EU DECLARATION OF CONFORMITY

Name and address of the manufacturer:

Jiangxi Kemei Medical Apparatus & Instruments Group Co., Ltd.
No. 316, National Road, Xinju, Changshan Town, Jinxian County,
Nanchang City, 331724 Jiangxi, P.R. China

Name and address of the European Authorized Representative

CMC Medical Devices & Drugs, S.L.
C/Horacio Lengo No 18, CP 29006, Málaga, Spain

We declare under our sole responsibility that

the medical device:

Latex Examination Gloves for Single Use

Product code
Basic UDI-DI

Basic UDI-DI	Type/Model
697226728008KX	Powder-free textured XS,S,M,L,XL

Intended purpose

Latex examination gloves is intended for use in medical examinations to protect patient and user from cross-contamination.

of class:

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according to Annex VIII of Regulation (EU) 2017/745

meets the provisions of the Regulation (EU) 2017/745 which apply to it. The declaration of conformity is valid in connection with the batch-related “final inspection report” of the device.

Conformity assessment procedure:

EU Declaration of Conformity referred to in MDR Article 19

Certificate:

NA

Notified Body:

NA

Jiangxi, 2022-11-08

Place, date (地址/日期)

Che Xuemei Management representative

Name and function （姓名/职务）