



Annex to Report (No. 01992)

Foshan Shenghai Medical Equipment Co., Ltd.

Product Name	Type
Manual Hospital Bed	SH-B001, SH-B001S, SH-B002, SH-B003, SH-B004, SH-B101, SH-B102, SH-B103, SH-B104, SH-B105, SH-B106, SH-B201, SH-B202, SH-B203, SH-B204, SH-B205, SH-B206, SH-B207, SH-B208, SH-B209, SH-B251, SH-B251S, SH-B301, SH-B303, SH-B304, SH-B501, SH-B502, SH-B503, SH-B504, SH-B505, SH-B551, SH-B552, SH-B553, SH-B554, SH-B555, SH-B557, SH-B558, SH-B601, SH-B602, SH-B603, SH-B604
Electric Hospital Bed	SH-B421, SH-B422, SH-B423, SH-B424, SH-B431, SH-B434, SH-B436, SH-B437, SH-B451, SH-B452, SH-B453, SH-B454
Medical Cabinet	SH-C001, SH-C002, SH-C101, SH-C102, SH-C103, SH-C201, SH-C202, SH-C203, SH-C204, SH-C205, SH-C305, SH-C306
Medical Trolley	SH-T001, SH-T002, SH-T003, SH-T004, SH-T005, SH-T007, SH-T008, SH-T009, SH-T010, SH-T011, SH-T012, SH-T013, SH-T015, SH-T016, SH-T018, SH-T019, SH-T101, SH-T149, SH-T150, SH-T151, SH-T152, SH-T152A, SH-T153, SH-T154, SH-T155, SH-T156, SH-T157, SH-T158, SH-T159, SH-T160, SH-T170, SH-T175, SH-T176, SH-T177, SH-T178, SH-T179, SH-T80, SH-T181, SH-T182
Medical Chair	SH-W101, SH-W102, SH-W103, SH-W104, SH-W301, SH-W302A, SH-W302B, SH-W201, SH-W202, SH-W203, SH-W204, SH-W205, SH-W206, SH-W207, SH-W208, SH-W209, SH-W210, SH-W210S, SH-W212, SH-W331, SH-W332, SH-W333

This annex is only valid if attached to the report mentioned above.



This is to certify that the Quality Management System of

Foshan Shenghai Medical Equipment Co., Ltd.

No.5-2, Row 3, Jiangbian Xiyue Industrial Area, Chancheng District, Foshan City,
Guangdong Province, P.R.China

applicable to

**The Production and Sales of Manual Bed, Stretcher, Cabinet, Trolley,
and Medical Chair(Only for Export)**

has been assessed and registered by NQA against the provisions of
ISO 13485: 2003

This registration is subject to the company maintaining a quality management
system, to the above standard, which will be monitored by NQA.
Please consult the website: www.snqa.com.cn

Adam Weir.

Certification Director



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Certificate Number **28887**

Date: 08 July 2013

Valid Until: 08 July 2016

EAC Code: 19



Compliance Report

Applicant: Foshan Shenghai Medical Equipment Co., Ltd.
Address: No. 5-2, Row 3, Xiyue Industrial Area, Jiangbian Village,
Chancheng District, Foshan City, Guangdong Province, China

Product: Manual Wheelchair, Emergency Stretcher

Type: See annex for details

Product Classification: Class I

The submitted technical files including test report of the above products have been reviewed against the self declaration requirements of conformity for CE marking according to Annex I & VII of the 93/42/EEC Medical Device Directive (including 2007/47/EC).

The review result of the technical files and test report support the self declaration for the devices listed above. The test report and the technical files are the annex of this report and should be used together.

Where the manufacturer affix's the CE marking to the product listed they must ensure that all the requirements of the appropriate EU directive(s) have and continue to be met.

This report is not a certificate of conformity.

No. 01990

Initial Issue Date: 13 Dec 2012

A handwritten signature in blue ink, appearing to be 'H. Santos', is written over a horizontal line.

General Manager (Signature)

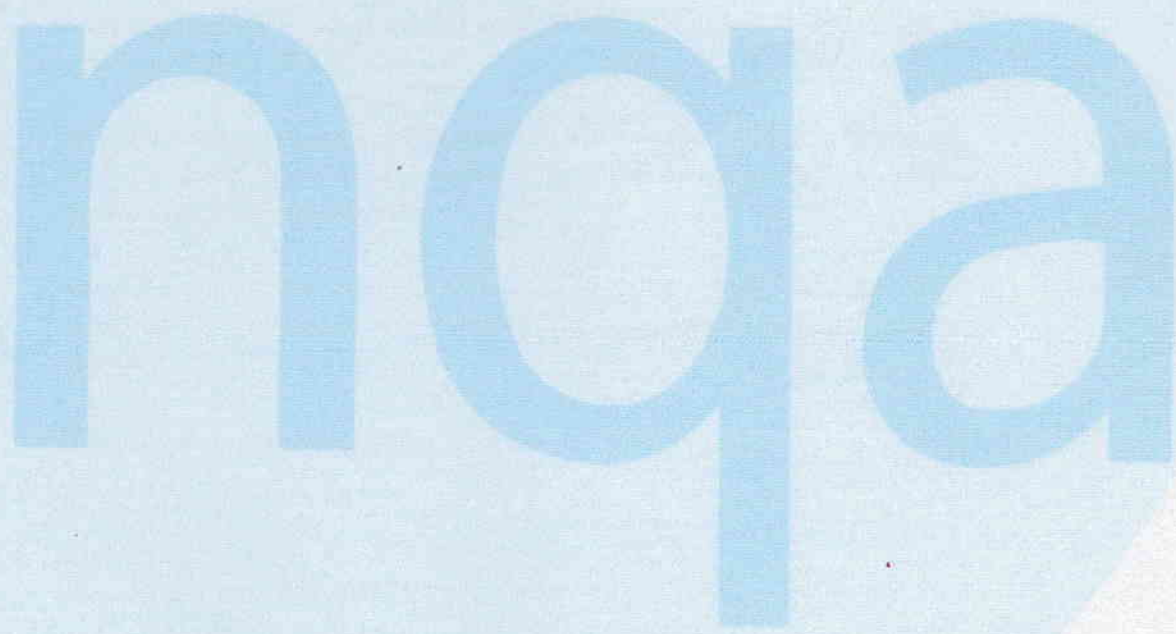


Annex to Report (No. 01990)

Foshan Shenghai Medical Equipment Co., Ltd.

Product Name	Type
Manual Wheelchair	SH-W001, SH-W004, SH-W005, SH-W006, SH-W007, SH-W221
Emergency Stretcher	SH-S001, SH-S002, SH-S003, SH-S004, SH-S005, SH-S006, SH-S007, SH-S008, SH-S009, SH-S010, SH-S011

This annex is only valid if attached to the report mentioned above.





Compliance Report

Applicant: Foshan Shenghai Medical Equipment Co., Ltd.
Address: No.5-2, Row 3, Xiyue Industrial Area, Jiangbian Village,
Chancheng District, Foshan City, Guangdong Province, China

Product: Manual Hospital Bed, Electric Hospital Bed, Medical Cabinet,
Medical Trolley, Medical Chair

Type: See annex for details

Product Classification: Class I

The submitted technical files including test report of the above products have been reviewed against the self declaration requirements of conformity for CE marking according to Annex I & VII of the 93/42/EEC Medical Device Directive (including 2007/47/EC).

The review result of the technical files and test report support the self declaration for the devices listed above. The test report and the technical files are the annex of this report and should be used together.

Where the manufacturer affix's the CE marking to the product listed they must ensure that all the requirements of the appropriate EU directive(s) have and continue to be met.

This report is not a certificate of conformity.

No. 01992
Initial Issue Date: 13 Dec 2012

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General Manager (Signature)