EU DECLARATION OF CONFORMITY

Responsible Manufacturer:	Laerdal Medical AS P.O. Box 377 Tanke Svilandsgate 30 4002 Stavanger Norway	
Manufacturing site:	Laerdal Medical (Suzhou) Co. Ltd No 19 Building, Huoju Road Science & Technology Industrial Park Suzhou, Jiangsu, P.R.China	
Product Name:	**Resusci Junior QCPR**	
Product Options:	181-00150 182-00150	Resusci Junior QCPR Resusci Junior QCPR AED AW w/IO Leg
Accessories	206-300xx 170-30050 181-80025	SimPad PLUS SkillReporter SkillGuide Cable, USB micro Am to Cm

to which this declaration relates is in conformity with
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)**
The standard Resusci Junior QCPR (Cat. No 181-00150) manikin when updated with the AED Service Upgrade Kit (Cat.No for the Service Part will be 182-60150) is electronically equivalent to the 182-00150 configuration.

All supporting documentation is retained by the manufacturer.

The conformity is based on the following standards:

EMC (Article 3.1(b) of RED):
- EN 301 489-1 V2.1.1
- EN 301 489-17 V3.1.1
- EN 61000-6-1:2007
- EN 61000-6-3:2007+A1:2011

Safety (Article 3.1(a) of RED):
- EN 62368-1:2014
- EN 60950-1:2006
- EN 62479:2010

Radio (Article 3.2 of RED):
- EN 300 328 V2.2.2

Stavanger, 22nd September 2022

Mercildes Dutra

Regulatory Affairs

CE

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