

ShockLink

Important Product Information



Warnings and Cautions

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 manikin.

Notes

Important information about the product or its operation.

Cautions

- Maximum energy in one shock: 360 J (Only give a cumulative total of 1,000 J per minute.) ShockLink will disconnect the defibrillator for one minute if a cumulative total of more than 4000 J has been given over the last 3 minutes
- Do not use ShockLink if the cable or the housing has visible damage or openings.
- Do not use ShockLink in wet conditions.

Waste Handling

The ShockLink unit contains electronic components. Dispose of it at the applicable recycling facility in accordance with local regulations.



WEEE

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, or on the documents accompanying 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. For more detailed information about treatment, recovery and recycling of this product, please contact your local city office, your household waste disposal service or the Laerdal representative where you purchased the product.

FCC

Federal Communications Commission Statement and Industry Canada Statements

This device complies with part 15 of the FCC rules and RSS-210 of the Industry Canada rules. Operation is subject to the following two conditions:

1. This device may not cause harmful interference, and
2. this device must accept any interference received, including interference that may cause undesired operation.

Ce dispositif est conforme à la norme CNR-210 d'Industrie Canada applicable aux appareils radio exempts de licence. Son fonctionnement est sujet aux deux conditions suivantes:

1. Le dispositif ne doit pas produire de brouillage préjudiciable, et
2. ce dispositif doit accepter tout brouillage reçu, y compris un brouillage susceptible de provoquer un fonctionnement indésirable.

This equipment has been tested and found to comply with the limits for a Class B digital device, pursuant to part 15 of the FCC Rules. These limits are designed to provide reasonable protection against harmful interference in a residential installation. This equipment generates, uses and can radiate radio frequency energy and, if not installed and used in accordance with the instructions, may cause harmful interference to radio communications. However, there is no guarantee that interference will not occur in a particular installation. If this equipment does cause harmful interference to radio or television reception, which can be determined by turning the equipment off and on, the user is encouraged to try to correct the interference by one or more of the following measures:

- Reorient or relocate the receiving antenna.
- Increase the separation between the equipment and receiver.
- Consult the dealer or an experienced radio/TV technician for help.

FCC ID: QHQ-185-20050

IC ID: 20263-18520050

The term "IC" before the equipment certification number only signifies that the Industry Canada technical specifications were met.









Caution

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

English

ShockLink Important Information

Symbol Glossary

	ShockLink is CE marked in accordance with Council Directive 1999/5/EC relating to Radio and Telecommunications Terminal Equipment (R&TTE), and Council Directive 2011/65/EU on Restriction on the use of certain hazardous substance (RoHS).
	Australian Radiocommunications and EMC Compliance Mark
	Dispose of in accordance with your country's recommendations
	FCC Statement and IC statements
	Reference order number
	Production lot number

Specifications

Dimensions	85 x 53 x 14 mm (3.3 x 2 x 0.5")
Weight	120 g (0.26 lbs) (including battery)
Cable length	Approximately 0.9 m (3ft)
Defibrillator impedance	115 Ohms
Operation temperature	0 °C to +35 °C (32 °F to 95 °F)
Storage temperature	-15 °C to +50 °C (5 °F to 122 °F)
Humidity	< 95% relative humidity non condensing
Communication	Bluetooth Smart
Battery	AAA size Alkaline battery 1.5V
Battery life	ON: 25 - 35 hours OFF: 3 years (shelf life with battery mounted)
Battery Low warning:	@ 1.1 V, typically 4 hours left
Unit turns off:	@ 0.95 V nominal
Shock detection:	5 – 360 J, +/- 15%

Warranty

Refer to the Laerdal Global Warranty for terms and conditions.
For more information visit www.laerdal.com

ShockLink® Patent Pending
Laerdal Medical, the Laerdal logo and ShockLink
are trademarks owned by Laerdal Medical AS.

© 2015 Laerdal Medical AS. All rights reserved.

Manufacturer: Laerdal Medical AS
P.O. Box 377
Tanke Svilandsgt. 30, 4002 Stavanger, Norway
T: (+47) 51 51 17 00

Printed in China.

20-09356 Rev A

www.laerdal.com



Laerdal
helping save lives