Transforming lives through science, innovation and collaboration

Pharmaceutical
Training Schedule 2024

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Auditing

COURSE	DAYS	JAN	FEB	MAR	APR	MAY	JUN	JUL	AUG	SEPT	OCT	NOV	DEC
Active Pharmaceutical Ingredients (API) Auditing	1						5 Virtual						
Auditing to Pharmaceutical Standards	1			12 Virtual									2 Virtual
GMP Lead Auditor – Pharmaceutical Quality Systems *(IRCA Ref: PR325)*	5	15-19 Virtual	26 Feb-1 Mar In person Cookham, UK			20-24 Virtual		1-5 In person Cookham, UK		23-27 In person Waterford, Ireland	14-18 Virtual		
QMS Lead Auditor – Pharmaceutical Supply Chain *(IRCA Ref: PR330)*	5						17-21 In person Cookham, UK			9-13 Virtual		18-22 In person Cookham, UK	
Internal Auditor	2		7-8 Virtual		22-23 Virtual		5-6 In person Reading, UK					4-5 In person Liverpool, UK	
GDP/RP Auditor	2	10-11 Virtual		26-27 Virtual				2-3 In person Reading, UK		3-4 In person Reading, UK			4-5 In person Reading, UK
Auditing against the new Annex 1 Steriles Guideline	90 mins			26 Virtual									
Virtual Auditing – enhancing your skills	90 mins					16 Virtual							
Auditing Alumni CPD Refresher	90 mins						11 Virtual						
Auditing ATMPs	90 mins									17 Virtual			

Group booking discount

Book three or more delegates on the same course, running on the same date to unlock our group discounts!

- First booking charged at full published rate
- Second booking charged at full published rate minus 20%
- Third or more bookings charged at full published rate minus 50%

Contact our team for details.

Information is correct at the time of going to print.
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For more details please contact the Training Team:
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Good Manufacturing Practice

COURSE	DAYS	JAN	FEB	MAR	APR	MAY	JUN	JUL	AUG	SEPT	OCT	NOV	DEC
Good Manufacturing Practice Advanced *RSC Approved*	3			25-27 Virtual	24-26 In person Waterford, Ireland	13-15 In person Liverpool, UK		15-17 In person Reading, UK		23-25 Virtual			4-6 Virtual
Good Manufacturing Practice – The Essentials	1			20 Virtual		8 Virtual		3 Virtual				5 Virtual	
Good Manufacturing Practice in the Laboratory	1				4 Virtual					2 Virtual			
Interpretation of Statistical Values Used in GMP Applications	1		12 Virtual				19 Virtual						
Management of GMP Inspections	2			7-8 Virtual									
Medical Device Regulations (MDR) 2017/745	1					8 Virtual						6 Virtual	
Pharmaceutical Product Development – GMP Requirements	2			4-5 Virtual						9-10 In person Reading, UK			
Pharmaceutical Quality Management System	3											6-8 In person Liverpool, UK	

Microbiology

COURSE	DAYS	JAN	FEB	MAR	APR	MAY	JUN	JUL	AUG	SEPT	OCT	NOV	DEC
Microbiology Environmental Monitoring *in Partnership with Cherwell Laboratories*	1		6 In person Oxford, UK								31 In person Oxford, UK		
Introduction to Pharmaceutical Microbiology	1				17 Virtual					12 Virtual			
Manufacturing Sterile Products	2				25-26 Virtual					19-20 Virtual		26-27 Virtual	
Pharmaceutical Microbiology Advanced	3				8-10 In person Wokingham, UK			29-31 In person Wokingham, UK			15-17 In person Wokingham, UK		
Water Systems and Microbiological Control	1				23 Virtual					5 Virtual			

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Qualified Person

ROYAL SOCIETY OF CHEMISTRY | APPROVED TRAINING

COURSE	DAYS	JAN	FEB	MAR	APR	MAY	JUN	JUL	AUG	SEPT	OCT	NOV	DEC
Active Pharmaceutical Ingredients	2				15-16 In person Reading, UK		5-6 In person Liverpool, UK					18-19 Virtual	
Analysis and Testing	3				22-24 In person Reading, UK			8-10 In person Reading, UK			21-23 In person Reading, UK		
Biotechnology Issues	1					20 Virtual						4 Virtual	
Investigational Medicinal Products	2	10-11 Virtual				16-17 In person Liverpool, UK							2-3 In person Reading, UK
Mathematics and Statistics	3		20-22 Virtual					3-5 In person Liverpool, UK				5-7 In person Reading, UK	
Medicinal Chemistry and Therapeutics	3			11-13 Virtual						2-4 Virtual			
Pharmaceutical Formulation and Processing Part 1*	2*	29-30 Virtual				28-29 In person Reading, UK				9-10 In person Reading, UK			
Pharmaceutical Formulation and Processing Part 1* – Optional Practical Day	1*					30 In person Reading, UK				11 In person Reading, UK			
Pharmaceutical Formulation and Processing Part 2*	3*			5-7 In person Reading, UK				22-24 In person Reading, UK				26-28 Virtual	
Pharmaceutical Law and Administration	2	22-23 Virtual					3-4 In person Reading, UK				1-2 In person Liverpool, UK		
Pharmaceutical Microbiology	3				8-10 In person Wokingham, UK			29-31 In person Wokingham, UK			15-17 In person Wokingham, UK		
Pharmaceutical Packaging	3		5-7 Virtual							16-18 In person Barnstaple, UK			10-12 In person Barnstaple, UK
Pharmaceutical Quality Systems	3	24-26 Virtual					25-27 In person Liverpool, UK				8-10 In person Reading, UK		
Role and Professional Duties	2	15-16 Virtual			3-4 Virtual	7-8 In person Reading, UK					30-31 In person Liverpool, UK		

*Please note to cover the criteria set out in the QP study guide, both Part 1 and Part 2 should be attended.

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Biopharmaceutical

COURSE	DAYS	JAN	FEB	MAR	APR	MAY	JUN	JUL	AUG	SEPT	OCT	NOV	DEC
Good clinical practice for ATMPs	1			11 Virtual				9 Virtual					
Biotechnology Issues	1					20 Virtual						4 Virtual	
ATMPs – Quality by Design from Bench to Clinic	1							2 Virtual		25 Virtual			
Biologically Derived Products – Manufacture, Testing and Compliance Challenges	2				29-30 Virtual			10-11 Virtual			28-29 Virtual		
ATMPs – What Are They and What Do The Laws Say	90 mins			14 Virtual									
ATMP Facility Design and Method Development	90 mins					20 Virtual							
ATMP Scale Out, Scale Up and Decentralised Manufacturing	90 mins							16 Virtual					

Supply Chain

COURSE	DAYS	JAN	FEB	MAR	APR	MAY	JUN	JUL	AUG	SEPT	OCT	NOV	DEC
Good Distribution Practice	1			20 Virtual			4 Virtual				14 Virtual		
Responsible Person and GDP *(Cogent Gold Standard Approved)*	3		12-14 In person Reading, UK			13-15 Virtual	24-26 In person Dublin, Ireland	15-17 Virtual		16-18 In person Liverpool, UK		6-8 Virtual	
Responsible Person Refresher	1				18 Virtual			25 Virtual				28 Virtual	
Responsible Person Forum	2 hours	19 Virtual		1 Virtual		10 Virtual		19 Virtual		13 Virtual		15 Virtual	

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Other Courses

COURSE	DAYS	JAN	FEB	MAR	APR	MAY	JUN	JUL	AUG	SEPT	OCT	NOV	DEC
Data Integrity, Electronic Records and Signatures	1			7 Virtual							22 Virtual		
Introduction to Pharmacovigilance	1						27 Virtual						
Leadership and Influencing Skills for QPs & New Managers	1		22 Virtual								29 Virtual		
Quality Risk Management – How to apply ICH Q9 in Practice	1				29 In person Reading, UK			9 In person Reading, UK				20 In person Reading, UK	
QC Chemistry Crash Course	1			25 Virtual						24 Virtual			
Root Cause Analysis and CAPA	2			20-21 Virtual									5-6 Virtual
Technical Report Writing	1		15 Virtual					11 Virtual					
Technology Transfer of Pharmaceutical Products	1		5 In person Reading, UK			7 In person Reading, UK							

Validation

COURSE	DAYS	JAN	FEB	MAR	APR	MAY	JUN	JUL	AUG	SEPT	OCT	NOV	DEC
Cleaning Validation	1			14 Virtual							15 Virtual		
Introduction to Validation	1					1 Virtual					2 Virtual		
GxP Computerised Systems Validation and Compliance	2		7-8 Virtual			1-2 Virtual					23-24 Virtual		
Process Validation and Qualification, including Validation Methods	2			26-27 In person Reading, UK						17-18 In person Reading, UK			
GAMP 5 Second Edition Workshop	1					14 In person Reading, UK						5 In person Reading, UK	

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Training for your team

We pride ourselves in offering flexible learning, perfectly tailored to your company's specific requirements.

For this reason, all the courses on our 'open' schedule can be run via our digital platform or face to face in a location of your choice.

Train several staff at the same time, saving on multiple course fees, travel and accommodation costs.

Popular courses in 2023

- Preparing for a Regulatory Inspection
- ICH Q8, ICH Q9 and ICH Q10
- Annex 1
- Data Integrity
- Good Manufacturing Practice
- Responsible Person
- GMP Lead Auditor
- Internal Auditor
- Cleaning Validation

To find out more or discuss your needs, please contact the Training Team

trainingsales@rssl.com

+44(0) 118 918 4000

Or visit:

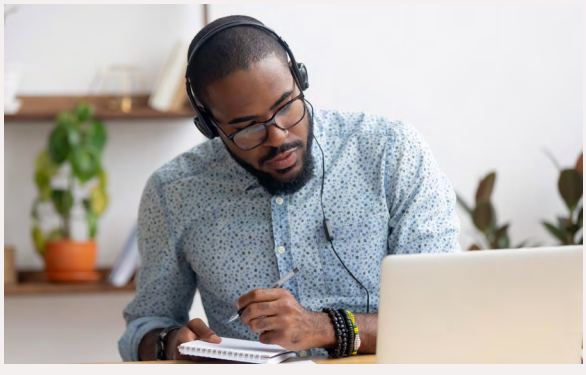
www.rssl.com/training-consultancy/training-courses-a-to-z

Self-paced eLearning

If you prefer to study at a time and pace that fits around your schedule, eLearning may be just the thing. Also useful if you have a large team to train with varying availability, you will have three months of unrestricted access in which to complete your course.

View our eLearning portfolio:

www.rssl.com/training-consultancy/e-learning



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About RSSL Training & Consultancy

We've been training and developing pharmaceutical, biopharmaceutical, medical device, and healthcare professionals across the complete life cycle of pharmaceutical products for over 30 years. We know what it takes to succeed and employ leading industry experts to deliver the most up-to-date training and CPD so individuals can achieve their career aspirations and companies have the skills they need to excel on the global stage and make a positive difference in the world.

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