong,
P.R. China

Contact

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Date January 05, 2024

Notified Body Confirmation Letter
Reference. : 10924096

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:

EXCELLENTCARE MEDICAL (HUIZHOU) LTD.
Shatou Industrial Zone,
Yuanzhou Town,
Boluo County, Huizhou,
516123, Guangdong,
P.R. China

SRN Number (if available): CN-MF-000009376

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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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**
Laryngeal Mask Airways Models: Ordinary type, Silicone type, Non-inflatable type Basic UDI-DI: **6957071LarynMA0401AX**	Class IIa	N/A	Certificate # HD 60144935 0001 NB #0197 Note: Non-inflatable type is not covered by MDD certificate.
Ventilation Mask Models: Ordinary type, Soft Cushion type, Endoscopic type, Non-inflatable type, CPAP face mask, NIV face mask Basic UDI-DI: **6957071VentiMa0101QK**	Class IIa	Aneasthetic Face Mask	Certificate # HD 60144935 0001 NB #0197 Note: Endoscopic type, Non-inflatable type, CPAP face mask, NIV face mask are not covered by MDD certificate.
Oxygen Masks Models: Normal Type, Airbag Type, Venturi type,Tracheostomy type	Class IIa	N/A	Certificate # HD 60144935 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**
Basic UDI-DI: **6957071OxygenM01017D**			
Nebulizers Models: A Type, T Type, High-capacity type Basic UDI-DI: **6957071Nebuliz0801VE**	Class IIa	N/A	Certificate # HD 60144935 0001 NB #0197
Suction Tube Models: Controllable Type Basic UDI-DI: **6957071SuctiTu30016K**	Class IIa	N/A	Certificate # HD 60144935 0001 NB #0197
Suction Tube Models: Closed type, Infant closed type Basic UDI-DI: **6957071SuctiTu30026M**	Class IIa	N/A	Certificate # HD 60144935 0001 NB #0197 Note: Infant closed type is not covered by MDD certificate.
Suction Tube Models: Collecting Type, Bronchoscopy Type Basic UDI-DI: **6957071SuctiTu30036P**	Class IIa	N/A	Certificate # HD 60144935 0001 NB #0197 Note: Bronchoscopy Type is not covered by MDD certificate.
Breathing System Filters Models: Anti-bacterial/viral Filter Basic UDI-DI: **6957071BreathF12016T**	Class IIa	N/A	Certificate # HD 60144935 0001 NB #0197
Breathing System Filters Models: HME and anti-bacterial/viral Filter (HMEFs), Pleated Hydrophobic Bacterial/Viral Filter, HEPA Filter Basic UDI-DI: **6957071BreathF12026V**	Class IIa	N/A	Certificate # HD 60144935 0001 NB #0197 Note: HEPA Filter is not covered by MDD certificate.
Breathing System Filters Models: Pulmonary Function Test Filter (PET) Basic UDI-DI: **6957071BreathF12036X**	Class IIa	N/A	Certificate # HD 60144935 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**
Breathing System Filters Models: HME for Tracheostomy patient Basic UDI-DI: **6957071BreathF12046Z**	Class IIa	N/A	Certificate # HD 60144935 0001 NB #0197
Breathing System Filters Models: Heat and Moisture Exchanger (HME) Basic UDI-DI: **6957071BreathF120573**	Class IIa	N/A	Certificate # HD 60144935 0001 NB #0197
Breathing Circuit Models: Anaesthesia type, Airbag type, Corrugated type, Smoothbore type, Basic type Basic UDI-DI: **6957071BreathC050162**	Class IIa	N/A	Certificate # HD 60144935 0001 NB #0197
Breathing Circuit Models: ： Non-invasive Type, Smoothbore type, Corrugated type, Basic typ, Single heated type, Double heated type Basic UDI-DI: **6957071BreathC050366**	Class IIa	N/A	Certificate # HD 60144935 0001 NB #0197
Breathing Circuit Models: Corrugated type, Smoothbore type, Basic typ, Single heated type, Double heated type Basic UDI-DI: **6957071BreathC050468**	Class IIa	N/A	Certificate # HD 60144935 0001 NB #0197
Breathing Circuit Models: Single heated type, Double heated type Basic UDI-DI: **6957071BreathC05056A**	Class IIa	N/A	Certificate # HD 60144935 0001 NB #0197
Breathing Circuit Models: Anaesthesia type, Airbag type, Corrugated type, Smoothbore type, Non-invasive Type, Basic type, Single heated type, Double heated type	Class IIa	N/A	Certificate # HD 60144935 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
Basic UDI-DI: **6957071BreathC05066C**			
Breathing Circuit Models: Anaesthesia type, Airbag type, Corrugated type, Smoothbore type, Non-invasive Type, Basic type, Single heated type, Double heated type Basic UDI-DI: **6957071BreathC05076E**	Class IIa	N/A	Certificate # HD 60144935 0001 NB #0197
Breathing Circuit Models: Suction type Basic UDI-DI: **6957071BreathC05086G**	Class IIa	N/A	Certificate # HD 60144935 0001 NB #0197
Breathing Circuit Models: Suction type Basic UDI-DI: **6957071BreathC05096J**	Class IIa	N/A	Certificate # HD 60144935 0001 NB #0197
Nasal Cannulas Models: Normal Type, U Type, Adjustable ear hanging type, Adjustable ear hanging III type, Adjustable fixed II type, T type,High flow type, Generator Basic UDI-DI: **6957071NasalCa080145**	Class IIa	N/A	Certificate # HD 60144935 0001 NB #0197 Note: Only Normal Type is covered by MDD certificate.

Table 2: Devices covered by this letter and for which the NB is <u>NOT</u> 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
2024/01/05	10924096	Initial issue