GOOD CLINICAL PRACTICES TRAINING OVERVIEW

The Good Clinical Practice (GCP) online training consists of 12 modules. Each module discusses a specific aspect of GCP guidelines.

Throughout the course, the instructional material includes ICH GCP guidelines as well as the Code of Federal Regulations (CFR) for clinical research trials in the U.S.

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Welcome

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.

To preview the new enhanced features, please **click here**.

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.

Login

email

password

Forgot password?

Sign in

Need an account?
Sign up here!

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National Drug Abuse Treatment Clinical Trials Network

Good Clinical Practice

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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*

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Study Role/Title*

Organization*

Location* United States

If Other Please Specify

NIH-Affiliated*

Yes No

Email*

Password*

Confirm Password*

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Study Role/Title*

Researcher

Organization*

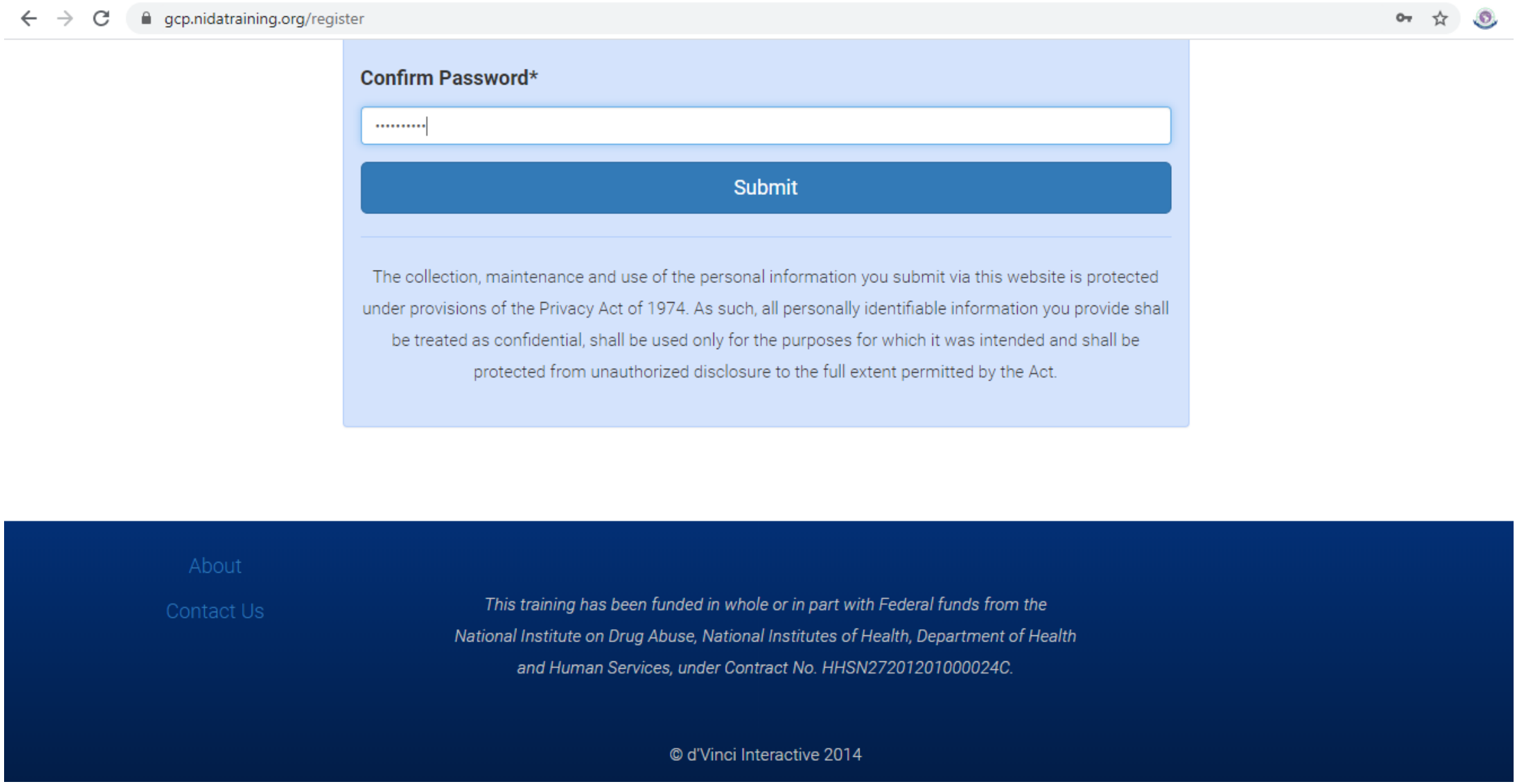
Chiang Mai University

Location* Thailand

If Other Please Specify

NIH-Affiliated*

Yes No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Confirm Password*
Submit
The collection, maintenance and use of the personal information you submit via this website is protected under provisions of the Privacy Act of 1974. As such, all personally identifiable information you provide shall be treated as confidential, shall be used only for the purposes for which it was intended and shall be protected from unauthorized disclosure to the full extent permitted by the Act.
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This training has been funded in whole or in part with Federal funds from the National Institute on Drug Abuse, National Institutes of Health, Department of Health and Human Services, under Contract No. HHSN272012010000024C.
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Welcome!

The **Good Clinical Practice** (GCP) online training consists of 12 modules. Each module discusses a specific aspect of GCP guidelines. Throughout the course, the instructional material includes ICH GCP guidelines as well as the Code of Federal Regulations (CFR) for clinical research trials in the U.S.

Every three years, NIH requires completion of the course to remain current with regulations, standards, and guidelines.

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Quizzes and Certification

To obtain a certificate, users must complete all module quizzes successfully. The quiz questions are based on the information provided directly in this course, including extracted summaries and definitions. However, the questions are not based on information provided in the externally linked documents or websites. Module quizzes are randomized from a bank of questions each time a quiz is taken. Therefore, it is highly recommended that users review the instructional material prior to completing the quizzes.

Users are required to complete a quiz following each module, except for the Introduction module. To receive a certificate, all quizzes must be completed with at least 80% accuracy. Upon successful completion of all quizzes, the user will be given access to the Certificate of Completion.


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Certificate of Completion
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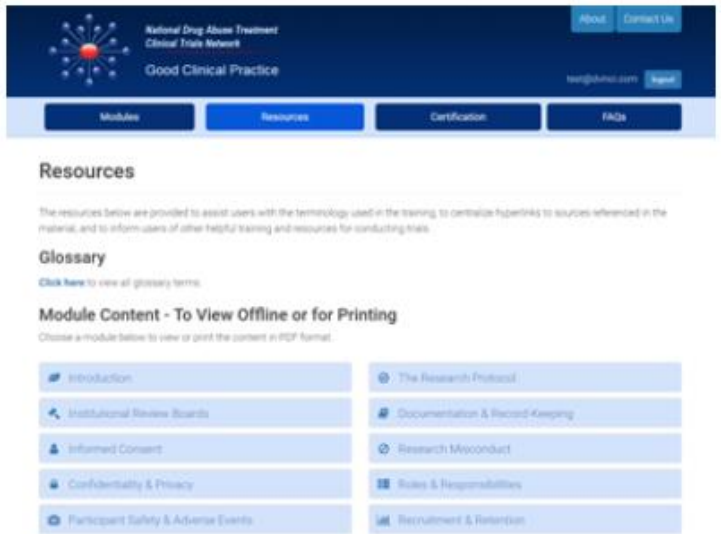
Resources

Please use the **Resources** tab to review terminology and definitions, course links, and additional resources and websites.

The course is designed to be comprehensive for users to review the content of required information in conjunction with an examination of other source documents and websites pertinent to Good Clinical Practice and research regulations.



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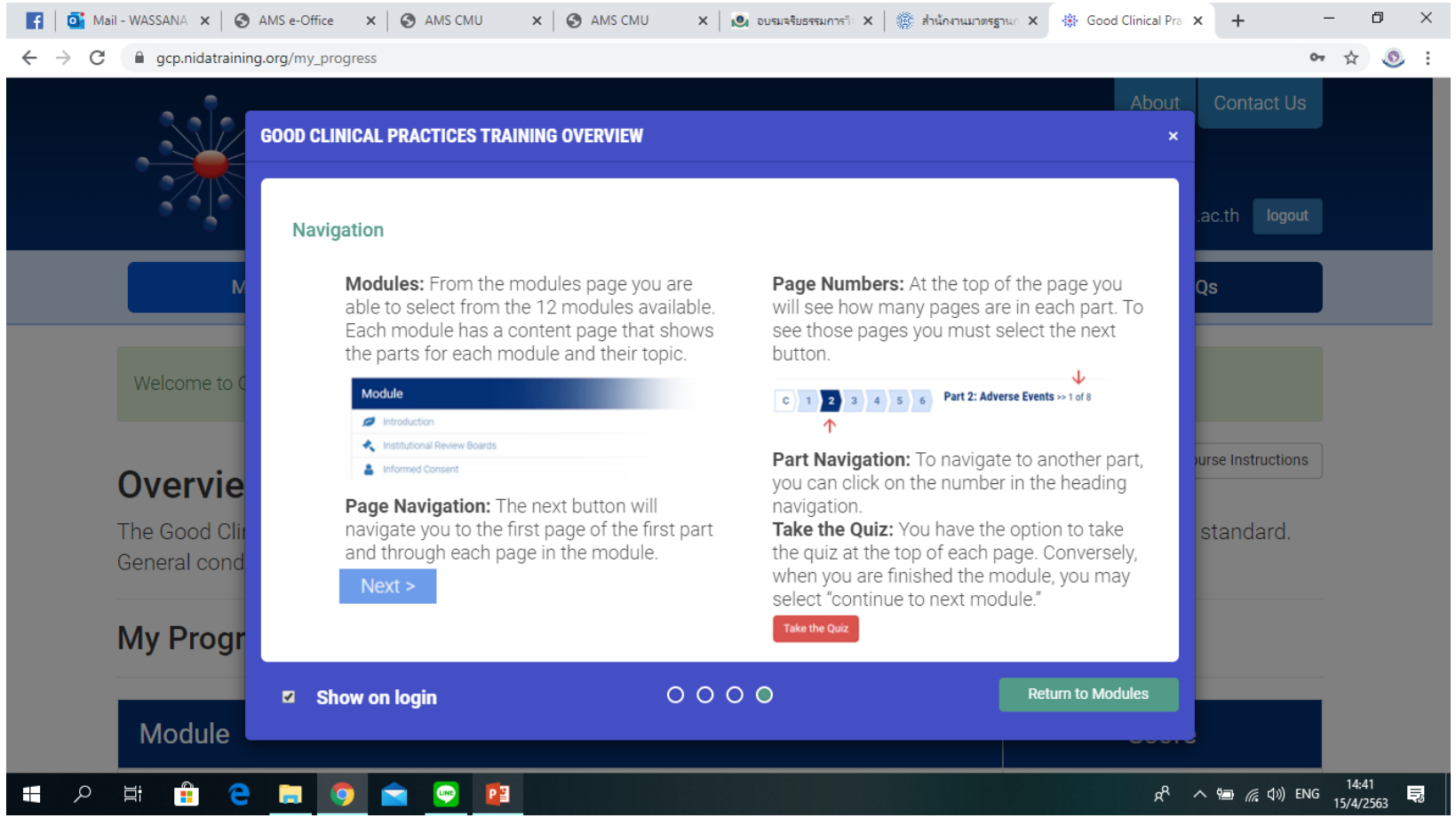
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Navigation
Modules: From the modules page you are able to select from the 12 modules available. Each module has a content page that shows the parts for each module and their topic.
Module
Introduction
Institutional Review Boards
Informed Consent
Page Navigation: The next button will navigate you to the first page of the first part and through each page in the module.
Next >
Page Numbers: At the top of the page you will see how many pages are in each part. To see those pages you must select the next button.
C 1 2 3 4 5 6 Part 2: Adverse Events >> 1 of 8
Part Navigation: To navigate to another part, you can click on the number in the heading navigation.
Take the Quiz: You have the option to take the quiz at the top of each page. Conversely, when you are finished the module, you may select "continue to next module."
Take the Quiz
Show on login
Return to Modules
Module
ENG
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Module	Score
Introduction	N/A
Institutional Review Boards	Take the Quiz
Informed Consent	Take the Quiz
Confidentiality & Privacy	Take the Quiz
Participant Safety & Adverse Events	Take the Quiz
Quality Assurance	Take the 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	___%