# OHRPP Post-Its

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Office of Human Research Protection Programme (OHRPP) Post–Its: Bringing you the latest updates on research policies, educational resources and event information

## Control Research Data Access & Monitoring

Include in Section T3 (ECOS): study data access control & monitoring plan. Maintain access logs & assign a person-in-charge for password management.

Learn more about what to include in Section T3 and other application form sections in our ECOS Guides.

SCAN HERE

REMINDER !

## Getting Consent Right For Adults with Capacity to Consent

**Before** any study activity begins, check that:

- ✓ The IRB-Approved ICF was used for the consent process.
- ✓ The **Informed Consent Form (ICF)** had been **personally signed & dated** (by the **participant** in **a language they understand**, by a **witness (if required)** and by the **person obtaining consent**).
- ✓ ICF is **complete and accurate**, and the **participant had received a signed copy**.

*Consent is not a formality - it protects participants' autonomy and their right to make an informed decision.*

Source: Proper Conduct of Research (PCR) SOP 501-C01, Informed Consent Form and Process

REGISTER NOW !

## Pick up Tools for Audits, Monitoring & Inspections

Sign up for PCR004 – Now incorporating ICH GCP E6(R3) & Tissue Bank quality tools.

NHG Staff: register on LENS. See website for details.

## DSRB Green Lane

Accelerating Clinical Trial reviews through collaborative excellence.
Find out more here.

## SARAH welcomes our friends from NUHS & CDA!

Supportive Automated Response Agent for Human research

Ask SARAH

Our AI-powered chatbot for human research conduct, is now accessible to our partners from NUHS and the Communicable Disease Agency (CDA).

**For feedback & more information**

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NHG Office of Human Research Protection Programme (OHRPP)

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