



Certificate of Compliance

We hereby declare that the technical file of product complied with the requirement of **Medical Device Directives (93/42/EEC)**

Manufacturer:

Name: BHAVNA FURNITURE

Address: REGD. OFFICE:-7, SIDDHIRAJ COMPLEX, BOPAL GHUMA ROAD,

BOPAL, AHMEDABAD, GUJARAT - 380058, INDIA

FACTORY ADDRESS:- 9 UMIYA INDUSTRIAL ESTATE, KHATWADA SINGARVA ROAD, OPP BAVCHAR PARTY PLOT, KHATWADA, AHMEDABAD - 382430, INDIA

Product: Laboratory furniture, Laboratory Island table, Pass box, fume hood, work bench, reagent rack, anti vibration table, library furniture, hostel furniture, hotel furniture, corporate interior designing, work station, cubical, wooden furniture, steel furniture. All type of revolving chair, visitor chair, fixed chair, sofas, college and school benches and

VISITOR CHAIR, FIXED CHAIR, SOFAS, COLLEGE AND SCHOOL BENCHES AND LIBRARY FURNITURE.

The certification body has performed an audit of the above product quality system covering the design, manufacture and final Inspection of certified product. The quality system has been assessed, approved and is subject to continuous surveillance according Medical Device Directives (93/42/EEC)

This certificate issued under the following conditions:

- 1. It applies only to the quality system maintained in the manufacturing 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 system are changed
- 3. The certificate validity is conditioned by positive results of surveillance audits After fulfilling relevant EEC legislation, the manufacturer shall affix to each device, of the referenced models The CE mark as shown above can be used under the responsibility of the manufacturer after completion of an EEC Declaration of conformity and compliance with all relevant EEC Directives. The statement is based on a single evaluation of one sample of above mentioned product. Not imply an assessment of the whole production.

:: Certificate No :: CE/23M02466

Date of initial registration: 24 August 2023

First Surveillance Audit on or before: 23 August 2024

Second Surveillance Audit on or before: 23 August 2025

Re-certification Due: 23 August 2026

This Certificate is property of MQA and remains valid Subject to satisfactory surveillance audits.

Authorized Signatory
MQA CERTIFICATION SERVICES

130 Thessaly Rd, Nine Elms, London SW8 5EJ, United Kingdom





CE

UKAF-CB-011

To check validity of the certificate please visit at www.mqacertification.com

This certification of registration is issued by MQA Certification Services accredited with UKAF CERT LIMITED Accreditation Board for Certification Bodies (www.ukafcert.org.uk). This certificate remains the property of MQA Certification Services having and must be returned upon request.