



# EU DECLARATION OF CONFORMITY

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<b>Responsible Manufacturer:</b>	Laerdal Medical AS P.O. Box 377 Tanke Svilandsgate 30 4002 Stavanger Norway
<b>Manufacturing site:</b>	Laerdal Medical (Suzhou) Co. Ltd No 19 Building, Huoju Road Science & Technology Industrial Park Suzhou, Jiangsu, P.R.China
<b>Product Name:</b>	<b>Resusci Anne QCPR</b>
<b>Product Options:</b>	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 177-00160 Resusci Anne QCPR HC Torso Torso/HPCPR 178-03160 Resusci Anne QCPR RQI Torso/HPCPR/with arms 178-03161 Res Anne QCPR AED RQI Torso/HPCPR/no arms 178-04160 Res Anne QCPR AED AW RQI Full Body/HPCPR 171-01235 Resusci Anne QCPR CHN Community - Full Body 171-00135 Resusci Anne QCPR CHN Community - Torso
<b>Accessories</b>	206-300xx SimPad PLUS SkillReporter 170-30050 SkillGuide 181-80025 Cable, USB micro AM to CM 171-15000 RAQCPR electr. upgrade Customer Kit 2018 173-15000 RAQCPR AED elect. upgrade Customer Kit 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 301 489-17 V3.1

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 V2.2.2

**Stavanger, 22<sup>nd</sup> September 2022**

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*Mecildes Dutra*  
Regulatory Affairs



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