Required Course Guide for IRB Members

Social-Behavioral-Educational Research & Biomedical Research

Basic Human Subjects Research (HSR) Modules

Basic Social-Behavioral-Educational (SBE) Modules		
1	Assessing Risk - SBE (ID: 503)	
2	Defining Research with Human Subjects - SBE (ID: 491)	
3	History and Ethical Principles - SBE (ID: 490)	
4	Informed Consent - SBE (ID: 504)	
5	Internet-Based Research - SBE (ID: 510)	
6	Privacy and Confidentiality - SBE (ID: 505)	
7	Research in Public Elementary and Secondary Schools - SBE (ID: 508)	
8	Research with Children - SBE (ID: 507)	
9	Research with Prisoners - SBE (ID: 506)	
10	The Federal Regulations - SBE (ID: 502)	
11	Unanticipated Problems and Reporting Requirements in Social and Behavioral Research (ID: 14928)	
	International Research - SBE (ID: 509)	
Additional Modules of Interest		
12	Conflicts of Interest in Human Subjects Research (ID: 17464)	
13	Hot Topics (ID: 487)	
	Are You Thinking About Being in a Research Study? (ID: 14562)	
	Cultural Competence in Research (ID: 15166)	
	International Studies (ID: 971)	
Clinical Trial Agreement (CTA) Modules		
	Clinical Trial Agreement (CTA) Negotiation for Researchers and Sites (ID: 17359)	
	Overview of the Clinical Trial Agreement (CTA) (ID: 17356)	
	Role of the Researcher and Site in Managing the Clinical Trial Agreement (CTA) (ID: 17358)	
	Understanding the Terms of the Clinical Trial Agreement (CTA) (ID: 17357)	
Community-Engaged Research (CEnR) Modules		

Consent Modules		
	Consent and Biobanks and Associated Databases (ID: 17254)	
	Consent and Cultural Competence (ID: 17263)	
	Consent and Subject Recruitment Challenges: Remuneration (ID:16881)	
	Consent and Subject Recruitment Challenges: Therapeutic Misconception (ID:17259)	
	Consent in the 21st Century (ID: 17060)	
	Consent Tools Used in Research (ID: 16944)	
	Consent with Subjects Who Do Not Speak English (ID: 17260)	
	Informed Consent and Incidental Findings in Research with Human Subjects (ID: 17342)	
IRB Focused Modules		
	External IRB Review (ID: 16711)	
	I Have Agreed to be an IRB Community Member. Now What? (ID: 13018)	
	The IRB Administrator's Responsibilities (ID: 13813)	
	The IRB Member Module - "What Every New IRB Member Needs to Know" (ID: 816)	
Phase I Research Modules		
	Phase I Research: Protecting Phase I Subjects (ID: 16874)	
	Phase I Research: Understanding Phase I Research (ID: 16873)	
Population-Specific Modules		
14	Vulnerable Subjects - Research Involving Workers/Employees (ID: 483)	
	Gender and Sexuality Diversity (GSD) in Human Research (ID: 16556)	
	Illegal Activities or Undocumented Status in Human Research (ID: 16656)	
	Research Involving Subjects at the End of Life (ID: 16658)	
	Research with Critically Ill Subjects (ID: 16592)	
	Research with Decisionally Impaired Subjects (ID: 16610)	
	Research with Older Adults (ID: 16502)	
	Research with Persons who are Socially or Economically Disadvantaged (ID: 16539)	
	Research with Subjects with Physical Disabilities & Impairments (ID: 16657)	
	Students in Research (ID: 1321)	
Stem Cell Research Modules		
	Stem Cell Research Oversight (Part I) (ID: 13882)	

Basic Biomedical (Biomed) Modules

	Avoiding Group Harms - International Research Perspectives (ID: 14081)
	Avoiding Group Harms - U.S. Research Perspectives (ID: 14080)
	Basic Institutional Review Board (IRB) Regulations and Review Process (ID: 2)
	FDA-Regulated Research (ID: 12)
	Genetic Research in Human Populations (ID: 6)
	History and Ethics of Human Subjects Research (ID: 498)
	Informed Consent (ID: 3)
	Populations in Research Requiring Additional Considerations and/or Protections (ID: 16680)
	Recognizing and Reporting Unanticipated Problems Involving Risks to Subjects or Others in Biomedical Research (ID: 14777)
	Records-Based Research (ID: 5)
	Research and HIPAA Privacy Protections (ID: 14)
	Social and Behavioral Research (SBR) for Biomedical Researchers (ID: 4)
	Vulnerable Subjects - Research Involving Children (ID: 9)
	Vulnerable Subjects - Research Involving Pregnant Women, Fetuses, and Neonates (ID: 10)
	Vulnerable Subjects - Research Involving Prisoners (ID: 8)

This training is required every 4 Years, at which time you may take the Refresher Course, an abbreviated version of t
course.

Please send Completion Reports to Gwen Parnell in the GRaSP Office

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