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CERTIFICATE



We hereby declare that the technical file of product complied with the requirement of Protective Equipment 89/686/EEC as Amended Regulation (EU) 2016/425 & Council Directive on Medical Devices 93/42/EEC as Amended 2007/47/EC.

UNITY HEALTHCARE

OFFICE: C-24, PREET VIHAR, DELHI 110092

Product: Medical Nitrile Gloves, Examination Latex Gloves, Pulse Oxymeters, Sterile Surgical Medical Masks, Medical Protective Coveralls, Sterile Medical Shoe Cover, Surgical Towels, Isolation Gowns

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 Protective Equipment 89/686/EEC as Amended Regulation (EU) 2016/425 & 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.

CertificateDate ofFirst SurveillanceSecond SurveillanceDate ofNumber:Certification:Audit Due:Audit Due:Expiry:P01M02H07116004-May-202103-May-202203-May-202303-May-2024









#CAB 130118

Lack of fulfillment of conditions as set out in Certification T&C may render this certificate invalid. Validity of this certificate is subject to annual surveillance audit. Verification of this certificate can be verified at TIQ Certification and Inspection Limited (TIQ) website at www.tiqcertification.org/verify or by scanning the QR code to the right. This certificate remains the property of TIQ and shall be returned immediately on request.

TIQ is accredited by UKCB (United Kingdom Certification Board), 27 Old Gloucester Street, London, United Kingdom. www.ukcb.org.uk WC1N 3AX (#CAB 130118). Website: www.tiqcertification.org | Email: info@tiqcertification.org | India Office: 117, Antriksh Bhawan, Barakhamba Road, New Delhi-110001

