Certificate of Compliance

This Certificate is awarded to the Quality Management System for Medical Devices of

MAHIKA MEDICAL PVT. LTD.

E-117, PREET VIHAR, DELHI – 110 092, INDIA

has been audited by the sgi management and found to be in compliance with the requirements of the standard

ISO 13485:2012

This Certificate is applicable to the following scope

"MANUFACTURER & SUPPLIER OF HOSPITAL FURNITURE, HOSPITAL HOLLOWWARE, ORTHOPAEDIC REHABILITATION & FRACTURE AIDS, LABORATORY EQUIPMENTS, AUTOCLAVES & STERILIZERS, SUCTION MACHINES, RESUSCITATORS, ANAESTHESIA EQUIPMENTS & ACCESSORIES, DIAGNOSTIC EQUIPMENTS & PRODUCTS"

Certificate No : 10Q/MMD/070013

Issue Date	**13 July 2016**	**Valid Until:**	**12 July 2021**
1st Surveillance on or before	**12 June 2017**	**Recertification Date**	**12 July 2021**
2nd Surveillance on or before	**12 June 2018**		

To Check Validity of this certificate please Visit www.sgicert.org

Digitally signed by Iarovoi Petru
Date: 2021.03.18 16:37:25 EET
Reason: MoldSign Signature
Location: Moldova

Auth. Sign

HO: E-48 , GROUND FLOOR , NEW MULTAN NAGAR , NEW DELHI-110056 , INDIA

Email:- info@sgicert.org , cert.sgi@gmail.com , Website :- www.sgicert.org

Digitally signed by Iarovoi Petru
Date: 2019.02.06 16:46:33 EET
Reason: MoldSign Signature
Location: Moldova

US Certification Services

Certificate of Conformity

We confirm that the technical documentation for the below mentioned product (Medical Devices of the Class I) according to the Council Directive 93/42/EEC as amended by 2007/47/EC

Hospital Furniture, Examination/ OPD Furniture, Gynaecology Equipments, Operation Theatre Equipments, Hospital Hollowware, O. T. Lights & Examinations Lights, Patient Transport System, Laboratory Equipments, Mortuary Equipments, Paediatric Ward Equipments, Hospital Linen, Hospital Equipments & Orthopaedic Rehabilitation and Fracture Aids

(For Detailed Specifications, Refer to Annexure of this Certificate, Pages 12)

Manufactured by Company

MAHIKA MEDICAL PVT. LTD.

E - 117, PREET VIHAR, DELHI – 110 092, INDIA

complies with the applicable requirements of the Council Directive 93/42/EEC as amended by 2007/47/EC.

Referring to the intended use, the Certification Body has conducted with successful results the review of the Manufacturer's Technical documentation of the certified product according to above mentioned Directive and appropriate Harmonized European Standards.

This Certificate is issued under the following conditions:

1. *The Manufacturer's Technical Documentation, as required for Class I Devices, has been reviewed and found to comply with the requirements in Annex VII, Section 3.*
2. *It applies only to the above mentioned Medical Devices.*
3. *The Manufacturer is obligated to assure conformity of all the Medical Devices of the respective model to the type assessed by the mean of this Certificate.*
4. *Any significant changes in the Design or Process used to manufacture the Product, or revision to the Directives or standards referred above may require special audit by U. S. Certification Services. The product liability rests with the Manufacturer or his Representative in accordance with Council Directive.*
5. *After fulfilling the relevant EU Legislation Requirements, the Manufacturer shall affix to each Medical Device, CE Marking according to the following example:*

CE

U.S. CERTIFICATION SERVICES

Certificate No. : ***USC/MDD/2016/124***
Issued On : ***July 30, 2016***
Valid Up To : ***July 29, 2021***

Authorized Signature

US Certification Services

The Certificate remains the property of US Certification Services to whom it must be returned on request.
Deliberate misuse of certificate will results in cancellation without notification. To check current validity of certificate, send mail at
URL: *www.uscertification.co.in* ***E-mail:*** *uscertificationservices@gmail.com*