CITATION REPORT List of articles citing

What information and the extent of information research participants need in informed consent forms: a multi-country survey

DOI: 10.1186/s12910-018-0318-x BMC Medical Ethics, 2018, 19, 79.

Source: https://exaly.com/paper-pdf/70636985/citation-report.pdf

Version: 2024-04-23

This report has been generated based on the citations recorded by exaly.com for the above article. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

#	Paper	IF	Citations
15	Informational needs for participation in bioequivalence studies: the perspectives of experienced volunteers. <i>European Journal of Clinical Pharmacology</i> , 2019 , 75, 1575-1582	2.8	O
14	Managing uncertain recovery for patients nearing the end of life in hospital: a mixed-methods feasibility cluster randomised controlled trial of the AMBER care bundle. <i>Trials</i> , 2019 , 20, 506	2.8	14
13	Researchersbviews on, and experiences with, the requirement to obtain informed consent in research involving human participants: a qualitative study. <i>BMC Medical Ethics</i> , 2020 , 21, 93	2.9	7
12	Supporting students in making informed data sharing decisions: from comprehension to consenting. 2020 ,		
11	Why bunderstandingbof research may not be necessary for ethical emergency research. <i>Philosophy, Ethics, and Humanities in Medicine</i> , 2020 , 15, 6	2.2	2
10	Enhancing consent forms to support participant decision making in multimodal learning data research. <i>British Journal of Educational Technology</i> , 2020 , 51, 1631-1652	4.3	7
9	Informed Consent: What Must Be Disclosed and What Must Be Understood?. <i>American Journal of Bioethics</i> , 2021 , 21, 46-58	1.1	18
8	Factors Influencing Successful Recruitment of Racial and Ethnic Minority Patients for an Observational HIV Cohort Study in Washington, DC. <i>Journal of Racial and Ethnic Health Disparities</i> , 2021 , 1	3.5	1
7	Citizens, Research Ethics Committee Members and ResearchersbAttitude Toward Information and Consent for the Secondary Use of Health Data: Implications for Research Within Learning Health Systems. <i>Journal of Empirical Research on Human Research Ethics</i> , 2021 , 16, 165-178	1.6	2
6	The AMBER care bundle for hospital inpatients with uncertain recovery nearing the end of life: the ImproveCare feasibility cluster RCT. <i>Health Technology Assessment</i> , 2019 , 23, 1-150	4.4	7
5	Getting the Most out of Consent: Patient-Centered Consent for an Acute Stroke Trial <i>Ethics & Amp; Human Research</i> , 2022 , 44, 33-40	2.1	1
4	Co-creation with research participants to inform the design of electronic informed consent. <i>Digital Health</i> , 2022 , 8, 205520762211090	4	0
3	What information and the extent of information to be provided in an informed assent/consent form of pediatric drug trials. 2022 , 23,		O
2	Using provocative design to foster electronic informed consent innovation. 2022, 22,		0
1	Fragmented understanding: exploring the practice and meaning of informed consent in clinical trials in Ho Chi Minh City, Vietnam. 2023 , 24,		О