



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 Huojia Road Science & Technology Industrial Park Suzhou, Jiangsu Province 215009 China
Product Name:	Stifneck Select Extrication Collar Stifneck Pedi-Select Extrication Collar
Basic UDI-DI:	0704543209932SV
Intended Purpose:	Intended to support cervical spine in a neutral position during transportation, either alone or in combination with other cervical and full body immobilization devices (CSIDs and FBSIDs).
Product Options:	980010 Stifneck Select Extrication Collar (US/AU) 98001005 Stifneck Select Extrication Collar(J) 98001033 Stifneck Select Extrication Collar 980012 Stifneck Select Green Extrication Collar (US/AU) 98001205 Stifneck Select Green Extrication Collar (J) 98001233 Stifneck Select Green Extrication Collar 980020 Stifneck Pedi-Select Extrication Collar (US/AU) 98002005 Stifneck Pedi-Select Extrication Collar(J) 98002033 Stifneck Pedi-Select Extrication Collar

to which this declaration relates is in conformity with the General Safety and Performance Requirements of EU Regulation 2017/745.

Classification:

The Stifneck Select Extrication Collars are 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 CS for Extrication Collars has been published at this time

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

Adrienne Farrington
Regulatory Affairs Specialist
On behalf of CEO Alf Christian Dybdahl
14th August 2025
Stavanger Norway

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Adrienne Farrington

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