



Certificate of Registration

This is to certify that

KEE DIAGNOSTICS PVT. LTD.

**A-8, 3RD FLOOR NARAINA INDUSTRIAL AREA, PHASE II
NEW DELHI -110028 INDIA**

has been independently assessed by QRO
and is compliant with the requirement of:

ISO 13485:2016

Quality Management System for Medical Devices

For the following scope of activities:

**MANUFACTURING AND MARKETING OF IN-VITRO DIAGNOSTICS KIT, RAPID DIAGNOSTICS TEST
KIT, ELISA KITS & LABORATORY TESTING EQUIPMENT**

Date of Certification: 7th August 2025

1st Surveillance Audit Due: 6th August 2026

2nd Surveillance Audit Due: 6th August 2027

Certificate Expiry: 6th August 2028

Certificate Number: 305025080742MD



Chunant...

Head of Certification

Validity of this certificate is subject to annual surveillance audits to be done successfully on or before 365 days from date of the audit.
(In case surveillance audit is not allowed to be conducted; this certificate shall be suspended / withdrawn).

The Validity of this certificate can be verified at www.qrocert.org

This certificate of registration remains the property of QRO Certification LLP, and shall be returned immediately upon request.

India Office : QRO Certification LLP

142, IInd Floor, Avtar Enclave, Near Paschim Vihar West Metro Station, Delhi-110063, (INDIA)

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