

The Past, Present, and Future of FDA Human Drug Regulation

Regulation of prescription drugs in the United States is a relatively new concept that has undergone many revisions since the early 1900s. In this skill module, you will learn about the progression to the current regulations.

To complete this skill module visit this website:

<http://www.fda.gov/Training/ForHealthProfessionals/default.htm>.

Select “The Past, Present, and Future of FDA Human Drug Regulation” from under the course list.

CDERLearn

Welcome to CDERLearn, the web page for educational tutorials offered by the Center for Drug Evaluation and Research. CDER's primary mission is to make certain that safe and effective drugs are available to the American people. There is, however, a strategic initiative to inform and educate people about the safe use of medicine, the drug regulatory process, the vital role health care professionals play to assist FDA in fulfilling its duties, and many other important issues. Online training is one way to share FDA expertise with many more people than face-to-face classroom sessions would allow, and we will offer additional CDER courses in the future.

Course List

- [The Past, Present, and Future of FDA Human Drug Regulation](#)
An updated version of "Drug Review and Related Activities in the United States," with continuing education (CE) credit for pharmacists, physicians, and nurses. The program gives health care professionals, industry, consumers, and all other interested participants an overview of the human drug regulatory process in the United States. CE Credit available through June 28, 2013. Course time: approximately 90 minutes

On the next screen select “Start the activity.”

The Past, Present, and Future of FDA Human Drug Regulation

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Registration: You can participate in this training without taking the post test. **However, registration information is required to enter this course and use the bookmarking feature.** To receive continuing education credit and/or certificate, you must take a post test and complete an evaluation.

- Activity Description, and Learning Objectives
- Accreditation
- Disclosure
- CE Statements of Credit/Certificate of Completion (non-CE)
- Biography
- Activity Time
- Replacement of CE Statement of Credit/Certificate of Completion (non-CE)
- Free Registration
- Privacy Policy

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You will then need to create a login and password.

Once you create your login, click “begin activity”.

Activity Overview and Information

Welcome Genevieve Allen!

Thank you for participating in the Food and Drug Administration, Center for Drug Evaluation and Research activity on the Past, Present, and Future of FDA Human Drug Regulation.

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