Approved Record Document Nr: 00072686 Rev: C DA-00083976

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Rev C DA-00083976

Title CPRcard User Guide (IE)

Document Type Doc - Print Specification

**Document Category** DMR



Description of Change:	Revise CPRcard3 User Guide specification to Rev C
Approved Record Date:	2021-05-12T13:27:37Z
Lifecycle:	Production (when released in PLM System)
Tool Information:	
Replaces Document:	
Site:	LMAS (Stavanger)

# **Attachments (Electronic Files)**

Filename	File Description	File Type
Att 2 to 00072686.pdf	CPRcard UG - Webfile	pdf
Att 5 to 00072686.pdf	CPRcard UG - Print Specifications	pdf
Att 3 to 00072686.zip	CPRcard UG - Native Indesign Folder	zip
Att 1 to 00072686.pdf	CPRcard UG - Prepressfile	pdf
Att 4 to 00072686.pdf	CPRcard UG - Folding Instructions	pdf

# **Electronic Signatures**

Status	Signoff User	Local Client Time
Create/update and approve document	Fosnes, Camilla Andersen	2021-05-12T12:18:19Z
Verify and approve document	Nguyen, Phuong Helen	2021-05-12T13:27:37Z
Verify and approve document	Risanger, John Sigve	2021-05-12T12:56:29Z
Verify and approve document	Eilevstjønn, Joar	2021-05-12T13:12:41Z

# Part References Valid at Time of Print (for current status see Agile PLM system)

Item Number	Description
20-18063	CPRcard User Guide

Approved Record Document Nr: 00072686 Rev: C DA-00083976

# **Additional Info**

Create User	Risanger, John Sigve
Create Date	2021-02-02T14:13:49Z
Replaces Document	

# **Specifics**

Type	01 User Guide	
Language	01= English (US)	
Page Number	NA NA	
Colour	4+4 CMYK	
Other	Accordian User Guide with print on both sides.	
	See Att 4 to 00072686 for folding instructions and size guidelines.	
	See Att 5 to 00072686 for print specifications	

## Get Familiar with CPRcard

CPR is best performed by CPR-trained rescuers. It is also recommended to get familiar with the CPRcard by practicing on a training manikin using standard CPR technique.



Practice following the feedback from the CPRcard:

Turn on CPRcard.

2. Place the card correctly without removing the liner from the adhesive. (See Correct Placement on page 6).

- 3. Focus on depth feedback (See Compression Depth on page 8). Gradually increase the depth of compression so that the recommended depth target is reached.
- 4. Focus on rate feedback (see Compression Rate on page 8). Compress slowly and gradually increase the rate of compression until each of the rate indicators lights up.
- 5. Focus on performing compressions at an adequate depth and rate.
- 6. Remember to release between each compression.

To stay familiar with the CPRcard, repeat as often as needed.

Do not practice on a person as this may cause injuries to the person.

## Troubleshooting

The CPRcard The liner was not removed Remove the liner and

moves around | before the CPRcard was | place the CPRcard on

indication while detected

compressions

connectivity issues

Too shallow

compressions (<1 cm)

 Too slow compressions (<40/min)

during CPR placed on the patient. the patient's bare chest as

Chest compressions not | Press harder and/or

quickly as possible to avoid

faster. Release completely

between compressions.

Do not interrupt CPR -

continue CPR.

interrupting CPR

Iroubleshooting			Specifications	Specifications		
Symptom	Possible cause	Possible solution	Dimensions	Height 86 mm x Width 54 mm x Depth 2 mm Weight < 7 g		
Card does not turn on (no LEDs	On/Off button is not sufficiently pressed	Press and hold the On/ Off button firmly to try	External Material	Polycarbonate (PC) and self-adhesive medical tap		
turning on)	Card temperature is below 0 °C (32 °F)     The device is broken	to turn on the CPRcard.  If the problem persists, do not use the CPRcard on a patient.	Battery	Non-rechargeable lithium battery 15 mAh nomin capacity* Typical battery life: >30 minutes of CPR		
Warning LED turns on at	Internal error detected     Depleted battery	Turn the card off and on again.		End-of-shelf-life battery life: minimum 10 minutes of CPR		
start-up and stays on for 1 minute	Depicted battery	If the problem persists,  do not use the CPRcard	Shelf life	3 years from the production date		
Not all LEDs light up at start-up	The device is broken.	on a patient.	Operating/Storage Conditions	Temperature: 0 – 40 °C (32 – 104 °F) Humidity: ≤ 90% RH Atmospheric Pressure: 620 – 1060 hPa		
CPRcard turns off during CPR	Compression inactivity     1 min     Accidental activation of On/Off button	Do not interrupt CPR – continue CPR without feedback.	Shipping Conditions	<del>  '</del>		
	Internal error detected.     Depleted battery		Bluetooth® Low energy transmitter	Frequency band: 2.400 – 2.4835 GHz Modulation: Gaussian frequency shift modulation		
Depth target (green LED) not achieved during	Too shallow compressions     Incomplete release	Press harder and release completely between compressions.	energy transmitter	Maximum radio-frequency shift modulation: 1 mV Effective radiated power: 0 dBm		
CPR	(leaning)	compressions.	Ingress protection rating	IP67. Dust tight, and protected against water submersion to 1 meter (3.3 feet) for 30 minutes		
The CPR card	The liner was not removed	Remove the liner and	raurig	Submersion to Timeter (3.3 leet) for 30 minutes		

### Specifications

sions	Height 86 mm × Width 54 mm × Depth 2 mm Weight < 7 g	Compress Depth
al Material	Polycarbonate (PC) and self-adhesive medical tape.	
у	Non-rechargeable lithium battery 15 mAh nominal capacity* Typical battery life: >30 minutes of CPR End-of-shelf-life battery life: minimum 10 minutes of CPR	
fe	3 years from the production date	
ting/Storage tions	Temperature: 0 – 40 °C (32 – 104 °F) Humidity: ≤ 90% RH	Compress
	Atmospheric Pressure: 620 – 1060 hPa	Compress
ng Conditions	Temperature: -40 – 40 °C (-40 – 104 °F) Humidity: ≤ 90% RH Atmospheric Pressure: 550 – 1060 hPa	
oth® Low	Frequency band: 2.400 – 2.4835 GHz	
transmitter	Modulation: Gaussian frequency shift modulation Maximum radio-frequency shift modulation: 1 mW Effective radiated power: 0 dBm	Compress accuracy

\*Battery performance varies with temperature.

tth 54 mm × Depth 2 mm	Compression Depth	[
and self-adhesive medical tape.		
hium battery 15 mAh nominal		
30 minutes of CPR ery life: minimum 10 minutes		(
duction date		t
°C (32 – 104 °F)	Compression depth accuracy	=
e: 620 – 1060 hPa	Compression Rate	F
°C (-40 – 104 °F)		
550 – 1060 hPa		
00 – 2.4835 GHz		
frequency shift modulation uency shift modulation: 1 mW wer: 0 dBm	Compression rate accuracy	-

Do not store your CPRcard above 40 °C as this may reduce the lifetime of the battery.

## Specifications

•		0
Compression Depth	Depth feedback is based on the median depth of the last 5 compressions.  < 5 cm (2.0 in)  5 - 6 cm (2 - 2.36 in)  > 6 cm (2.36 in)	The device is inte equipment, and it No particular act to electromagnet  Warnings  Use of this equal avoided because
	Compressions with a depth of less than 1 cm or more than 10 cm will not be detected	equipment and operating norm • Portable RF cor cables and exte
Compression depth accuracy	±5 mm or ±10%, whichever is greater	part of the CPF could result.
Compression Rate	Rate feedback is based on the median rate of the last 5 compressions. 100 – 120 /min	Electromagne
	< 100 /min > 120 /min	Emissions Test
Compression rate	±5%	RF emissions C

## Electromagnetic Conformity

tended for use outdoors and indoors except for near HF surgical wireless communication nd the RF shielded room for magnetic resonance imaging. actions are required to maintain safety and performance with regard gnetic disturbances for the expected service life.

Proximity fields from RF | IEC 61000-4-3 | 380-390 MHz: 27 V/m

uipment adjacent to or stacked with other equipment should be Rated power frequency | IEC 61000-4-8 | 30 A/m ause it could result in improper operation. If such use is necessary, this nd the other equipment should be observed to verify that they are

Electrical fat transients | IEC 61000-4-4 | ±2 kV communications equipment (including peripherals such as antenna xternal antennas) should be used no closer than 30 cm (12 in) to any CPRcard. Otherwise, degradation of the performance of this equipment Surges: Line-to-line | IEC 61000-4-5 | ±0.5 kV. ±1 kV

## netic Emissions Tests

Emissions Test	Standard or test method	Compliance	Conducted disturbances induced by RF fields	IEC 61000-4-6	3 V ; 0.15 MHz – 80 MHz
RF emissions CISPR 11	Group 1 Class B	Group 1 Class B			6 V in ISM and amateur radio bands between 0.15 MHz
Harmonic emissions	IEC 61000-3-2	Class A			and 80 MHz 80% AM at 1 kHz
Voltage fluctuations/ flicker	IEC 61000-3-3	Complies	Voltage dips	IEC 61000-4-11	0% U <sub>т</sub> ; 0.5 cycle

## Electromagnetic Immunity Tests

Immunity Test	Standard or test Compliance Level and			Single phase: at 0	
	method	Immunity Test Level	Voltage interruptions	IEC 61000-4-11	0% U.; 250/300 c
Electrostatic discharge	rostatic discharge IEC 61000-4-2 ±8 kV contact ±2kV, ±4kV, ±8kV ±15 kV air	voltage interruptions	120 01000 111	070 O <sub>T</sub> , 230/300 C	
			$\boldsymbol{U}_{\scriptscriptstyle T}$ is the a.c. mains voltage prior to application of the test lev		
Radiated RF EM fields	IEC 61000-4-3	10 V/m 80 MHz – 2.7 GHz 80% AM at 2 Hz			

## Federal Communications Commission (FCC) and Industry Canada (IC) Statements

This device complies with part 15 of the FCC Rules and Industry Canada's

- licence-exempt RSSs. Operation is subject to the following two conditions: This device may not cause harmful interference, and
- 2. This device must accept any interference received, including interference that may cause undesired operation.

Le présent appareil est conforme aux CNR d'Industrie Canada applicables aux appareils radio exempts de licence. L'exploitation est autorisée aux deux

- . L'appareil ne doit pas produire de brouillage, et
- 2. L'appareil doit accepter tout brouillage radioélectrique subi, même si le brouillage est susceptible d'en compromettre le fonctionnement.

Changes or modifications not expressly approved by Laerdal Medical could void the user's authority to operate the equipment.

FCC ID: QHQ-20-10468

IC: 20263-2010468

80% AM at 1 kHz IEC 61000-4-11 0% U., : 0.5 cvcle At 0°. 45°. 90°. 135°. 180°. 225°, 270° and 315° EU on restriction of the use of certain hazardous substances (RoHS). 0% U.; 1 cycle and 70% U. 25/30 cycles Single phase: at 0° IEC 61000-4-11 0% U<sub>\*\*</sub>; 250/300 cycle

430-470 MHz: 28 V/m

800-960 MHz: 28 V/m

1700-1990 MHz: 28 V/m

2400-2470 MHz: 28 V/m

5100-5800 MHz: 29 V/m

100 kHz repetition frequency

50 Hz or 60 Hz

Surges: Line-to-ground | IEC 61000-4-5 | ±0.5 kV. ±1 kV. ±2 kV

704-787 MHz: 9 V/m

## Symbol Glossary

## Symbol Definition CE mark Single Use. Do not re-use. Follow Instructions for Use WEEE Symbol Manufacturer Date of Manufacture YYYY MM DD Defibrillation-proof type BF applied part. The entire CPRcard is the applied part. Warning/Caution Device catalogue number reference Unique Device Identification IP67 Ingress protection rating Temperature limitations

This product is in compliance with the essential requirements of EU Council directive 93/42/ EEC as amended by EU Council directive 2007/EC, Council

Humidity limitations Directive 2014/53/EU on Radio Equipment (RED) and Council Directive 2011/65/ Pressure limitations

Machine readable Unique Device Identification (UDI).

Not to be used in a bed or on soft surfaces

Datamatrix with UDI numbers (XXX = last three UDI digits)

(X) Not for patient under 8 years

If you need assistance or to report any issues, contact a local Laerdal representative

## Waste Handling



Waste Electrical and Electronic Equipment (WEEE).

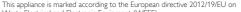
negative consequences for the environment and human health, which could otherwise be caused by inappropriate waste handling of this product.

household waste. Instead, it shall be handed over to the applicable collection point for the recycling of electrical and electronic equipment.

Warranty for terms and conditions.

For more information visit www.laerdal.com.

or visit www.laerdal.com/CPRcard for more information.



By ensuring this product is disposed of correctly, you will help prevent potential

The symbol on the product indicates that this appliance may not be treated as

Disposal must be carried out in accordance with local environmental regulations

Do not cut the device as this may damage the battery and expose harmful chemicals

### Service and Warranty

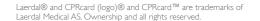
CPRcard does not have any replaceable or serviceable parts.

The CPRcard has a one-year limited warranty. Refer to the Laerdal Medical

4002 Stavanger, Norway



www.laerdal.com/CPRcard



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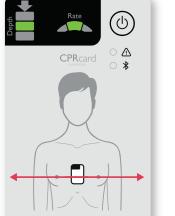
Manufactured in China for:

Laerdal Medical AS,

P.O. Box 377

Tanke Svilandsgate 30,

Printed in China

















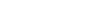




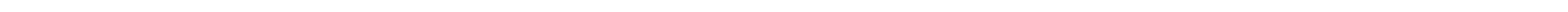






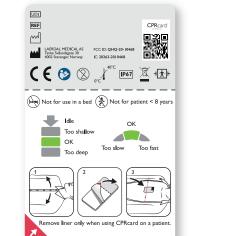






## Important Information and User Instructions

(on the back of the card)



### Intended Use

CPRcard is a single-use device intended to provide chest compression feedback to a CPR trained rescuer performing CPR on a suspected cardiac arrest patient, 8 years or older, lying flat on the back on a firm warnings, cautions and instructions in this User Guide. Retain this guide

### Indication for Use

When performing CPR on a suspected sudden cardiac arrest patient. despite any available care.

## Operating Principle

A battery powered device placed between the patient's bare chest and the rescuer's hands for giving feedback on compression depth and rate to a rescuer performing CPR. Using an accelerometer, the device measures independent of the device. its own relative movement perpendicular to the device surface and uses this to give input to an algorithm for calculation of correct feedback on compression rate and depth. The device is equipped with wireless communication functionality for transfer of performance data.

## Warnings and Cautions

A Warning states a condition, hazard, or unsafe practice that can result in serious bersonal injury or death.

A Caution states a condition, hazard, or unsafe practice that can result in minor personal injury or damage to the product

A Note states important information about the product or its operation.

( ) Important Information Before using CPRcard, read these instructions thoroughly. Observe all

> for future reference. CPR cannot ensure survival regardless of how well chest compressions are performed. For some patients, the cardiac arrest is not reversible

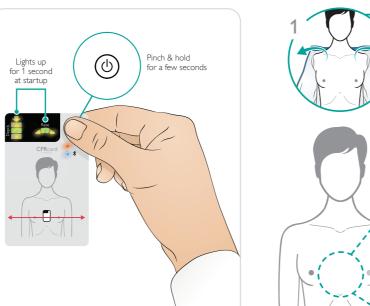
Common side effects of properly performed CPR include skin abrasion. bruises, rib and sternum fractures, and occasionally injuries to internal

The CPRcard does not guide the decision of whether or not to perform CPR on a suspected cardiac arrest victim. This decision must be made

- Do not use CPRcard on a patient on a "soft surface" (e.g. bed/mattress. stretcher or transportation cot) as it will provide inaccurate feedback that can result in too shallow combressions.
- Do not delay CPR to search for the CPRcard or if you experience any problem using it. Begin CPR without the card.
- Do not use the CPRcard in a moving environment (e.g. during patient transport in a car, boat or aircraft) as it may provide inaccurate feedback.
- Do not use CPRcard on an open wound or recent incision site as this may lead to cross-contamination and cause further injury.

Using CPRcard under sunlight might affect the visibility of the feedback from the card. Continue CPR even if you don't see the feedback from the card.

Turn On/Off



Correct Placement



hands are placed correctly on the CPRcard, the feedback

panel should be visible.

The patient tabe helps the CPRcard to stay in place on the patient's

chest during CPR. The patient tabe's liner should only be removed for

Compression Rate Place the card with the red placement line in the center of the patients bare chest, and the feedback panel towards the

> 100 – 120 compressions per minute Use standard CPR hand placement technique. When the

> > Release pressure between compressions to allow the chest to recoil

The CPRcard does not provide feedback on incomplete release (leaning) between compressions.

## Idle Indicator

When no compressions are detected for more than 1.5 seconds, the idle indicator will blink,



completely to maintain efficient compressions.

Do not interrupt CPR if compression feedback stops. Continue CPR without



If there are no compressions detected for one minute, the CPRcard shuts

Do not place the defibrillation bads on top of the CPRcard as this may

obstruct the CPR feedback and interfere with the defibrillation.

## Warning Indicator

If a device error is detected, the red Warning Indicator will turn on and compression feedback will stop. Such an error may be caused by a depleted battery or a technical malfunction. The CPR card should be replaced.

## Use with a Defibrillator

Follow defibrillator voice guidance when using the CPRcard together with a defibrillator. There is no need to remove the CPRcard before delivering a shock.



Bluetooth® Wireless Technology

than 50 compressions

Possible uses with compatible apps are:

turned on and before chest compressions are initiated.

Refer to www.laerdal.com/CPRcard for more information.

Do not delay CPR if you are not able to establish a Bluetooth connection.

\*Bluetooth is a trademark owned by Bluetooth SIG, Inc.

## CPRcard allows wireless connectivity using Bluetooth® Low Energy\*.

The CPRcard is intended for single-use only and should not be re-used. After use on a patient, the CPRcard may be contaminated and must be disposed of in accordance with local protocol.

If data transfer is required after patient use, the CPRcard can be placed in a plastic bag. The compatible app or software can perform the transfer through the bag.

### Maintenance and Inspection

Perform functional test quarterly to ensure that the CPRcard is functioning as it should.

1. Inspect the CPRcard for physical damage (e.g. tears and cracks).

### Turn on the device.

Single Use

- 3. Observe and verify that all the lights function (see illustration), the lights should display for one second.
- 4. Turn off the device.



Replace the card if it doesn't work properly or is severely damaged.

# Do not attempt to modify the CPRcard in any way before use as it can affect

its functionality.



