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CVR. nr. 63644919

RoHS EU Declaration of Conformity No.: REG-001425

Product Name: Ambu® Neuroline Inoject

REF and Configuration

744 25-30/NN
744 30-36/NN
744 35-40/NN
744 38-45/NN
744 50-50/NN
744 75-55/NN

NN states number of electrodes in a box

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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