

## 1. เข้า google พิมพ์ good clinical Practice training คลิกตาม

The screenshot shows a Google search results page for the query "good clinical Practice training". The search bar at the top contains the text "good clinical Practice training". Below the search bar, there are several search filters and a "ลงชื่อเข้าระบบ" (Sign in) button. The search results are displayed in a list format. The first result is "บทความทางตติศึกษาเกี่ยวกับ good clinical practice training" (Articles about good clinical practice training). The second result is "Good Clinical Practice" from <https://gcp.nihtraining.com/>. The third result is "GCP - CITI - Collaborative Institutional Training Initiative" from <https://www.citiprogram.org/index.cfm?pageID=90>. The fourth result is "ICH GCP (Good Clinical Practice) Training Course" from [www.onlinegcp.com/](http://www.onlinegcp.com/). A black arrow points from the text "คลิกตาม" in the section header to the "Good Clinical Practice" result.

## 2. กดคลิก Click here ด้านล่างสุด

The screenshot shows the homepage of the Good Clinical Practice (GCP) training website. The URL in the browser is <https://gcp.nihtraining.com>. At the top, there is a green notification box that says "You are now logged out." Below this, the word "Welcome" is displayed in a large, bold font. The main content area contains a paragraph of text: "The Good Clinical Practice (GCP) course is designed to prepare research staff in the conduct of clinical trials with human participants. The 12 modules included in the course are based on ICH GCP Principles and the Code of Federal Regulations (CFR) for clinical research trials in the U.S. The course is self-paced and takes approximately six hours to complete." Below this paragraph, there is a link: "To preview the new enhanced features, please [click here](#)." Another paragraph follows: "To begin, please *sign in* using the link to the right if you have already created an account. If you do not have an account, click [here](#) to register." On the right side of the page, there is a "Login" form with input fields for "email" and "password", a "Forgot password?" link, and a "Sign in" button. Below the "Sign in" button, there is a link: "Need an account?". A black arrow points from the text "กดคลิก Click here ด้านล่างสุด" in the section header to the "click here" link in the main content area.

### 3. กรอกข้อมูลส่วนตัว

Good Clinical Practice

หน้าเว็บนี้เป็น **อังกฤษ** คุณต้องการแปลหรือไม่ **แปล** **ไม่** **ไม่ต้องแปลอังกฤษ** **ตัวเลือก**

## Create an Account

Please enter your first and last name as you would like it to appear on your certificate. You will **NOT** be able to change your name later.

**First Name\***

**Last Name\***

**Node/University Name**

start | srithanyaintranet - W... | Doc1.docx - Microsof... | Good Clinical Practice ... | EN | 15:31

### 4. เลือกหัวข้อ RS-Research Staf

Good Clinical Practice

หน้าเว็บนี้เป็น **อังกฤษ** คุณต้องการแปลหรือไม่ **แปล** **ไม่** **ไม่ต้องแปลอังกฤษ** **ตัวเลือก**

**Protocol Role(s)\***

- IN - Investigator
- QA - QA Staff
- RS - Research Staff**
- RG - Regulatory Staff
- RX - Research Superv. Staff
- DM - Data Management Staff
- TH - Therapist
- DX - Data Management Staff
- TX - Therapy Supervisor
- BO - Business Operations Staff
- MD - Medical Staff
- RO - Research Staff: Other

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## 5.กรอกข้อมูลให้ครบและกด submit

The screenshot shows a web browser window with the URL <https://gcp.nihtraining.com/register>. The page contains a registration form with the following fields:  IA - Independent Assessor, Email\*, Password\*, Confirm Password\*, Phone, and Study Site Name. A blue 'Submit' button is located at the bottom of the form. The browser's taskbar at the bottom shows the 'start' button and several open applications, including 'srithanyaintranet - W...', 'Doc1.docx - Microsof...', and 'Good Clinical Practice ...'. The system tray shows the time as 15:33.

## 6. เลือกทำหัวข้อ Institutional Review Boards เป็นต้นไป

The screenshot shows the 'Overview' page on the GCP training website. The URL is <https://gcp.nihtraining.com/overview>. The page features a section titled 'Modules' with the following text: 'The 12 modules may be completed one at a time or in one sitting as users may login and logout, as needed. Users have the option to complete module quizzes after reviewing the instructional material, or choose to complete the module quizzes only, particularly for returning users. Accessing quizzes is easy. Select the **Take the Quiz** button in each module or choose one of the links on the **My Progress** page.'

Below the text is a grid of 12 module buttons, each with an icon and a title:

- Introduction
- Institutional Review Boards
- Informed Consent
- Confidentiality & Privacy
- Participant Safety & Adverse Events
- Quality Assurance
- The Research Protocol
- Documentation & Record-Keeping
- Research Misconduct
- Roles & Responsibilities
- Recruitment & Retention
- Investigational New Drugs

The browser's taskbar at the bottom shows the 'start' button and several open applications, including 'srithanyaintranet - W...', 'Doc1.docx - Microsof...', and 'Good Clinical Practice ...'. The system tray shows the time as 15:38.

Good Clinical Practice x

← → ↻ <https://gcp.nihtraining.com/modules/1> ☆ ☰

หน้าเว็บนี้เป็น อังกฤษ คุณต้องการแปลหรือไม่ แปล ไม่ ไม่ต้องแปลอังกฤษ ตัวเลือก

# Institutional Review Boards

Take the Quiz

## Contents

- Part 1: What is an Institutional Review Board (IRB)?
- Part 2: Purpose of an IRB
- Part 3: Membership of an IRB
- Part 4: Responsibilities of an IRB
- Part 5: Criteria for IRB Approval of Research
- Part 6: Expedited Review
- Part 7: Investigators' Responsibilities to the IRB
- Part 8: IRBs and Cooperative Research
- Summary of Key Points

[Return to Previous Module](#)

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8. ทำไปเรื่อยๆ จบครบทุก Part

Good Clinical Practice x

← → ↻ <https://gcp.nihtraining.com/modules/1/1> ☆ ☰

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# Institutional Review Boards

Take the Quiz

## Part 1: What is an Institutional Review Board?

An Institutional Review Board (IRB) is an independent committee established to protect the rights and welfare of human research participants. Under Title 45 Part 46 of the Code of Federal Regulations (**45 CFR 46**), any research that is federally funded must be reviewed and approved by an IRB.

Any clinical investigation involving a product regulated by the U.S. Food and Drug Administration (FDA) must also be reviewed and approved by an IRB (**21 CFR 56**). Individual institutions or sponsors may require that all research, no matter how it is funded, be reviewed and approved by an IRB.

An IRB has specific authority over the conduct of research under its jurisdiction. No clinical study may begin enrolling participants until it has received IRB approval. The IRB has the authority to:



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## 9. ตัวอย่างคำถาม

The screenshot shows a web browser window with the URL <https://gcp.nihtraining.com/quiz/1>. The page header includes the text "Good Clinical Practice" and a user email "yuyuu30@hotmail.com" with a "logout" button. Below the header are four navigation tabs: "Overview", "My Progress", "Resources", and "Certification". The main content area is titled "Quiz: Institutional Review Boards" and includes a "Return to Module" button. The quiz consists of two questions:

- 1 The primary purpose of the Institutional Review Board (IRB) is to:
  - A. Investigate allegations of research misconduct.
  - B. Administer compensation for participation to study volunteers.
  - C. Protect the rights and welfare of research participants.
  - D. All of the above.
- 2 The three key principles that serve as the criteria for IRB approval of research can be found in which of the following documents:
  - A. Belmont Report
  - B. Code of Federal Regulations (CFR)
  - C. ICH Good Clinical Practice guidelines (ICH GCPs)

## 10. หลังจากทำทุก Part ครบ ด้านขวา หัวข้อ STATUS จะขึ้น Passed ปรีนมาส่งได้เลย

The screenshot shows a web browser window with the URL [https://gcp.nihtraining.com/my\\_progress](https://gcp.nihtraining.com/my_progress). The page displays a table of quiz results for various topics. The table has two columns: the topic name and the status/percentage. The "The Research Protocol" row is highlighted in blue.

Informed Consent	80%
Confidentiality & Privacy	Retake Quiz
Participant Safety & Adverse Events	Retake Quiz
Quality Assurance	Retake Quiz
The Research Protocol	Take the Quiz
Documentation & Record-Keeping	Take the Quiz
Research Misconduct	Take the Quiz
Roles & Responsibilities	Take the Quiz
Recruitment & Retention	Take the Quiz
Investigational New Drugs	Take the Quiz
Final Grade	___%

If you have completed certification and need to renew, please select the **Certification** tab and choose the button for