



DECLARATION OF CONFORMITY

ACCORDING TO (EU) 2017/745 MEDICAL DEVICE REGULATION

EU Representative

SUNGO Europe B.V.
Olympisch Stadion 24, 1076DE
Amsterdam, Netherlands
SRN: NL-AR-000000247

Conformity Assessment

Conformity Assessment Procedure
Annex II+III of Regulation (EU) 2017/745

Applicable Standards

EN ISO 14971:2012
EN ISO 15223-1:2016
EN ISO 20417: 2021
EN ISO 10993-1:2020
EN ISO 10993-5:2009
EN ISO 10993-10:2013
EN 60601-1-2:2015+A1:2020
EN 60601-1:2006+A1:2013
EN 12182: 2012

Remark

The declaration of conformity is valid in connection
with the release technical document
CE/MDR-V080502-01.

All the supporting documentation is retained at the
premises of the manufacturer.

The Declaration of Conformity is exclusively under
the sole responsibility of the manufacturer. *SUNGO Europe office, I confirmed we are*
EU REP of the company who issue this document.



Authorized Signature (S)

Manufacturer

Name: Xiang Fa Li Technology (Xiamen) Co., Ltd
Address: 503, 5th floor, No.5 factory building, No. 63,
Pingcheng North Road, Haicang District, Xiamen city,
China

Product Information

Name: Electric Lift Transfer Chair
Model: XFL-QX-YW01

XFL-QX-YW01-1
XFL-QX-YW02
XFL-QX-YW03
XFL-QX-YW05
XFL-QX-YW06

EMDN: V080502

Basic UDI-DI: /

Classification: Class I, According to Rule 13, Annex
VIII, Regulation (EU) 2017/745

Declaration

We herewith declare that the above-mentioned
products meet the requirements of Medical Device
Regulation (EU) 2017/745 and the applicable
standards above.

Signature: *SHUI FA LI*

Date: *2021.12.21*



Position: GM