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 Medical AS, Stavanger Manufacturing Vestre Svanholmen 3 4313 Sandnes, Norway
Product Name:	**SimMan Essential** **SimMan Essential Bleeding**

Product Options:

213-02155	SimMan Essential Medium Manikin with active eyes
213-02255	SimMan Essential Dark Manikin with active eyes
213-02355	SimMan Essential Light Manikin with active eyes
214-02155	SimMan Ess. Bleed Medium Manikin with active eyes
214-02255	SimMan Ess. Bleed Dark Manikin with active eyes
214-02355	SimMan Ess. Bleed Light Manikin with active eyes

to which this declaration relates is in conformity with

EU Council Directives

EU Radio Equipment Directive (RED) 2014/53/EU, 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/A11:2020
EN 55035:2017/A11:2020

Safety:

IEC 62368-1:2014

Radio:

EN 18031-1:2024
EN 18031-2:2024

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