

From: [OC GCP Questions](#)
To: [REDACTED]
Subject: GCP & HSP Training Requirement
Date: Friday, February 07, 2020 1:51:18 PM
Attachments: [REDACTED]

Good afternoon –

Thank you for your email. Neither FDA's regulations nor guidance provide guidelines on how often GCP training should be completed and by whom and who can perform the training. FDA does not require or provide certification. This would be up to the sponsor. The regulations require that sponsors choose investigators qualified by training and experience (see 21 CFR 312.53(a)). The regulations also require that investigators commit themselves to personally conduct or supervise the investigation (see 21 CFR 312.53(c)(1)(vi)(c)). FDA's guidance on Investigator Responsibilities - Protecting the Rights, Safety, and Welfare of Study Subjects (available at <https://www.fda.gov/media/77765/download>) states that, "The investigator should ensure that any individual to whom a task is delegated is qualified by education, training, and experience (and state licensure where relevant) to perform the delegated task."

FDA regulations regarding qualifications for those involved with the conduct of clinical studies are very broad. Both Title 21, Code of Federal Regulations (21 CFR) Part 312 (for drugs and biologics) and 812 (for medical devices) simply state that sponsors are required to choose clinical investigators and study staff that are qualified by training and experience. Logically, clinical investigators and study staff need to be knowledgeable about applicable regulations as well as specific areas essential to the conduct of the particular study.

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. As noted, the 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. This is the sponsor's responsibility.

Additionally what training is needed and how it is documented depends to some degree on the nature of the study. Some protocols need extensive training and others may need minimal, also dependent upon the background and experience of study staff.

Kind regards,

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From: [REDACTED]

Sent: Friday, February 07, 2020 1:08 PM

To: OC GCP Questions <gcpquestions@fda.hhs.gov>

Cc: [REDACTED]

Subject: GCP & HSP Training Requirement

Hello

I am trying to find in writing how often the Good Clinical Practice and/or Human Subject Protection Training should occur.

I hear every 2 years and I've seen after googling the subject every 3 years.

Can you assist me in finding out what the regulations states and where I can find it?

Thanks,

[REDACTED]