

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

RESEARCH TRAINING & CONDUCT

ICH E6 (R3) Good Clinical Practice (GCP) Guideline has effected on 1 January 2026

🎉 Well Done to all who have completed ICH E6 (R3) GCP Training! 🎉

To keep up your knowledge on ICH E6 (R3) GCP Guideline, refer to these resources on HSA website:

- [HSA-SCRI Public Webinar on the ICH E6 \(R3\) GCP guideline \(28 Feb 2025\)](#)
- [HSA IOCTB Learn webpage \(Clinical Trials Learning and Educational Hub\)](#)
- [Regulatory Guidances](#)

Proper Conduct of Research (PCR) SOP Reminder

Informed Consent Form for Non-English-speaking subjects

Researchers should provide written consent forms in the subject's preferred language. If a **full translation** is unavailable, use the DSRB-approved English consent form alongside a **Short Consent Form (SCF)** in the language understood by the subject.

For cross-cluster studies under IRB mutual recognition agreement, always follow your Institutional requirements, regardless of which IRB is conducting the review.

Reference: [PCR SOP 501-C01: Informed Consent Form and Process](#)

Resource: SCF templates in local languages available on [NHG Health research website](#).

ECOS HELPDESK

Can't see Queries from Research Institution in ECOS?

Your role might need an update!

Quick Fix: Ask your PI or Site PI to change your access to "Study Team Member" in the **CRMS** User Authorisation List.

Refreshed ECOS Quick Start User Guides & FAQ!

Updated [ECOS User Guides](#) and [ECOS FAQs](#) now available on ECOS website.

- ❖ Refer to the Quick Start Guide to navigate ECOS, submit IRB applications and respond to queries!

NEW YEAR, FRESH OPPORTUNITIES

Join OHRPP!

Shape Research Ethics. Safeguard Participants.

Join the NHG Health research ethics community and directly impact participant safety through policy development and regulatory guidance. Help advance our national healthcare research whilst ensuring ethical standards protect every volunteer.

Multiple **roles** available: **IRB Specialist**, **IRB Analyst** positions. Full-time & part-time options. Interested?

Contact Ms Ng Hwee Hian, Director of OHRPP: hwee.hian.ng@nhghealth.com.sg

RESOURCES

Supportive Automated Response Agent for Human Research

Meet SARAH
YOUR NEW HUMAN RESEARCH CONDUCT ASSISTANT
Research Rules and Procedures Made Simple
Skip the Wait, Get Research Guidance [Now!](#)

[Chicken Soup For The Busy Coordinator](#)
Bite-size resource for Clinical Research Coordinators

[Proper Conduct of Research \(PCR\) Courses](#)
Learn how to conduct your research properly

[Join OHRPP on Viva Engage](#)
Get updates on research guidelines, best practices, OHRPP news & happenings!

[Responsible Conduct of Research \(RCR\)](#)
Doing Science Right, find out more about RCR