EU DECLARATION OF CONFORMITY

RESPIRONICS®

Doc Number REG 2101464
Revision 08

This Declaration of Conformity is issued under the sole responsibility of the manufacturer. The device(s)/accessories covered by this Declaration are in conformity with all regulations or directives below.

1. Object of the declaration:

Product Name:	Pico Traditional Nasal Mask
Product Type:	Nasal Mask
Intended Purpose:	The Pico Nasal Mask is intended to provide an interface for application of CPAP or bi-level therapy to patients. The mask is for single patient use in the home or multi-patient use in the hospital/institutional environment. The mask is to be used on patients (>66lbs/30kg) for whom CPAP or bi-level therapy has been prescribed.
Product Part Number(s) and Descriptions:	Part Number(s) listed in this section comply with all directives indicated in DoC unless otherwise noted. The following part numbers are compliant with 93/42/EEC Medical Devices Directive, as amended up to and inclusive of Council Directive 2007/47/EC:

Part Number	Device Name/Description
1104940	Fitpack S/M, L, XL Pico Traditional Nasal Mask w/HGR
1104915	S/M Pico Traditional Nasal Mask w/HGR
1104916	L Pico Traditional Nasal Mask w/HGR
1104917	XL Pico Traditional Nasal Mask w/HGR
1104918	S/M Pico Traditional Nasal Mask W/O HGR
1104919	L Pico Traditional Nasal Mask W/O HGR
1104920	XL Pico Traditional Nasal Mask W/O HGR
1104921	S/M Pico Traditional Nasal Mask w/HGR, INTL
1104922	L Pico Traditional Nasal Mask w/HGR, INTL
1104923	XL Pico Traditional Nasal Mask w/HGR, INTL
1104921MY	Small/Medium Pico Nasal Mask with Headgear Malaysia
1104922MY	Large Pico Nasal Mask with Headgear Malaysia
1104923MY	Extra Large Pico Nasal Mask with Headgear Malaysia
1104940MY	Small/Medium, Large, Extra Large Pico Fitpack, with Headgear, Malaysia

EU DECLARATION OF CONFORMITY

RESPIRONICS®

Doc Number REG 2101464
Revision 08

Product Options/Accessories Part Number(s) and Descriptions:	Refer to REG 22849 for accessory headgear and chin straps.
Basic UDI-DI:	N/A
Control Indicator:	

Initial Issue Date:	REF (Part Number):
17-Jun-14	1104940, 1104915, 1104916, 1104917, 1104918, 1104919, 1104920
26-Sep-14	1104921, 1104922, 1104923
11-Jan-21	1104921MY, 1104922MY, 1104923MY, 1104940MY

Global Medical Device Nomenclature code (GMDN) and Description	57815 CPAP/BiPAP Face Mask Reusable

The object of the Declaration (product part numbers and if applicable the component part numbers) that are described above as included in this DoC is / are in conformity with the following regulations / directives. If optional accessories that are used in combination with the product (medical device), but are CE marked on their own are referenced in this DoC, they are excluded further on as these optional accessories have their own DoC.

EU Directive	**Council Directive 93/42/EEC of 14 June 1993 concerning medical devices amended up to and inclusive of Council Directive 2007/47/EC (MDD)** **<This DoC may not be used to declare conformity to both the MDD and the MDR at the same time.>**
Risk Classification	Class IIa based on Annex IX and Rule 2
Conformity Assessment Route	Annex II excluding (4)
Notified Body Name, Address, and ID	TÜV SÜD Product Service GmbH Ridlerstrasse 65 80339 München, Germany 0123
Certificate(s) Issued	TÜV SÜD EC Certificate Number: G1 015581 0611
Standards	The products listed on this Declaration of Conformity have been assessed and/or tested in a typical configuration as described in

	the Manufacturer's accompanying documentation in accordance with the product standards listed below. Refer to Attachment A.

2. Mandatory information:

Manufacturer	Respironics, Inc. 1001 Murry Ridge Lane Murrysville, PA 15668 USA
EU Authorized Representative (AR):	Respironics Deutschland GmbH & Co. KG Gewerbestrasse 17 82211 Herrsching, Germany Tel: +49 8152 93060
ISO Quality Certificates Issued:	The Manufacturer is certified by TÜV SÜD to the following: TÜV SÜD EN ISO 13485:2016 Certificate Number: Q5 015581 0609 TÜV SÜD MDSAP Certificate Number: QS6 112601 0001

Signature (signed for and on behalf of)

Date of Issue: 09 November 2022

Printed Name: Jennifer Richardson

Place of Issue: Murrysville, PA, USA

Title: Senior Manager, Regulatory Affairs

This declaration is valid until: 26 May 2024

3. Attachment A Standards and/or Common Specifications

Standard	Standard Title
Quality System	
EN ISO 13485:2016	Medical devices – Quality management systems – Requirements for regulatory purposes
Particular Safety Standards	
Patient Interface	
EN ISO 17510-2:2009	Sleep apnoea breathing therapy – Part 2: Masks and application accessories
Biocompatibility	
EN ISO 10993-1:2020	Biological evaluation of medical devices – Part 1: Evaluation and testing within a risk management process
EN ISO 10993-3:2009	Biological evaluation of medical devices – Part 3: Tests for genotoxicity, carcinogenicity and reproductive toxicity
EN ISO 10993-5:2009	Biological evaluation of medical devices – Part 5: Tests for in vitro cytotoxicity
EN ISO 10993-6:2009	Biological evaluation of medical devices – Part 6: Tests for local effects after implantation
EN ISO 10993-10:2013	Biological evaluation of medical devices – Part 10: Tests for irritation and skin sensitization
ISO 18562-1: 2017	Biocompatibility evaluation of breathing gas pathways in healthcare applications – Part 1: Evaluation and testing within a risk management process
ISO 18562-2: 2017	Biocompatibility evaluation of breathing gas pathways in healthcare applications – Part 2: Tests for emissions of particulate matter
ISO 18562-3: 2017	Biocompatibility evaluation of breathing gas pathways in healthcare applications – Part 3: Tests for emissions of volatile organic compounds (VOCs)
ISO 18562-4: 2017	Biocompatibility evaluation of breathing gas pathways in healthcare applications – Part 4: Tests for leachables in condensate
Other Standards	
Accompany Documents and Labeling	
EN 1041:2008/ A1:2013	Information supplied by the manufacturer of medical devices
EN ISO 15223-1: 2017	Medical devices – Symbols to be used with medical device labels, labelling and information to be supplied – Part 1: General requirements
Risk Management	
ISO 14971:2019	Medical devices – Application of risk management to medical devices
Usability	
IEC 62366-1:2015	Medical devices – Application of usability engineering to medical devices
Tubing and Connections	
ISO 5356-1:2015	Anaesthetic and respiratory equipment – Conical connectors – Part 1: Cones and sockets
Cleaning and Disinfection	
EN ISO 17664: 2017	Processing of health care products - Information to be provided by the medical device manufacturer for the processing of medical devices