



EC DECLARATION OF CONFORMITY

For the following equipment:

MEDICAL CART (Model Designation)

ITEM NO: MX series

Is herewith confirmed to comply with the requirements set out in the Council Directive on the harmonization of the laws of Member States concerning Medical Devices Directive 93/42/EEC & 2007/47/EC.

For the evaluation regarding the safety limits, the following standards were applied:

ISO14971:2007 ; EN980:2003

The following representative/ importer in Europe is responsible for this declaration:

AUTO CHROBOK Brunon Chrobok

(Representative Name)

Ul. M.C. Skłodowskiej 2, 41-947 Piekary Śląskie, Poland

(Representative Address)

Person responsible for making this declaration:

Machan International Co., Ltd.

(Manufacturer Name)

No. 354, Tama Rd., Waipu, Taichung, 43856 Taiwan, R.O.C.

(Place, Manufacturer Address)

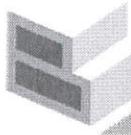
Ellen Lu

(Sales supervisor)

(Signature)

2015/03/26

(Date)



BAILIDA

Machan International Co.,Ltd

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MEDICAL / HOSPITAL EQUIPMENT



EC DECLARATION OF CONFORMITY

Medical Devices Directive 93/42/EEC

We herewith declare that medical carts are produced in our subcontractor manufacturing place according to harmonize standards and according to Medical Devices Directive Annex VII EC Declaration of Conformity

Products: MEDICAL CART, CABINETS

Model: Bailida products

Classification: Class I (93/42/EEC Annex VII)

Directive : Medical Devices Directive - 93/42/EEC

Conformity Assessment Procedure : Annex VII EC Declaration of Conformity

Manufacture: Machan International Co., Ltd.

No. 354, Dama Rd., Waipu, Taichung, 43856 Taiwan, R.O.C.

Yours faithfully,



Signature of manufacturer/supplier

Date: 9 April, 2018