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CVR. nr. 63644919

RoHS EU Declaration of Conformity No.: REG-001422

Product Name: Ambu® Neuroline

REF and Configuration

700-XX-Y/NN ; 700-XX-K/C/NN
710-XX-Y/NN ; 710-XX-K/C/NN
715-XX-K/C/NN
720-XX-Y/NN ;720-XX-K/C/NN ;720-00-S/NN
725-XX-Y/NN

XX states the lead wire length: 10, 20, 50, 80, 100, 120, 150, 200 cm
Y states the connector type: A, J, K, SC
(Not all the connector types and wire length apply for the mentioned cat. No.)
NN states number of electrodes in a pouch

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

23 June 2014

Laila Strange Lundtoft, Senior Manager Corporate Regulatory Affairs

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