CERTIFICATE

This is to certify that

CE

We hereby declare that the technical file of product complied with the requirement of directives Council Directive on Medical Devices 93/42/EEC as Amended 2007/47/EC.

Certificate Number: 2022071631

Manufacturer

Name : **NARANG HOSPIMEDEQP EXPORTS**

Address : **7/33, ANSARI ROAD, DARYA GANJ, NEW DELHI – 110002, INDIA**

Products : **Manufactures, Contractors & Exporters of Hospital Furniture, (Beds, Trolleys and SS Ward Items), Hospital Holloware, Waste Management, Operation Theatre Furniture, Operation Theatre Tables, Operation Theatre Lights, Delivery Tables, Patient Handling Equipment's for Ambulance & Hospital, Doctors & Surgeon Room Equipment's, Surgical Instruments & Equipment's Obstetric Instruments, Gynaecological Equipment's & Instruments Arthroscopy Set, Laparotomy Set, General Surgery Instruments & Sets, Physiotherapy & Rehabilitation Equipment's, Walking Aids, Wheel Chair, Dental Equipment's & Instruments, Plastic Lab Ware, Ophthalmology & ENT Equipment's, Surgical Rubber Goods, Home Health Care Devices, Diagnostics Devices, Medical Eqp. & Anaesthesia Products, Laboratory Equipments, Scientific Equipments, Autoclaves & Steam Sterilizers {Portable, Horizontal, Rectangular & Vertical}, Heating Equipments, Cooling Equipments, Clean Air Equipments, Blood Bank Equipments, Mortuary Equipments, R & D Lab Equipments, Microbiology Lab Equipments, Pharmaceutical Laboratory Equipments, Laboratory Research Microscopes, Testing Instruments, Poultry Appliances, Analytical Equipments, Scrub Stations, Grossing Tables, Kitchen Equipments (Commercial) (more detail as per appendix-I)**

Complies with the requirements applicable to it

The quality system file has been assessed, approved and is subject to continuous surveillance according to the Council Directive on Medical Devices 93/42/EEC as Amended 2007/47/EC.

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 changed.
3. The certificate validity is conditioned by positive results or 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.

Date of Certification: 16th July 2022
1st Surveillance Audit Due: 15th July 2023
2nd Surveillance Audit Due: 15th July 2024
Certificate Expiry: 15th July 2025

Head of Certification

UKAF
UKAF-CB-009

Management System Certification
DBS CERT

DBS CERTIFICATIONS PVT. LTD.
142, IInd Floor, Avtar Enclave, Paschim Vihar, Delhi-110063, (INDIA) info@dbscertification.com, www.dbscertification.com

ACCREDITED BY :
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