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46. Leveraging IT Insight in IRB Review: Why Technology-Based Expertise is Critical to Human Subjects **(3 CREC)**
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50. Noncompliance and the IACUC: Basic Approaches for Success
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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
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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)**
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70. Supervision for Supervisors
71. The Challenge of Medicare Advantage Plans and Local Coverage Determinations
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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)**

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78. U.S. Department of Energy Interim Policy on Conflicts of Interest
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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
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