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: 69450401CMS800GFM

PRODUCT AND TRADE NAME: Fetal Monitor

EMDN CODE: Z12080101

CATALOGUE NUMBER/MODEL: CMS800G

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

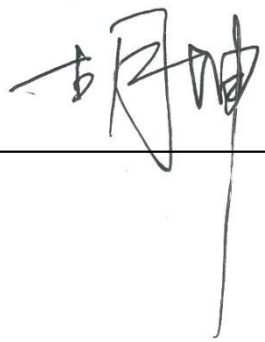
NOTIFIED BODY IDENTIFICATION NUMBER: 0123

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.	Standard	Title and Description
1	EN ISO 13485:2016/A11:2021	Medical devices - Quality management systems - Requirements for regulatory purposes (ISO 13485:2016)
2	EN 60601-1:2006/A2:2021	Medical electrical equipment - Part 1: General requirements for basic safety and essential performance
3	EN 60601-1-2:2015/A1:2021	Medical electrical equipment - Part 1-2: General requirements for basic safety and essential performance - Collateral Standard: Electromagnetic disturbances - Requirements and tests
4	EN 60601-1-6:2010/A2:2021	Medical electrical equipment - Part 1-6: General requirements for basic safety and essential performance - Collateral standard: Usability
5	EN 62366-1:2015/A1:2020	Medical devices - Part 1: Application of usability engineering to medical devices
6	EN 62304:2006/A1:2015	Medical device software - Software life-cycle processes
7	EN 60601-2-37:2008/A1:2015	Medical electrical equipment - Part 2-37: Particular requirements for the basic safety and essential performance of ultrasonic medical diagnostic and monitoring equipment
8	EN 61157:2007/A1:2013	Standard means for the reporting of the acoustic output of medical diagnostic ultrasonic equipment
9	EN ISO 14971:2019/A11:2021	Medical devices - Application of risk management to medical devices (ISO 14971:2019)
10	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)
11	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)
12	EN ISO 20417:2021	Medical devices - Information to be supplied by the manufacturer (ISO 20417:2021, Corrected version 2021-12)