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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 Medical AS, Stavanger Manufacturing Vestre Svanholmen 3 4313 Sandnes Norway
Product Name:	**SimMan ALS**
Product Options:	
235-02350	SimMan ALS Light
235-02150	SimMan ALS Medium
235-02250	SimMan ALS Dark
235-03350	SimMan ALS LiveShock L
235-03150	SimMan ALS LiveShock M
235-03250	SimMan ALS LiveShock D

to which this declaration relates is in conformity with
EU Council Directives
Directive 2014/53/EU on Radio Equipment (RED), including the cybersecurity requirements introduced by Commission Delegated Regulation (EU) 2022/30, 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 55032:2015//A1:2020
- EN 55035:2017/A11:2020
- EN 301 489-1 V2.2.3
- EN 301 489-17 v3.2.4

Safety:
- IEC 62368-1:2018
- EN 62311: 2008

Radio:
- EN 300 328 V2.2.2
- EN 301 893 V2.1.1
- EN 18031-1:2024
- EN 18031-2:2024

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