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

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EU DECLARATION OF CONFORMITY

Responsible Manufacturer:	Laerdal Medical AS Seehusens gate 1, P.O. Box 377 4002 Stavanger, Norway	
Manufacturing sites:	Laerdal Medical (Suzhou) Co., Ltd. Building 18, 19, 20 No. 57 Huoju Road, SND, Suzhou, Jiangsu Province China 215009	Printec H.T. electronics Corp. Liyan Street No.38 Zhong-He District New Taipei City 23557, Taiwan
Product Names:	**Little Baby QCPR** **Little Baby QCPR 4-pack** **Little Family QCPR**	

Product Options:	**133-01050**	**Little Baby QCPR**
	133-03050	**Little Baby QCPR Dark**
	134-01050	**Little Baby QCPR 4-pack**
	134-03050	**Little Baby QCPR 4-pack Dk**
	136-01050	**Little Family QCPR**
	136-03050	**Little Family QCPR Dark**
Accessories:	**170-30050**	**SkillGuide**

to which this declaration relates is in conformity with
Essential Requirements of the Council Directive 2014/53/EU on radio equipment (RED); and **Council Directive 2011/65/EU on Restriction on the use of certain hazardous substance (RoHS), as amended by EU 2015/863**

All supporting documentation is retained by the manufacturer.

The conformity is based on the following standards:

Safety (Article 3.1(a) of RED):
- EN 62368-1:2014+A11

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
- EN 62479:2010

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

Stavanger, 2026-08-27

DocuSigned by:
Mari Kaada
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Corporate Director Q&R

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