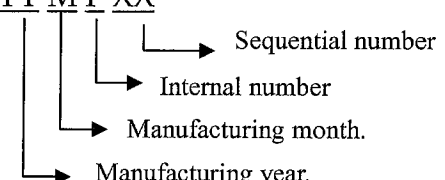


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Effective Date	2014/02/14

1. EC Declaration of Conformity

Manufacturer Address	: Rossmax Swiss GmbH Widnauerstrasse 1, CH-9435 Heerbrugg, Switzerland
Notified Body Address EU Identification No.	: SGS Belgium NV : SGS House Noorderlaan 87 2030 Antwerp Belgium : 1639
Certificate No.	: TW19/20056
Representative in Europe Address	: CMC Medical Devices & Drugs S.L. C/ Horacio Lengo No 18, CP 29006, Málaga, Spain
Product type	: Nebulizer (including nebulizer pack, nebulizer bottle set) for respiratory therapy
Type Designation	: nose piece(NB_AC_020_00, NB_AC_040_00), mouth piece(NB_AC_021_00, NB_AC_031_00) Nasal Aspirator(NS1)
Conformity Assessment	: EU Council Directive 93/42/EEC amended by 2007/47/EC Annex II (excluding Section 4) and Regulation (EU) 2023/607 amendment of Regulations (EU) 2017/745 as regards the transitional provisions for certain medical devices
Classification	: Class IIa (According to EU Council Directive 93/42/EEC amended by 2007/47/EC, Annex IX, Rule 5)
Lot No.	: <u>YY M P XX</u> <div style="margin-left: 40px;">  </div>

The above-mentioned devices are in full compliance with the relevant provisions of EU Council Directive 93/42/EEC amended by 2007/47/EC, Regulation (EU) 2023/607 amendment of Regulations (EU) 2017/745, Annex I-Essential Requirements and applied harmonized standards, national standards or other normative documents.

EN ISO 10993-1:2009/AC2010, EN ISO 10993-5:2009, EN ISO 10993-10:2013, EN ISO 13485:2016, EN ISO 14971:2012

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 and/or the expiration date (31.12.2028) of the certificate of 93/42/EEC.

Yolanda Lin

Signature: Yolanda Lin, Management Representative

Date: Jun. 13, 2023