

# **EC Certificate**Full Quality Assurance System

Certificate No.: 9805-2017-CE-KOR-NA-PS Rev. 2.0

Project No.: PRJC-558665-2017-MSL-KOR

Valid Until: **07 May 2022** 

This is to certify that the quality system of:

## **CU Medical Systems Inc.**

130-1, Donghwagongdan-ro, Munmak-eup, Wonju-si, Gangwon-do, Republic of Korea

For design, production and final product inspection/testing of:

Defibrillator, Defibrillator/monitor with defibrillation electrodes.

Has been assessed with respect to:

The conformity assessment procedure described in Annex II excluding section 4 of Council Directive 93/42/EEC on Medical Devices, as amended

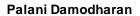
and found to comply.

Further details of the product(s) and conditions for certification are given overleaf.

Place and Date: Høvik, 23 May 2019



For: **DNV GL PRESAFE AS** 



The Certificate has been digitally signed.

See www.presafe.com/digital\_signatures for more info



Notice: The Certificate is subject to terms and conditions as set out in the Certification Agreement. Failure to comply may render this Certificate invalid.



# **EC Certificate**Full Quality Assurance System

Certificate No.: 9805-2017-CE-KOR-NA-PS Rev. 2.0

Project No.: PRJC-558665-2017-MSL-KOR

Valid Until: **07 May 2022** 

#### **Jurisdiction**

Application of Council Directive 93/42/EEC of 14 June 1993, adopted as "Forskrift om Medisinsk Utstyr" by the Norwegian Ministry of Health and Care Services.

#### Certificate history:

Revision	Description	Issue Date
0	Replace the Nemko certificate EU1110405 (NB0470) following the transfer of Notified Body functions to DNV GL NEMKO Presafe AS (NB 2460) issued after re-certification	2017-07-05
1	Model add	2018-07-02
2	Pediatric Defibrillation Electrode add (bold font)	2019-05-17

#### Products covered by this Certificate:

Product Description	Product Name	Class
Defibrillator	<ul> <li>CU-SP1</li> <li>CU-SP1 PLUS</li> <li>NF1201</li> <li>NF1200</li> <li>NFK200</li> </ul>	llb
Defibrillator/monitor	■ CU-HD1 ■ CU-SP2	IIb
Pediatric Defibrillation Electrode	- CUA0512P, CUA0711P, CUA0809PA	llb

The complete list of devices is filed with the Notified Body

#### Sites covered by this certificate

130-1, Donghwagongdan-ro, Munmak-eup, Wonju-si, Gangwon-do, Republic of Korea

### **EU Representative**

Medical Device Safety Service, GmbH Schiffgraben 41, 30175 Hannover, Germany

### **Terms and conditions**



# **EC Certificate**Full Quality Assurance System

Certificate No.:

9805-2017-CE-KOR-NA-PS Rev. 2.0

Project No.:

PRJC-558665-2017-MSL-KOR

Valid Until: **07 May 2022** 

The certificate is subject to the following terms and conditions:

- Any producer (see 2001/95/EC for a precise definition) is liable for damage caused by a defect in his product(s), in accordance with directive 85/374/EEC, as amended, concerning liability of defective products.
- The certificate is only valid for the products and/or manufacturing premises listed above.
- The Manufacturer shall fulfil the obligations arising out of the quality system as approved and uphold it so that it remains adequate and efficient.
- The Manufacturer shall inform Presafe of any intended updating of the quality system and Presafe will assess the changes and decide if the certificate remains valid.
- Periodical audits will be held, in order to verify that the Manufacturer maintains and applies the quality system. Presafe reserves the right, on a spot basis or based on suspicion, to pay unannounced visits.

The following may render this Certificate invalid:

- Changes in the quality system affecting production.
- Periodical audits not held within the allowed time window.

### Conformity declaration and marking of product

When meeting with the terms and conditions above, the producer may draw up an EC declaration of conformity and legally affix the CE mark followed by the Notified Body identification number of Presafe.

**End of Certificate**