DECLARATION OF CONFORMITY TO COUNCIL REGULATION (EU) 2017/745 CONCERNING MEDICAL DEVICES

MANUFACTURER: Shenzhen Creative Industry Co., Ltd.

1001, Building West, Lepu Tower, No.66 Xingke Road,

Xili Community, Xili Street, Nanshan District,

518055 Shenzhen, PEOPLE'S REPUBLIC OF CHINA

SRN OF the MANUFACTURER: CN-MF-000009430

MEDICAL DEVICE: SpO2 Probe Cable Extender

MODEL: 2909-0000004/2909-0000034/2909-0000035/2909-0000038

15040012

INTENDED USE: The Cable Extender is used to connect a compatible

patient monitor or a pulse oximeter device and a SpO2 Probe. It ensures that there is enough length for the SpO2 Probe to acquire oxygen saturation(SpO2)

and pulse rate(PR) from a patient.

CLASSIFICATION - ANNEX VIII: Class I, Rule 1

BASIC UDI-DI: 69419006CableExtender3U

CONFORMITY ASSESSMENT ROUTE: Annex II+ Annex III

We, **Shenzhen Creative Industry Co., Ltd.**, herewith declare that the stated medical devicesmeet the transposition into national law, the provisions of Council REGULATION (EU) 2017/745 concerning medical devices; All supporting documentation are retained under the premises of the manufacturer. We are exclusively responsible for the declaration of conformity.

STANDARDS APPLIED:

EN ISO 15223-1:2021	EN ISO 20417:2021	EN ISO 10993-1:2020	
EN ISO 10993-5:2009	EN ISO 10993-10:2013	ISO 10993-23:2021	

EC REP

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

Eiffestraße 80, 20537 Hamburg, Germany

SRN OF the European Representative: DE-AR-000000001

PLACE, DATE OF DECLARATION: Shenzhen, Oct. 17, 2025

SIGNATURE:

NAME: Zhang Xiang

POSITION: Management Representative