

# Certificate of Compliance



We hereby declare that the technical files of all the items in each product group of complied with the requirements of the Medical Device Directive EU Directive 93/42/EEC

**Certificate No.: CE-1211**

## Manufacturer

Name : MOKSHIT CORPORATION

Address : **Registered Office: Ganjpara Durg (C.G) 491001, India**  
Works : HSIIDC IGC, First Floor, Saha Ambala –  
Haryana -133104, India

Products : **Design, Supply, Installation, Fabrication, Operation and Maintenance of Medical Gas Pipe Lines Systems, Modular and prefabricated OTS , ICU, CCU, ICCU, SICU, Oxygen Generator / concentrator plants, Air Ceiling Management system, Nurse Call System, Wall Guard and Corner Guard System, Surgical and Medical Equipment and Instruments, CSSD, Cubicle Partition Track System Bed Head Panels OT Lights, OT tables, Clean Room Luminaries / Peripheral Lights , Hospital beds and Hospital Furniture, Critical Care Ambulance / mobile Hospitals on Complete Turnkey Project**

## Complies with the requirements applicable to it

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 Medical Device Directive EU Directive 93/42/EEC

## 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 of surveillance audits.  
The CE mark as shown above 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.

**Validity of this certificate can be verified at [www.ukcertifications.co.uk/verify](http://www.ukcertifications.co.uk/verify)**

Date of Certification	08 <sup>th</sup> December 2018
1 <sup>st</sup> Surveillance Audit Due	07 <sup>th</sup> December 2019
2 <sup>nd</sup> Surveillance Audit Due	07 <sup>th</sup> December 2020
Certificate Expiry (subject to the company maintaining its system to the required standard)	07 <sup>th</sup> December 2021

*Daniel..*

Authorised Signatory

