



Laerdal

helping save lives

00019022 Rev G

EU DECLARATION OF CONFORMITY

Responsible Manufacturer:	Laerdal Medical AS P.O. Box 377 Tanke Svilandsgate 30 4002 Stavanger Norway	
Manufacturing site: <i>(If other than mentioned above)</i>	Manufacturing site for 170-30050 SkillGuide: PRINTEC H.T. ELECTRONICS CORP. Liyan Street No.38 Zhong-He District, New Taipei City 23557 Taiwan	
Product Name:	Resusci Baby QCPR	
Product Options:	162-01201	Resusci Baby QCPR HeartCode Full Body Suitcase
	163-01001	Resusci Baby QCPR RQI
	163-01301	Resusci Baby QCPR RQI-Go
Accessories	170-30050	SkillGuide

to which this declaration relates is in conformity with
Essential Requirements of the Council Directive 2014/30/EU; and
Council Directive 2011/65/EU on Restriction on the use of certain hazardous substances (RoHS).

All supporting documentation is retained by the manufacturer.

The conformity is based on the following standards:

EN 61000-6-1:2007	Generic standards – Immunity for residential, commercial and light-industrial environments commercial and light-industrial environments
EN 61000-6-3:2007 +A1:2011+A2:2012	Generic standards – Emission standard for residential, commercial and light-industrial environments residential, commercial and light-industrial environment
EN/IEC 61000-3-2:2006 +A1+A2	Electromagnetic compatibility (EMC) - Part 3-2: Limits- Limits for harmonic current emissions (equipment input current ≤ 16 A per phase)
EN/IEC 61000-3-3:2008	Electromagnetic compatibility (EMC) - Part 3-3: Limits - Limitation of voltage changes, voltage fluctuations and flicker in public low-voltage supply systems, for equipment with rated current ≤ 16 A per phase and not subject to conditional connection

Stavanger, 5th September 2022


Regulatory Affairs Specialist




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