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.