

00064764 Rev F

EU DECLARATION OF CONFORMITY

Responsible Manufacturer:	Laerdal Medica P.O. Box 377 Tanke Svilands 4002 Stavange Norway	sgate 30
Manufacturing site:	Laerdal Medical AS P.O. Box 377 Tanke Svilandsgate 30 4002 Stavanger Norway	
Product Name:	Resusci Baby QCPR	
Product Options:	161-01260 162-01260 163-01260	Resusci Baby QCPR Wireless Resusci Baby QCPR Airway Head Wireless Resusci Baby QCPR RQI Wireless
Accessories	206-300xx 170-30050 181-80025 161-15000	SimPad PLUS SkillReporter SkillGuide Cable, USB micro Am to CM Resusci Baby QCPR Elect. Upgrade

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 V3.1.1 EN 61000-6-1:2007 EN 61000-6-3:2007+A1:2011 Safety (Article 3.1(a) of RED): EN 62368-1:2014 EN 62479:2010 Radio (Article 3.2 of RED): EN 300 328 V2.2.2

Stavanger, 30th August 2022

<u>Muildus Dutra</u> Regulatory Affairs Specialist

CE

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