EU DECLARATION OF CONFORMITY

Responsible Manufacturer:	Laerdal Medical AS P.O. Box 377 Tanke Svilandsgate 30 4002 Stavanger Norway
Single Registration Number:	NO-MF-000000012
Manufacturing site:	Laerdal Medical (Suzhou) Co. Ltd. Building 18,19,20, No. 57 Huoju Road Science & Technology Industrial Park Suzhou, Jiangsu Province 215009, China
Product Name:	300 ml Disposable Canister
Basic UDI-DI:	0704543209955T9
Intended Purpose:	The 300 ml Disposable Canister is intended for use with the Laerdal Compact Suction Unit 4 to collect suctioned bodily fluids from the patient's airways.
Product Options:	886100 300 ml Disposable Canister with tubing 886100L10 10-pack 300 ml canister

to which this declaration relates is in conformity with the General Safety and Performance Requirements of EU Regulation 2017/745; and **Council Directive 2011/65/EU on Restriction on the use of certain hazardous substance (RoHS)**

Classification:

The 300 ml Disposable Canister is class I according to rule 1 of Annex VIII of the EU Medical Device Regulation.

Conformity Assessment is based on the principles described in Article 52 of Regulation 2017/745.

Conformity is declared in relation to common Specification(s):

No Common Specifications are available at this time for Canisters.

This EU Declaration of Conformity is issued under the sole responsibility of Laerdal Medical AS.

DocuSigned by:
Claudia Pajares Ibáñez
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Claudia Pajares Ibanez

Regulatory Affairs Manager – Medical Device
On behalf of CEO Alf Christian Dybdahl
18 February 2026
Stavanger, Norway