



Certificate of Compliance

We hereby declare that the product complied with the requirement of
Medical Device Directive 93/42/EEC as amended (2007/47/EU)

Manufacturer

Name : A ONE HOSPICARE

Address : 353/1/1, PATWARI HALKA NO. 52, PALDA, INDORE - 452001,
MADHYA PRADESH, INDIA

Products : HOSPITAL AND MEDICAL EQUIPMENTS

The Certification body has performed an audit of the above product covering the design, manufacture and final inspection of the certified product. The technical documentation / inspection / test results comply with the requirements of Medical Device Directive 93/42/EEC as amended (2007/47/EU)

Hence the manufacturer has issued a Declaration of Conformity according and places the CE marking with his own responsibility.

Certificate No : 25/CE/AON09257
Initial Registration Date : 25/09/2025

Issuance Date : 25/09/2025
Date of Expiry* : 24/09/2028

Auth. Signatory



Times Certification Services UK Ltd.

Regd Office : 71-75 Shelton Street, Covent Garden, London, WC2H 9JQ, United Kingdom

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Validity of this certificate can be verified at : www.timescert.co.uk or via email at info@timescert.co.uk





Certificate of Compliance

This is to Certify that

A ONE HOSPICARE

**353/1/1, PATWARI HALKA NO. 52, PALDA, INDORE - 452001,
MADHYA PRADESH, INDIA**

FOR THE PRODUCT :

**MANUFACTURING OF HOSPITAL FURNITURE AND
TRADING OF MEDICAL EQUIPMENTS**

**RoHS Directive(2024/232/EU)
(Restriction of use of certain Hazardous Substance)**

RoHS Directive (2011/65/EU) as Amended (EU) 2024/232

This is to state that the above mentioned product is in compliance with RoHS Directive(2015/863/EU) of the European Parliament and Commission Decision 2005/618/EC on the restriction of use of certain Hazardous Substance [Led (Pb), Mercury (Hg), Cadmium (Cd), Hexavalent Chromium (Cr6+), Polybrominated Biphenyls (PBBs) and Polybrominated Diphenyl, Bis (2-Ethylhexyl) phthalate (DEHP), Benzyl butyl phthalate (BBP), Dibutylphthalate (DBP), Diisobutyl phthalate (DIBP)] ethers (PBDEs) in Electrical and Electrical Equipments.

Statement: This Certificate Declares That the Product type/model described above complies with the mentioned above European Standards(s).

Remarks: This certificate of complies is based on the evaluation of a sample of the above 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 Report. The 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

Certificate No : 25/R/AON09255

Initial Registration Date : 25/09/2025

Issuance Date : 25/09/2025

Date of Expiry : 24/09/2028

Auth. Signatory



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Verification of Compliance of BIFMA

A ONE HOSPICARE

353/1/1, PATWARI HALKA NO. 52, PALDA, INDORE - 452001,
MADHYA PRADESH, INDIA

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 not changed.
3. The certificate validity is conditioned by positive results of surveillance audits.
4. 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.

For the Products:

**MANUFACTURING OF HOSPITAL FURNITURE AND
TRADING OF MEDICAL EQUIPMENTS**

Certificate No : 25/B/AON09256
Initial Registration Date : 25/09/2025

Issuance Date : 25/09/2025
Date of Expiry : 24/09/2028

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