



Certificate of Compliance

We hereby declare that the technical file of product class 1 complied with the requirement of directives Medical Council Directive 93/42/EEC of June 1993

Manufacturer

Name : MEDITECH (INDIA)

Address : # 487/67, GLASS STREET, NATIONAL MARKET, PEERAGARHI,
NEW DELHI-110087, INDIA

Product : PRODUCT AS PER ANNEXURE

The Certification body has performed an audit of the above product quality system covering the design, manufacture and final inspection of the certified product. The quality system has been assessed, approved and is subject to continuous surveillance according to the directive 2004/108/EC Electromagnetic Compatibility Directive

This certificate is issued under the following conditions:

1. It applies only to the quality system maintained in the manufacture of above referenced models and it does not substitute the design or type-examination procedures, if requested.
2. The certificate remains valid until the manufacturing conditions or the quality systems are not changed.
3. The certificate validity is conditioned by positive results or surveillance audits.
4. After fulfilling the relevant EU legislation, the manufacturer shall affix to each device, of the referenced models.
5. The CE mark as shown Below can be used, under the responsibility of the manufacturer, after completion of an EC Declaration of conformity and compliance with all relevant EC Directives. The statement is based on a single evaluation of one sample of above mentioned product. It does not imply an assessment of the whole production

Date of Registration

06 January 2021

1st Surveillance Due

05 January 2022

2nd Surveillance Due

05 January 2023

Recertification Due (subject to the company maintaining its system to the required standard)

05 January 2024

Certificate No.: 301060121

Validity of this certificate can be verified at www.otabucert.co.uk



Authorised Signatory

Otabu Certification Limited (UK)

Validity of this Certificate is subject to Annual Surveillance audits done successfully

This Certificate of Registration remains the Property of Otabu Certification Limited and shall be returned immediately upon request.

Email:- info@otabucert.co.uk Website:- www.otabucert.co.uk

Suite 48, 88-90 Hatton Garden, London, EC1N 8PN, UK



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Appendix Certificate No. 301060121

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This certificate referred to above covers the following products:

Product	Model
Operation Theatre Table	Astra 103+, Astra 101+, Celebrita, MI 103, MI 102, MI 101, MI 111, MI 2003
Neuro Attachment	Three Head Pin Mayfield Skull Clamp, Horse Shoe Shape Attachment
Orthopaedic Attachment	Hanging & Floor
Operation Theatre Lights	Medlight 50+50, Medlight 40+40, Medlight 50+40, Medlight 50+40+40, Medlight 50, Medlight 40, XN 243, XN 233, XN 104, XN 103, XN 4, XN 3, Matrix 2, Matrix 1, Matrix M, Z 500+500, Z 400+400, Z 300+300, Z 500+400, Z 500+300, Z 400+300, Z 500, Z 400, Z 300, Z 500M, Z 400M, Z 300M,
Hospital Bed & Furniture	
Examination Lamps	EL 570, EL 580, EL 590



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