Docusign Envelope ID: 85BB5844-DE96-410B-8DB8-D06A1523AD72

00099612 Rev A

Supplier's Declaration of Conformity

under FCC Part 15 SDoC procedure

Product

Product Description	Preterm Patient Simulator
Model Number	PHE02 Paul 02
Trade Name	Laerdal

Test Report

Test Report No.	00099038 (EMV-E 55/23)
Standard	FCC 47 CFR Part 15 Subpart B, Class A

Testing Laboratory

Name	Seibersdorf Labor GmbH
Address	2444 Austria
FCC Test Site Registration Number	835627
FCC Accredited CAB Designation Number	AT0002

Responsible Party

Name	Laerdal Medical Corporation
Address	167 Myers Corners Road Wappingers Falls, New York 12590 USA
Telephone	877-LAERDAL (523-7325)

FCC Compliance Statement

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions: (1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

For and on behalf of Responsible Party,

Signed by: Gina Bashai SIGNATURE	QA/RA Manager POSITION IN ORGANISATION
QA/RA Manager PRINT NAME	2025-10-08 DATE