



Certificate of Compliance



We hereby declare that the technical files of all the items in each product group complies with the requirements of the Council Directive on **Medical Device Regulation - (Council Regulation 2017/745)**

Certificate No.: CE-7750

Manufacturer : CINTEX INDUSTRIAL CORPORATION

Address : 237, 242, JAYGOPAL INDUSTRIAL ESTATE, BHAVANI SHANKAR X ROAD, DADAR WEST, MUMBAI-400028, MAHARASHTRA, INDIA

Products : LAB, PHARMA & HOSPITAL EQUIPMENTS

Complies with the requirements applicable to it

The quality system file has been assessed, approved and is subject to continuous surveillance according to the Council Directive on **Medical Device Regulation - (Council Regulation 2017/745)**

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 changed.
3. The certificate validity is conditioned by positive results or surveillance audits.

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 test report 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.ukcertifications.org.uk/verify

Date of initial registration	14th April 2026
Date of this Certificate	14th April 2026
Certificate Expiry	13th April 2029
Recertification Due	13th April 2029

Authorised Signatory

