

From: [OC GCP Questions](#)
To: [REDACTED]
Subject: GCP Training
Date: Thursday, November 20, 2014 3:01:23 PM

Good afternoon --

Both the Center for Drug Evaluation (CDER) and the Center for Devices and Radiological Health (CDRH) provide online training relevant to clinical trials on their respective websites. These training programs are accessible at CDER Learn <http://www.fda.gov/Training/ForHealthProfessionals/default.htm> and CDRH (Center for Devices) Learn <http://www.fda.gov/Training/CDRHLearn/default.htm> respectively.

The expectation is that investigators, sub-investigators, and study staff will be knowledgeable about good clinical practice, including human subject protection, data integrity, recordkeeping, etc. Note that sponsors have discretion in determining what qualifications will be needed to conduct a study and may identify in the protocol a required frequency of GCP training, in which case the investigator and sub-investigators would be expected to meet that frequency of training in order to comply with the sponsors requirements. Every effort should be made to train the investigational sites.

While we cannot endorse/recommend non-government training entities, you may also find the courses provided by a number of the professional societies to be useful. These include: the Association of Clinical Research Professionals (ACRP) (<http://www.acrpnet.org/>), the Society of Clinical Research Associates (SoCRA) (<http://www.socra.org/>) the Regulatory Affairs Professionals Society (RAPS) (<http://www.raps.org/personifyebusiness/>), the Drug Information Association (DIA) (<http://www.diahome.org/DIAHome/Home.aspx>), and the Society of Quality Assurance (SQA) (<http://www.sqa.org/>). Several of these associations also have certification programs for clinical trial staff.

I hope this information is helpful. Please contact us again at gcp.questions@fda.hhs.gov should you have additional questions.

Kind regards,

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Office of Good Clinical Practice
Office of the Commissioner, FDA

This communication does not constitute a written advisory opinion under 21 CFR 10.85, but rather is an informal communication under 21 CFR 10.85(k) which represents the best judgment of the employee providing it. This information does not necessarily represent the formal position of FDA, and does not bind or otherwise obligate or commit the agency to the views expressed.

From: [redacted]
Sent: Wednesday, November 19, 2014 2:55 PM
To: OC GCP Questions
Subject: GCP Training

Hi

My name is [redacted]. I am a

new hire that started three weeks ago and I need to do the GCP training. Could you please send me a link or let me know how to do this.

Thank You for your time

[redacted]