Certificate of Compliance

We Hereby Declare That the Technical File of Product Complied with The Requirement of Machine Directive 2006/42/CE

Certificate No.: CE-97801

COMPANY NAME :- **GEL CRAFT HEALTHCARE PVT.LTD**

REGD. OFFICE :- **A2/48, SITE-IV, SAHIBABAD INDUSTRIAL AREA GHAZIABAD, UTTAR PRADESH – 201010**

PRODUCTS :- **C3I GEL® (MEDIGEL®) TECHNOLOGY BASED FOOTCARE PRODUCTS, HEALTHCARE PRODUCTS, CUSHIONING / BED SORE PREVENTION PRODUCTS, OPERATION THEATRE POSITIONING DEVICES, FOOTWEAR AND SPORTS PRODUCTS, DRDO TECHNOLOGY VISCOELASTIC GEL WITH ANTI-MICROBIAL COVID COAT , DRDO CBRN ULTRA DEVICES, ULTRA SWACHH PPE DISINFECTION UNIT, ATI SWACHH – HEAT SENSITIVE MEDICAL DEVICE DISINFECTION UNIT, OZONATED RADICAL CONFINED SPACE DISINFECTION UNIT(POORAN SWACHH), TRI-NETRA HAND SANITIZATION UNIT, TAARAN PATIENT TRANSFER SYSTEM, SAMGRAH SWACHH, VAYU SWACHH, VISANKRA OT STERILIZATION UNIT, TRIYOGANI HAND SANITIZER, TRIYOGANI FUMIGATION ,COVID COAT PLUS , ULTRA SWACHH – PERSINAL PROTECTIVE EQUIPMENT DISINFECTION UNIT , TRI-NETRA HAND SANITIZATION UNIT, TAARAN (SAFE PASSAGE) PATIENT TRANSFER SYSTEM , ORCS SANITIZATION UNIT (POORN SWACHH) , ATI SWACHH HEAT SENSITIVE MEDICAL DEVICE DISINFECTION UNIT , TRIYOGANI HAND SANITIZATION SOLUTION , TRIYOGANI FUMIGATION SOLUTION , COVID COAT 90 DAYS DISINFECTION SOLUTION , SAMAGRAH SWACHH FRUIT & VEGETABLE WASH & OZONATED SPACE STERILIZERS**

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 Machine Directive 2006/42/CE

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 with applicable CE Requirement 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 and CE Requirement, the manufacturer shall affix to each device, of the referenced models.**
5. **The CE Certificate 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 with EN standard requirement. 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.ukglobals.uk/Verify

Date of Certification	09 April. 2021
1 st Surveillance Audit	08 April. 2022
2 nd Surveillance Audit	08 April. 2023
Certificate Expiry (subject to the company maintaining its system to the required standard)	08 April. 2024

Authorised Signatory

CE

This certificate is the property of UK Global Certification & Inspection Limited and shall be returned immediately on request.
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