Docusign Envelope ID: A5FD9A2D-6B61-8CCF-82D3-F38D22B7B2D2

00064764 Rev I

Declaration of Conformity

Responsible Manufacturer:	Laerdal Medical AS Seehusens gate 1, P.O. Box 377 4002 Stavanger, Norway
Manufacturing site:	Laerdal Medical Monterrey SA de CV Avenida Las Torres 905 Parque Industrial Guadalupe Guadalupe N.L. 67190 Mexico
Product Name:	Resusci Baby QCPR, wireless
Product Options:	
161-01260	Resusci Baby QCPR, wireless
162-01260	Resusci Baby QCPR AW, wireless
163-01260	Resusci Baby QCPR RQI, Light wireless
163-01360	Resusci Baby QCPR RQI-Go, Light wireless
Accessories:	
206-300xx	SimPad PLUS SkillReporter
170-30050	SkillGuide
181-80025	Cable, USB micro Am to CM 1.8m, USB 2.0
161-15000	Resusci Baby QCPR Elect. Upgrade

to which this declaration relates is in conformity with

EU Council Directives

Council Directive EU 2014/53/EU on Radio Equipment Directive (RED) and Council Directive 2011/65/EU on Restriction on the use of certain hazardous substance (RoHS), as amended by EU 2015/863

UK Regulations

Radio Equipment Regulations SI 2017/1206 and RoHS Regulations SI 2012/3032

All supporting documentation is retained by the manufacturer.

The conformity is based on the following standards:

EMC:

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:

EN 62368-1:2014
EN 62479:2010

Radio:

EN 300 328 V2.2.2

DocuSigned by:
Mari Kaada
60D9901BD6AA495...

SIGNATURE

Corporate Director Q&R
POSITION IN ORGANISATION

Mari Kaada
PRINT NAME

2026-08-27
DATE