

Vasiliki Rahimzadeh

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/8874437/publications.pdf>

Version: 2024-02-01

38
papers

274
citations

1162889

8
h-index

1058333

14
g-index

43
all docs

43
docs citations

43
times ranked

623
citing authors

#	ARTICLE	IF	CITATIONS
1	The Serious Factor in Expanded Prenatal Genetic Testing. American Journal of Bioethics, 2022, 22, 23-25.	0.5	7
2	Ethical and epistemic issues in the design and conduct of pragmatic stepped-wedge cluster randomized clinical trials. Contemporary Clinical Trials, 2022, 115, 106703.	0.8	1
3	Promoting Ethical Deployment of Artificial Intelligence and Machine Learning in Healthcare. American Journal of Bioethics, 2022, 22, 4-7.	0.5	2
4	Institutional Review Board Use of Outside Experts: A National Survey. AJOB Empirical Bioethics, 2022, 13, 251-262.	0.8	8
5	Insights for Teaching During a Pandemic: Lessons From a Pre-COVID-19 International Synchronous Hybrid Learning Experience. Family Medicine, 2022, 54, 471-476.	0.3	0
6	Regulatory Angels and Technology Demons? Making Sense of Evolving Realities in Health Data Privacy for the Digital Age. American Journal of Bioethics, 2022, 22, 68-70.	0.5	1
7	Automating Justice: An Ethical Responsibility of Computational Bioethics. American Journal of Bioethics, 2022, 22, 30-33.	0.5	6
8	Leveraging Algorithms to Improve Decision-Making Workflows for Genomic Data Access and Management. Biopreservation and Biobanking, 2022, 20, 429-435.	0.5	6
9	Pros and cons of present as an alternative to traditional consent in medical research. Journal of Medical Ethics, 2021, 47, 251-252.	1.0	5
10	Frontline Ethico-Legal Issues in Childhood Cancer Genetics Research. , 2021, , 387-414.		1
11	Communication of Pharmacogenomic test results and treatment plans in pediatric oncology: deliberative stakeholder consultations with parents. BMC Palliative Care, 2021, 20, 15.	0.8	3
12	A policy Delphi study to validate the key implications of data sharing (KIDS) framework for pediatric genomics in Canada. BMC Medical Ethics, 2021, 22, 71.	1.0	1
13	Assessing the quality of deliberative stakeholder consultations involving allied health professionals in pediatric palliative care and hematology/oncology in Canada. BMC Palliative Care, 2021, 20, 189.	0.8	0
14	Ethical, Legal, and Social Issues (ELSI) of Responsible Data Sharing Involving Children in Genomics: A Systematic Literature Review of Reasons. AJOB Empirical Bioethics, 2020, 11, 233-245.	0.8	7
15	Interactive role-playing and health-related quality of life assessment in children with neurocognitive sequelae: A global neuroethics research approach. , 2020, , 165-187.		0
16	Biobanking for Genomic and Personalized Health Research: Participant Perceptions and Preferences. Biopreservation and Biobanking, 2020, 18, 204-212.	0.5	15
17	A Policy and Practice Review of Consumer Protections and Their Application to Hospital-Sourced Data Aggregation and Analytics by Third-Party Companies. Frontiers in Big Data, 2020, 3, 603044.	1.8	3
18	Understanding how professionals cultures impact implementation of a pediatric oncology genomic test. Journal of Health Organization and Management, 2019, 33, 919-928.	0.6	2

#	ARTICLE	IF	CITATIONS
19	Involving children with neurodevelopmental disorders in biomedical research. <i>The Lancet Child and Adolescent Health</i> , 2019, 3, 143-144.	2.7	4
20	Open science precision medicine in Canada: Points to consider. <i>Facets</i> , 2019, 4, 1-19.	1.1	7
21	Key Implications of Data Sharing in Pediatric Genomics. <i>JAMA Pediatrics</i> , 2018, 172, 476.	3.3	29
22	Deliberative stakeholder consultations: creating insights into effective practice-change in family medicine. <i>Family Practice</i> , 2018, 35, 749-752.	0.8	4
23	The sIRB System: A Single Beacon of Progress in the Revised Common Rule?. <i>American Journal of Bioethics</i> , 2017, 17, 43-46.	0.5	9
24	Policies and practices of data-intensive primary care in the precision-medicine era. <i>Internal Medicine Review (Washington, D C: Online)</i> , 2017, 3, .	0.3	2
25	P3G: Towards an International Policy Platform for Population Genomics. , 2017, , 155-167.		0
26	The Raison d'Être of Mutual Recognition: An Analysis of the 2015 Reform to Research Ethics Review Policies, Processes and Problems in Québec. <i>Health Reform Observer - Observatoire Des Riformes De Santé</i> , 2017, 5, .	0.4	0
27	Legal and Ethical Implications of Data Sharing in International Biobanking Research: Toward a Global Response. <i>Biopreservation and Biobanking</i> , 2016, 14, 193-194.	0.5	4
28	How mutually recognizable is mutual recognition? An international terminology index of research ethics review policies in the USA, Canada, UK and Australia. <i>Personalized Medicine</i> , 2016, 13, 101-105.	0.8	5
29	An International Framework for Data Sharing: Moving Forward with the Global Alliance for Genomics and Health. <i>Biopreservation and Biobanking</i> , 2016, 14, 256-259.	0.5	35
30	Uncertainty and innovation: Understanding the role of cell-based manufacturing facilities in shaping regulatory and commercialization environments. <i>Applied & Translational Genomics</i> , 2016, 11, 27-39.	2.1	15
31	Addressing ethical challenges at the intersection of pharmacogenomics and primary care using deliberative consultations. <i>Pharmacogenomics</i> , 2016, 17, 1795-1805.	0.6	13
32	Promoting an ethic of engagement in pediatric palliative care research. <i>BMC Palliative Care</i> , 2015, 14, 50.	0.8	12
33	Streamlining review of research involving humans: Canadian models: Table 1. <i>Journal of Medical Genetics</i> , 2015, 52, 566-569.	1.5	7
34	Statement of principles on the return of research results and incidental findings in paediatric research: a multi-site consultative process. <i>Genome</i> , 2015, 58, 541-548.	0.9	25
35	To disclose, or not to disclose? Context matters. <i>European Journal of Human Genetics</i> , 2015, 23, 279-284.	1.4	15
36	Phasing Out Voluntary Donation in Clinical Trials: The Ethics of Mandating Biospecimen Collection. <i>Journal of Clinical Trials</i> , 2015, 05, .	0.1	0

#	ARTICLE	IF	CITATIONS
37	Genetics and primary care: where are we headed?. Journal of Translational Medicine, 2014, 12, 238.	1.8	20
38	Primary Care and the Ethics of Integrating Genomic Medicine. International Journal of Whole Person Care, 2014, 1, .	0.0	0