

CE Technical Documentation Review Report

Manufacturer: **JERRY MEDICAL INSTRUMENT (SHANGHAI) CO., LTD.**
Building 12, No.615 Fengdeng Rd., Malu Town,
Jiading District, Shanghai, China

Report Number: 15081428 001

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): Power Wheelchair

Type(s)/Model(s): JRWD501, JRWD502, JRWD503, JRWD601, JRWD301, JRWD602, JRWD603 , JRWD1801, JRWD1002, JRWD1803

Classification: Class I, rule 12
(according to manufacturer's declaration)

Review result: During the examination of the provided Technical Documentation (No.: MDDJR2015CE, Version A/0, Date: 2015.06.24), no Non-compliance according to the requirements of the Medical Devices Directive 93/42/EEC Annex VII was detected.

Shanghai, 2016-12-30

TÜV Rheinland (Shanghai) Co., Ltd.



Allen CHEN
Lead Auditor, Product Assessor
Medical Device Services

