Reform of Clinical Research Regulations, Finally

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Citation Report

#	Article	IF	Citations
1	Clinical Research Informatics for Big Data and Precision Medicine. Yearbook of Medical Informatics, 2016, 25, 211-218.	1.0	24
2	A Flawed Revision of the Common Rule. Annals of Internal Medicine, 2016, 165, 143.	3.9	4
3	The ERS Research Agency: the beginning. European Respiratory Journal, 2016, 47, 1017-1023.	6.7	13
4	Reform of Clinical Research Regulations. New England Journal of Medicine, 2016, 374, 1693-1694.	27.0	3
5	CT Fluoroscopyâ€“Guided Blood Patching of Ventral CSF Leaks by Direct Needle Placement in the Ventral Epidural Space Using a Transforaminal Approach. American Journal of Neuroradiology, 2016, 37, 1951-1956.	2.4	20
6	Confronting Biospecimen Exceptionalism in Proposed Revisions to the Common Rule. Hastings Center Report, 2016, 46, 4-5.	1.0	14
7	Human Subjects Protection and Technology in Prevention Science: Selected Opportunities and Challenges. Prevention Science, 2016, 17, 765-778.	2.6	24
8	Lessons from HeLa Cells: The Ethics and Policy of Biospecimens. Annual Review of Genomics and Human Genetics, 2016, 17, 395-417.	6.2	90
9	Unravelling the human genomeâ€“phenome relationship using phenome-wide association studies. Nature Reviews Genetics, 2016, 17, 129-145.	16.3	222
10	Improving biobank consent comprehension: a national randomized survey to assess the effect of a simplified form and review/retest intervention. Genetics in Medicine, 2017, 19, 505-512.	2.4	34
11	At Last! Aye, and There's the Rub. American Journal of Bioethics, 2017, 17, 4-7.	0.9	34
12	Rethinking the Belmont Report?. American Journal of Bioethics, 2017, 17, 15-21.	0.9	59
13	A qualitative exploration of the informed consent process in hematopoietic cell transplantation clinical research and opportunities for improvement. Bone Marrow Transplantation, 2017, 52, 292-298.	2.4	6
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16	Revisions to the Common Rule: A proposal in search of evidence. Research Ethics, 2017, 13, 92-96.	1.7	6
17	Untapped Potential of Observational Research to Inform Clinical Decision Making: American Society of Clinical Oncology Research Statement. Journal of Clinical Oncology, 2017, 35, 1845-1854.	1.6	87
18	Challenges in the Regulation of Autologous Stem Cell Interventions in the United States. Perspectives in Biology and Medicine, 2018, 61, 25-41.	0.5	17
19	A checklist for clinical trials in rare disease: obstacles and anticipatory actionsâ€”lessons learned from the FOR-DMD trial. Trials, 2018, 19, 291.	1.6	26

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21	Public Comments on the Proposed Common Rule Mandate for Singleâ€IRB Review of Multisite Research. Ethics & Human Research, 2019, 41, 15-21.	0.9	6
22	Rethinking the role of Research Ethics Committees in the light of Regulation (EU) No 536/2014 on clinical trials and the COVIDâ€19 pandemic. British Journal of Clinical Pharmacology, 2022, 88, 40-46.	2.4	15
23	A population-based approach for implementing change from opt-out to opt-in research permissions. PLoS ONE, 2017, 12, e0168223.	2.5	19
24	Reimagining Human Research Protections for 21st Century Science. Journal of Medical Internet Research, 2016, 18, e329.	4.3	30
25	SurveyÂon Centralization of Research Ethics Review for Multicenter Clinical Trials in Japan . Japanese Journal of Clinical Pharmacology and Therapeutics, 2018, 49, 159-167.	0.1	0
27	Paying Research Participants: Regulatory Uncertainty, Conceptual Confusion, and a Path Forward. Yale Journal of Health Policy, Law, and Ethics, 2017, 17, 61-141.	1.5	42