EN

Laerdal-SonoSim

Procedure Trainer
&
Ultrasound Solution

Important Product Information

www.laerdal.com

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helping save lives

This Important Product Information covers Laerdal-SonoSim Procedure Trainer and Laerdal-SonoSim Ultrasound Solutions. Unless otherwise specified the information applies to all product configurations.

Read these instructions thoroughly. Observe all warnings, cautions and instructions in the User Guide and in this Important Product Information booklet. Retain this booklet for future reference.

⚠ 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 product.

Notes
A note states important information about the product or its operation.

Needle & Syringe

⚠ Caution
The tip of the Needle & Syringe is sharp. Take care when handling.

Storage

Store the components of the Laerdal-SonoSim Procedure Trainer in the Accessory box when not in use.

Care, Maintenance and Cleaning

Note
Unplug all components before cleaning.

Use mild soap and water on a soft cloth to clean the equipment immediately after simulation sessions.

Federal Communications Commission Statement

This device complies with part 15 of the FCC 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.

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.
- Connect the equipment into an outlet on a circuit different from that to which the receiver is connected.
- Consult the dealer or an experienced radio/TV technician for help.

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

Industry Canada Statement

This device complies with RSS-210 of the Industry Canada Rules. Operation is subject to the following two conditions:
1. This device may not cause harmful interference.
2. This device must accept any interference received, including interference that may cause undesired operation.

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

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.

Mise en garde: *Tout changement ou toute modification n'ayant pas fait l'objet d'une approbation expresse de Laerdal Medical peut annuler le droit dont dispose l'utilisateur de se servir de l'équipement.*

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

Le terme « IC » qui précède le numéro d'agrément de l'équipement signifie uniquement que les caractéristiques techniques spécifiées par Industrie Canada sont respectées.

USA

Laerdal SonoSim Probe contains FCC ID: 2AEMESSLS

Needle & Syringe (Laerdal SonoSim Procedure Trainer) contains FCC ID: 2AEME-1002040

Canada

IC: 20197-1002040

Laerdal- SonoSim Probe IC Certificate: 20197-SSLS

Needle & Syringe (Laerdal SonoSim Procedure Trainer)
IC Certificate: xxxxx-xxxx

EU

CE:
This product is in compliance with the essential requirements of Council Directive. 2014/53/EU on Radio Equipment (RED) and Council Directive 2011/65/EU on restriction of the use of certain hazardous substances (RoHS).

Waste Handling

Dispose of in accordance with your country's recommendations

This appliance is marked according to the European directive 2012/19/EC 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 Laerdal representative.

Symbol Glossary

CE	CE mark
	WEEE symbol
	Australian Radiocommunications and EMC Compliance Mark
	CSA Certification Mark
	Warning/Caution
	Note
	Manufacturer
REF	Reference Number
SN	Serial Number

Temperature and Conditions	
Operating / Storage temperature	Needle: -20 °C to 25 °C (-4 °F to 77 °F) Probe: -40 °C to 85 °C (-40 °F to 185 °F) TrackPad: -20 °C to 25 °C (-4 °F to 77 °F)
Dimensions	
Measurements	Torso: 94.3 x 20.4 x 35.1 cm (36.8 x 8 x 138.2 in) Needle: 17.4 x 1.6 x 3.5 cm (6.9 x 0.6 x 1.4 in) Probe: 11.2 x 2.8 x 4.7 cm (4.4 x 1.1 x 1.9 in) TrackPad: 19.7 x 9.5 x 0.3 cm (7.75 x 3.75 x 0.1 in)
Weight	Torso: 3.42 kg (120.6 oz) Needle: 1.1 g (0.94 oz) Probe: 146 g (5.15 oz) TrackPad: 152.5 g (5.3792 oz)

Warranty

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

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