

COURSE OVERVIEW LE0290 Advanced Good Laboratory Practice (GLP)

CEUS

Course Title

Advanced Good Laboratory Practice (GLP)

Course Date/Venue

July 13-17, 2025/Tamra Meeting Room, Al Bandar Rotana Creek, Dubai UAE

Course Reference LE0290

Course Duration/Credits Five days/3.0 CEUs/30 PDHs

Course Description









This practical and highly-interactive course includes various practical sessions and exercises. Theory learnt will be applied using "MS-Excel" application.

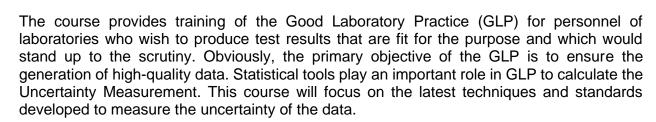
Good laboratory practice means different things to different people. This course covers the advanced topics in Good Laboratory Practice (GLP). All the practices that ensure that laboratory testing is planned, performed, monitored, recorded and reported in an organised and controlled manner, will be discussed in general and the Uncertainty Measurement (GUM) in particular.

The course addresses the requirements of EN ISO / IEC 17025: 2005 General Requirements for the Competence of Testing and Calibration Laboratories as well as the common requirements of GLP Compliance (OECD, USA, applicable Canada. Europe & Japan) as to QC/QA/Production Control Laboratories. The different GLP standards set out the requirements for technical competence and proper management of the analytical laboratories. The standards usually cover a variety of topics, that will be covered during this course: Laboratory Organization; Personnel; Safety Assurance; Laboratory Facility; Testing and Measuring Equipment; Apparatus Reagents; Test Methods and and Procedures: Calibration; Samples and Test Data Storage, Inspection, Assessment and Uncertainty Measurement (UM). The course will also discuss the requirements to be met by a laboratory supporting its organisations quality system to the ISO 9000 series of standards.



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The Guideline for Uncertainty Measurement (GUM) will be introduced in a simple way. Every effort is made in this course to eliminate unnecessary complications, to apply the GUM at its simplest level and to take away apparent mystery. Participants who have never drawn up uncertainty budgets before usually develop the required skill well before the end of the course. Others who seek explanations of GUM complexities obtain clarifications expressed in simple terms. Measurement uncertainty problems are solved by brainstorming methods so as to generate interaction by all participants. The course will use the Statistical Process Control (SPC) module as a tool to validate the uncertainty measurement. It covers control charts and their applicability to uncertainty before covering a step-by-step process of calculating uncertainty for a typical application. Methods of reporting uncertainty, the uncertainty budget and applicable standards and guidelines for expressing measurement uncertainty are covered in some detail. Successful completion of this course will enable participants to understand, evaluate and express measurement uncertainty.

Course Objectives

Upon the successful completion of this course, each participant will be able to:-

- Apply an in-depth knowledge in the Good Laboratory Practice (GLP) in accordance with the international rules and standards
- Practice uncertainty measurement, data validation, method validation and statistical process control in compliance with the latest revision of ISO 17025 standard
- Implement the Good Laboratory Practice (GLP) and gain advanced knowledge in modern laboratories GLP system
- Distinguish the various GLP standards and regulations that set out the requirements for technical competence and proper management of analytical laboratories
- Identify the common requirements of GLP compliance as per OECD, the USA, Canada, Europe and Japan
- Explain the differences between the USA and OECD regulations and identify which system is suitable for your laboratory
- Discuss the critical GLP compliance issues
- Describe the GLP requirements related to organization, personnel, facilities, equipment, operations, testing, control, protocols, records and reports
- Recognize the scope, field of application, components & sources of uncertainty and carry out analytical measurement and experimental studies of method performance
- Review and improve the process of measurement uncertainty estimation and be able to analyze the specification of the measurand
- Identify uncertainty sources and employ the process of quantifying uncertainty
- Carryout the process of calculating the combined uncertainty and be able to review the procedure on reporting uncertainty
- Explain Statistical Process Control (SPC) and identify its applications
- Apply method validation techniques in accordance with ISO17025



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Who Should Attend

This course provides an overview of all significant aspects and considerations of GLP in accordance with the international standards for those involved in laboratory management. This includes laboratory managers, superintendents, supervisors, scientists, doctors, physicians, chemists, researchers, analysts, officers, coordinators and other technical staff. R&D and government statutory employees will benefit from the international standards and regulation part of the course. Further, this course is very important for QA/QC employees and Third-Party Inspection and certification companies.

Exclusive Smart Training Kit - H-STK®



Participants of this course will receive the exclusive "Haward Smart Training Kit" (**H-STK**[®]). The **H-STK**[®] consists of a comprehensive set of technical content which includes **electronic version** of the course materials conveniently saved in a **Tablet PC**.

Training Methodology

All our Courses are including **Hands-on Practical Sessions** using equipment, State-ofthe-Art Simulators, Drawings, Case Studies, Videos and Exercises. The courses include the following training methodologies as a percentage of the total tuition hours:-

- 30% Lectures
- 20% Practical Workshops & Work Presentations
- 30% Hands-on Practical Exercises & Case Studies
- 20% Simulators (Hardware & Software) & Videos

In an unlikely event, the course instructor may modify the above training methodology before or during the course for technical reasons.

Course Fee

US\$ 5, 500 per Delegate + **VAT**. This rate includes H-STK[®] (Haward Smart Training Kit), buffet lunch, coffee/tea on arrival, morning & afternoon of each day.

Accommodation

Accommodation is not included in the course fees. However, any accommodation required can be arranged at the time of booking.



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Course Certificate(s)

(1) Internationally recognized Competency Certificates and Plastic Wallet Cards will be issued to participants who completed a minimum of 80% of the total tuition hours and successfully passed the exam at the end of the course. Certificates are valid for 5 years.

Recertification is FOC for a Lifetime.

Sample of Certificates

The following are samples of the certificates that will be awarded to course participants:-







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(2) Official Transcript of Records will be provided to the successful delegates with the equivalent number of ANSI/IACET accredited Continuing Education Units (CEUs) earned during the course.

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Course Certificate(s)

Internationally recognized certificates will be issued to all participants of the course who completed a minimum of 80% of the total tuition hours.

• *** BAC

British Accreditation Council (BAC)

Haward Technology is accredited by the **British Accreditation Council** for **Independent Further and Higher Education** as an **International Centre**. BAC is the British accrediting body responsible for setting standards within independent further and higher education sector in the UK and overseas. As a BAC-accredited international centre, Haward Technology meets all of the international higher education criteria and standards set by BAC.



The International Accreditors for Continuing Education and Training (IACET - USA)

Haward Technology is an Authorized Training Provider by the International Accreditors for Continuing Education and Training (IACET), 2201 Cooperative Way, Suite 600, Herndon, VA 20171, USA. In obtaining this authority, Haward Technology has demonstrated that it complies with the **ANSI/IACET 2018-1 Standard** which is widely recognized as the standard of good practice internationally. As a result of our Authorized Provider membership status, Haward Technology is authorized to offer IACET CEUs for its programs that qualify under the **ANSI/IACET 2018-1 Standard**.

Haward Technology's courses meet the professional certification and continuing education requirements for participants seeking **Continuing Education Units** (CEUs) in accordance with the rules & regulations of the International Accreditors for Continuing Education & Training (IACET). IACET is an international authority that evaluates programs according to strict, research-based criteria and guidelines. The CEU is an internationally accepted uniform unit of measurement in qualified courses of continuing education.

Haward Technology Middle East will award **3.0 CEUs** (Continuing Education Units) or **30 PDHs** (Professional Development Hours) for participants who completed the total tuition hours of this program. One CEU is equivalent to ten Professional Development Hours (PDHs) or ten contact hours of the participation in and completion of Haward Technology programs. A permanent record of a participant's involvement and awarding of CEU will be maintained by Haward Technology. Haward Technology will provide a copy of the participant's CEU and PDH Transcript of Records upon request.



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Course Instructor(s)

This course will be conducted by the following instructor(s). However, we have the right to change the course instructor(s) prior to the course date and inform participants accordingly:



Dr. Yousef Al-Mashni, PhD, MSc, BSc, is an International Expert in Analytical Laboratory with over 30 years of extensive experience. He is an authority in Laboratory Equipment, Laboratory Quality Management Systems (ISO 17025 and ISO 15189), Lab Safety & Health, Good Laboratory Practice (GLP) and Safety Procedure in Laboratories. His wide expertise also covers Water Analysis & Reporting, Water Sampling & Testing, Water Analyzer, Medical Laboratory Auditing, ISO 15489, Infection Control, Internal Quality

Control for Microbiologists, Analytical Techniques, Biochemical, Hematological, Parasitological, Biochemical, Microbiological & Serological Analysis of Clinical Specimens, Helmith Ova & Salmonella in Waste Water & Sludge, Microbiological Aspects & Analysis of Wastewater, Microbiology of Wetlands, Microbiological Indoor Air Quality, Entrococcus, Pseudomonas & Aeromonas, Sulfate Reducing Bacteria, Fluorescense Microscopy, Planktology of Ambient Environment, Oral, Medical & Diagnostic Microbiology and Oral & Dental Hygiene. Further, he is also well-versed in the areas of Food Hygiene and HACCP, Food Safety, Food Poisoning, First Aid & CPR and Fire Safety. He is currently the Deputy Principal & Chief Technical Instructor of UNRWA wherein he is responsible in developing and managing operations at the college/centre including building workshops and laboratories capacity, curriculum development and introducing new courses.

During his long career life, Dr. Yousef worked for many international companies handling key positions such as ICDL Centre Manager, Deputy Principal, Chief Technical Instructor, Acting Principal, Laboratory Supervisor, Technical Instructor, Technical & Vocational Instructor, Senior Medical Laboratory Technician and Medical Laboratory Technician.

Dr. Yousef has a PhD degree in Natural Health Sciences from the University of Florida (USA), Master degree in Clinical Microbiology and Bachelor degree with Honours in Microbiology. Further, he has Diploma in Vocational Education (UNRWA & UNESCO) and received several certifications like ICDL and Training of Trainers (TOT) in Cambridge University (England). He is a Certified Internal Verifier/Assessor/Trainer by the Institute of Leadership & Management (ILM), a Certified Instructor/Trainer and an active member of Jordan Medical Laboratories Society, Technical Accreditation Committee of Medical Laboratories (Jordan Institution & Metrology) and the Technical Accreditation Committee for Granting ISO 15189 Certificate. Furthermore, he has also published numerous technical papers and books including Medical & Diagnostic Microbiology, Practical Competencies in Medical Laboratory Technology, Safety in Medical Laboratory Science and Quality Control in Medical Laboratory Science just to name a few.



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Course Program

The following program is planned for this course. However, the course instructor(s) may modify this program before or during the course for technical reasons with no prior notice to participants. Nevertheless, the course objectives will always be met:

Day 1:	Sunday, 13 th of July 2025
0730 – 0800	Registration & Coffee
0800 - 0815	Welcome & Introduction
0815 - 0830	PRE-TEST
0830 - 0930	<i>Guide to Good Laboratory Practice</i> Introduction • Quality Assurance in Laboratory • GLP & The Transfer of Technology • The Cost of GLP Non-Compliance • International Regulations Governing GLP • Chronology & Current Regulations (U.S.A, OECD, Canada, Europe & Japan 3.2 Comparison Between US & OECD Regulations)
0930 - 0945	Break
0945 – 1100	Guide to Good Laboratory Practice (cont'd)Critical GLP Compliance Issues• Organization & Personnel• FacilitiesEquipment• Testing Facilities Operation• Test & Control Articles• Protocolfor & Conduct of a Study• Records & Reports• Conclusions
1100 - 1230	<i>Essential Statistics for Analytical Science</i> <i>Sample</i> • <i>Population</i> • <i>Normal Distribution</i> • <i>Summarizing Data</i> • <i>Confidence Intervals</i>
1230 - 1245	Break
1245 - 1420	Essential Statistics for Analytical Science (cont'd) Outliers • Errors • Accuracy • Precision • Significant Figures
1420 - 1430	Recap
1430	Lunch & End of Day One

Day 2: Monday, 14th of July 2025

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Scope & Field of Application
Uncertainty
Definition of Uncertainty • Uncertainty Sources • Uncertainty Components •
Error & Uncertainty
Break
Analytical Measurement & Uncertainty
Method Validation • Conduct of Experimental Studies of Method Performance •
Traceability
The Process of Measurement Uncertainty Estimation
Specification of the Measurand
Break
Identifying Uncertainty Sources
Quantifying Uncertainty (GUM)
Introduction • Uncertainty Evaluation Procedure • Relevance of Prior Studies
• Evaluating Uncertainty by Quantification of Individual Components • Closely
Matched Certified Reference Materials
Recap
Lunch & End of Day Two



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Day 3:	Tuesday, 15 th of July 2025
0730 - 0930	Quantifying Uncertainty (GUM) (cont'd) Using Prior Collaborative Method Development & Validation Study Data • Using In-House Development & Validation Studies • Evaluation of Uncertainty for Empirical Methods • Evaluation of Uncertainty for Ad-Hoc Methods • Quantification of Individual Components
0930 - 0945	Break
0945 - 1045	<i>Quantifying Uncertainty (GUM) (cont'd)</i> <i>Experimental Estimation of Individual Uncertainty Contributions</i> • <i>Estimation Based on other Results or Data</i> • <i>Modelling from Theoretical Principles</i> • <i>Estimation Based on Judgement</i> • <i>Significance of Bias</i>
1045 - 1230	Calculating the Combined Uncertainty Standard Uncertainties • Combined Standard Uncertainty • Expanded Uncertainty
1230 – 1245	Break
1245 - 1420	Reporting Uncertainty General • Information Required • Reporting Standard Uncertainty • Reporting Expanded Uncertainty • Numerical Expression of Results • Compliance Against Limits
1420 - 1430	Recap
1430	Lunch & End of Day Three

Day 4:	Wednesday, 16 th of July 2025
0730 – 0930	Statistical Process Control (SPC)The Measurement Process Model • Setting Up the Statistics • Evaluating theProcess • Setting Control Limits • Using Control Limits • PracticalConsiderations • Rules
0930 - 0945	Break
0945 - 1045	Statistical Process Control (SPC) (cont'd) Pareto Charts • Fishbone Diagram • Flowcharting • Scatter Plots • Charts
1045 - 1230	<i>Method of Validation</i> Scope and Method Specifications • Setting Validation Criteria that will Satisfy the Needs of Customers • Validation Planning • Accuracy by the Analysis of a CRM • Precision/Repeatability • Working range • Linearity
1230 - 1245	Break
1245 - 1420	<i>Method of Validation (cont'd)</i> <i>Testing for Performance Characteristics</i> • <i>Limit of Detection (LOD) &</i> <i>Quantitation (LOQ)</i> • <i>Quality Control Plan for Routine Analysis</i>
1420 - 1430	Recap
1430	Lunch & End of Day Four

Day 5:	Thursday, 17 th of July 2025
0730 – 0830	<i>Method of Validation (cont'd)</i> <i>Precision/Ruggedness</i> • <i>Robustness</i> • <i>Stabilities</i> • <i>Validation Report</i>
0830 - 0930	<i>Method of Validation (cont'd)</i> <i>Traceability</i> • <i>Proof of the Performance of A Method and/or Assuring the Quality</i> <i>of a Measurement According to ISO 17025</i> • <i>Analysis of QC Samples</i> • <i>Documentation</i>
0930 - 0945	Break



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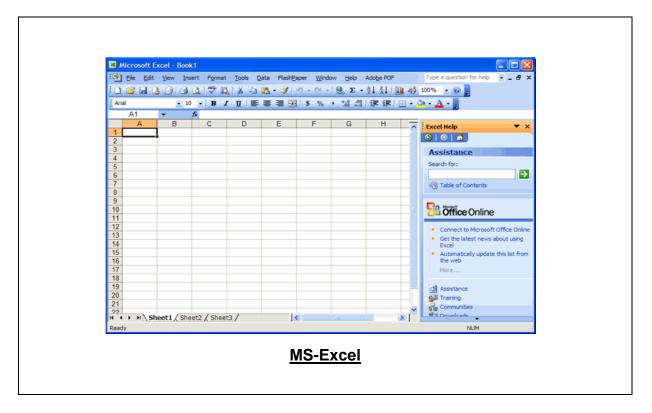




0945 – 1230	Method of Validation (cont'd)Analysis of CRM's • Reproducibility Studies Like Proficiency Testing &Interlaboratory Studies • Customer Management
1230 - 1245	Break
1245 – 1300	Case Study: How to write a Method Validation Document. Students will Work in Groups or Individually to Solve a Case Study
1300 - 1315	Course Conclusion
1315 – 1415	COMPETENCY EXAM
1415 – 1430	Presentation of Course Certificates
1430	Lunch & End of Course

Simulator (Hands-on Practical Sessions)

Practical sessions will be organized during the course for delegates to practice the theory learnt. Delegates will be provided with an opportunity to carryout various exercises using "MS-Excel" application.



Course Coordinator

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