

For the following equipment:

Hospital Bed & Medical Furniture

(Product Name)

Manual bed model: 525H1/528H1/530H1/640H2 /CK830/ SS330-C1/SS001/001(A)/001(B)/SS-010

Electric bed model: 500H1/535H1/600H2/625H2-H/868H1/868H1/SS888-FM /SS888-FMK/SS888-U1/CK-835

Furniture model: SF-014/SF020/SR-010C/SR-010M/SR-010M/SR-010J/ SA-010/SA-019/

SM031/SY019/SY-017/SY-017-1/SR-018/CK-DFT-1C/CK-DFT-2C/CK-DFT-2D/CK-DFT-2E

Hereby confirmed to comply with the requirements set out in the Council Directive on the harmonization of the laws of Member States concerning <u>Medical Devices Directive (93/42/EEC)</u>.

For the evaluation regarding the safety hazard limits of Class I device, the following standards were applied:

EN 1441:1998, EN60204-1:1997

EN 60601-1:1990, +A1:1993, +A2:1995, +A11:1995, +A12:1993, A13:1996

EN 60601-1-2:1993 + EN60601-2-52:2010

Whose responsible for making this declaration:

CHEN KUANG INDUSTRIES CO., LTD.,

(Manufacturer Name)

No.11, Ln. 295, Sec.6, Hanxi W. Rd., Tanzi Dist. Taichung, Taiwan

(Manufacturer Address)

CHEN KUANG INDUSTRIES CO., LTD.

AUTHORIZED SIGNATURE

CEO Mr. Chen, Chu Chang

May 1, 2019

Position/Title

Legal Signature

Date