Certificate of Compliance

We hereby declare that the technical file of product complied with the requirement of directives Low voltage directive (LVD) (2014/35/EU) & Medical directive 2006/42/EC

Applicants

Name : MONARCPLUS MANUFACTURERS PRIVATE LIMITED

Address : SHOP NO S-1, P NO-123, SHYAM MITRA MANDAL NAGAR, ROAD NO 5 VKI AREA, MURLIPURA, JAIPUR, RAJASTHAN -302039, INDIA.

Product : MANUFACTURERS, SUPPLIER & IMPORT-EXPORT BUSINESS VARIOUS TYPE OF MEDICAL, HOSPITAL, OFFICE & HOME FURNITURE, VARIOUS TYPE OF MEDICAL EQUIPMENT AND DEVICES, SURGICAL & DISPOSABLE MEDICAL PRODUCTS, SURGICAL INSTRUMENTS, ORTHOPEDIC AND DENTAL IMPLANTS, ICU, OT & LAB EQUIPMENTS, C ARM MACHINE, X RAY MACHINE & FILM, CT SCAN, MRI MACHINES, PHARMACEUTICAL PRODUCTS, COSMETIC AND WELLNESS PRODUCTS, FOOD SUPPLEMENTS, HOSPITAL SERVICES.

Complies with the requirements applicable to it

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 the Low voltage directive (LVD) (2014/35/EU) & Medical directive 2006/42/EC

This certificate is issued under the following conditions:

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.

1. The certificate remains valid until the manufacturing conditions or the quality systems are changed.
2. The certificate validity is conditioned by positive results or surveillance audits.
3. After fulfilling the relevant EU legislation, the manufacturer shall affix to each product, of the above referenced models.
4. 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.

Certificate No. QVA-MSPL-23-0128905

Certificate can be verified at www.gaafs.us

Date of Certification	**01ST DEC 2023**
1st Surveillance Due	**30TH NOV 2024**
2nd Surveillance Due	**30TH NOV 2025**
Certificate Expiry (Subject to the company maintaining its system To the required standard)	**30TH NOV 2026**

Registered

Authorized Signatory

CERTIFIED QVA Certification

CE

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 returned immediately on request
QVA Certification is an independent Systems Products and Personal assessment Body, QVA Certification is a accredited by GAAFS.US