

cGMP, Quality, Productivity, Validation and Positive Attitude as a Organization Culture for Corporate Growth

Sr.No	Time	Topic
1	9.00 to 9.15 a.m	Objective and Importance of the training programme.
2	9.15 to 10.45 a.m	GMP, Quality, Productivity, Positive Attitude: Definition and concept of Quality, Definition, Pharma manufacturers view on Quality, Pharma Regulatory's view on Quality, Doctors and Patients view on Quality, Quality and Attitude, various meaning of Quality, Resources and systems to build Quality, Meaning of word GMP, concepts of GMP, Productivity and its importance.
3	10.45 to 11.00 a.m	TEA BREAK
4	11.00 to 12.00 a.m	Schedule M: and its requirements in pharmaceutical Industry.
5	12.00 to 01.00 p.m	Validation: Objectives of validation, Benefits of validation, validation guidelines, Types of validation, Applicability of validation, D.Q. I.Q. O.Q.,P.Q.
6	01.00 to 02.00 p.m	LUNCH BREAK
7	02.00 to 03.00 p.m	Current Good manufacturing Practices : Why cGMP, Ten basic rules of GMP, documentation and its importance, Reasons of documentation, Rules about making records, House Keeping related to area.
8	03.00 to 3.45 p.m	Current Good Manufacturing Practices : House Keeping related to Materials, Records, and Documents, Personnel and Men, Materials, Machines and online set-ups, General Factory discipline, systems to be followed.
9	3.45 to 4.00 p.m	Video clip on GMP
10	4.00 to 4.15 p.m	TEA BREAK
11	4.15 to 4.30 p.m	Objective Test
12	4.30 to 5.00 p.m	Question and Answer Session

**Number of Participants Required
Who can participate ?**

**: Minimum 60
:Third Year & Final year B.Pharm. students, M.Pharm.
Pharmaceuticals and Quality Assurance students.**

Fees- Rs. 500 per participant

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