

The Middle East Research Ethics Training Initiative (MERETI)

at the University of Maryland, USA

in collaboration with

The Salim El-Hoss Bioethics and Professionalism Program at the American University of Beirut

and

Jordan Society of Forensic Science

Announces Two Workshops in Research Ethics to be held at Landmark Hotel in Amman, Jordan

<i>Training for IRB/REC Members</i>	<i>International Research Ethics</i>
<i>Dates: March 18-22, 2012 Registration fees: \$350</i>	<i>Dates: March 24-26, 2012 Registration fees: \$250</i>

*Registration fees for both workshops is \$500
Certificates will be given for each workshop*

Closing date for applications is 15 February 2012

Space is Limited

Registration fees cover coffee breaks and lunch, but not travel and hotel expenses

Limited Number of Scholarships are available to cover registration fees, travel, and hotel expenses

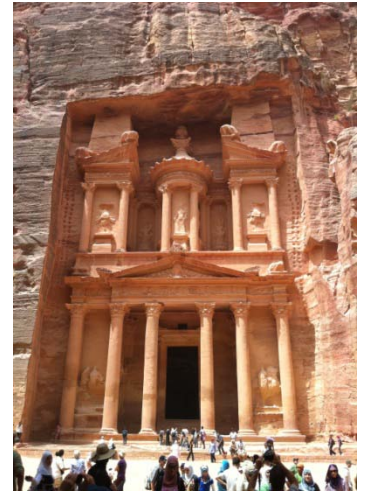
Who Should Attend?

- Chairs and Members of Research Ethics Committees
- Investigators and research coordinators
- Members of Contract Research Organizations

Learning Objectives:

By the end of these courses, participants will be able to:

- Discuss the ethical foundations for research ethics
- Review the ethical aspects of research protocols
- Explain vulnerability
- Explain the importance of informed consent
- Discuss barriers faced by Research Ethics Committees
- Discuss issues in international research



Hotel Information: Landmark Amman Hotel and Conference Center
Tel: (962) 6 560 7100
www.landmarkamman.com
Rates: starting at 80 JD (special conference rate)
Includes breakfast

For Inquiries and Registration, please contact:

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For more information: www.mereti.net



www.mereti-network.net



Landmark Hotel

Workshop Agendas

Certificate for Members of IRBs/RECs (March 18 – March 22)

	March 18 Sunday	March 19 Monday	March 20 Tuesday	March 21 Wednesday	March 22 Thursday
Session I	<p>Introduction Why Research Ethics? Theory and Principles</p> <p>Ethical Requirements for Protocol Review: How to use a framework</p>	<p>Therapeutic Misconception Conflicting roles of physician and investigators</p> <p>Therapeutic Misconception Lecture</p>	<p>Privacy and Confidentiality Lecture Case Study</p> <p>Coercion and Undue Inducement Lecture</p>	<p>Risks and Benefits Lecture Risks and Benefits Case Study</p> <p>IRB II Monitoring of Clinical Trials</p> <p>Responsible Conduct of Research I</p>	<p>Genetic I Ethics of Stem Cell Research Lecture</p> <p>Genetics II Stored Sample Lecture Case Study</p>
Session II	<p>Research Ethics in the Arab Region Status of Research Ethics Capacity in the Arab Region</p> <p>Perspectives of Investigators in the Arab Region Regarding Research Ethics</p>	<p>Ethical Theory I Islamic Bioethics</p> <p>Informed Consent I Case Study</p>	<p>Vulnerability Lecture Hepatitis Vaccine Case Study</p> <p>Ethics of Children Research Lecture</p>	<p>Clinical Trials I Phases of Clinical Trials Lecture</p> <p>Clinical Trials II Ethics of Phase I Studies Lecture</p>	<p>IRB III Structure and Function of IRBs</p> <p>IRB IV: Collaborative IRB Review: The Role of a Central IRB</p>
Lunch					
Session III	<p>IRB I Categories of Research Review</p> <p>Challenges and Capacity of IRBs in the Arab Region</p>	<p>Informed Consent II Presentation</p> <p>Waiver of Consent Case Studies</p>	<p>Ethical Theory II Philosophical Bioethics</p> <p>Social Behavioral Research Lecture</p>	<p>Fair Recruitment of Subjects Participants' Views on research ethics</p> <p>Ethics of Payment to participants</p>	<p>Presentations by IRBs from different Arab Countries: Egypt, Jordan, Saudi Arabia, Sudan, Yemen</p>
Session IV	<p>Protocol Review I Outpatient Treatment of Pneumonia</p>	<p>Informed Consent III Writing an Informed Consent</p>	<p>Protocol Review II Case Study: Domestic Violence</p>	<p>Protocol Review III</p>	<p>Protocol Review IV</p>

Certificate in International Research Ethics (March 24 – 26)					
	March 24 Saturday	March 25 Sunday	March 26 Monday		
Session I	International Research Ethics I Standard of Care Case Study: AZT Trial The Role of Equipoise in International Trials	International Ethics II Achieving Fair Benefits Lecture and Case Studies The concept of social value in research in developing countries	Clinical Trials III Ethics of Placebo Trials Controversial Cases in International Research Trovan, Tenofovir, and TeGenero		
Session II	The Challenge of Informed Consent In International Trials Case study Movie Clip: Constant Gardener Discussion: Helsinki Document CIOMS Document	International Research Ethics III Exploitation <ul style="list-style-type: none"> • Frostbite Case Study • The SAPIT (Starting Antiretroviral Therapy at Three Points in Tuberculosis Therapy) 	Responsible Conduct of Research II Research Integrity in the Context of Global Cooperation: What are the Challenges? Responsible Conduct of Research III Conflict of Interest and Publication Issues among International Partners		
Lunch					
Session III	Regulations and Existing Laws on Clinical Trials in the Arab Countries: What is missing to ensure protection for participants? Research and Global Health: Issues of Justice	Community Participatory Research: Issues regarding community engagement and community consent. Challenges of Research Collaborative Partnership in International Trials	What are Ancillary Care Responsibilities of Investigators and Sponsors? Ethical challenges in the release and publication of ethnicity related research results		
Session IV	Case Studies: Group Work	Case Studies: Group Work	Case Studies: Group Work		