Promoting Regulatory Quality and Compliance

Indian Chapter of the Society of Quality Assurance (ICSQA)

(A Not-for-profit Organisation Registered under the Government of India)

Increasing and Improving Awareness for Assuring Compliance

Webinar for the Benefit of All stakeholders

Background

OECD Council Decision of 1997 has specific requirements for test item use in GLP studies. Similar requirements are applicable for conduct of studies in compliance with USFDA 21 CFR Part 58 and USEPA 40 CFR Part 160.

Requirements

- **Test item Characterization**
- **Dose Formulation Analysis**

Responsible Parties

- **Sponsor**
- **Test Facility Management**
- **Study Director**

Real-world Challenges

- **Lack of recognition**
- **Non-compliance / Deviation**
- **Study rejection**

Webinar Date and Time

14 March 2026 (Saturday)

From14:30 to 16:00 hours

Topic

Test Item Characterisation and Its Analysis in Vehicle

Responsibilities and Challenges for Meeting Compliance

Speaker: Dr. Natesan S., PhD, having extensive experience in conduct of studies, management of test facility and assuring quality.

Target Audience

- Quality Assurance professionals
- Study personnel (Study Directors, Principal Investigators, Study Personnel and others)
- Management, Monitoring Scientists and Sponsors
- Personnel managing regulatory data
- Consultants, Academics and Students

Benefits of the Webinar

- Knowledge sharing
- Interactive discussion
- Participation certificate

Registration

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

ICSQA member*	:	Free. CLICK HERE to Register for the Webinar
Non-member of ICSQA (Choose any one option)	:	Option 1: Become a member (₹1,000/-) and register for the Webinar. Option 2: Directly Register by paying ₹500/ for the webinar. CLICK HERE to become a member and/or register for the Webinar
Student	:	Free. CLICK HERE to register for the Webinar
*All webinars in 2026 are free for members		

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