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

00035704 Rev F

Declaration of Conformity

Responsible Manufacturer:	Laerdal Medical AS Seehusens gate 1, P.O. Box 377 4002 Stavanger, Norway
Manufacturing site:	Laerdal Medical AS, Stavanger Manufacturing Vestre Svanholmen 3 4313 Sandnes, Norway
Product Name:	PrematureAnne
Product Options:	
295-00050	PrematureAnne Standard

to which this declaration relates is in conformity with

EU Council Directives

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

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:

ETSI EN 301 489-1 V2.2.3
ETSI EN 301 489-17 V3.2.6
EN 61000-4-3: 2006 A1: 2008 A2: 2010
EN 61000-4-2: 2009
EN 55032:2015

Safety:

EN 62368-1:2014
EN 62479:2010

Radio:

EN 300 328 V2.2.2

DocuSigned by: Mari Kaada 50D99013D6AA495...	Corporate Director Q&R
SIGNATURE	POSITION IN ORGANISATION
Mari Kaada	2026-08-27
PRINT NAME	DATE