00038930 Rev D

DECLARATION OF CONFORMITY

Responsible Manufacturer:	Laerdal Medical AS P.O. Box 377 Tanke Svilandsgate 30 4002 Stavanger Norway	
Product Name	**Resusci Anne**	
Product Options:	150-22000	RA Simulator AED Link, IV arm left
	150-23000	RA Simulator AED Link, IV Arm right
	150-27000	RA Simulator Paddle, IV arm left
	150-28000	RA Simulator Paddle, IV arm right
	151-22000	RA Adv SkillTr. AED Link, IV arm left
	151-23000	RA Adv SkillTr. AED Link, IV arm right
	151-27000	RA Adv SkillTr. Paddle, IV arm left
	151-28000	RA Adv SkillTr. Paddle, IV arm right
Accessories:	204-301XX	SimPad PLUS LLEAP *XX = Country code*

to which this declaration relates is in conformity with the **Essential Requirements of EU Radio Equipment Directive (RED) 2014/53/EU** and
Council Directive 2011/65/EU on Restriction on the use of certain hazardous substance (RoHS)
All supporting documentation is retained by the manufacturer.

The conformity is based on the following standards:

Safety (Article 3.1 (a) of RED):
- EN 62368-1:2014
- EN 62479:2010
- EN 62311:2008

EMC (Article 3.1 (b) of RED)
- EN 301 489-1 V1.9.2
- EN 301 489-17 V2.2.1
- EN 61000-6-1:2007
- EN 61000-6-3:2007/A1:2011/AC:2012

Radio (Article 3.2 of RED):
- EN 300 328 V1.9.1
- EN 301 893 V1.8.1

Stavanger, 7th November 2022

Mercildes Dutra
Regulatory Affairs

CE

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