# EU DECLARATION OF CONFORMITY

According to Art. 19 of Regulation (EU) 2017/745 on Medical Devices

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|---|---|
| **Manufacturer:** | Medsource Factory, Inc.<br>#160 Jinsheng Dong Lu, Jintan District,Changzhou City,Jiangsu Province, 213200 China |
| **Trademark:** | Unitedmed 珂 迈 迪 |
| **SRN:** | Not available yet |
| **European Representative:** | MedPath GmbH<br>Mies-van-der-Rohe-Strasse 8<br>80807 Munich, Germany |
| **SRN:** | **DE-AR-000000087** |
| **Trade name:** | FISRT AID KITS |
| **Product Name:** | FISRT AID KITS |
| **Models:** | HS500BK |
| **Basic UDI：** | 69752936914016K48 |
| **Classification acc. to MDR Ax. VIII:** | Class I, rule I |
| **Applied Common Specifications / Standards:** | EN ISO 14971： 2019 |
| **Conformity assessment procedure:** | MDR Annex II + Annex III |
| **CE certificate No.:** | N.A. |
| **Name and ID of the Notified Body:** | N.A. |

We, the manufacturer, herewith declare under our sole responsibility that the above-mentioned products meet the provisions of the Regulation (EU) 2017/745 on Medical Devices (MDR). All supporting documentations are retained under the premises of the manufacturer.

Signature of issue person: Lu GuoFang

Jiangsu, June. 16,2022

Name： Lu Guofang

Position： General Manager