Well Done To All Who Have Completed ICH E6 (R3) GCP Training!

The ICH E6 (R3) GCP guideline will take effect from **01 January 2026** in Singapore.

Enhanced GCP Concepts

What Should I Consider Before Ethics Submission?

- **What phase is my trial and how complex is the design?**
- **What level of risk do participants face?**
- **How critical is each piece of data I'm collecting (essential vs. "good to have")?**

FIT FOR PURPOSE

What Does It Mean?

Appropriate tools, processes, and systems matched to trial objectives and risks.

Effort matches the risks to participants and importance of the data being collected.

QUALITY BY DESIGN (QbD)

"Building quality into clinical trials from the start"

Proportionate, Risk-based Approaches

"Align effort with risk level and data criticality"

IMPORTANT **[For NHG Health Staff Only]**

Haven't Completed Your ICH E6 (R3) GCP Training?

From 01 Oct 2025 (Wed) NHG Health Principal Investigators (PI), Site PIs and Co-Investigators WITHOUT an updated ICH E6 (R3) GCP training record will NOT be able to submit new/in-flight IRB applications or study amendments for clinical trials regulated by Health Science Authority on ECOS.

ACT NOW

Click here for Step-by-Step guide on how to update your ICH GCP Training.

If you have any questions regarding ICH E6 (R3) GCP Training, please contact your Institutional Minimum Training Secretariat.

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