

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.

## Venue - CIM Moor Hall Conference Centre, The Moor, Cookham SL6 9QH, Cookham

## Course Outline

- Overview of MD
classifications EU and US
- When do you need to do a clinical trial?
- Update on the MDR and the impact it will have on clinical investigations
- Review of other relevant
directives and guidance
including ISO 14155
- Designing your clinical investigation
- Key Points for Project

Managing a clinical investigation

- Key Points for Monitoring a clinical investigation
- Medical Devices and Safety

Monitoring in Clinical
Investigations

- What about drug device combinations?

Who would benefit
Anyone looking for an overview of conductive clinical
investigations with medical devices.

## Course Fees

| Guest | $£ 550.00$ |
| :--- | ---: |
| ICR Member | $£ 450.00$ |
| ICR Member Academic | $£ 350.00$ |

## Victoria Cavendish

A highly skilled clinical affairs specialist who has worked in clinical research for over 15 years, including 11 years in the medical device arena. Her experience includes work within academic, manufacturing, consulting and CRO environments. Dr. Cavendish has extensive knowledge of medical device development and clinical affairs in regulatory, post-market surveillance, marketing and registry environments within Europe. She has proven ability in the set-up, study document preparation, incountry submissions, vendor negotiations and leadership of global and European projects to ensure they are conducted in accordance with the highest ethical, clinical and scientific standards. Victoria has held increasingly senior positions at various medical device companies including DePuy Synthes, Novartis, and Reckitt Benckiser.

Her therapeutic areas of expertise include orthopedics, software, sexual health, Over The Counter (OTC) and combination products. For the last 18 months, Dr. Cavendish has owned a medical device consultancy, Orca Solutions Ltd, providing strategic direction and management of the global clinical development portfolio for a UK based medical device software manufacturer, alongside providing training courses for medical device professionals.Victoria also has experience of the implementation and management of a quality system to obtain ISO 13485 certification and is able to conduct internal and site audits. Dr. Cavendish obtained a BSc(Hons) in Immunology at theUniversity of Glasgow, an MSc in Diabetes at the University of Manchester, a PhD in Respiratory Medicine at the University of Bristol, and holds a Post-graduate Diploma in Public Health.


## Course Title: M17 Clinical Investigations for Medical Devices - Foundation

Date: 05 June 2023

## Name:

Company / Hospital: $\qquad$
Position / Job Title: $\qquad$

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

## What are you hoping to get out of the day?

$\qquad$
$\qquad$
$\qquad$

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

## Dietary Requirements

$\qquad$
$\qquad$
$\qquad$

* 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.

Address: CIM Moor Hall Conference Centre, The Moor, Cookham SL6 9QH, CIM Moor Hall Conference Centre, The Moor, Cookham, Berkshire, United Kingdom, SL6 9QH

## Local Taxi Companies

## Accommodation

The ICR does not specifically recommend any accommodation - however the following are within easy travelling distance of the training venue
contact Lydia.Macdonald@cim.co.uk

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.

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## Registration Form

## Please photocopy this form for further registrations



## 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 ofI enclose a cheque payable to "The Institute of Clinical Research"
OR
$\square$ I wish to pay byVISAMASTERCARDDELTAEUROCARD

Card Number


Start Date


Expiry Date


Name (as it appears on the card) $\qquad$
Signature of card holder
ORPlease invoice my company using Purchase Order Number $\square$ Invoices can only be raised when a PO no. is provided

Correspondence Address

Address:

