

Increasing and Improving Awareness for Assuring Compliance

Professional Webinar for GLP Stakeholders and Quality Leaders

Background

OECD Principles on Good Laboratory Practice (GLP) and other GLP regulations (e.g. USFDA and USEPA) mandate conduct of non-clinical health and environmental safety studies in compliance with those principles/regulations by test facilities. Inadequate implementation of systems and processes may lead to study rejection by regulatory agencies.

Key Reasons for Study Rejection

- Potential non-compliance
- Data integrity
- QA failures

Responsible Parties

- Test Facility Management
- Quality Assurance Unit
- Study Director

Real-World Challenges

- Ensuring compliance
- Assuring compliance
- Varying expectations and interpretations

Webinar Date and Time

19 September 2026 (Saturday)

2:00 PM to 4:00 PM

Topic

**Regulatory Insights Towards
Rejection of GLP Studies**

Speaker: Anand Pandya, RQAP-GLP

Head, Preclinical QA
Microcrispr Pvt. Ltd., Vapi, Gujarat

Target Audience

- Test Facility Management
- Quality Assurance professionals
- Study Directors and personnel
- Test Item Controllers and Archivists
- Monitoring Scientists and Sponsors
- Consultants, academics, and students

Benefits of the Webinar

- Practical knowledge sharing
- Interactive discussion with an experienced speaker
- Participation certificate

Registration: All registrants will receive a separate email containing a link to join the webinar

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ICSQA Circular: Webinar on 19 September 2026