

**UMDNJ- Office of the VP for Research**  
**HUMAN SUBJECTS PROTECTION PROGRAM**  
*(Your Partner in the Responsible Conduct of Research)*

**Hosting an Webinar**

**“Biobanking Basics: An Overview of Current Legal and  
Compliance Issues Related to Human Specimens  
Collected in Research”**

**Tuesday, April 5, 2011**  
**1:00 – 2:30 p.m.**

**LOCATIONS:**                   New Brunswick Campus  
  125 Paterson Street  
  Clinical Academic Building, Room 1302

  Newark Campus  
  185 So. Orange Avenue  
  Medical Science Building, Room B-556

**Webinar Instructors:**

**Shana Fried, Reed Smith, LLP**  
**Celeste Letourneau, Reed Smith, LLP**

**By the end of this program, participants will be able to:**

- Identify the differences between HHS and FDA Informed Consent requirements and their applicability to different scenarios such as clinical trials and material transfer arrangements with industry sponsors
- Describe the application of HIPAA to the use of protected health information
- Describe and discuss the applicability of case law, as well as trends and developments in state and federal legislation (including GINA and ARRA), with regard to "ownership" of biological specimens, with a view toward understanding current industry sponsor and academic medical center concerns in clinical research and development

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