

FDA Guidance Documentation and Warning Letters... What it means for you?

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The VCU Clinical Trials Office is holding a lecture discussing the FDA guidance documents and subsequent warning letters, and provide tools to meet these requirements.

At the conclusion of this event, participants will be able to:

- review processes applicable to the FDA guidance requirements.
- identify documentation processes that must be maintained to fulfill FDA expectations.
- use tools provided to implement additional source documentation needs.



DATE: Thursday, July 15, 2010

TIME: 12-12.50p

WHERE: Sanger Hall, 8-036

WHO: Clinical Trial Coordinators, Regulatory Coordinators, and anyone who handles clinical trial regulatory documentation are encouraged to attend.

This continuing education activity meets the criteria of Virginia Commonwealth University and the Southern Association of Colleges and Schools. 0.1 CEUs will be awarded and recorded with the University.