

EU Declaration of Conformity

No.: REG-002421

Product Name: Ambu® AmbuMan® Wireless
Ambu® AmbuMan® Defib
Ambu® AmbuMan® Advanced IV
Ambu® AmbuMan® Advanced
Ambu® AmbuMan® Airway Wireless

REF and Configuration	Ambu® AmbuMan® Wireless	A234407000
	Ambu® AmbuMan® Defib	A265407000
	Ambu® AmbuMan® Advanced IV	A284407000
	Ambu® AmbuMan® Advanced	A284417000
	Ambu® AmbuMan® Airway Wireless	A284427000

ROHS Category of the device Category 11 – Other EEE

Device classification Class 1, according to the RED directive, the product falls within Class 1 (sub-class 22) equipment of Commission Decision 2000/299/EC of 6 April 2000.

Identification: All products manufactured after issue date.

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

Restriction of Hazardous Substances Directive 2011/65/EU as amended by Commission Delegated Directive (EU) 2015/863 on the restriction of the use of certain hazardous substances in electrical and electronic equipment (RoHS recast) as it specified in Article 4.

Radio Equipment Directive 2014/53/EU

The following harmonized standards and technical specification have been applied: DSC-029143 List of Applied Standards

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

For and on behalf of Ambu A/S

Gitte Juel Friis
Head of Regulatory Affairs, Vigilance, ANE & PM, Market Expansion

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