

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



SHANGHAI INTERNATIONAL HOLDING CORP. GMBH(EUROPE)
EIFFESTRASSE 80, 20537 HAMBURG GERMANY

SRN of AUTHORISED REPRESENTATIVE: DE-AR-000000001

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: 69450401IGN0008C2; 69450401IGN0009C4;
69450401IGN0010BM; 69450401IGN0058CH;
69450401IGN0050BZ; 69450401IGN0060C4;
69450401IGN0004BS; 6945401IGN000594;
6945401IGN00579P; 69450401IGN0003BQ;
69450401IGN0006BW;06945040101239;
06945040101246

PRODUCT AND TRADE NAME: CUFF

CATALOGUE NUMBER/MODEL: IGN0008 ; IGN0009;IGN0010 ;IGN0058;IGN0050; IGN0060;
IGN0004; IGN0005;IGN0057; IGN0003; IGN0006;IGN0001
IGN0002; IGN0066

RISK CLASS OF THE DEVICE: Class I according to rule 13 Annex VIII ,as well as its intended purpose

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.

CONFORMITY ASSESSMENT PROCEDURE: Regulation (EU) 2017/745, Annex II + III

PLACE, DATE OF ISSUE: QINHUANGDAO, 2022/10/21

HUKUN, Chairman/ manufacturer

NAME AND FUNCTION, SIGNATURE: