Research Policy & Compliance

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Topics

- UCLA Procedure 925.3
- ClinicalTrials.gov adverse event reporting requirement
- NSF America COMPETES Act Implementation
UCLA Procedure 925.3

- “Disclosing Financial Interests Relevant to Federally Sponsored Contracts and Grants for Research”

- **NSF**: Disclosure required for **all** sponsored projects

- **Other federal agencies**: Disclosure required for sponsored **research** only
ClinicalTrials.gov Adverse Event Reporting

- Effective September 27, 2009
- "Responsible Party" must report adverse events when reporting or updating the results of a clinical trial
ClinicalTrials.gov Adverse Event Reporting

- "Responsible Party"
  - For PI-initiated clinical trial
    - Principal Investigator
  - For sponsor-initiated clinical trial
    - Sponsor or Sponsor’s designated Principal Investigator
NSF America COMPETES Act Implementation

Requires training in the *Responsible Conduct of Research* for all undergraduates, graduate students, and postdoctoral scholars who will be supported by NSF to conduct research.
NSF America COMPETES Act Implementation

- UCLA must have training plan in place by January 4, 2010
- Vice Chancellor for Research, Dean of Students, Graduate Division working on plan