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

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Declaration of Conformity

Responsible Manufacturer:	Laerdal Medical AS Seehusens gate 1, P.O. Box 377 4002 Stavanger, Norway
Manufacturing site:	Laerdal Labs Vienna GmbH Lehargasse 1/5 1060 Vienna Austria
Product Name:	Paul
Product Description	Preterm Patient Simulator
Product Options:	
PHE02	Paul 02
Accessories:	
HA02	Stethoscope

to which this declaration relates is in conformity with
EU Council Directives
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:
EMC:
- EN 301 489-1 V2.2.3
- EN 301 489-17 v3.2.4
- EN IEC 61000-6-1:2019
- EN IEC 61000-6-3:2021

Safety:
- EN IEC 62368-1:2020+A11:2020
- EN 62311:2008

Radio:
- EN 300 330 V2.1.1
- EN 300 328 v2.2.2
- EN 301 893 v2.1.1
- EN 300 440 v2.1.1
- EN 50665:2017

DocuSigned by:
Mari Kaada
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SIGNATURE	POSITION IN ORGANISATION
Mari Kaada	Corporate Director Q&R
PRINT NAME	DATE
	2026-08-27