


**DECLARATION OF CONFORMITY  
TO COUNCIL DIRECTIVE 93/42/EEC  
CONCERNING MEDICAL DEVICES**

<b>MANUFACTURER:</b>	<b>CONTEC MEDICAL SYSTEMS CO., LTD</b> No.112 Qinhuang West Street, Economic & Technical Development Zone, Qinhuangdao, Hebei Province, PEOPLE'S REPUBLIC OF CHINA
<b>MEDICAL DEVICE:</b>	SPIROMETER , SP10
<b>CLASSIFICATION - ANNEX IX:</b>	Class II a, Rule 10
<b>CONFORMITY ASSESSMENT ROUTE:</b>	Annex II excluding chapter 4
WE, ( CONTEC MEDICAL SYSTEMS CO., LTD ) HEREWITH DECLARE THAT THE STATED MEDICAL DEVICES. MEET THE TRANSPOSITION INTO NATIONAL LAW, THE PROVISIONS OF COUNCIL DIRECTIVE 93/42/EEC OF 14 JUNE 1993 CONCERNING MEDICAL DEVICES; INCLUDING, AT 21 MARCH 2010, THE AMENDMENTS BY COUNCIL DIRECTIVE 2007/47/EC ALL SUPPORTING DOCUMENTATION IS RETAINED AT THE PREMISES OF THE MANUFACTURE.	
STANDARDS APPLIED: SEE ATTACHED LIST OF (HARMONISED - EN) STANDARDS FOR WHICH DOCUMENTED EVIDENCE OF COMPLIANCE CAN BE PROVIDED.	
<b>NOTIFIED BODY:</b>	TÜV SÜD PRODUCT SERVICE GMBH RIDLERSTR 65, D-80339 MÜNCHEN, GERMANY
<b>IDENTIFICATION NUMBER:</b>	<b>CE</b> 0123
<b>(EC) CERTIFICATE(S):</b>	<u>G1 050972 0050 Rev.04</u>
<b>EUROPEAN REPRESENTATIVE:</b>	Shanghai International Holding Corp. GmbH(Europe) Eiffestrasse 80, 20537 Hamburg Germany

**START OF CE-MARKING:** 2011-01-14 (Date or Lot or serial number)

<b>PLACE, DATE OF DECLARATION:</b>	QINHUANGDAO, 2020-06-18
<b>SIGNATURE:</b>	 _____ President

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**Appendix: list of (harmonised - EN) standards**

<b>NO.</b>	<b>Reference</b>	<b>Title</b>
1	EN60601-1:1990+A1:1993+A2:1995	Medical electrical equipment- Part 1: General requirements for safety
2	EN 60601-1-2:2007	Medical electrical equipment- Part 1-2: General requirements for safety - Collateral standard: Electromagnetic compatibility - Requirements and tests
3	EN60601-1-4:1996+A1:1999	Medical electrical equipment - Part 1-4: General requirements for safety - Collateral standard: Programmable electrical medical systems
4	EN 60601-1-6:2007	Medical electrical equipment-Part 1-6: General requirements for basic safety and essential performance-Collateral Standard: Usability
5	EN ISO23747:2009 (ISO 23747:2007)	Anaesthetic and respiratory equipment - Peak expiratory flow meters for the assessment of pulmonary function in spontaneously breathing humans
6	EN 62304:2006	Medical device software-Software life-cycle processes