

Tower Mains Training

Spring 2020 Course Dates

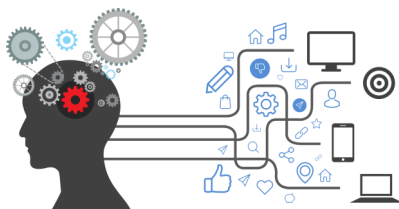
For course bookings and enquiries please email training@towermains.com

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 CLINICAL PRACTICE (GCP)	This course provides an overview of how to conduct commercial and non-commercial clinical trials in compliance with GCP.	27th April 2020 1 day £500
INTRODUCTION TO GOOD MANUFACTURING PRACTICE (GMP)	An overview of Good Manufacturing Practice requirements suitable for all staff.	1st May 2020 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.	22nd May 2020 1 day £500
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.	29th May 2020 1 day £500

See next page for our Refresh and Update Courses
Prices exclude VAT and accommodation



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Spring 2020 Courses

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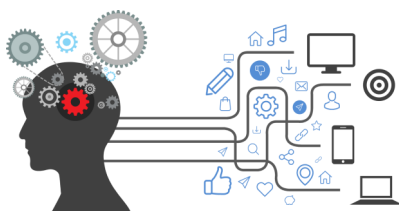
Refresh and Update

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>
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.	28th April 2020 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.	29th April 2020 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).	30th April 2020 half day £300
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.	7th May 2020 half day £300

Prices exclude VAT and accommodation

See next page for our Clinical Research Masterclasses



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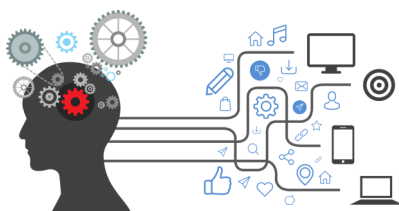
Clinical Research Masterclasses

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.	11th May 2020 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.	12th May 2020 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.	13th May 2020 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	14th May 2020 half day £300
MONITORING AND MONITORING OVERSIGHT	What is Monitoring? Role of Monitors and oversight. Frequent Audit findings	15th May 2020 half day £300

Prices exclude VAT and accommodation

See next page for our Quality Management Courses



Tower Mains Training

Spring 2020 Courses

Quality Management

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

Course Title	Summary	Course Information
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.	20th May 2020 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.	21st May 2020 1 day £500
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.	27th and 28th May 2020 2 days £800

Prices exclude VAT and accommodation

Want to get in touch?



training@towermains.com

Tower Mains Training – Learn from the Experts



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