NIHR | Clinical Research Network East of England

Informed Consent in Research

Available Dates

15th September 1:00-1:30pm & 13th October 2-4pm

1st November 1:30-2:00pm & 29th November 2-4pm

Up to 10 spaces per session

Who is this training for?

- Research delivery staff who have a patient-facing role; have been in practice for 6 months or less; who are working on or with NIHR Portfolio studies and have completed ICH-GCP training.
- This 4 week training package is designed in a modular format and combines interactive content, tasks to complete, reflection on practice and opportunities to experience receiving consent.
- You must attend a pre and post course zoom session to complete the training.

Learning objectives:

- Recognise the ethical and legislative frameworks that underpin research consent
- Understand the principles of consent
- Examine the research consent process
- Gain an insight into the added protection required for vulnerable group
- Build confidence in participating in the research consent
- Demonstrate high quality practices to support the process

To apply, visit:

Informed Consent Booking Form