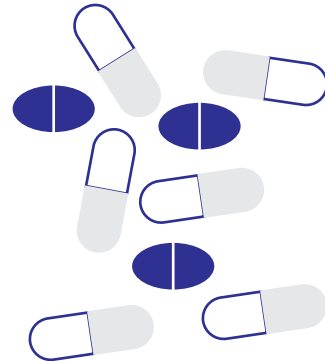
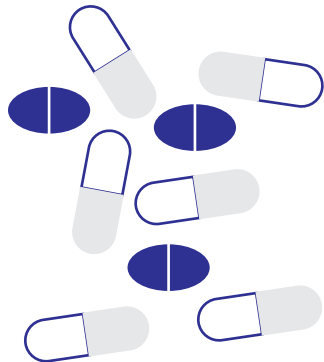




**TWO DAYS TRAINING PROGRAM  
(NON RESIDENTIAL)  
ON  
CONCEPTS OF cGMP  
(GOOD MANUFACTURING PRACTICES)**

**Faculty : Dr. Milind Pathak**



**Organized By:**

**QUALITY CIRCLE FORUM OF INDIA**

Ankleshwar Chapter & Surat Sub Chapter.

Opp. State Bank of India (Main Branch)

Station Road, ANKLESHWAR - 393 001

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Tel. : 02646 - 247461

Mobile : 9974062641 (Mr. Suresh Sharma)

Mobile : 93774 34079 (Dr. Milind Pathak)

**:: Venue**

**Hotel Lords Plaza,  
Station Road, Surat.**

**27th & 28th, MAY - 2016**

**Time : 9:00 A.M. to 5.00 P.M.**

## **INTRODUCTION :**

FDA insists compliance of cGMP while manufacturing of pharmaceuticals. Also, compliance of cGMP ensures satisfaction to stakeholders (patients).

It has been a mandatory requirement that persons involved in manufacturing of pharmaceuticals should be adequately trained for cGMP, as compliance of cGMP is a responsibility of all the persons involved in manufacturing.

## **OBJECTIVE :**

Objective of this training programme is to guide participants to current Good Manufacturing Practices during manufacturing of pharmaceuticals.

## **CERTIFICATE :**

Certificate will be awarded to the participants after successful compliance of the test at the end of the training programme.

## **METHODOLOGY :**

Current Good Manufacturing Practices during manufacturing of pharmaceuticals will be explained in detail. It will be a participatory session and followed by an evaluation on second day.

## **FACULTY**

**Dr. Milind Pathak - (☎) 09377434079**

\* Having 29 years of experience in reputed APIs manufacturing industries at various capacities in QA and Regulatory Department.

## SCHEDULE :

### DAY-1

9.00 AM	Registration
9.30 AM	Inauguration / Prayer / Introduction
10.00 AM	Concepts of cGMP Purpose Scope Tools
11.30 AM	Tea Break
11.45 AM	Who is responsible for cGMP? Responsibility of Ware-house Responsibility of Administration
1.00 PM	Lunch Break
2.00 PM	Responsibility of Production, Responsibility of Eng./Utility/Adm.
3.30 PM	Tea Break
3.45 PM	Responsibility of QC and QA
5.00 PM	Day Off

### DAY-2

9.00 AM	Qualification and Calibration of equipment.
11.00 AM	Tea Break
11.45 AM	Concepts of validation Documents and Records
1.00 PM	Lunch
2.00 PM	House-keeping SOPs Benefits of cGMP
3.00 PM	Group Play
4.00 PM	Evaluation Test
4.30 PM	Valedictory Function Certificate distribution
5.00 PM	Closure

**DATE & TIME :**

27th - 28th, May - 2016 : 9.00 A.M. to 5.00 P.M.

**FEES :**

Rs. 4000/- per participant + Service Tax @ 14.00 + 0.5 % SBC = Rs. 4580/- Course Fee includes training material, Tea / Snacks and Working Lunch.

**DISCOUNT**

10% Discount for QCFI-Life Members. 5% Discount for 3 or more participants from QCFI - Institutional Member organizations.

**VENUE**

Hotel Lords Plaza, Station Road, Surat., Gujarat.

**ENROLLMENT**

Please e-mail the list of names of participants along with local cheque/Demand Draft in favour of “**Quality Circle Forum of India**” at

Mr. Suresh Sharma, QCFI, Surat sub Chapter: Mo. 9974062641, E-mail : suresh\_sharmas@rediffmail.com

Dr. Milind Pathak, Joint Secretary & Faculty, QCFI, Ankleshwar Chapter: Mo. 9377434079, E-mail : nikan35@yahoo.co.in

QCFI, Ankleshwar Chapter: Ph. 02646 - 247461, E-mail : qcfi.ank@gmail.com

Last Date For Registration : **17th May - 2016**