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RoHS EU Declaration of Conformity No.: REG-001426

Product Name: Ambu® Neuroline Subdermal
Ambu® Neuroline Twisted Pair Subdermal

REF and Configuration
745 12-YY/NN
746 12-YY/Z/NN
YY states the pre-attached lead wire length: 50, 100, 150, 200 and 250 cm.
NN states number of pouch in a box
Z states lead wire color combination
Subdermal: Color-coded red, blue, green, yellow, black and white
Twisted Pair Subdermal: Ten colours twisted with white and ten colours twisted with black

Accessory: None

ROHS2 Category of the device Category 8 – Medical devices

Identification: All products manufactured after issue date.

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

DIRECTIVE 2011/65/EU OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 8 June 2011
on the restriction of the use of certain hazardous substances in electrical and electronic equipment (RoHS recast) as it specified in Article 4.

This Declaration of Conformity is issued under the sole responsibility of Ambu A/S.

For and on behalf of Ambu A/S, Denmark

26 June 2014

Laila Strange Lundtoft, Senior Manager Corporate Regulatory Affairs

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