



**The Human Subjects Protection Program and  
Newark Institutional Review Board Office**

**Educational Session**

**“Basics for Informed Consent in Research”**

This session will cover several key topics instrumental to understanding the consent process in human subjects' research:

- Overview of the ethical principles underlying informed consent
- Current regulatory requirements
- Common findings by auditors
- The subject's view of the consent process

**Date 1: April 4, 2012  
9:30 a.m. – 11:30 a.m.**

UMDNJ- New Brunswick Campus  
Cancer Institute of New Jersey  
195 Little Albany Street  
Auditorium A

**Date 2: April 26, 2012  
12:30 p.m. – 2:30 p.m.**

UMDNJ- Newark Campus  
New Jersey Medical School  
Medical Science Building  
Lecture Hall B-552

**Speakers:**

Paula Bistak, D.M.H., RN, MS, CIP, CHRC  
Executive Director, Human Subjects Protection Program

Donna Hoagland, LPN, BS, CIP, CCRC  
IRB Director, New Brunswick/Piscataway Campus

Carlotta Rodriguez, BS, CIP, CHRC  
IRB Director, Newark Campus

Cheryl A Forst, RN, BSN, CCRP  
Director, Human Subjects Protection Analysts

Chris Asmann-Finch, DMH, MS  
Program Development Specialist, HSPP

There is no continuing education credit awarded for this session

RSVP to Shelia Evans: [evanssg@umdnj.edu](mailto:evanssg@umdnj.edu)