rossmax	EU Declaration of Conformity	Doc No.	RD-4-136
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		Effective Date	2021/6/2

EU Declaration of Conformity

Manufacturer Address SRN	: :	Rossmax Swiss GmbH Widnauerstrasse 1, CH-9435 Heerbrugg, Switzerland CH-MF-000011245
Representative in Europe Address	:	CMC Medical Devices& Drugs S.L. C/ Horacio Lengo No 18, CP 29006, Málaga, Spain
Basic UDI-DI	:	4715139CuffBC
Product name	:	Cuff
Product code	:	Cone cuff, Medical cuff, DK cuff
Conformity Assessment	:	Regulation (EU) 2017/745 of the European Parliament and of the Council ANNEX II, ANNEX III, ANNEX IV
Classification	:	Class I (According to g Regulation (EU) 2017/745, Annex VIII, Rule 1)
Lot No.	:	YY M XXX Sequential number Manufacturing month. Manufacturing year.

The above-mentioned devices are in full compliance with the relevant provisions of Regulation (EU) 2017/745 of the European Parliament and of the Council, Annex I-General Safety and Performance Requirements and applied harmonized standards, national standards or other normative documents and, if applicable, common specification.

EN ISO15223-1:2021; EN 1041:2008; EN ISO 10993-1:2009/ AC2010; EN ISO 10993-5:2009; EN ISO 10993-10:2013; EN1060-1:1995+A2:2009

Rossmax Swiss GmbH is exclusively responsible for the declaration of conformity.

This declaration is limited by the issuing of a revised declaration of conformity after change of the product.

Yolanda Lin

Signature: **Yolanda Lin, Management Representative**

Date: Feb. 20, 2023