

# EC-CONFORMITY DECLARATION

We hereby declare that **"Laerdal Stifneck Set - 980000 and Combo Pack - 980016"** which are

manufactured by:

Laerdal Medical (Suzhou) Co. Ltd.  
 No.19 Building, Huoju Rd.  
 Science & Technology Industry Park  
 Suzhou New District  
 215009, Suzhou  
 P.R. China

and manufactured for:

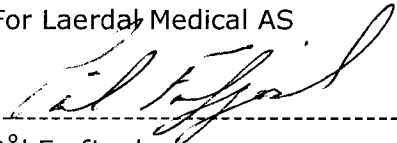
Laerdal Medical AS  
 P.O.Box 377  
 N-4002 Stavanger  
 Norway

are 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.

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


  
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 Pål Frafjord  
 Technical Product Manager

21.06.2004

Date



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		
			DATE AND SIGN	DATE AND SIGN	DATE AND SIGN		
A	Release	9092	DATE AND SIGN 21/06-04 [Signature]	DATE AND SIGN 21/06-04 [Signature]	DATE AND SIGN 21/06-04 [Signature]		
THE INFORMATION CONTAINED HEREIN IS PROPRIETARY TO LAERDAL AND SHALL NOT BE USED FOR ANY PURPOSE DETRIMENTAL TO OR BE REPRODUCED WITHOUT PERMISSION OF LAERDAL		DMR <input checked="" type="checkbox"/> DHF <input type="checkbox"/>					
REFERENCE	N/A	REPLACES	N/A	REV	TITLE		
					<b>Stifneck Set &amp; Combo Pack EC- Conformity Declaration (CE Marking)</b>		
PRODUCT GROUP	TOOL	PART NO	REV	DOCUMENT NO	PAGE	ATTACH	REV
98	N/A	See above		PRO-RP01-0311	1 of 1	0	A