

## Continuing Research Education Credit [\(CREC\)](#) Webinars In CITI

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Add a webinar | [Instructions for new CITI users](#) | [Instructions for current CITI users](#)

1. A 21st Century Approach to ADA Compliance: Equity and Access **(3 CREC)**
2. A Beginner's Guide to Being a Sponsor-Investigator **(3 CREC)**
3. Accreditation 101 for New and Adjunct Faculty
4. Addressing and Understanding Mental Health Challenges for Faculty and Staff
5. Artificial Intelligence (AI) and Human Subject Protections **(2 CREC)**
6. Best Practices for Global Research Partnerships: Benefits and Challenges
7. Blockchain and Higher Education
8. Bring Your Own Device (BYOD) Studies **(2 CREC)**
9. Building a ClinicalTrials.gov Compliance Program – Tips for Investigators and Institutions **(3 CREC)**
10. Clinical Trial Staff Diversity and Cultural Humility **(3 CREC)**
11. ClinicalTrials.gov Enforcement: An Update
12. Cost Allowability on Federally Sponsored Awards
13. COVID-19 and Human Research Protection Programs **(3 CREC)**
14. COVID-19: Supporting Ethical Care and Responding to Workforce Concerns in a Public Health Emergency
15. CRA Soft Skills, Time Management, and Effective Site Relationships
16. CRISPR Genome Editing: Rewriting DNA and the Future of Humanity **(2 CREC)**
17. Data Management and Security for Student Researchers: An Overview
18. Decentralized Clinical Trials (DCTs) and Your Workforce
19. Drones in Academia
20. Ethics & Policy Issues in CRISPR Gene Editing New Content **(2 CREC)**

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21. Export Compliance: An Overview for Staff, Students, and Faculty
22. Facial Recognition Considerations for Researchers **(3 CREC)**
23. FCPA and University Research: What Faculty and Administrators Need to Know
24. FDA Inspections of GMP Facilities: How to Be Inspection Ready
25. FERPA and Online Learning in the Time of COVID-19
26. FERPA: A Quick Review of the Law for Researchers and IRBs **(3 CREC)**
27. From Cancer to COVID-19, Does Science Self-Correct?
28. GDPR & Human Subject Research in the U.S. **(3 CREC)**
29. GDPR: Top Noncompliance Risks and Mitigation Strategies
30. Gender and Sexual Minorities (GSM) in Human Subjects Research **(2 CREC)**
31. Getting Started in Grant Writing: An Introduction for Graduate Students, Postdocs, and New Faculty
32. Health Disparities: Promoting Equity and Diversity in Clinical Research **(3 CREC)**
33. Higher Education Accelerated Credit
34. How to Conduct an Audit of a Ceded Study
35. How to Effectively Manage a Research Administration Team
36. Human Enhancement and its Ethical Implications
37. Importance of Peer Review and Data Validation in Research **(3 CREC)**
38. Improving the Clinical Trial Participant's Experience: From Recruitment through Study Closure **(2 CREC)**
39. Informed Consent and Clinical Investigations: A Focus on the Process **(3 CREC)**
40. Informed Consent and Research with Wearable Tech **(3 CREC)**

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41. Intellectual Property and Working With Your Technology Transfer Office
42. International Students in Focus at U.S. Higher Education Institutions (HEIs)
43. IRB Administrator Professional Development and Self-Advocacy
44. IRB Protocol Noncompliance: When Research Goes Rogue, What Next? **(3 CREC)**
45. IRB Review of Observational Research **(3 CREC)**
46. Leveraging IT Insight in IRB Review: Why Technology-Based Expertise is Critical to Human Subjects **(3 CREC)**
47. Managing Conflict with Your Dissertation Chair
48. Managing Your Grants as Systems: A Guide for Grant Management Success
49. Medical Marijuana: A Budding Field of Research
50. Noncompliance and the IACUC: Basic Approaches for Success
51. "Nuts & Bolts" of Running a Virtual IRB Meeting **(2 CREC)**
52. Open Access Publishing: An Introduction
53. Partnering with Technology Companies
54. Preparing for Single IRB (sIRB) under the Common Rule **(3 CREC)**
55. Preparing to Publish in Traditional and Hybrid Journals
56. Principles and Practices for Managing Undue Foreign Influence in an Academic Environment
57. Privacy and Ethical Considerations for Connected and Automated Vehicles (CAVs)
58. Quality Improvement Activities and the Common Rule **(3 CREC)**
59. Race in Clinical Research: Ethics and IRB Decision Making **(3 CREC)**
60. Remote Informed Consent: The Same, but Different, but Still the Same **(3 CREC)**

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61. Research Equity and the Part We Play **(3 CREC)**
62. Research in Wound Care
63. Research with Audio-Visual Mobile Data Collection Tools: Ethics and Regulations **(3 CREC)**
64. Research with Native American Communities: Important Considerations When Applying Federal Regulation **(2 CREC)**
65. Revised Common Rule: Overview of Revisions **(3 CREC)**
66. Revised Common Rule: Revisions to Definitions **(2 CREC)**
67. Revised Common Rule: Revisions to Informed Consent **(2 CREC)**
68. Running a Virtual IRB Meeting **(2 CREC)**
69. Social Media and Research Recruiting **(3 CREC)**
70. Supervision for Supervisors
71. The Challenge of Medicare Advantage Plans and Local Coverage Determinations
72. The Playbook: Successfully Developing and Deploying Digital Clinical Measures **(3 CREC)**
73. Tips for Research Administrators: Working with Faculty and Research Teams
74. Title IX and the New Regulations
75. Title IX: 50 Years and Modern Challenges
76. Transitioning Research to the Revised Common Rule: The What, How, and Why **(3 CREC)**
77. U.S. Department of Defense (DoD) Regulations & Requirements for Human Subject Research **(3 CREC)**
78. U.S. Department of Energy Interim Policy on Conflicts of Interest
79. Understanding 483s and Surviving Them

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- 80. Understanding and Addressing Mental Health on Campus: Opportunities and Challenges in Higher Education
- 81. Understanding Consent Requirements and “Key Information” Under the Revised Rule **(3 CREC)**
- 82. Understanding Decentralized Clinical Trials (DCTs) and Virtual Study Visits **(2 CREC)**
- 83. Understanding ISO 14155:2020 Revision **(3 CREC)**
- 84. Working with the FDA: Medical Devices and Regulatory Touchpoints
- 85. Working with Your IRB **(2 CREC)**