Rebekah E Mcwhirter

List of Publications by Year in descending order

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687363 580821 37 740 13 25 citations g-index h-index papers 39 39 39 2493 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	General practitioner perceptions of assessment and reporting of absolute cardiovascular disease risk via pathology services: a qualitative study. Family Practice, 2021, 38, 172-179.	1.9	2
2	General practitioners maintain a focus on blood pressure management rather than absolute cardiovascular disease risk management. Journal of Evaluation in Clinical Practice, 2021, 27, 1353-1360.	1.8	7
3	Australian human research ethics committee members' confidence in reviewing genomic research applications. European Journal of Human Genetics, 2021, 29, 1811-1818.	2.8	2
4	Recurrence patterns identify aggressive form of human papillomavirusâ€dependent vulvar cancer. Australian and New Zealand Journal of Obstetrics and Gynaecology, 2020, 60, 231-237.	1.0	1
5	Equitable Expanded Carrier Screening Needs Indigenous Clinical and Population Genomic Data. American Journal of Human Genetics, 2020, 107, 175-182.	6.2	24
6	Self-directed multimedia process for delivering participant informed consent. BMJ Open, 2020, 10, e036977.	1.9	3
7	Cerebral small vessel disease genomics and its implications across the lifespan. Nature Communications, 2020, 11, 6285.	12.8	89
8	A Scenario-Based Methodology for Analyzing the Ethical, Legal, and Social Issues in Genomic Data Sharing. Journal of Empirical Research on Human Research Ethics, 2020, 15, 355-364.	1.3	4
9	Integration of absolute cardiovascular disease risk assessment into routine blood cholesterol testing at pathology services. Family Practice, 2020, 37, 675-681.	1.9	4
10	Enhancing early detection of cognitive impairment in the criminal justice system: feasibility of a proposed method. Current Issues in Criminal Justice, 2019, 31, 60-74.	1.4	3
11	Genetic and lifestyle risk factors for MRI-defined brain infarcts in a population-based setting. Neurology, 2019, 92, .	1.1	30
12	Disclosure of Genetic Results to At-risk Relatives without Consent: Issues for Health Care Professionals in Australia. Journal of Law & Medicine, 2019, 27, 108-121.	0.0	1
13	Moving Forward on Consent Practices in Australia. Journal of Bioethical Inquiry, 2018, 15, 243-257.	1.5	11
14	Regulation of unregistered birth workers in Australia: Homebirth and public safety. Women and Birth, 2018, 31, 134-142.	2.0	3
15	Australia: regulating genomic data sharing to promote public trust. Human Genetics, 2018, 137, 583-591.	3.8	14
16	Provenance and risk in transfer of biological materials. PLoS Biology, 2018, 16, e2006031.	5.6	7
17	Identifying public expectations of genetic biobanks. Public Understanding of Science, 2017, 26, 671-687.	2.8	25
18	Adjusting for Familial Relatedness in the Analysis of GWAS Data. Methods in Molecular Biology, 2017, 1526, 175-190.	0.9	7

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19	Performance data and informed consent: a duty to disclose?. Medical Journal of Australia, 2017, 207, 100-101.	1.7	O
20	Key challenges in bringing CRISPR-mediated somatic cell therapy into the clinic. Genome Medicine, 2017, 9, 85.	8.2	17
21	Implementing values-based governance for a new bioresource model. Journal of Law and the Biosciences, 2017, 4, 404-411.	1.6	0
22	The complex genetics of gait speed: genome-wide meta-analysis approach. Aging, 2017, 9, 209-246.	3.1	21
23	Understanding public reactions to commercialization of biobanks and use of biobank resources. Social Science and Medicine, 2016, 162, 79-87.	3.8	53
24	Novel genetic loci underlying human intracranial volume identified through genome-wide association. Nature Neuroscience, 2016, 19, 1569-1582.	14.8	213
25	Precision medicine: drowning in a regulatory soup?. Journal of Law and the Biosciences, 2016, 3, 281-303.	1.6	18
26	Regulation of Non-consensual Genetic Testing in Australia: Use of Samples from Deceased Persons. Journal of Law & Medicine, 2016, 24, 150-65.	0.0	0
27	Genomics in research and health care with Aboriginal and Torres Strait Islander peoples. Monash Bioethics Review, 2015, 33, 203-209.	0.8	11
28	All in the Blood. Journal of Empirical Research on Human Research Ethics, 2015, 10, 347-359.	1.3	34
29	Community Engagement for Big Epidemiology: Deliberative Democracy as a Tool. Journal of Personalized Medicine, 2014, 4, 459-474.	2.5	49
30	Runs of homozygosity and a cluster of vulvar cancer in young Australian Aboriginal women. Gynecologic Oncology, 2014, 133, 421-426.	1.4	14
31	Genetic and epigenetic variation in vulvar cancer: Current research and future clinical practice. Australian and New Zealand Journal of Obstetrics and Gynaecology, 2014, 54, 406-411.	1.0	6
32	New avenues within community engagement: addressing the ingenuity gap in our approach to health research and future provision of health care. Journal of Responsible Innovation, 2014, 1, 321-328.	4.9	12
33	Body ownership and research. Journal of Law & Medicine, 2013, 21, 323-9.	0.0	2
34	Genome-wide homozygosity and multiple sclerosis in Orkney and Shetland Islanders. European Journal of Human Genetics, 2012, 20, 198-202.	2.8	16
35	Ethical genetic research in Indigenous communities: challenges and successful approachesâ [†] . Trends in Molecular Medicine, 2012, 18, 702-708.	6.7	18
36	The History of Bioethics: implications for current debates in health research. Perspectives in Biology and Medicine, 2012, 55, 329-338.	0.5	10

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37	Evidence for a common genetic aetiology in highâ€risk families with multiple haematological malignancy subtypes. British Journal of Haematology, 2010, 150, 456-462.	2.5	7