H3 Impact of the EU Regulation (536/2014)

8th May 2019

The Institute of Clinical Research

The day will start at 08:45 with registration and coffee for a prompt start at 09:15. We aim to finish by 17:30.

ICR Training

H3 Impact of the EU Regulation (536/2014)

10 CPD POINTS

8th May 2019

H3

Venue - The Institute of Clinical Research Training Suite, White Waltham

Course Information

The Course

The 2001 Clinical Trials Directive (2001/20/EC) has been criticised by patients, researchers and industry alike for its disproportionate regulatory requirements. High costs and a lack of harmonisation of the applicable rules necessary for multinational clinical trials have contributed to a significant decline in the number of clinical trials in the EU – a reduction of about 25% in the last few years. The new EU Clinical Trials Regulation is due to be enacted in 2016 and will replace the 2001 Clinical Trials Directive (CTD) (2001/20/EC). The Regulation aims to restore the EU's competitiveness in clinical research and the development of new and innovative treatments and medicines by cutting red-tape and bringing patient-oriented research back to Europe. The regulation includes rules for clinical trials which are conducted outside the EU but referred to in a clinical trial application within the EU. For such trials, the rules call for compliance with regulatory requirements at least equivalent to those in the EU, including rules on transparency. In this highly interactive course, we explore the changes proposed by the regulation and how they will impact our current practices in conducting global clinical trials.

Learning Objectives

- Distinguish between a Regulation and a Directive
- Identify what will change under Regulation 536/2014
- Identify what will not change under Regulation 536/2014
- Define Low-Impact Interventional Trials
- Discuss the impact on:

– Application processes

– Informed consent

– Transparency requirements

- Describe next steps for implementation of Regulation 536/2014

Course Outline

- Review of the current regulatory situation in EU
- Directive vs Regulation
- Clinical Trial Authorization Process
- Trials in Emergency Situations
- Trials with authorized medicinal products
- Risk Based Evaluations
- Non-EU Sponsors
- Transparency
- Insurance Issues
- Co-Sponsorship Concept
- Safety Reporting
- Inspections in EU and Beyond

Who would benefit

Clinical researchers with at least one year of experience, who wish to explore the impact and wider application of the new EU Regulation for Clinical Trials on their current practices.

Course Fees

Guest	**£550.00**
ICR Member	**£450.00**
ICR Member Academic	**£350.00**

Your Trainer

Alison Messom

A molecular geneticist by training, Dr. Alison Messom has nearly 20 years of industry experience, working both within Pharmaceutical Companies and Clinical Research Organisations and is the current Chairman of the Board of Directors of the ICR.

With detailed experience of directing global clinical trials, Dr. Messom has worn many different hats during her career. She has had the opportunity to travel far and wide on business; lived in the UK, France and Switzerland, and gained operational experience in over 50 countries and line management in over 30.

During the course of her career she has line managed at all levels from CTAs through to Directors and VPs. At both i3 Research and ICON she helped build a new department of dedicated line managers for CRAs and also Start Up teams. While within the Pharma environment she line managed teams within Study and Project Management, Clinical Supplies, Translational Medicine, Data Management, Statistics, Informatics and IT – showing that line management skills are transferrable across functional areas.

Pre-Course Questionnaire - To be completed by all delegates

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Please complete and sent to training@icr-global.org or fax to +44 01628 501 709

Course Title: H3 Impact of the EU Regulation (536/2014)

Date: 08 May 2019

Name: .....

Company / Hospital: .....

Position / Job Title: .....

How much experience of clinical trials do you have? (Years)

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What are you hoping to get out of the day?

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State one issue/problem you would like discussed at the meeting

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Dietary Requirements

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* The ICR will work with the venue catering team and endeavour to accommodate specific dietary requirements e.g. Vegetarian/Vegan/Gluten Free - however it may not be possible to cover all requests for dietary preferences.

Training Location

Address: The Institute of Clinical Research Training Suite, Suite 1, Cedar Court Grove Business Park White Waltham, Berkshire, SL6 3LW

The ICR's training suite is situated near the charming village of White Waltham. Located just off the A404, with connections by road or rail.

The Institute of Clinical Research Training Suite

Suite 1, Cedar Court

Grove Business Park

White Waltham

Maidenhead

SL6 3LW

By Car: Join the A404 either via at J5 of the M40, or J8/9 of the M4. Follow the A404 until you reach J9A signposted to Maidenhead/Reading. Turn right at the first roundabout, then continue on that road, going straight across any subsequent roundabouts. Take the second right past the airfield. If you reach the village, you've gone too far.

Once inside Grove Park, turn left at the first mini roundabout, then right at the T junction. Turn right at the red brick building, and we are in the one furthest from the road. Park in the large car park on your right. Please contact the Secretariat using the buzzer by the door - type in 12 and then press the small bell.

By Train: The nearest station is Maidenhead, located on the Paddington - Reading line. We recommend getting a taxi from the station, as bus routes are sporadic. It should take roughly 15 minutes.

Local Taxi Companies

Dot to Dot + 44 01628 200 999
Golden Executive + 44 1628 622955
Imperial +44 1628 771777

Accommodation

The ICR does not specifically recommend any accommodation - however the following are within easy travelling distance of the training venue

Hotel or Bed & Breakfast Name	Contact number	Website	Approximate Distance to The Institute of Clinical Research
Premier Inn Maidenhead	08715 279520	http://www.premierinn.com	4 miles 14 minutes
Taplow House Hotel	01628 670056	https://taplowhousehotel.com	6 miles 16 miles
Bel & The Dragon	01628 521263	https://belandthedragon-cookham.co.uk/	7 miles 18 minutes
CIM Moor Hall	+44 1628 427500	https://moorhall.cim.co.uk/	7 miles 18 minutes

There are also a number of travel websites which may allow you to identify local accommodation and special offers - e.g. expedia

The Small Print

As a matter of policy we do not issue electronic copies of the slides used.

All ICR materials are copyrighted.

All delegates receive a delegate book.

Payment must be received in advance of a training course commencing. The ICR has the right to refuse entry for non-payment. Payment by invoice must be settled within 14 days from the date of invoice.

We understand that occasionally circumstances may change and that you will be unable to attend your chosen course. Notification of cancellation must be made in writing. If you cancel **more than 14 days prior to the event**, we will refund the course less £50 to cover administration costs. If you cancel within 14 days, no refund will be payable, but we will allow you to transfer to another course of your choice.

We will accept a change of delegate at any time without you incurring a penalty. The Institute of Clinical Research reserves the right to cancel any course that is under-subscribed but will give you 7 days notice in writing and will refund your course fees without any liability for any consequential or indirect loss.

At anytime, you may transfer to the same course within 12 months, or to another course of your choice within 6 months; a £25 administration fee will be charged for such transfers.

We may also need to change the venue but will give you 7 days notice in writing of the new location.

Programmes as published are correct, however due to circumstances beyond our control, trainers, speakers and/or the programme may need to be altered occasionally.

The ICR will work with the venue catering team and endeavour to accommodate specific dietary requirements e.g. Vegetarian/Vegan/Gluten Free – however it is not possible to cover all possible requests for dietary preferences.

Registration Form

Please complete and sent to training@icr-global.org or fax to +44 01628 501 709

Registration Form

Please photocopy this form for further registrations

Course Title: H3 Impact of the EU Regulation (536/2014) **Course Date:** 08 May 2019

Membership No.: ..... **Title(Dr,Mr,Mrs,etc):** ..... **First Name:** .....

Surname: ..... **Job Title:** .....

Company Name: .....

Email Address: .....

Confirmation of booking will be sent by email, unless you request here that it is sent by post ☐

Correspondence Address

Address: .....

Postcode: ..... **Country:** ..... **Telephone Number:** .....

Special Dietary Requirements

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Declaration

I agree to the terms and conditions of booking **Signature:** .....

Method of Payment

Please not that your place will only be confirmed when payment has been received (please tick as required)

I wish to pay the fee of .....

☐ I enclose a cheque payable to "The Institute of Clinical Research"

OR

☐ I wish to pay by

☐ VISA ☐ MASTERCARD ☐ DELTA ☐ EUROCARD

Card Number ☐☐☐☐ ☐☐☐☐ ☐☐☐☐ ☐☐☐☐

Start Date ☐☐ ☐☐ **Expiry Date** ☐☐ ☐☐

Name (as it appears on the card) .....

Signature of card holder .....

OR

☐ Please invoice my company using Purchase Order Number ☐☐☐☐☐☐☐☐ Invoices can only be raised when a PO no. is provided

Correspondence Address

Address: .....

Postcode: ..... **Country:** .....