



Tower Mains Training

Autumn 2019 Course Dates

For course bookings and enquiries please email training@towermains.com

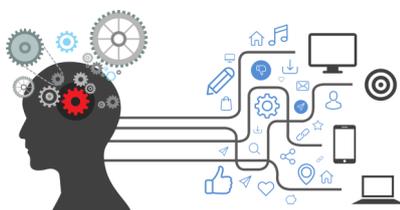
Discovery

Our one-day **Discovery** courses aim to provide introductions to the Good Practices (GxP see definitions below). They are intended for participants who would like to implement these guidelines into their organisations or they wish to expand their knowledge.

<u>Course Title</u>	<u>Summary</u>	<u>Course Information</u>
INTRODUCTION TO GOOD LABORATORY PRACTICE (GLP)	What is GLP and when does it apply? This is an introductory course for staff at all levels involved in laboratory work.	3 rd October 2019 1 day £500
INTRODUCTION TO GOOD CLINICAL LABORATORY PRACTICE (GCLP)	We discuss the special requirements that apply to the analysis of clinical samples. This is an introductory course for staff at all levels involved in clinical laboratories.	4 th October 2019 1 day £500
INTRODUCTION TO GOOD CLINICAL PRACTICE (GCP)	This course provides an overview of how to conduct commercial and non-commercial clinical trials in compliance with GCP.	22 nd October 2019 1 day £500
INTRODUCTION TO GOOD MANUFACTURING PRACTICE (GMP)	An overview of Good Manufacturing Practice requirements suitable for all staff.	23 rd October 2019 1 day £500

Prices exclude VAT and accommodation

See next page for our Refresh and Update Courses



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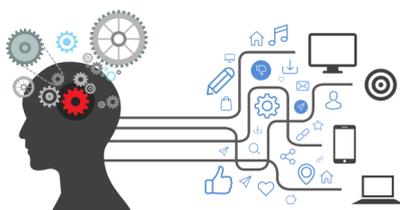
Refresh and Update

The **Refresh and Update** half-day modules are designed to keep staff up-to-date with developments in the regulations and meet regulatory authorities latest requirements and expectations.

<u>Course Title</u>	<u>Summary</u>	<u>Course Information</u>
DATA INTEGRITY	Course is designed to provide guidance on how data should be controlled and managed from initial generation and recording, through processing, use, retention, archiving and retrieval.	7 th October 2019 half day £300
GLP REFRESHER	Provides an update in what is happening in the GLP world in the previous 12 months such as new guidance or position papers and the implications and impact on facilities.	8 th October 2019 half day £300
MULTI-SITE STUDIES	This course will provide participants comprehensive understanding of the requirements and roles and responsibilities for being involved in multi-site studies being conducted in compliance with Good Laboratory Practice (GLP).	9 th October 2019 half day £300
GCP REFRESHER	Key aspects of Good Clinical Practice (GCP), main content of the ICH-GCP and key principles, highlight main changes from the previous version to R2 and present how Regulations and ICH-GCP are cross referenced.	28 th October 2019 half day £300

Prices exclude VAT and accommodation

See next page for our Clinical Research Masterclasses



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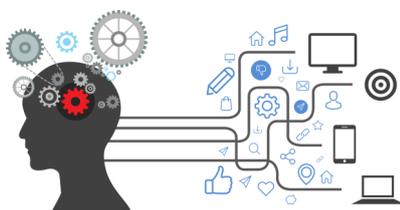
Clinical Research Masterclasses

The **Clinical Research Masterclasses** are aimed at clinical research professionals who need to refine and develop their professional skills.

<u>Course Title</u>	<u>Summary</u>	<u>Course Information</u>
DATA INTEGRITY AND DATA PROTECTION IN CLINICAL TRIALS	Data must be credible and reliable to have any value. Use of personal confidential information requires special consideration. Both these are discussed in this course.	24 th October 2019 1 day £500
PEOPLE MANAGEMENT IN CLINICAL TRIALS	Why do we need people skills? PPA conduct and analysis, how to identify the DISC types and how to manage them effectively in order to achieve what you need.	19 th November 2019 1 day £500
RISK MANAGEMENT IN CLINICAL TRIALS	This course aims to create a practical, process driven, risk management system that can be adapted to fit any requirement.	20 th November 2019 1 day £500
ENSURE YOUR TRIAL MASTER FILE (TMF) IS INSPECTION READY	What does the TMF comprise of? Paper, electronic and hybrid TMF. What to expect during the inspection? Regulations and best practices	10 th December 2019 half day £300
MONITORING AND MONITORING OVERSIGHT	What is Monitoring? Role of Monitors and oversight. Frequent Audit findings	11 th December 2019 half day £300

Prices exclude VAT and accommodation

See next page for our Quality Management Courses



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Autumn 2019 Courses

Quality Management

Quality Management courses aim to provide an understanding of how to develop, manage and audit Research Quality Management Systems.

<u>Course Title</u>	<u>Summary</u>	<u>Course Information</u>
AUDITING RESEARCH QUALITY SYSTEMS	The course is aimed at those who are currently working in or wish to consider moving to Research Quality Assurance. It is applicable to anyone who is involved in audit or inspection either within a regulated system such as GLP or GCP or in 'non-regulated' research and development.	3 rd and 4 th December 2019 2 days £800
MANAGING A QUALITY ASSURANCE PROGRAMME	This course will give participants an understanding and the tools to identify critical aspects of the quality system to be audited through the use of risk evaluations. In addition will cover aspects of the audit process including assessing audit effectiveness, the audit team, audit follow up and correction and corrective action.	5 th December 2019 1 day £500
QUALITY MANAGEMENT FOR NON-REGULATED RESEARCH	This course will give participants an understanding of the key elements of a quality system and how to develop, deliver and maintain a quality system within a working research facility.	6 th December 2019 1 day £500

Prices exclude VAT and accommodation

Want to get in touch?



training@towermains.com

Tower Mains Training – Learn from the Experts



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