Declaration of Conformit

(€ 0123

EU declaration of conformity

Manufacturer information:

Name: Shanghai Caremate Medical Device Co. Ltd

Registered trade name(or registered trade mark):Caremate

Address: Building 4,No.281 HongAn Road.Zhuijng Town Jinshan

SRN: CN-MF-000037470

EU Quality Assurance Certificate(MDR) NO.:G21 067667 0015 Rev.00

Authorised representative information:

Name: MedPath GmbH SRN:DE-AR-000000087

Address: Mies-van-der-Rohe-Strasse 8, 80807 Munich, Germany

Product covered by the EU declaration of conformity:

Product name: Aneroid Sphygmomanometer

Intended use: Measure blood pressure of human being

Model: CM-1621, CM-2621, CM-2021 , CM-1329, CM-2243, CM-3003, CM-3410, CM-3001, CM-3412, CM-3411, CM-3413. CM-1013, BPM-AN-50, BPM-CH-50-HH, BPM-ST-SPL, CLOCK-Sphyg, BPM-AN-50-HH, CM-BPM, CM-PBPM

Basic UDI-DI: 697132998200701VJ

EMDN Code: C9006

MDA Code:0203

MDS Code:1010

MDT Code I:2001 MDT Code II:2011 MDT Code III:2002

Risk class: ClassI(With measuring function). According to the intended use and requirement in REGULATION (EU) 2017/745, the classification and definition are as followings: ANNEX VIII CLASSIFICATION RULES 4.1 Rule 1

All non-invasive devices are classified as class I, unless one of the rules set out hereinafter applies.

Applied Standard & Common Specification: EN ISO 81060-1.

Notified body:

Name: TÜV SÜD Product Service GmbH

Identification number: 0123

Conformity Assessment Procedure: Annex XI Part A

We herewith declare that the device is covered by the present declaration is in conformity with REGULATION (EU) 2017/745 of THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of Facil 2017 on medical devices and the

EU declaration of conformity is issued under the sole responsibility of the manufacture

All supporting documentations are retained under the premises of the manufactured

Name: Fensen Zhu

Function or Title General manager

Signature:

Place: Shanghai

Monk. Zhu

Date: 2075.11.25

Issue on behalf of Shanghai Caremate Medical Device Co. Ltd.