## Karmela Krleza-Jeric

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/2096585/publications.pdf

Version: 2024-02-01

840776 8,849 17 11 citations h-index papers

17 g-index 17 17 17 18335 docs citations times ranked citing authors all docs

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#	Article	IF	CITATIONS
1	SPIRIT 2013 Statement: Defining Standard Protocol Items for Clinical Trials. Annals of Internal Medicine, 2013, 158, 200.	3.9	4,463
2	SPIRIT 2013 explanation and elaboration: guidance for protocols of clinical trials. BMJ, The, 2013, 346, e7586-e7586.	6.0	3,435
3	Outcome reporting bias in randomized trials funded by the Canadian Institutes of Health Research. Cmaj, 2004, 171, 735-740.	2.0	380
4	Principles for international registration of protocol information and results from human trials of health related interventions: Ottawa statement (part 1). BMJ: British Medical Journal, 2005, 330, 956-958.	2.3	186
5	Sharing and reuse of individual participant data from clinical trials: principles and recommendations. BMJ Open, 2017, 7, e018647.	1.9	116
6	The Use of Electronic Data Capture Tools in Clinical Trials: Web-Survey of 259 Canadian Trials. Journal of Medical Internet Research, 2009, 11, e8.	4.3	74
7	Clinical Trial Registration: The Differing Views of Industry, the WHO, and the Ottawa Group. PLoS Medicine, 2005, 2, e378.	8.4	43
8	Reporting of Methodologic Information on Trial Registries for Quality Assessment: A Study of Trial Records Retrieved from the WHO Search Portal. PLoS ONE, 2010, 5, e12484.	2.5	38
9	Do trialists endorse clinical trial registration? Survey of a Pubmed sample. Trials, 2007, 8, 30.	1.6	37
10	Evaluation of repositories for sharing individual-participant data from clinical studies. Trials, 2019, 20, 169.	1.6	36
11	International Dialogue on the Public Reporting Of Clinical Trial Outcome and Results – PROCTOR Meeting. Croatian Medical Journal, 2008, 49, 267-268.	0.7	13
12	Sharing individual participant data from clinical studies: a cross-sectional online survey among Italian patient and citizen groups. BMJ Open, 2019, 9, e024863.	1.9	13
13	IMPACT Observatory: tracking the evolution of clinical trial data sharing and research integrity. Biochemia Medica, 2016, 26, 308-317.	2.7	8
14	Setting the IMPACT (IMProve Access to Clinical Trial data) Observatory baseline. Biochemia Medica, 2018, 28, 010201.	2.7	3
15	Clinical Trials Registries and Results Databases. Computers in Health Care, 2012, , 389-408.	0.3	2
16	First BH Cochrane Symposium held. Materia Socio-medica, 2016, 28, 74.	0.7	1
17	Clinical Trial Registries, Results Databases, and Research Data Repositories. Computers in Health Care, 2019, , 453-480.	0.3	1