Certificate of Compliance

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

Manufacturer

Name : **YASHIKA ENTERPIRSES**

Address : **PLOT NO. 49, SHOP NO. 1, JAY NAGAR, KESAR CIRCLE, ISKON ROAD, MANSAROVAR, JAIPUR- 302020, RAJASTHAN, INDIA**

Product : **O.T LIGHT, O.T. TABLE, PATHOLOGY MICROSCOPE, RESEARCH MICROSCOPE, DENTAL MICROSCOPE, INDUSTRIAL MICROSCOPE, MULTI PARA MONITOR, DEFIBRILLATOR, SYRINGE INFUSION PUMP, HORIZONTAL AUTOCLAVE,VERTICAL AUTOCLAVE, DENTAL AUTOCLAVE, HOSPITAL FURNITURE, SINGLE BLOOD BAG, DOUBLE BLOOD BAG, TRIPLE BLOOD BAG, QUADRUPLE BLOOD BAG, ANESTHETIC AND RESPIRATORY EQUIPMENT – TRACHEOSTOMY TUBES & CONNECTORS, MEDICAL DISPOSABLE DEVICES, CAUTERY MACHINE, RAPID CARD, BLOOD COLLECTION TUBES, GLOVES, SURGERY INSTRUMENTS, LABORATORY PRODUCTS**

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 Medical Council Directive 93/42/EEC of June 1993.

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.
4. After fulfilling the relevant EU legislation, the manufacturer shall affix to each device, of the above referenced models.
5. 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 Registration	: 03 February 2020
1st Surveillance Due	: 02 February 2021
2nd Surveillance Due	: 02 February 2022
Certificate Expiry (subject to the company maintaining its system to the required standard)	: 02 February 2023

Certificate No:- 105030220

To Verify this certificate please visit at www.qvcert.co.uk

CE

Authorised Signatory

Quality Veritas Certification Limited

Validity of this certificate is subject to annual surveillance audits done successfully

This Certificate Of Registration Remains The Property of Quality Veritas Certification Limited and Shall be Returned Immediately Upon Request

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

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