EC-CONFORMITY DECLARATION

We hereby declare that **"Laerdal BaXstrap model 982500/982600 and Laerdal BaXstrap Custom SP model 982599/982699"**,

manufactured for:
Laerdal Medical AS
P.O.Box 377
N-4002 Stavanger
Norway

is manufactured in conformity with the product described in Product Configuration Document and the Product Device Master Record Index and other information found in the technical file.

The product has been tested against the following standards and specifications:
NS-EN 1865:2000 Specifications for stretchers and other patient handling equipment used in road ambulances.

Compliance is claimed for the essential requirements in council directive **93/42/EEC Medical Device Directive** regarding the harmonisation of the member-states' regulations regarding medical devices.

The product is in risk class **I**, according to Rule II in Appendix IX of the Directive.

For Laerdal Medical AS

Pål Frafjord
Technical Product Manager

23.04.2004
Date

CE

Laerdal Medical AS
P.O.Box 377
N-4002 Stavanger
Telephone: 51511700
Fax: 51511788

REV	DESCRIPTION OF CHANGE	CHANGE REFERENCE	ORIGINATOR	VERIFIED	APPROVED
			DATE AND SIGN	DATE AND SIGN	DATE AND SIGN
B	Added standard NS-EN 1865:2000	LN 04-115	DATE AND SIGN 23.04.04	DATE AND SIGN 23.04.04	DATE AND SIGN 23.04.04
A	Release	LN 03-170 LN 03-171	DATE AND SIGN	DATE AND SIGN	DATE AND SIGN

DMR ☒
DHF ☐

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REFERENCE	REPLACES	REV	TITLE
N/A	N/A		**BaXstrap EC- Conformity Declaration (CE Marking)**

PRODUCT GROUP	TOOL	PART NO	REV	DOCUMENT NO	PAGE	ATTACH	REV
98	N/A	See above		PRO-RP01-0266	1 of 1	0	B