The EU Clinical Trials Regulation: key priorities, purpos public health

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#	Article	IF	Citations
1	Application challenges of the new EU Clinical Trials Regulation. European Journal of Clinical Pharmacology, 2017, 73, 795-798.	0.8	17
2	Informed consent to research trials on Alzheimerâ€™s disease: How to foster research without infringing upon the patientâ€™s right to self-determination. Pharmacological Research, 2018, 132, 69-71.	3.1	2
3	Proportionality and shared responsibility for Dutch trial reviews. Nature Human Behaviour, 2018, 2, 99-99.	6.2	1
4	International Perspectives on Substantiating the Efficacy of Herbal Dietary Supplements and Herbal Medicines Through Evidence on Traditional Use. Comprehensive Reviews in Food Science and Food Safety, 2019, 18, 910-922.	5.9	17
5	Prospective registration and reporting of trial number in randomised clinical trials: global cross sectional study of the adoption of ICMJE and Declaration of Helsinki recommendations. BMJ, The, 2020, 369, m982.	3.0	44
6	O Regulamento (UE) NÂº 536/2014 relativo aos ensaios clÃnicos de medicamentos para uso humano: oportunidades de inovaÃ§Ã£o e desafios Ã©ticos. Cadernos Ibero-americanos De Direito SanitÃ¡rio, 2021, 10, 211-224.	0.1	0
7	Research Volunteer. , 2019, , 497-504.		0
8	CONTRACTUAL REGISTRATION OF ORGANIZATIONAL AND LEGAL RELATIONS BETWEEN SUBJECTS INVOLVED IN THE CONDUCT OF CLINICAL TRIALS OF MEDICINAL PRODUCTS. WiadomoÅ›ci Lekarskie, 2020, 73, 2840-2847.	0.1	0
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12	Serious adverse event reporting and compensation. , 2024, , 141-156.		0