Certificate of Compliance

:: Certificate No :: WQ/22Q57365

This is to certify that

INDIA MEDICO INSTRUMENTS

WORKS-S-46, BADLI INDUSTRIAL ESTATE, PHASE-1, DELHI-110042, INDIA.

Has been assessed and found to be conforming the requirements of

FDA

(FDA Regulatory Guideline for Food and Drug Administration)

MANUFACTURER AND SUPPLY OF PHYSIOTHERAPY EQUIPMENT, OCCUPATIONAL THERAPY EQUIPMENT, REHABILITATION AID AND PANCHA KARMA EQUIPMENT.

This certificate is issued under the following conditions:

1. It applies only to the quality system maintained in the manufacture of above referenced scope / activities.
2. The certificate remains valid until the manufacturing conditions or the quality systems are changed and is subject to continuous surveillance according to the FDA Guidelines
3. The certificate validity is conditioned by positive results or surveillance audits

Validity of this certificate can be verified at www.mqacertification.com

Date of initial registration:	**22 May 2024**
First Surveillance Audit on or before:	**21 May 2025**
Second Surveillance Audit on or before:	**21 May 2026**
Re-certification Due:	**21 May 2027**

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

UKAF-CB-011

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