CE Technical Documentation Review Report

Applicant:	**Foshan Dongfang Medical Equipment Manufactory (Ltd.)** No.1 Pulan Road, Foshan City, Guangdong Province 528000,P.R.China
Order Number:	63010931
Examination intent:	Examination the completeness of the Technical Documentation according to the requirements of the Medical Devices Directive 93/42/EEC Annex VII
Product(s):	Mechanical Wheelchairs
Type(s)/Model(s):	FS-series
Classification:	Class I (according to manufacturer's declaration)
Examination period:	2004 Sep. 01-02
Review result:	During the examination of the provided Technical Documentation (dated 2004-08-15), no Non-compliance according to the requirements of the Medical Devices Directive 93/42/EEC Annex VII was detected.

TÜV Rheinland (Shanghai) Co., Ltd.

Xiaoli REN
Manager (Asia Region)
Medical Services

Product Safety · Quality Assurance
TÜV Rheinland
Approved

TÜV Rheinland (Shanghai) Co., Ltd.
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EC-Declaration of Conformity

According to MDD 93/42/EEC

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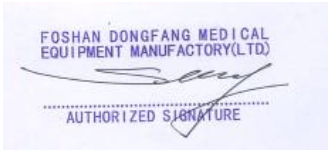
Declare under our sole responsibility that the products:
Wheelchair
Power wheelchair
Shower chair
Walker

CLASSIFICATION: CLASS I MEDICAL DEVICE

are produced in conformity with the technical documentation of Annex VII clause 3 and meets the provisions of the Medical Device Directive 93/42/EEC which apply to them.
This declaration is valid for the above mentioned product with the CE-Mark.



FOSHAN October 11, 2019
(place and date of issue)

(name and signature or equivalent marking of authorized person)

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