Documents Information preparation for ISO 9001:2015

COURSE DESCRIPTION

Documentation and records generated within the scope of the Quality System are critical. It is very important that all quality documentation / records are generated in a correct, consistent and complete manner.

This course Provides practical guidance on how develop and document a system for quality assurance effectively in accordance with ISO 9001:2015 and avoid regulatory enforcement actions.

Participants learn how to write essential documents, maintain documentation, audit documentation, ensure change control and implement proper review and approval processes.

The involvement of trainees in the open discussions and case studies on various aspects associated with the authentication of quality system.

COURSE OUTLINE

- Recognize the value of good documentation in Quality Management Systems
- Basics of quality systems documentation
- Identify the required ISO 9001 documented Information
- Learn how to write effective documentation
- Identify steps for creating your documented Information
- Know how to adapt existing documents
- Understand control of documented information
- Practice writing quality system documents
- Final review of documented information
- Recognize how auditors look at documented information

Who Should Attend

- *Professionals who face the challenges of helping their organization focus and deploy common goals, strategies, plans, and customer requirements*
- *Management representatives*
- *Persons involved in defining, planning, or implementing an ISO 9001:2015 quality management systems*
- *New and experienced people in quality who want to use the QM framework and proven approaches and tools to be more effective on the job*

COURSE DURATION

4 Days 16 Hours