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Laerdal
helping save lives

00038930 Rev I

Declaration of Conformity

Responsible Manufacturer:	Laerdal Medical AS Seehusens gate 1, P.O. Box 377 4002 Stavanger, Norway
Manufacturing site:	Laerdal Medical AS, Stavanger Manufacturing Vestre Svanholmen 3 4313 Sandnes, Norway
Product Name:	Resusci Anne Simulator Resusci Anne Advanced Skill Trainer
Product Options:	
150-22000	Resusci Anne Simulator AED
150-23000	Resusci Anne Simulator AED
150-27000	Resusci Anne Simulator Paddle
150-28000	Resusci Anne Simulator Paddle
151-22000	Resusci Anne Advanced SkillTrainer AED
151-23000	Resusci Anne Advanced SkillTrainer AED
151-27000	Resusci Anne Advanced SkillTrainer Paddle
151-28000	Resusci Anne Advanced SkillTrainer Paddle

to which this declaration relates is in conformity with

EU Council Directives

EU Radio Equipment Directive (RED) 2014/53/EU, including the cybersecurity requirements introduced by Commission Delegated Regulation (EU) 2022/30, and Council Directive 2011/65/EU on Restriction on the use of certain hazardous substance (RoHS), as amended by EU 2015/863

All supporting documentation is retained by the manufacturer.

The conformity is based on the following standards:

EMC:

EN 55032:2015 + A11:2020
EN 55035:2017 + A11:2020

Safety:

EN 62368-1:2014
EN 62368-1:2018

Radio:

ETSI EN 300 328 V2.2.2
ETSI EN 301 893 V2.1.1
EN 18031-1:2024
EN 18031-2:2024

DocuSigned by:
Mari Kaada
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SIGNATURE	POSITION IN ORGANISATION
Corporate Director Q&R	
PRINT NAME	DATE
Mari Kaada	2026-08-27