

For more information
and apps go to:

laerdal.com/littleanne



2022 Laerdal Medical AS.
All rights reserved.

P.O. Box 377
Tanke Svilandsgate 30
4002 Stavanger, Norway
T:(+47) 51 51 17 00




Little Anne PLUS

Important Product Information



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.

Care, Maintenance and Cleaning

Keep the manikin clean and in a hygienic condition.

Replace the airway filter after every class, especially if mouth to mouth resuscitation is practiced.

Change the airway assembly if there are visible signs of wear or damage to lung or metal patch.

We recommend that you use a separate Laerdal Manikin Face for every student. If several students use one manikin face, thoroughly sanitize the manikin face after every use.

Clean all skin parts regularly. Use warm soapy water or Laerdal manikin wipes. Use of Manikin Face Shields provides a clean barrier between your lips and the manikin face.

These can discolor the manikin:

- Pigments from lipstick and pens
- Latex gloves
- Using clothes other than what comes with the product originally.

 Cautions

- Do not use liquids to clean inside the manikin.
- Do not submerge any parts.
- Do not remove the spring plate rope.

Battery

Only use consumer grade, non-rechargeable, carbon zinc or alkaline batteries, LR6 1.5V AA. Batteries must be replaced with correct type. Remove batteries if product is not to be used for an extended period.

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

This device complies with part 15 of the FCC Rules and RSS-210 of IC 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.
- 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.

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

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.

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

This device contains licence-exempt transmitter(s)/receiver(s) that comply with Innovation, Science and Economic Development Canada's licence-exempt RSS(s). Operation is subject to the following two conditions:

1. This device may not cause interference.
2. This device must accept any interference, including interference that may cause undesired operation of the device.

L'émetteur/récepteur exempt de licence contenu dans le présent appareil est conforme aux CNR d'Innovation, Sciences et Développement économique Canada applicables aux appareils radio exempts de licence. L'exploitation est autorisée aux deux conditions suivantes :

1. L'appareil ne doit pas produire de brouillage;

L'appareil doit accepter tout brouillage radioélectrique subi, même si le brouillage est susceptible d'en compromettre le fonctionnement.

FCC ID: QHQ-LA

IC: 20263-LA

Japan

MIC certification 012-220014

Korea

R-CMM-Lm1-LA

China

Basic Torso: CMIIT ID: xxxxxxxxxx

EU

This product is in compliance with the essential requirements of Council Directive 2014/53/EU on Radio Equipment (RED),

The product is in compliance with Council Directive 2011/65/EU on restriction of the use of certain hazardous substances (RoHS).

UK

UKCA (U.K. Conformity Assessed): This product complies with the requirements of U.K. legislation of Radio Equipment Regulations (S.I. 2017/1206).
UKCA (U.K. Conformity Assessed): This product complies with the requirements of U.K. legislation of Restriction of the Use of Certain Hazardous Substances in Electrical and Electronic Equipment Regulations (RoHS) Regulations (S.I. 2012/3032).

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

Symbol	Definition
	CE mark
	UKCA Mark
	Australia Radiocommunications and EMC Compliance Mark
	KC (Korea Certification) certification mark
	Japanese MIC Certification
	Waste Electrical and Electronic Equipment symbol
	Manufacturer
	Date of Manufacture
	Reference Number
	Serial Number

Specifications

Size and Weight	
Dimensions	64 cm x 37 cm x 21,5 cm (25.2" x 14.5" x 8.5")
Weight	2 kg (± 0.2 kg)
Environment	
Operating temperatures	5 °C to 35 °C (41 °F to 95 °F) Degradation in product performance may be experienced when used at low temperatures (below 15 °C / 59 °F)
Storage temperatures	-10 °C to 50 °C (14 °F to 122 °F)
Relative humidity	5 % - 95 % relative humidity

Warranty

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

English

Little Anne PLUS Important Information

© 2022 Laerdal Medical AS. All rights reserved.



Manufactured in China for Laerdal Medical AS,
P.O. Box 377
Tanke Svilandsgate 30, 4002 Stavanger,
Norway
T: (+47) 51 51 17 00

Printed in China

20-xxxxx DRAFT

www.laerdal.com



Laerdal
helping save lives