

Declaration of Conformity

Manufacturer: Mobility Source Medical Technology Co., Ltd.

Address: 160 Heheng Road, Yanghe Town, Gaoming District, Foshan City, Guangdong Province, 528000, China

SRN: Not assigned yet

Product name: Bed Rail

LQX-110004, LQX-110004-01, LQX-110013, LQX-110018, LQX-110027, LQX-110027-02, LQX-110033, LQX-110034, LQX-110035, LQX-110036, LQX-

Model(s): 110037, LQX-110038, LQX-110039, LQX-110040, LQX-110047, LQX-110048, LQX-110049, LQX-110050, LQX-110051, LQX-110052, LQX-110053, LQX-110054, LQX-110055, LQX-110056, LQX-110057, LQX-110058

Type: Class I, rule 1

Basic UDI-DI: 6975996301100H4

The manufacturer declares that the device complies with the Medical Device Regulation (EU) 2017/745, Standard(s) used for showing compliance with the essential requirements in the specified directive(s):

EN ISO 14971:2019, EN ISO 20417:2021, EN ISO 15223-1:2021, EN 62366-1:2015,

EN ISO 10993-1:2020, EN ISO 10993-5:2009, EN ISO 10993-10:2023

The referred report(s) show that the device complies with standard(s) recognized as giving presumption of compliance with the essential requirements in the specified EU Directive(s). The manufacturer has marked the device with the CE mark.

European Authorized Representative

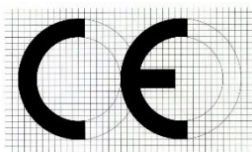
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SRN: NL-AR-000001490



This declaration is issued under the sole responsibility of the manufacturer.



Signature of Authorized Person

Gigeng

Title: Quality Manager

Place: Foshan City