IS YOUR RESEARCH READY FOR ICH E6 (R3) GCP?

ICH E6 (R3) GCP guideline will take effect from **01 Jan 2026** in Singapore. All investigators and study teams in HSA-regulated trials should conduct a **gap analysis** comparing current practices with new GCP requirements to ensure participant safety and data reliability.

STEP 1 : LEARN THE CHANGES

- **Understand** the new requirements & principles.
- **Complete ICH E6 (R3) GCP training** & **upload the completion report to ECOS** as soon as possible.

Refer to PCR SOPs summary of changes to understand how NHG Health is adopting the ICH E6 (R3) GCP requirements.

STEP 2 : ASSESS YOUR CURRENT STATE

- Review your protocols & procedures against the Guideline and identify gaps.

STEP 3 : TAKE ACTION

- **Prioritize** critical gaps (e.g. participant safety & data reliability)
- **Update** your processes and documents
- **Train** your team
- **Maintain** records of the gap analysis & implemented changes.

Complete Your CITI GCP Course Today!

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

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NHG Health Office of Human Research Protection Programme (OHRPP)