



# Certificate of Compliance

RoHS Directive (2011/65/EU) of the European Parliament and of the Council on the restriction of use of certain Hazardous Substances in Electrical and Electronic Equipments

Certificate No. ROHS-MHLS-21-222615

## Manufacture

Name : **MODULAR HEALTHCARE SYSTEM**

Address : REGD. ADDRESS: 202, GULMOHAR ESTATE, POCKET- E, SECTOR- PI-1,  
GREATER NOIDA, GAUTAM BUDHH NAGAR- 201310, INDIA  
FACTORY ADDRESS: PLOT NO.37, UDYOG KENDRA - I ECOTECH - III, GREATER  
NOIDA, GAUTAM BUDDHA NAGAR-201308, UTTAR PRADESH, INDIA

Product : "MANUFACTURER & SERVICE, DESIGN, SUPPLY, INSTALLATION, TESTING AND COMMISSIONING FOR MODULAR OPERATION THEATRE, ICU, NICU, IVF ROOMS, CLEAN ROOM, MGPS, BED HEAD PANEL, OT CONTROL PANEL, SURGICAL SCRUB STATION, HOSPITAL FURNITURE, HVAC & MEDICAL EQUIPMENT'S , HERMETICALLY SEALED DOORS,SURGEON PENDENT, ICU TRACK & CURTAIN, AIR HANDLING UNIT, TOTAL HOSPITAL SOLUTIONS"

This is to state that the above mentioned products is in compliance with RoHS Directive (20/95/EC) of the European Parliament and Commission Decision 2005/618/EC on the restriction of use of certain Hazardous Substances [Lead (Pb), Mercury (Hg), Cadmium (Cd), Hexavalent Chromium (Cr6+), Polybrominated biphenyls (PBBs) and Polybrominated Diphenyl ethers (PBDEs)] in Electrical and Electronic Equipments.

### Statement:

This certificate declares that the product type/model described above complies with the mentioned above European Standard(s).

### Remarks:

This certificate of complies is based on the evaluation of a sample of the above mentioned products. It does not imply and assessment of the mass-production of the product. This certificate holder may use this certificate in connection with the test certification body should be informed (revision of technical file)for any modification or alterations made to the aforementioned product type(s), including design and manufacture and/or extension to the existing scope of application.

The certificate is valid for three years if the company applies the technical construction file which has been stored IMC office. This certificate includes declaration of manufacturer.

Certificate remains property of IMC to whom it must be returned upon request. The certificate validity is conditioned by positive results or surveillance audits.

**Validity of this certificate can be verified at [www.gaafs.us](http://www.gaafs.us)**

**Date of Certification**

**1<sup>st</sup> Surveillance Due**

**2<sup>nd</sup> Surveillance Due**

**Certificate Expiry** (Subject to the company maintaining its system  
To the required standard)

**22<sup>ND</sup> JULY 2021**

**21<sup>ST</sup> JULY 2022**

**21<sup>ST</sup> JULY 2023**

**21<sup>ST</sup> JULY 2024**

Registered

Authorized Signatory



## QVA Certification

CAB Address : Maryland Avenue, SW Washington, D.C. 20202

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

This certificate is the property of QVA Certification and shall be return immediately on request

QVA Certification is and independent Systems Products and Personal assessment Body, QVA Certification is a accredited by GAAFS.US