

# Human Research Protection Programs 1a: How to Navigate Human Subject Protection Regulations

Chair: Mark E. Sobel, M.D., Ph.D.  
American Society for Investigative Pathology

**Saturday, April 17, 2004**

**9:00AM - Room 101**

**Experimental Biology 2004  
Washington DC Convention Center**

This workshop will provide an introduction to principles of bioethics and current United States federal regulations for the protection of human subjects involved in research. There will be particular emphasis on use of human biological materials in research, the roles of Institutional Review Boards, when informed consent is required, interpretation of HIPAA, and how institutions can develop comprehensive programs to effectively protect human subjects and simultaneously conduct good scientific research.

- ▶ 9:00 a.m. **Introduction to Principles of Biomedical Ethics**, Mark E. Sobel, M.D., Ph.D., ASIP
- ▶ 9:30 a.m. **Regulations**, J. Kancshiro, Office of Human Research Protections (OHRP)
- ▶ 10:00 a.m. **Approaches to Accreditation of Human Subjects Protection Programs**, Marjorie Speers, Ph.D., Association for the Accreditation of Human Research Protection Programs
- ▶ 10:30 a.m. **Panel Discussion and Questions from the Audience**

**Pre-Registration Required - Deadline: March 31, 2004**



**American Society for  
Investigative Pathology**

9650 Rockville Pike  
Bethesda, MD 20814-3993 (USA)

Contact:  
Tara Zeitner  
Director of Meetings and  
Membership Services  
Tel: 301.634.7950  
Email: [meetings@asip.org](mailto:meetings@asip.org)  
Fax: 301.634.7990  
[www.asip.org](http://www.asip.org)

Name \_\_\_\_\_

Department \_\_\_\_\_

Institution \_\_\_\_\_

Address \_\_\_\_\_

City, State, Zip \_\_\_\_\_

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Email \_\_\_\_\_

**Fax completed form to 301-634-7990**