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

Manufacturer: Shenzhen Urion Technology Co., Ltd.

ADD: Floor 4-6th of Building D, Jiale Science & Technology Industrial Zone, No.3, ChuangWei Road, Heshuikou Community, MaTian Street, GuangMing New District, 518106 Shenzhen, PEOPLE'S REPUBLIC OF CHINA

MEDICAL DEVICE:

Infrared thermometer

MODLE:

UFR101,UFR102,UFR103,UFR105,UFR106,UFR116,

UFR118,UFR201,UFR203,UFR205

CLASSIFICATION - ANNEX IX:

CLASS IIA, RULE10

CONFORMITY ASSESSMENT ROUTE:

ANNEX II(EXCLUDE II. 4)

WE, THE MANUFACTURER, HEREWITH DECLARE THAT THE STATED MEDICAL DEVICES
MEET THE TRANSPOSITION INTO NATIONAL LAW, THE PROVISIONS OF COUNCIL DIRECTIVE 93/42/EEC
CONCERNING MEDICAL DEVICES;

ALL SUPPORTING DOCUMENTATION IS RETAINED AT THE PREMISES OF THE MANUFACTURER. WE ARE EXCLUSIVELY RESPONSIBLE FOR THIS DOC.

NOTIFIED BODY:

TÜV SÜD PRODUCT SERVICE GMBH

RIDLERSTRABE 65, 80339 MUNICH, GERMANY

IDENTIFICATION NUMBER

0123

(EC) CERTIFICATE(S):

No. G1 078672 0014 Rev. 01

EC REP

EUROPEAN REPRESENTATIVE: Shanghai International Holding Corp. GmbH (Europe)

ADD:

Eiffestrasse 80, 20537 Hamburg, Germany

START OF CE-MARKING:

MARCH 30, 2018

PLACE, DATE OF DECLARATION:

SHENZHEN, APRIL 27, 2021

SIGNATURE:

NAME: MR. MALLE ZHU

POSITION: (GENERAL MANAGER)