Mandatory Training for FDA Regulated Researchers
(updated 6-30-11)

To review the course content

1. Put cpd.uc.edu in your Internet browser
2. Click Compliance Training or Competency Testing
3. Look in the blue section in the left side of the screen and click either
   Member Login (if you already have an account in CPD) OR
   New User? Register! (if you are new to CPD)
   Follow the instructions to log in to CPD
4. On the Compliance Training Catalog page
   Look under Compliance Courses / **Compliance Training**.
   Click **Human Subjects Research Compliance Training**.
5. On the Compliance Training page, look for the course titles that begin with HRP (dated May 2011).
   Note that the course titles are alphabetical. They may be completed in any order.
   Click the course title to access the slides and the quiz.
6. On the page for the selected course, look below the goals for the course.
7. Click on the link to view the course materials. This will open a new window where the course content may be viewed and printed. Closing the window will reveal the page for the selected course.
8. Click the Back button to return to the Compliance Training page and select another course.

To take the course quiz

1. Enter CPD as above and follow all steps to go to the page for the selected course.
   Note that viewing the course materials online is not required before taking the quiz. However, it is strongly recommended.
2. The very last statement on the page has a link ("click here") to take the quiz for the selected course.
   Each time the quiz is opened, the order of the questions is shuffled. The questions themselves and the possible answers remain the same.
3. After the quiz has been completed successfully (at least 80% correct), a link will be provided to print the Certificate of Completion for the course.
4. Even after successfully completing the quiz, it is possible to go back to the page for the selected course, go to the last statement on the page, and re-take the quiz.
5. Click the Back button to return to the Compliance Training page and select another course.
   Note that a course's Certificate of Completion can be printed from the Compliance Training page also.

Center for Continuous Professional Development
for Researchers and Healthcare Professionals

Compliance Training Catalog

Use this page to view the categories of training, testing, or news and information available or to make a selection. Also use this page to view assigned requirements. Use the following methods to locate specific information.

- Compliance Home
- Logout
- My Account
- Transcript
- Administration
- Technical Support
- Feedback
- Contact Us

Continuing Education Program

Find Course by Keyword

If you are looking for a specific topic, you may search for it below:

Search Criteria

Keywords: Match ANY Search

Category: Any Category

Find Course by Category

If you are looking for topics under a category, click on it below:

Compliance Courses

Compliance Training

Animal Acquisitions
Animal Research
Blood-Borne Pathogens Training
Biosafety Compliance Training
COM Industry Relationships
Grant Submissions - SRS
HIPAA Compliance Training
Human Subjects Research Compliance Training
IACUC Training & Verification
LAMS CE Class
LAMS SOPs & Forms
Radiation Safety Training
UC Information Security
UCP HIPAA Compliance Training - Not On-line
UCP HIPAA Compliance Training - OnLine
UCP Infection Control
UCP Other Compliance Training

Competency Testing

Biomed Research HRP Knowledge - expired 10/13/08
Clinical Research Orientation - IRB Roles and Responsibilities
Clinical Research Orientation - Session 1
Clinical Research Orientation - Session 2
Clinical Research Orientation - Session 3
Clinical Research Orientation - Submissions to the IRB
Grant Submissions Competency Exam
Human Subjects Research Compliance Quizzes
Key Instit Officials Record - expired 10/13/08

University of Cincinnati

Center for Continuous Professional Development
for Researchers and Healthcare Professionals

Compliance Training
Human Subjects Research Compliance Training

(Page 1 of 2) « < Previous | Next > »

Having trouble finding the course you are looking for?
Search in "Human Subjects Research Compliance Training" -or- Search ALL Categories

- Compliance Home
- Logout
- My Account
- Transcript
- Administration
- Technical Support
- Feedback
- Contact Us

Continuing Education Program

Course Title	Start Date	End Date
Coercion in Recruiting Human Subjects		
Conflict of Interest		
Ethical Considerations in Investigator Initiated Study Design		
HRP - Adverse and Other Events in Human Research May 2011	06/20/2011	12/31/2020
HRP - Case Report Forms Source Records Data Entry May 2011	06/20/2011	12/31/2011
HRP - Clin Research Overview May 2011  Course Completed: 06/30/2011 Print Certificate	06/20/2011	12/31/2020
HRP - Device Accountability in Human Research May 2011	06/20/2011	12/31/2020
HRP - Drug Accountability in Human Research May 2011	06/20/2011	12/31/2020
HRP - How to Avoid Deviations and Violations May 2011	06/20/2011	12/31/2011
HRP - Informed Consent for Human Research May 2011	06/20/2011	12/31/2020
HRP - Resp and Oblig of Investigators Part 1 May 2011	06/20/2011	12/31/2020
HRP - Resp and Oblig of Investigators Part 2 May 2011	06/20/2011	12/31/2020
HRP - Sponsor Resp and Oblig of Research Sponsor-Investigators May 2011	06/20/2011	12/31/2020
HRP - Submissions and Reports per Fed Authority May 2011	06/20/2011	12/31/2020
Human Subjects Research 1 (meets NIH requirement)		

(Page 1 of 2) « < Previous | Next > »

Center for Continuous Professional Development
for Researchers and Healthcare Professionals

HRP - Clin Research Overview May 2011

This is the initial course in a series of 11 modules for human subjects researchers both at UC and at other institutions who receive oversight from the UC IRB.

The following is a description of the goals for this Compliance Training Course

Compliance Home
Logout
My Account
Transcript
Administration
Technical Support
Feedback
Contact Us

Continuing Education Program

- Define human subjects research and list roles that researchers have.
- Describe the types of relationships Investigators have with Sponsors.
- Describe the role of the UC Sponsored Research Office.
- Provide an overview of good clinical practices in research.
- Describe AAHRPP and its Accreditation.
- List the follow-in modules in this HRP series.

Please Click on the link below to view the course materials

- Clin Res Oview May 2011.pdf IND/IDE Assistance Program, UC

You have successfully completed this course.
If you would like to take the test again click on the link below.

After you have reviewed the content, click here to complete the on-line test.

Back