




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

 MANUFACTURER:	CONTEC MEDICAL SYSTEMS CO., LTD No.112 Qinhuang West Street, Economic & Technical Development Zone, Qinhuangdao, Hebei Province, PEOPLE'S REPUBLIC OF CHINA		
MEDICAL DEVICE:	Pulse Oximeter Probe GSE0030		
CLASSIFICATION - ANNEX IX:	Class II b, Rule 10		
CONFORMITY ASSESSMENT ROUTE:	Annex II excluding chapter 4		
<p>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/EEC ALL SUPPORTING DOCUMENTATION IS RETAINED AT THE PREMISES OF THE MANUFACTURE.</p>			
STANDARDS APPLIED: SEE ATTACHED LIST OF (HARMONISED - EN) STANDARDS FOR WHICH DOCUMENTED EVIDENCE OF COMPLIANCE CAN BE PROVIDED.			
NOTIFIED BODY:	TÜV SÜD PRODUCT SERVICE GMBH RIDLERSTR 65, D-80339 MÜNCHEN, GERMANY		
IDENTIFICATION NUMBER:	 0123		
(EC) CERTIFICATE(S):	<u>G1 050972 0050 Rev.04</u>		
<table border="1" style="display: inline-table; border-collapse: collapse;"><tr><td style="padding: 2px;">EC</td><td style="padding: 2px;">REP</td></tr></table> Prolinx GmbH:	EC	REP	Brehmstr. 56,40239,Duesseldorf, Germany
EC	REP		

START OF CE-MARKING: 2024-05-20 (Date or Lot or serial number)

PLACE, DATE OF DECLARATION:	QINHUANGDAO, 2020-06-18
SIGNATURE:	 _____ President

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Appendix: list of (harmonised - EN) standards

NO.	Reference	Title
1	EN 60601-1:2006	Medical electrical equipment - Part 1: General requirements for basic safety and essential performance
2	EN 60601-1-2:2007 (IEC60601-1-2:2007)	Medical electrical equipment- Part 1-2: General requirements for basic safety and essential performance - Collateral standard: Electromagnetic compatibility - Requirements and tests
3	EN 60601-1-6:2010 (IEC 60601-1-6:2010)	Medical electrical equipment-Part 1-6:General requirements for basic safety and essential performance-Collateral Standard: Usability
4	EN 62366:2008	Medical devices - Application of usability engineering to medical devices
5	ISO 80601-2-61: 2011	Medical electrical equipment —Part 2-61: Particular requirements for basic safety and essential performance of pulse oximeter equipment