Docusign Envelope ID: 60E4F360-EDB7-41E9-B635-9F010A901F10

00038930 Rev H

Supplier's Declaration of Conformity

under FCC Part 15 SDoC procedure

Product

Product Description	Resusci Anne Simulator, AED or Paddle Resusci Anne Advanced Skill Trainer, AED or Paddle
Model Number	150-22000 150-23000 150-27000 150-28000 151-22000 151-23000 151-27000 151-28000
Trade Name	Laerdal

Test Report

Test Report No.	00095800
Standard	FCC 47 CFR Part 15 Subpart B, Class A

Testing Laboratory

Name	Intertek Semko Korea Ltd.
Address	Intertek Building 3, Gongdan-ron 160beon-gil, Gunpo-si, Gyeonggi-do, 15845 Korea
FCC Test Site Registration Number	
FCC Accredited CAB Designation Number	KR0163

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 Beshai SIGNATURE	QA/RA Manager POSITION IN ORGANISATION
Gina Beshai PRINT NAME	2025-07-08 DATE