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00072888 Rev I

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 3G PLUS**
Product Options:	
212-03150	SimMan 3G PLUS Medium Manikin and Accessories
212-03250	SimMan 3G PLUS Dark Manikin and Accessories
212-03350	SimMan 3G PLUS Light Manikin and Accessories
Accessories:	
212-14550-L	Arm Kit SM3G Light Left + Right + RFID kit
212-14550-M	Arm Kit SM3G Medium Left + Right + RFID kit
212-14550-D	Arm Kit SM3G Dark Left + Right + RFID kit

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 50D90013D6AA495...	Corporate Director Q&R
SIGNATURE	POSITION IN ORGANISATION
Mari Kaada	2026-08-27
PRINT NAME	DATE