UCDAVIS HEALTH | Clinical and Translational Science Center

Informed Consent Boot Camp

Vulnerable Populations

Title: CRC 2.0 Series – Informed Consent Boot Camp, Part 3: Vulnerable Populations – Obtaining and Documenting Consent

Date: Tuesday, June 18, 2019

Time: 12-2 :00 p.m.

Location: Education Bldg, Lecture Hall, Rm 2222

In this interactive course, we will focus on the process of conducting and documenting informed consent for vulnerable populations (children, cognitively impaired adults, pregnant woman, prisoners, neonates, students/employees/subordinates).

Topics will include –

- IRB's expectation/requirements for obtaining consent
- Process of obtaining consent and considerations of the population
- Non-compliance issues that have been identified by IRB
- Best practices/tips

LMS registration is required and 2.0 self-study CEUs will be available for those who attend: Click to register, LMS Code: DAHS-CTSC-ICF3

Previous attendance in the series is not required.

For more information, contact Dannelle Jimenez, Clinical Research Education Program Manager, Clinical Trials Office at dijimenez@ucdavis.edu.

June

18

The course is beneficial to anyone who conducts the ICF process.

Speakers:

Stephanie Falwell, CRC
Pathology and Lab Medicine

Nicole Walters,
IRB Education
Program Manager

Sponsored by
Clinical Trials Office (CTO)