00018999 Rev Q

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 Anne QCPR**	
Product Options:	171-00160	Resusci Anne QCPR Torso - Rechargeable
	171-01260	Resusci Anne QCPR Full Body - Rechargeable
	172-00160	Resusci Anne QCPR AW Torso - Rechargeable
	172-01260	Resusci Anne QCPR AW Full Body - Rechargeable
	173-00160	Resusci Anne QCPR AED Torso - Rechargeable
	173-01260	Resusci Anne QCPR AED Full Body - Rechargeable
	174-00160	Resusci Anne QCPR AED AW Torso - Rechargeable
	174-01260	Resusci Anne QCPR AED AW Full Body - Rechargeable
	178-03160	Resusci Anne QCPR RQI Torso/HPCPR
	178-03161	Res Anne QCPR AED RQI Torso/HPCPR/no arms
	178-04160	Res Anne QCPR AED AW RQI Full Body/HPCPR
Accessories	206-300xx	SimPad PLUS SkillReporter
	170-30050	SkillGuide
	181-80025	Cable, USB micro AM to CM 1.8m, USB 2.0
	171-15000	RAQCPR electr. upgrade Customer Kit 2018
	173-15000	RAQCPR AED elect. upgrade Customer Kit 2018
	171-15010	RAQCPR Upgr. Bundle 2018
	173-15010	RAQCPR AED Upgr. Bundle 2018

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)**
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 61000-6-1:2007
EN 61000-6-3:2007+A3
Safety (Article 3.1(a) of RED):
EN 62368-1:2014 + A11
EN 60950-1/A11:2009/A1:2010/
A12:2011/A2:2013
EN 62479:2010
Radio (Article 3.2 of RED):
EN 300 328 V2.1.1

Stavanger, 3 May 2019

Corporate Regulatory Affairs Manager

CE