

In House Training



**FOCUS
PHARMA
WORKSHOPS**

FOCUS Pharma Workshops focus on technical aspects which covers the following aspects

- Quality Assurance & Quality Control
- Projects & Engineering
- Manufacturing & Production
- R&D & Clinical Research
- Systems & Documentation Practices

Focus Pharma Workshops are In house Training Workshops that are specially drawn out and custom-designed, in order to bring FOCUSsed results.

Why Should A Company Do Focus Pharma Workshops???

Cost Benefits

- **Pricing** :- Per day and not per person; an entire team can be trained at the same cost as sending a few persons to an external training.
- **Savings** :- Time and money; No travelling & accommodation expenses nor absence from workplace.
- **Teamwork**:- Get all your staff trained at the same level at the same time.

Professional Benefits

- **Tailor-made Programs** :- Designed for you; addressing issues that are most important to you and your business, using real-time examples.
- **Flexibility** :- Time, date, venue of your choice, duration of your convenience.
- **Audit Benefits** :- Auditors have made it mandatory to train your employees and keep them updated; set SOPs and make sure they are implemented.

Focus Benefits

- **Confidentiality** :- Assurance of not sharing your company specifics with others. Employees can openly share and discuss sensitive issues during the workshop, and find solutions to critical practical issues.
- **Services** :- During the workshops, Trainers of the Focus Group conduct Case Studies & Participant Assessment, before certification. This is followed by a Feedback process, to continue the training for the next batch of your employees.

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Focus Pharma Trainings have a series of course modules mentioned below. You can pick any one to suit your need of the hour.....

- Good Manufacturing Practices (GMPs)
- Good Laboratory Practices (GLPs)
- Good Clinical Practices (GCPs)
- Good Documentation Practices

- Drug Discovery and Drug Development
- Clinical Data Management
- Medical Writing for Clinical Research
- Regulatory Affairs in Clinical Research

- Technology Transfer
- Facility Designing
- Qualification of Machinery and Equipment

- Pharma Water Systems
- HVAC Systems
- Cleanrooms and Controlled Environments
- Isolators and Isolation Technologies

- Regulatory Inspections and Audits for GMP - Preparation
- Regulatory Compliance and Submissions to US / EU
- Regulatory Requirements – WHO, ISO, ICH, USFDA, MHRA, EDQM, TGA, etc.
- CTD and eCTD, Dossier Filing, and Regulatory Documentation
- IPR – Intellectual Property Rights, Patents, Copyrights, Trademarks

- Quality Risk Management
- Risk based approach
- CMC – Chemistry, Manufacturing, and Controls

- Stability Studies for Pharma and Biotech products
- Impurity Profiling
- Bio-Availability / Bio-Equivalence Studies

- Advanced Technologies
- Sterilization Processes and Sterility Assurance
- Process Simulations – Media Fills
- Laboratory Controls & Compliance

- Validation : Analytical, Cleaning, Process and Method
- Validation Master Plan
- Computerized Systems Validation

For more information, please contact:

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