QUALITY CIRCLE FORUM OF INDIA
ANKLESHWAR CHAPTER
PHARMACEUTICAL WING

Organizing
Two days' Training on:
Revised Schedule M for API
(Part I & Part XII)

Date: 2nd & 3rd February, 2026

Experts:

Dr. Milind Pathak

Mr. Bharat Bhamre

Program Facilitator:

Ms. Alka Kale

Schedule M:

- Schedule M of the Drugs and Cosmetics Rules, 1945, prescribes Good Manufacturing Practices (GMP) and specifies requirements for premises, plant, and equipment used in the manufacture of pharmaceutical products. Adherence to these standards ensures product quality, safety, and efficacy.

Training Objectives:

- Interpret each clause of Schedule M using relevant real-world examples.
- Identify gaps and deviations in current manufacturing practices.
- Prepare effectively for regulatory inspections and audits.
- Implement robust and sustainable compliance systems.

Audience:

- Senior & Middle Management professional engaged in pharmaceutical manufacturing industries.

Our Experts:

Dr. Milind Pathak

Associated with Pharmaceutical industries since 45 years

Mr. Bharat Bhamre

Mechanical Engineer Associated with Pharmaceutical industries since 40 years

DAY - 1

PART - 1

08:45 to 09:00	Breakfast & Registration
09:00 to 09:30	Inauguration
09:30 to 10:15	Pharmaceutical Quality System (PQS)
10:15 to 11:00	Quality Risk Management (QRM)
11:00 to 11:30	GMP, Sanitation and Hygiene, Qualification and Validation
11:30 to 11:45	Tea - Break
11:45 to 13:00	Complaints and adverse Reaction, Recall, Change Control, Contract Manufacturing, Self-inspection, Quality Audits and Suppliers' Audits and Approval
13:00 to 13:45	Lunch Break
13:45 to 15:30	Personnel, Premises
15:30 to 15:45	Tea Break
15:45 to 17:30	Equipment, Materials, Reference Standard, Waste Materials, Documentation
17:30	Closure

DAY - 2

PART - 12

08:45 to 09:00	Breakfast
08:45 to 09:00	Recape
09:15 to 11:00	Good Practices in Production, Quality Control & Computerized System
11:00 to 11:30	Introduction, Quality Management, Personnel, Building and Facilities
11:30 to 11:45	Tea Break
11:45 to 13:00	Process Equipment, Documentation and Records
13:00 to 13:45	Lunch Break
13:45 to 14:30	Materials Management, Production Controls
14:30 to 15:15	Packaging and Labelling, Storage and Distribution, Laboratory Controls
15:15 to 15:30	Tea Break
15:30 to 17:00	Validation and Change Control, Rejection and reuse of materials, Complaints and Recalls, Contract Manufacturers
17:00 to 17:30	Valedictory Function and Certificate Distribution
17:30	Closure

Schedule & Venue:

- **Date: 2nd & 3rd February, 2026 (two days)**
- **Time: 09:00 am to 05:30 pm**
- **Venue: Mastiff Hotel, Near Railway Station, Ankleshwar - 393002.**

Certificate:

- **Certificate will be awarded to the participants at the end of the training programme.**

Fees:

- **INR 7,500/- per participant + 18 % GST (Total 8,850 /- INR).**
- **Course fee includes training material, tea-snacks and working lunch.**
- **This training is a non-residential training.**

Registration Process:

- **Registration can be acknowledged on qcfi.ank@gmail.com**
- **Call on 9377434079 /9375233039 for any additional information and registration.**
- **Request to prepare Demand draft or Cheque in favour of "Quality Circle Forum of India" and can be submitted at following address.**

QUALITY CIRCLE FORUM OF INDIA,
Ankleshwar Chapter
Opp. State Bank of India (Main Branch), Station Road,
ANKLESHWAR-393001, Dist.: Bharuch (Gujarat)

- **GST number and Bank details shall be provided for payment through NEFT at qcfi.ank@gmail.com, if requested.**

Our Associates

SAI SEVA
HVAC PRIVATE LIMITED
Hygiene Air Partner

VISHWA CLEANROOM
Cleanroom Project

THE PD-VD CONSULTANCY

Marvenix Pharma Tech Solution

ACE Laboratories