

Richard Milne

List of Publications by Year in descending order

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41
papers

752
citations

623188

14
h-index

610482

24
g-index

54
all docs

54
docs citations

54
times ranked

934
citing authors

#	ARTICLE	IF	CITATIONS
1	“Ready for What?” Timing and Speculation in Alzheimer’s Disease Drug Development. <i>Science Technology and Human Values</i> , 2022, 47, 597-622.	1.7	3
2	Assessing and disclosing test results for “mild cognitive impairment”: the perspective of old age psychiatrists in Scotland. <i>BMC Geriatrics</i> , 2022, 22, 50.	1.1	5
3	Digital phenotyping and the (data) shadow of Alzheimer's disease. <i>Big Data and Society</i> , 2022, 9, 205395172110707.	2.6	7
4	Return of genomic results does not motivate intent to participate in research for all: Perspectives across 22 countries. <i>Genetics in Medicine</i> , 2022, 24, 1120-1129.	1.1	8
5	Towards equitable and trustworthy genomics research. <i>EBioMedicine</i> , 2022, 76, 103879.	2.7	34
6	Understanding “passivity” in digital health through imaginaries and experiences of coronavirus disease 2019 contact tracing apps. <i>Big Data and Society</i> , 2022, 9, 205395172210911.	2.6	2
7	What can data trusts for health research learn from participatory governance in biobanks?. <i>Journal of Medical Ethics</i> , 2021, , medethics-2020-107020.	1.0	10
8	Public trust and genomic medicine in Canada and the UK. <i>Wellcome Open Research</i> , 2021, 6, 124.	0.9	1
9	Demonstrating trustworthiness when collecting and sharing genomic data: public views across 22 countries. <i>Genome Medicine</i> , 2021, 13, 92.	3.6	39
10	Public trust and genomic medicine in Canada and the UK. <i>Wellcome Open Research</i> , 2021, 6, 124.	0.9	2
11	Italian public’s views on sharing genetic information and medical information: findings from the “Your DNA, Your Say” study. <i>Wellcome Open Research</i> , 2021, 6, 180.	0.9	4
12	Lived time and the affordances of clinical research participation. <i>Sociology of Health and Illness</i> , 2021, , .	1.1	2
13	Societal and equity challenges for Brain Health Services. A user manual for Brain Health Services” part 6 of 6. <i>Alzheimer’s Research and Therapy</i> , 2021, 13, 173.	3.0	14
14	Engaged genomic science produces better and fairer outcomes: an engagement framework for engaging and involving participants, patients and publics in genomics research and healthcare implementation. <i>Wellcome Open Research</i> , 2021, 6, 311.	0.9	6
15	The rare and the common: scale and the genetic imaginary in Alzheimer's disease drug development. <i>New Genetics and Society</i> , 2020, 39, 101-126.	0.7	8
16	Members of the public in the USA, UK, Canada and Australia expressing genetic exceptionalism say they are more willing to donate genomic data. <i>European Journal of Human Genetics</i> , 2020, 28, 424-434.	1.4	29
17	Alzheimer’s disease and the development of a post-genomic science. <i>New Genetics and Society</i> , 2020, 39, 1-12.	0.7	1
18	Disruption and dislocation in post-COVID futures for digital health. <i>Big Data and Society</i> , 2020, 7, 205395172094956.	2.6	13

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19	Global Public Perceptions of Genomic Data Sharing: What Shapes the Willingness to Donate DNA and Health Data?. <i>American Journal of Human Genetics</i> , 2020, 107, 743-752.	2.6	76
20	Professional duties are now considered legal duties of care within genomic medicine. <i>European Journal of Human Genetics</i> , 2020, 28, 1301-1304.	1.4	5
21	Societal considerations in host genome testing for COVID-19. <i>Genetics in Medicine</i> , 2020, 22, 1464-1466.	1.1	14
22	We need to think about data governance for dementia research in a digital era. <i>Alzheimer's Research and Therapy</i> , 2020, 12, 17.	3.0	6
23	Trust in genomic data sharing among members of the general public in the UK, USA, Canada and Australia. <i>Human Genetics</i> , 2019, 138, 1237-1246.	1.8	69
24	Should doctors have a legal duty to warn relatives of their genetic risks?. <i>Lancet, The</i> , 2019, 394, 2133-2135.	6.3	9
25	Attitudes of publics who are unwilling to donate DNA data for research. <i>European Journal of Medical Genetics</i> , 2019, 62, 316-323.	0.7	53
26	Strategies to improve recruitment of people with dementia to research studies. <i>Dementia</i> , 2019, 18, 2494-2504.	1.0	50
27	At, with and beyond risk: expectations of living with the possibility of future dementia. <i>Sociology of Health and Illness</i> , 2018, 40, 969-987.	1.1	31
28	From people with dementia to people with data: Participation and value in Alzheimer's disease research. <i>BioSocieties</i> , 2018, 13, 623-639.	0.8	10
29	Advances in Alzheimer's imaging are changing the experience of Alzheimer's disease. <i>Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring</i> , 2018, 10, 285-300.	1.2	27
30	Research participants as collaborators: Background, experience and policies from the PREVENT Dementia and EPAD programmes. <i>Dementia</i> , 2018, 17, 1045-1054.	1.0	13
31	Perspectives on Communicating Biomarker-Based Assessments of Alzheimer's Disease to Cognitively Healthy Individuals. <i>Journal of Alzheimer's Disease</i> , 2018, 62, 487-498.	1.2	38
32	On the personal utility of Alzheimer's disease-related biomarker testing in the research context. <i>Journal of Medical Ethics</i> , 2018, 44, 830-834.	1.0	27
33	Expanding engagement with the ethical implications of changing definitions of Alzheimer's disease. <i>Lancet Psychiatry</i> , 2017, 4, e6-e7.	3.7	8
34	[P4377]: EXPECTATIONS RELATED TO THE IMPLICATIONS OF LEARNING ALZHEIMER'S DISEASE RISK AMONG COGNITIVELY HEALTHY RESEARCH PARTICIPANTS. <i>Alzheimer's and Dementia</i> , 2017, 13, P1437.	0.4	0
35	In search of lost time: age and the promise of induced pluripotent stem cell models of the brain. <i>New Genetics and Society</i> , 2016, 35, 393-408.	0.7	8
36	Arbiters of Waste: Date Labels, the Consumer and Knowing Good, Safe Food. <i>Sociological Review</i> , 2012, 60, 84-101.	0.9	41

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37	Pharmaceutical prospects: Biopharming and the geography of technological expectations. <i>Social Studies of Science</i> , 2012, 42, 290-306.	1.5	24
38	Of Markets and Medicines. <i>Science As Culture</i> , 2011, 20, 121-126.	2.4	0
39	Fraught cuisine: food scares and the modulation of anxieties. <i>Distinktion</i> , 2011, 12, 177-192.	0.8	13
40	A focus group study of food safety practices in relation to listeriosis among the over-60s. <i>Critical Public Health</i> , 2011, 21, 485-495.	1.4	19
41	Drawing Bright Lines: Food and the Futures of Biopharming. <i>Sociological Review</i> , 2010, 58, 133-151.	0.9	6