

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

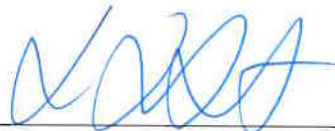
Identification: All products manufactured after issue date.

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

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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