Laboratory Quality Management System

ISO/IEC 17025:2017



OBJECTIVE

- To get an overview of ISO/IEC 17025:2017 and the benefits to implement it
- To know about the resource requirements and process requirements for ISO/IEC 17025:2017
- To understand the processes involved in auditing and to learn auditing techniques based on ISO 19011:2018
- To learn how to prepare and maintain internal audit records

ENTRY REQUIREMENT

Delegates are better to complete "Understanding of Laboratory quality management based on ISO/IEC 17025: 2017" (Course Code: TLM1) or Implementation of Laboratory quality control and documentation based on ISO/IEC 17025: 2017 (Course Code: TLM3) or equivalent.

GENERAL INFORMATION

Venue: ACI Training Venue

Duration: 1 Day

Language: Cantonese supplemented

with English Material

Methodology: Presentation and experience

sharing

Certificate: Certificate of successful

Completion will be awarded to delegates who have attended

full course

WHO SHOULD ATTEND?

SUCCESS

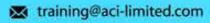
- Anybody who is involved in the implementation of the laboratory management system (LMS) within the organization
- Anybody who want to become LMS Internal Auditor and having a basic knowledge of standard or working in the organization
- Anybody wants to enhance their auditing knowledge and skills

COURSE CONTENT

- Overview of ISO 17025:2017 Laboratory Management system
- Requirements of ISO 17025:2017 Laboratory Management System
- ISO 17025:2017 Documentation
- ISO 17025:2017 Internal Audit Process
- ISO 17025:2017 Audit Records and Audit Checklist
- Impartiality, Risk Management, Decision Rule and Process Approach – Key changes in ISO 17025:2017

REMARK

- After completion of all TLM1, TLM3, TLM5 & TLM6, he or she is eligible for applying Professional Certificate of Laboratory Technical Officer.
- After completion of all TLM1, TLM3, TLM4, he or she is eligible for applying Professional Certificate of Laboratory Auditor.



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實驗室品質管理系統

ISO/IEC 17025:2017



培訓目的

- 全面了解ISO/IEC 17025:2017系統及其實施的好處
- 了解ISO/IEC 17025:2017的程序及其基本要求
- 了解基於ISO 19011:2018的審核方式及其審核流程
- 了解如何準備和維持內部審核記錄

培訓要求

建議代表們先完成「實驗室品質管理系統ISO/IEC 17025: 2017理解培訓」(課程編號: TLM1)或「實驗室品質管理系統ISO/IEC 17025: 2017 控制、執行與文件處理培訓」(課程編號: TLM3)或相關課程。

培訓詳情

地點: ACI 訓練中心

時間: 一天

課程語言: 廣東話授課輔以英語教材

上課模式: 講解、工作小組討論與練習

證書: 高出席率之學員將獲頒發課程

完成證書乙張

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培訓對象

SUCCESS

- 任何參與企業實施內部實驗室管理系之人士
- 任何想成為實驗室管理系統內部審核員並具有 基礎知識或在其企業中工作之人士
- 任何想提高自己的審核知識和技能之人士

培訓內容

- ISO 17025:2017實驗室管理體系的概述
- ISO 17025:2017實驗室管理體系的要求
- ISO 17025:2017的文檔
- ISO 17025:2017內部審核程序
- ISO 17025:2017的主要更改包括公正、風險管理、決策規定和流程方法

備註

- 在完成所有 TLM1、TLM3、TLM5 和 TLM6 後,他/ 她有資格申請實驗室技術人員專業證書。
- 完成所有TLM1, TLM3, TLM4後, 他/她有資格申請實驗室審核員專業證書。

