DECLARATION OF CONFORMITY TO REGULATION(EU) 2017/745 ON MEDICAL DEVICES

CONTEC MEDICAL SYSTEMS CO., LTD.
No.112 Qinhuang West Street, Economic ＆Technical Development Zone, Qinhuangdao, Hebei Province, PEOPLE’S REPUBLIC OF CHINA

SRN of Manufacturer :CN-MF-000007715

EU REP Prolinx GmbH
Brehmstr. 56, 40239, Duesseldorf, Germany

SRN of Authorised Representative:DE-AR-000005129

This EU declaration of conformity is issued under the sole responsibility of the manufacturer.
We keep all supporting documentation and ensure that the authorised representative has the necessary documentation permanently available.

Basic UDI-DI:	69450401SP70BLZ
Product and Trade Name:	SPIROMETER
EMDN Code:	Z12150101
Catalogue Number/model:	SP80B
Risk Class of the Device:	Class IIa according to Annex VIII Rule 10

We, （CONTEC MEDICAL SYSTEMS CO., LTD.）herewith declare that the stated medical devices meet REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 5 April 2017 on medical devices.
The harmonised standards and/or CS used and in relation to this conformity see annex.

NOTIFIED BODY:	TÜV SÜD Product service GmbH Ridlerstr 65, D-80339 München, Germany
NOTIFIED BODY IDENTIFICATION NUMBER:	0123
CONFORMITY ASSESSMENT PROCEDURE:	Regulation (EU) 2017/745, Annex IX excluding chapter II
(EC) CERTIFICATE(S):	G15 050972 0058 Rev. 02
PLACE, DATE OF ISSUE:	Qinhuangdao, 2026/04/07
NAME AND FUNCTION, SIGNATURE:	HUKUN,Chairman/ manufacturer



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Appendix: list of (harmonised - EN) standards

No.	Standards	Title and Description
1	EN ISO 13485:2016/A11:2021	Medical devices - Quality management systems - Requirements for regulatory purposes (ISO 13485:2016)
2	EN ISO 14971:2019/A11:2021	Medical devices - Application of risk management to medical devices (ISO 14971:2019)
3	EN 60601-1:2006/A2:2021	Medical electrical equipment-Part 1: General requirements for basic safety and essential performance
4	EN 60601-1-2:2015/A1:2021	Medical electrical equipment-Part 1-2: General requirements for safety and essential performance - Collateral standard: Electromagnetic disturbances - Requirements and tests
5	EN 60601-1-11:2015/A1:2021	Medical electrical equipment - Part 1-11: General requirements for basic safety and essential performance - Collateral Standard: Requirements for medical electrical equipment and medical electrical systems used in the home healthcare environment
6	EN ISO 23747:2015	Anaesthetic and respiratory equipment - Peak expiratory flow meters for the assessment of pulmonary function in spontaneously breathing humans (ISO 23747:2015)
7	EN ISO 26782:2009	Anaesthetic and respiratory equipment - Spirometers intended for the measurement of time forced expired volumes in humans (ISO 26782:2009)
8	EN 60601-1-6:2010/A2:2021	Medical electrical equipment - Part 1-6: General requirements for basic safety and essential performance - Collateral Standard: Usability
9	EN 62366-1:2015/A1:2020	Medical devices - Part 1: Application of usability engineering to medical devices
10	EN 62304:2006/A1:2015	Medical device software - Software life-cycle processes
11	EN ISO 10993-1:2020	Biological evaluation of medical devices - Part 1: Evaluation and testing within a risk management process (ISO 10993-1:2018, including corrected version 2018-10)
12	EN ISO 15223-1:2021/A1:2025	Medical devices - Symbols to be used with information to be supplied by the manufacturer - Part 1: General requirements - Amendment 1: Addition of defined term for authorized representative and modified EC REP symbol to not be country or region specific (ISO 15223 1:2021/Amd 1:2025)
13	EN ISO 20417:2021	Medical devices - Information to be supplied by the manufacturer (ISO 20417:2021, Corrected version 2021-12)