3 Declaration of Conformity

Manufacturer:

Name： Yiwu Ori-Power Medtech Co.,Ltd.

Address：#86 Xiahe Rd., Beiyuan Industrial Park 322000 Yiwu, CHINA

SRN: CN-PR-000039035

European Representative:

Name：MedPath GmbH (Europe)

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

SRN: DE-AR-000000087

Importer:XDConnectsB.V.

Address：LangeKleiweg6-28,2288GKRijswijk,TheNetherlands

Product Name： FIRST AID KIT

Model/size: V26500

Basic UDI-DI: 697468605FAK2852315H

Intended use: Provide initial, timely on-site emergency care to the injured in the event of sudden i llness or accidental injury.

Risk Class of the Device: The medical device has been assigned to class I, rule 1 according to ANNEX VIII, Medical Device REGULATION (EU) 2017/745.

GMDN code：44047

Conformity Assessment Route: Annex II, Annex III of EU 2017/745

We herewith declare that the abovementioned products meet the provisions of the following EC Council Directives and Standards. All technical documentations are retained under the promises of the manufacturer.

Yiwu Ori-Power Medtech Co.,Ltd. is exclusively responsible for the declaration of conformity.

DIRECTIVES

General applicable directives:

EN ISO 14971:2019, EN ISO15223-1:2021，EN ISO10993-1, EN ISO 10993-5, EN ISO 10993-10

Signature of issue person：

Position： General Manager **Name:** Li SongFu

Date： 2026-6-11 **Place：** YIWU