

# Mahsa Shabani

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/1018527/publications.pdf>

Version: 2024-02-01

38  
papers

982  
citations

430874

18  
h-index

501196

28  
g-index

39  
all docs

39  
docs citations

39  
times ranked

1164  
citing authors

#	ARTICLE	IF	CITATIONS
1	Credit and Recognition for Contributions to Data-Sharing Platforms Among Cohort Holders and Platform Developers in Europe: Interview Study. <i>Journal of Medical Internet Research</i> , 2022, 24, e25983.	4.3	10
2	Will the European Health Data Space change data sharing rules?. <i>Science</i> , 2022, 375, 1357-1359.	12.6	24
3	Data sharing platforms: instruments to inform and shape science policy on data sharing?. <i>Scientometrics</i> , 2022, 127, 3007-3019.	3.0	7
4	The Data Governance Act and the EU's move towards facilitating data sharing. <i>Molecular Systems Biology</i> , 2021, 17, e10229.	7.2	12
5	Ethics review of big data research: What should stay and what should be reformed?. <i>BMC Medical Ethics</i> , 2021, 22, 51.	2.4	39
6	Data Sharing in Biomedical Sciences: A Systematic Review of Incentives. <i>Biopreservation and Biobanking</i> , 2021, 19, 219-227.	1.0	20
7	Editorial: Personal Genomes: Accessing, Sharing, and Interpretation. <i>Frontiers in Genetics</i> , 2021, 12, 687584.	2.3	1
8	Factors that influence data sharing through data sharing platforms: A qualitative study on the views and experiences of cohort holders and platform developers. <i>PLoS ONE</i> , 2021, 16, e0254202.	2.5	15
9	The Impact of the GDPR on the Governance of Biobank Research. <i>Law, Governance and Technology Series</i> , 2021, , 45-60.	0.4	11
10	Toward better governance of human genomic data. <i>Nature Genetics</i> , 2021, 53, 2-8.	21.4	31
11	Genomic Sequencing Capacity, Data Retention, and Personal Access to Raw Data in Europe. <i>Frontiers in Genetics</i> , 2020, 11, 303.	2.3	9
12	Review of policies of companies and databases regarding access to customers' genealogy data for law enforcement purposes. <i>Personalized Medicine</i> , 2020, 17, 141-153.	1.5	23
13	Reporting, recording, and communication of COVID-19 cases in workplace: data protection as a moving target. <i>Journal of Law and the Biosciences</i> , 2020, 7, lsa008.	1.6	2
14	Data sharing platforms and the academic evaluation system. <i>EMBO Reports</i> , 2020, 21, e50690.	4.5	9
15	The Use of Samples Originating From Doping Control Procedures for Research Purposes: A Qualitative Study. <i>Journal of Empirical Research on Human Research Ethics</i> , 2019, 14, 254-261.	1.3	1
16	Transparency and objectivity in governance of clinical trials data sharing: Current practices and approaches. <i>Clinical Trials</i> , 2019, 16, 547-551.	1.6	11
17	Re-identifiability of genomic data and the <sc>GDPR</sc>. <i>EMBO Reports</i> , 2019, 20, .	4.5	49
18	Consent insufficient for data release. <i>Science</i> , 2019, 364, 445-446.	12.6	9

#	ARTICLE	IF	CITATIONS
19	DNA Data Marketplace: An Analysis of the Ethical Concerns Regarding the Participation of the Individuals. <i>Frontiers in Genetics</i> , 2019, 10, 1107.	2.3	37
20	Blockchain-based platforms for genomic data sharing: a de-centralized approach in response to the governance problems?. <i>Journal of the American Medical Informatics Association: JAMIA</i> , 2019, 26, 76-80.	4.4	69
21	Variant data sharing by clinical laboratories through public databases: consent, privacy and further contact for research policies. <i>Genetics in Medicine</i> , 2019, 21, 1031-1037.	2.4	12
22	Forensic Epigenetic Age Estimation and Beyond: Ethical and Legal Considerations. <i>Trends in Genetics</i> , 2018, 34, 489-491.	6.7	24
23	The challenges of the expanded availability of genomic information: an agenda-setting paper. <i>Journal of Community Genetics</i> , 2018, 9, 103-116.	1.2	45
24	Rules for processing genetic data for research purposes in view of the new EU General Data Protection Regulation. <i>European Journal of Human Genetics</i> , 2018, 26, 149-156.	2.8	95
25	Raw Genomic Data: Storage, Access, and Sharing. <i>Trends in Genetics</i> , 2018, 34, 8-10.	6.7	18
26	Registered access: authorizing data access. <i>European Journal of Human Genetics</i> , 2018, 26, 1721-1731.	2.8	33
27	Oversight of Genomic Data Sharing: What Roles for Ethics and Data Access Committees?. <i>Biopreservation and Biobanking</i> , 2017, 15, 469-474.	1.0	25
28	Research ethics review for the use of anonymized samples and data: A systematic review of normative documents. <i>Accountability in Research</i> , 2017, 24, 483-496.	2.4	12
29	Registered access: a "Triple-A" approach. <i>European Journal of Human Genetics</i> , 2016, 24, 1676-1680.	2.8	33
30	Who should have access to genomic data and how should they be held accountable? Perspectives of Data Access Committee members and experts. <i>European Journal of Human Genetics</i> , 2016, 24, 1671-1675.	2.8	15
31	Ethics review for international data-intensive research. <i>Science</i> , 2016, 351, 1399-1400.	12.6	44
32	"You want the right amount of oversight" interviews with data access committee members and experts on genomic data access. <i>Genetics in Medicine</i> , 2016, 18, 892-897.	2.4	32
33	From the principles of genomic data sharing to the practices of data access committees. <i>EMBO Molecular Medicine</i> , 2015, 7, 507-509.	6.9	51
34	Challenges of web-based personal genomic data sharing. <i>Life Sciences, Society and Policy</i> , 2015, 11, 3.	3.2	26
35	Controlled Access under Review: Improving the Governance of Genomic Data Access. <i>PLoS Biology</i> , 2015, 13, e1002339.	5.6	42
36	Attitudes of research participants and the general public towards genomic data sharing: a systematic literature review. <i>Expert Review of Molecular Diagnostics</i> , 2014, 14, 1053-1065.	3.1	82

#	ARTICLE	IF	CITATIONS
37	An agenda-setting paper on data sharing platforms: euCanSHare workshop. Open Research Europe, 0, 1, 80.	2.0	3
38	An agenda-setting paper on data sharing platforms: euCanSHare workshop. Open Research Europe, 0, 1, 80.	2.0	1