Turn on CPRcard.

- 2. Place the card correctly without removing the liner from the adhesive. (See Correct Placement on page 6).
- 3. Place your hands on the card making sure the feedback indicator lights are not covered.
- 4. Focus on depth feedback (See Compression Depth on page 8). Gradually increase the depth of compression so that the recommended depth target is reached.
- 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. 6. Focus on performing compressions at an adequate depth and rate.
- Remember to release between each compression.

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

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

Too slow compressions

connectivity issues | antenna on the card | feedback indicators

Hands are covering the Move hands away from the

(<40/min)

Symptom	Possible cause	Possible solution	Dimensions	
Card does not turn on (no LEDs turning on)	On/Off button is not sufficiently pressed     Card temperature is	Press and hold the On/ Off button firmly to try to turn on the CPRcard.	External Material I Battery I	
	below 0 °C (32 °F) • The device is broken	If the problem persists, do not use the CPRcard on a patient.		
Warning LED turns on at	Internal error detected     Depleted battery	Turn the card off and on again.	Shelf life	
start-up and stays on for 1 minute	. ,	If the problem persists, do not use the CPRcard		
Not all LEDs light up at start-up	The device is broken.	on a patient.	Operating/Storage Conditions	-
CPRcard turns off during CPR	Compression inactivity     2 min     Accidental activation of     On/Off button     Internal error detected.	Do not interrupt CPR – continue CPR without feedback.	Shipping Conditions  Bluetooth® Low energy transmitter	-
Depth target (green LED) not achieved during	Depleted battery      Too shallow compressions     Incomplete release	Press harder and release completely between compressions.		
CPR The CPRcard	(leaning) The liner was not removed	Remove the liner and	Ingress protection rating	1
moves around during CPR	before the CPRcard was placed on the patient.	place the CPRcard on the patient's bare chest as quickly as possible to avoid interrupting CPR	⚠ Caution	
Inactivity indication while performing	Chest compressions not detected     Too shallow	Press harder and/or faster: Release completely between compressions.	Do not store your CPRcard	1

Specifications

\_\_\_\_

	•
Height 86 mm × Width 54 mm × Depth 2 mm Weight < 7 g	Compression Depth
Polycarbonate (PC) and self-adhesive medical tape.	
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	
3 years from the production date	
Temperature: 0 – 40 °C (32 – 104 °F) Humidity: ≤ 90% RH	Compression dep accuracy
Atmospheric Pressure: 620 – 1060 hPa	Compression Rate
Temperature: $-40 - 40$ °C ( $-40 - 104$ °F) Humidity: $\le 90\%$ RH Atmospheric Pressure: $550 - 1060$ hPa	
Frequency band: 2.400 – 2.4835 GHz Modulation: Gaussian frequency shift modulation Maximum radio-frequency shift modulation: 1 mW Effective radiated power: 0 dBm	Compression rate accuracy
IP67. Dust tight, and protected against water submersion to 1 meter (3.3 feet) for 30 minutes	

nm x Depth 2 mm		Compression Depth	Depth feedback i last 5 compressio
adhesive medical tape.			
ttery 15 mAh nominal			5 -
ites of CPR ninimum 10 minutes			
			Compressions with than 10 cm will no
date			triair 10 tri wiir ii
– 104 °F)		Compression depth accuracy	±5 mm or ±10%
- 1060 hPa		Compression Rate	Rate feedback is b
- 104 °F)			5 compressions.
60 hPa			<
335 GHz			
ncy shift modulation nift modulation: 1 mW		Compression rate accuracy	±5%

Specifications

## \*Battery performance varies with temperature.

ard above 40 °C as this may reduce the lifetime of the battery.

### Electromagnetic Conformity

Harmonic emissions

Immunity Test

Voltage fluctuations/ flicker

Electrostatic discharge

Radiated RF FM fields

n n 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)  Compressions with a depth of less than 1 cm or more than 10 cm will not be detected  ±5 mm or ±10%, whichever is greater	The device is intended for use outdoors a equipment, and the RF shielded room for No particular actions are required to main to electromagnetic disturbances for the experiment of the equipment of the other equipment of the other equipment of the other equipment (cables and external antennas) should be part of the CPRCard. Otherwise, degradatic could result.
n Rate	Rate feedback is based on the median rate of the last 5 compressions.  100 – 120 /min	Electromagnetic Emissions Tests
	< 100 /min > 120 /min	Emissions Test Stan
	150/	RF emissions CISPR 11 Gro
n rate	±5%	

for use outdoors and indoors except for near HF surgical wireless communication shielded room for magnetic resonance imaging. re required to maintain safety and performance with regard urbances for the expected service life.

- adjacent to or stacked with other equipment should be Rated power frequency | IEC 61000-4-8 | 30 A/m uld result in improper operation. If such use is necessary, this ther equipment should be observed to verify that they are Electrical fat transients | IEC 61000-4-4 | ±2 kV
- ications equipment (including peripherals such as antenna tennas) should be used no closer than 30 cm (12 in) to any Otherwise, degradation of the performance of this equipment Surges: Line-to-line IFC 61000-4-5 +0.5 kV +1 kV

### Standard or test Compliance Conducted disturbances | IEC 61000-4-6 | 3 V : 0.15 MHz = 80 MHz method induced by RF fields Group 1 Class B Group 1 Class B IEC 61000-3-2 Class A IEC 61000-4-11 0% U., : 0.5 cvcle

# Electromagnetic Immunity Tests

tandard or test nethod	Compliance Level and Immunity Test Level
EC 61000-4-2	±8 kV contact ±2kV, ±4kV, ±8kV ±15 kV air
EC 61000-4-3	10 V/m 80 MHz – 2,7 GHz 80% AM at 2 Hz

# Federal Communications Commission (FCC) and Industry Canada

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

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

At 0°. 45°. 90°. 135°. 180°

ons IEC 61000-4-11 0% U<sub>\*\*</sub>; 250/300 cycle voltage prior to application of the test level.

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

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

430-470 MHz: 28 V/m

704-787 MHz: 9 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 frequence

6 V in ISM and amateur radio

bands between 0.15 MHz

and 80 MHz

80% AM at 1 kHz

Single phase: at 0°

50 Hz or 60 Hz

Follow Instructions for Use

Date of Manufacture YYYY MM DD

Temperature limitations

Not for patient under 8 years

Not to be used in a bed or on soft surfaces

Humidity limitations

WEEE Symbol

Manufacturer

2. This device must accept any interference received, including interference that may

est susceptible d'en compromettre le fonctionnement.

user's authority to operate the equipment.

FCC ID: QHQ-20-10468

IC: 20263-2010468

directive 93/42/ EEC as amended by EU Council directive 2007/EC, Council Directive 2014/53/FU on Radio Equipment (RED) and Council Directive 2011/65/

Symbol Definition ( € CE mark Single Use. Do not re-use.

L'appareil doit accepter tout brouillage radioélectrique subi, même si le brouillage

Defibrillation-proof type BF applied part.
The entire CPRcard is the applied part. Changes or modifications not expressly approved by Laerdal Medical could void the

Warning/Caution

Device catalogue number reference

Unique Device Identification IP67 Ingress protection rating

This product is in compliance with the essential requirements of EU Council

225°, 270° and 315° 0% U,; 1 cycle and 70% U,; 25/30 cycles

EU on restriction of the use of certain hazardous substances (RoHS). Pressure limitations

Machine readable Unique Device Identification (UDI). Datamatrix with UDI numbers (XXX = last three UDI digits)

If you need assistance or to report any issues, contact a local Laerdal representative or visit www.laerdal.com/CPRcard for more information.

## Waste Handling

Support



This appliance is marked according to the European directive 2012/19/EU on Waste Electrical and Electronic Equipment (WEEE).

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

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

The symbol on the product indicates that this appliance may not be treated as household waste, Instead, it shall be handed over to the applicable collection point for the recycling of electrical and electronic equipment.

Disposal must be carried out in accordance with local environmental regulations for waste disposal.



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 Warranty for terms and conditions.

For more information visit www.laerdal.com.

## www.laerdal.com/CPRcard



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Manufactured in China for:

Laerdal Medical AS.

P.O. Box 377

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Printed in China

**CPRcard** User Guide



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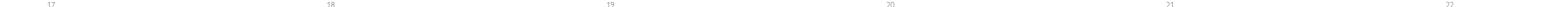






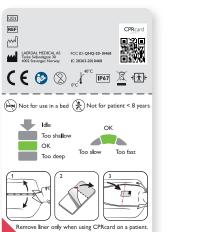






### Important Information and User Instructions

(on the back of the card)



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

### Indication for Use

Intended 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

## ∴ Warnings and Cautions

communication functionality for transfer of performance data.

A Warning states a condition, hazard, or unsafe practice that can result in serious personal 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

for future reference.

Before using CPRcard, read these instructions thoroughly. Observe all warnings, cautions and instructions in this User Guide. Retain this guide

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

- Do not use CPRcard on a batient on a "soft surface" (e.g. bed/mattress. stretcher or transportation cot) as it will provide inaccurate feedback that
- 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.

the card, Continue CPR even if you don't see the feedback from the card.

Turn On/Off

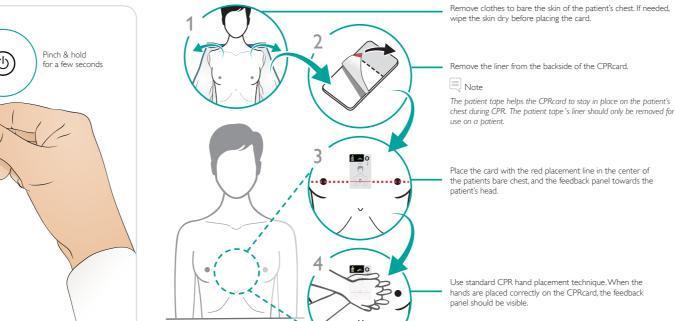
CPR cannot ensure survival regardless of how well chest compressions are performed. For some patients, the cardiac arrest is not reversible for 1 second at startup

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

- can result in too shallow combressions.

Using CPRcard under sunlight might affect the visibility of the feedback from

### Correct Placement



### Chest Compression Feedback

international CPR guidelines.

Feedback on compression depth and rate are provided by indicator lights. The green lights indicate the targets according to the 2020 indicator will blink,

## Compression Depth



### Compression Rate

Too slow Too fast

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

If there are no compressions detected for two minutes, the CPRcard shuts

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

Warning Indicator

100 – 120 compressions per minute

Release pressure between compressions to allow the chest to recoil completely to maintain efficient compressions.

# 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.

# Firmware upgrade

- Live streaming of CPR feedback during use
- CPRcard is only available for Bluetooth connection after the card is

turned on and before chest compressions are initiated.

Refer to <a href="https://www.laerdal.com/CPRcard">www.laerdal.com/CPRcard</a> 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.



Do not cover the Bluetooth antenna, located in the feedback banel, when

blacing the hands on CPRcard.

Do not place the defibrillation pads on top of the CPRcard as this may Do not interrupt CPR if compression feedback stops. Continue CPR without obstruct the CPR feedback and interfere with the defibrillation.

# Bluetooth® Wireless Technology

CPRcard allows wireless connectivity using Bluetooth® Low Energy\*. Possible uses with compatible apps are:

- Transfer of stored data. Summary data is stored for events with more than 50 compressions



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

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

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

Perform functional test quarterly to ensure that the CPRcard is

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

3. Observe and verify that all the lights function (see illustration), the

disposed of in accordance with local protocol.

lights should display for one second.

through the bag.

Maintenance and Inspection

functioning as it should.

Turn on the device.

Turn off the device.

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