ICR EVENTS

MONDAY 29<sup>TH</sup> OCTOBER 2012



Health Research Authority





# Ethics & GCP Forum

### **Overview**

The ICR Ethics and GCP Forum provides an excellent opportunity to:

- Keep up to date with current ethical and clinical research issues
- Learn about new industry and regulatory initiatives
- Hear key industry figures speak about their specialist fields
- Network with fellow clinical research professionals

#### **Learning Outcomes**

The learning outcomes for delegates attending this meeting are to:

- Learn about the latest information from the Health Research Authority
- Understand the challenges of emergency research
- Discover the best practices in order to obtain ethical approval
- Learn about the proposed EU Clinical Trials Regulation and what it will mean to UK research
- Understand the latest Phase I guidance
- Be aware of recent regulatory changes in clinical research

# Organisers

Ethics Forum Chair: Joan Perou GCP Forum Steering Committee: Nicky Dodsworth, Janice Hedgecock, Stuart Harris, Julia DeCesare UCL Joint Research Office: Helen Cadiou

Enquiries (ICR): +44 (0)845 521 0056

## Venue

UCL Institute of Neurology in Queens Square is just a short walk from Russell Square tube Stations

http://www.ucl.ac.uk/ion/contact

**Ethics & GCP Forum** 

Adapting to structural and regulatory changes

Venue: University College London, Institute of Neurology Lecture Theatre, Lower Ground Floor, 33 Queen Square, WC1N 3BG

## Programme

10.00	Welcome
10.10	<b>Health Research Authority</b> An update from the Health Research Authority Joan Kirkbride, Operations Director, National Research Ethics Service
11:10	<b>Proposed EU Clinical Trials Regulation – the MHRA perspective</b> Martyn Ward, MHRA Clinical Trials Unit
<b>11.5</b> 0	Ethics Applications – Best Practices Hugh Davies, Research Ethics Advisor, Health Research Authority
<b>12.20</b>	Lunch
13:20	The Code of Practice for the Pharmaceutical Industry 2012 Edition tbc, Association of the British Pharmaceutical Industry
13:50	<b>Phase I Guidelines</b> An update on the latest ABPI Guidelines Dr Louise Leong, Head of Research & Development, Association of the British Pharmaceutical Industry
14:20	Tea Break
14:50	Key GCP Updates A summary of changes in GCP-related regulations and guidelines since the last Ethics/GCP Forum meeting GCP Steering Committee
15:10	<b>IMP Management at Trial Sites</b> A summary of the ICR IMP Management Survey followed by a presentation from the National Pharmacy Clinical Trials Network Janice Hedgecock, Director, Greatspur Clinical Development Ltd & Claire Khalil, Lead Pharmacist, Clinical Trials and Audit, Great Ormond Street Hospital
16.00	Close
	<b>Registration</b> Log into the members' area of the ICR website to book: Fee: ICR Members £30 Non Members £40 <u>http://www.icr-global.org/community/calendar-of-events/</u>
	Places are limited to 150 so please book early!

The programme is subject to change. Please check the ICR website for the latest version.