

# DECLARATION OF CONFORMITY

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**Responsible  
Manufacturer:** Laerdal Medical AS  
P.O. Box 377  
Tanke Svilandsgate 30  
4002 Stavanger  
Norway

**Manufacturing site:** Laerdal Medical (Suzhou) Co. Ltd  
No 19 Building, Huoju Road  
Science & Technology Industrial Park  
Suzhou, Jiangsu, P.R. China

**Product Name:** **Little Anne AED**

**Product Options:** 122-01050 Little Anne AED  
122-50750 LA AED Upgrade Kit

to which this declaration relates is in conformity with  
**Essential Requirements of the Council Directive 2014/30/EU relating to  
Electromagnetic Compatibility (EMC); 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:

EN 61000-6-1:2007	Generic standards – Immunity for residential, commercial, and light industrial environments
EN 61000-6-3:2007 +A1:2011+A2:2012	Generic standards – Emission standard for residential, and light industrial environments

**Stavanger, 13<sup>th</sup> September 2022**

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*Mecildes Dutra*  
**Regulatory Affairs**



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