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CVR. nr. 63644919

EC Declaration of Conformity No.: REG-000139

Product Name:	Ambu® SPUR® II, Single Patient Use Resuscitator Adult, Pediatric and Infant		
REF and Configuration	Adult	52x xxx xxx	325 0xx 000
	Pediatric	53x xxx xxx	330 0xx 000
	Infant	54x xxx xxx	335 xxx 000
	Customized items: 325021000RH, 330021000RH, 335002000RH, 3350003000RH		
Accessory:	Ambu® Disposable Face masks Ambu® PEEP valve		

Identification: All products manufactured after issue date.

We the manufacturer hereby declare that this product is in conformity with the requirements in:

Council directive 93/42/EEC, Annex II enforced in Danish law.
Equipment class: IIa, non-sterile, Annex IX rule 2

and in accordance with the authorization given by BSI Certificate No. CE 68734.

For and on behalf of Ambu A/S, Denmark

16 Jun 2014

Kaja Tengbjerg, Senior Regulatory Affairs Professional

First Issued: 29 June 2005

EC Declaration of Conformity – Annex I: GMDN Code

No.: REG-000139

Product Name:

Ambu® SPUR® II Single Patient Use Resuscitator Adult, Pediatric and Infant

The Ambu® SPUR® II Single Patient Use Resuscitator Adult, Pediatric and Infant are covered by the following GMDN Code:

GMDN Code: 36086
Term: Manual pulmonary resuscitator, single-use