

# Bartha M Knoppers

## List of Publications by Year in descending order

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Version: 2024-02-01

239  
papers

8,994  
citations

47409

49  
h-index

66518

82  
g-index

245  
all docs

245  
docs citations

245  
times ranked

13347  
citing authors

#	ARTICLE	IF	CITATIONS
1	Ethical challenges of precision cancer medicine. <i>Seminars in Cancer Biology</i> , 2022, 84, 263-270.	4.3	8
2	Genetic counselors and legal recognition: A made-for-Canada approach. <i>Journal of Genetic Counseling</i> , 2022, 31, 49-58.	0.9	9
3	Artificial Intelligence in Cardiovascular Imaging: "Unexplainable" Legal and Ethical Challenges?. <i>Canadian Journal of Cardiology</i> , 2022, 38, 225-233.	0.8	14
4	The international data governance landscape. <i>Journal of Law and the Biosciences</i> , 2022, 9, Isac005.	0.8	11
5	The Serious Factor in Expanded Prenatal Genetic Testing. <i>American Journal of Bioethics</i> , 2022, 22, 23-25.	0.5	7
6	Re-contact Following Withdrawal of Minors from Research. <i>Canadian Journal of Bioethics</i> , 2022, 5, 45.	0.0	1
7	Genomics4RD: An integrated platform to share Canadian deep-phenotype and multiomic data for international rare disease gene discovery.. <i>Human Mutation</i> , 2022, , .	1.1	4
8	Gynecologic Cancer Risk and Genetics: Informing an Ideal Model of Gynecologic Cancer Prevention. <i>Current Oncology</i> , 2022, 29, 4632-4646.	0.9	1
9	Raising standards for global data-sharing"Response. <i>Science</i> , 2021, 371, 134-135.	6.0	0
10	Frontline Ethico-Legal Issues in Childhood Cancer Genetics Research. , 2021, , 387-414.		1
11	Women's Views on Multifactorial Breast Cancer Risk Assessment and Risk-Stratified Screening: A Population-Based Survey from Four Provinces in Canada. <i>Journal of Personalized Medicine</i> , 2021, 11, 95.	1.1	28
12	Parental Access to Children's Raw Genomic Data in Canada: Legal Rights and Professional Responsibility. <i>Frontiers in Genetics</i> , 2021, 12, 535340.	1.1	2
13	Coming Out to Play: Privacy, Data Protection, Children's Health, and COVID-19 Research. <i>Frontiers in Genetics</i> , 2021, 12, 659027.	1.1	8
14	The Equitable Implementation of Cystic Fibrosis Personalized Medicines in Canada. <i>Journal of Personalized Medicine</i> , 2021, 11, 382.	1.1	2
15	Three decades of genetic privacy: a metaphoric journey. <i>Human Molecular Genetics</i> , 2021, 30, R156-R160.	1.4	8
16	Personalized Risk Assessment for Prevention and Early Detection of Breast Cancer: Integration and Implementation (PERSPECTIVE I&I). <i>Journal of Personalized Medicine</i> , 2021, 11, 511.	1.1	59
17	A policy Delphi study to validate the key implications of data sharing (KIDS) framework for pediatric genomics in Canada. <i>BMC Medical Ethics</i> , 2021, 22, 71.	1.0	1
18	Longitudinal Health Studies: Secondary Uses Serving the Future. <i>Biopreservation and Biobanking</i> , 2021, 19, 404-413.	0.5	3

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19	Risk-Stratified Approach to Breast Cancer Screening in Canada: Women's Knowledge of the Legislative Context and Concerns about Discrimination from Genetic and Other Predictive Health Data. <i>Journal of Personalized Medicine</i> , 2021, 11, 726.	1.1	5
20	Of Screening, Stratification, and Scores. <i>Journal of Personalized Medicine</i> , 2021, 11, 736.	1.1	13
21	Ethical and Analytic Challenges With Genomic Sequencing of Relapsed Hematologic Malignancies Following Allogeneic Hematopoietic Stem-Cell Transplantation. <i>JCO Precision Oncology</i> , 2021, 5, 1339-1347.	1.5	2
22	A marathon, not a sprint – neuroimaging, Open Science and ethics. <i>NeuroImage</i> , 2021, 236, 118041.	2.1	14
23	Should Age-Dependent Absolute Risk Thresholds Be Used for Risk Stratification in Risk-Stratified Breast Cancer Screening?. <i>Journal of Personalized Medicine</i> , 2021, 11, 916.	1.1	8
24	Toward better governance of human genomic data. <i>Nature Genetics</i> , 2021, 53, 2-8.	9.4	31
25	GA4GH: International policies and standards for data sharing across genomic research and healthcare. <i>Cell Genomics</i> , 2021, 1, 100029.	3.0	94
26	Computational tools for genomic data de-identification: facilitating data protection law compliance. <i>Nature Communications</i> , 2021, 12, 6949.	5.8	3
27	A centralized rare disease database and whole-genome sequencing as a standard of care: two essential implementations for the future of health. <i>Facets</i> , 2021, 6, 1831-1834.	1.1	1
28	When information is the treatment? Precision medicine in healthcare. <i>Healthcare Management Forum</i> , 2020, 33, 120-125.	0.6	0
29	How to fix the GDPR's frustration of global biomedical research. <i>Science</i> , 2020, 370, 40-42.	6.0	29
30	The Right to Benefit from Science and Its Implications for Genomic Data Sharing. <i>European Journal of International Law</i> , 2020, 31, 665-691.	0.2	8
31	Ethical, Legal, and Social Issues (ELSI) of Responsible Data Sharing Involving Children in Genomics: A Systematic Literature Review of Reasons. <i>AJOB Empirical Bioethics</i> , 2020, 11, 233-245.	0.8	7
32	Of the Rights and Best Interests of Future Generations. <i>American Journal of Bioethics</i> , 2020, 20, 38-40.	0.5	0
33	Modeling consent in the time of COVID-19. <i>Journal of Law and the Biosciences</i> , 2020, 7, Isaa020.	0.8	9
34	The Genetic Family as Patient?. <i>American Journal of Bioethics</i> , 2020, 20, 77-80.	0.5	8
35	International mHealth Research: Old Tools and New Challenges. <i>Journal of Law, Medicine and Ethics</i> , 2020, 48, 178-186.	0.4	6
36	How Can We Not Waste Legacy Genomic Research Data?. <i>Frontiers in Genetics</i> , 2020, 11, 446.	1.1	11

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37	Doping controls and the "Mature Minor"™ elite athlete: towards clarification?. <i>International Journal of Sport Policy and Politics</i> , 2020, 12, 179-187.	1.0	9
38	The Provision of Genetic Testing and Related Services in Quebec, Canada. <i>Frontiers in Genetics</i> , 2020, 11, 127.	1.1	13
39	Pandemics, privacy, and public health research. <i>Canadian Journal of Public Health</i> , 2020, 111, 454-457.	1.1	9
40	Genomics: data sharing needs an international code of conduct. <i>Nature</i> , 2020, 578, 31-33.	13.7	31
41	Oversight, governance, and policy for making decisions about return of individual genomic findings. , 2020, , 29-41.		1
42	Biobanking for Genomic and Personalized Health Research: Participant Perceptions and Preferences. <i>Biopreservation and Biobanking</i> , 2020, 18, 204-212.	0.5	15
43	"Serious"™ factor"™ a relevant starting point for further debate: a response. <i>Journal of Medical Ethics</i> , 2020, 46, 153-155.	1.0	4
44	Bringing Code to Data: Do Not Forget Governance. <i>Journal of Medical Internet Research</i> , 2020, 22, e18087.	2.1	11
45	Genome-wide sequencing in acutely ill infants: genomic medicine™s critical application?. <i>Genetics in Medicine</i> , 2019, 21, 498-504.	1.1	42
46	Development of a consent resource for genomic data sharing in the clinical setting. <i>Genetics in Medicine</i> , 2019, 21, 81-88.	1.1	20
47	Privacy-Preserving Linkage of Genomic and Clinical Data Sets. <i>IEEE/ACM Transactions on Computational Biology and Bioinformatics</i> , 2019, 16, 1342-1348.	1.9	18
48	Model consent clauses for rare disease research. <i>BMC Medical Ethics</i> , 2019, 20, 55.	1.0	16
49	Regulatory Landscape of International Direct-to-Participant (DTP) Genomic Research: Time to Untie the Gordian Knot?. <i>Journal of Law, Medicine and Ethics</i> , 2019, 47, 336-341.	0.4	2
50	The Human Right to Science and the Regulation of Human Germline Engineering. <i>CRISPR Journal</i> , 2019, 2, 134-142.	1.4	12
51	A Tale of Two Capacities: Including Children and Decisionally Vulnerable Adults in Biomedical Research. <i>Frontiers in Genetics</i> , 2019, 10, 289.	1.1	16
52	Canada's Assisted Human Reproduction Act: Pragmatic Reforms in Support of Research. <i>Frontiers in Medicine</i> , 2019, 6, 157.	1.2	7
53	Heritable Genome Editing: Who Speaks for "Future"™ Children?. <i>CRISPR Journal</i> , 2019, 2, 285-292.	1.4	14
54	Reciprocity and the Quest for Meaningful Disclosure. <i>American Journal of Bioethics</i> , 2019, 19, 36-38.	0.5	3

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55	Return of individual genomic research results: are laws and policies keeping step?. <i>European Journal of Human Genetics</i> , 2019, 27, 535-546.	1.4	73
56	Whose Commons? Data Protection as a Legal Limit of Open Science. <i>Journal of Law, Medicine and Ethics</i> , 2019, 47, 106-111.	0.4	12
57	Importance of Participant-Centricity and Trust for a Sustainable Medical Information Commons. <i>Journal of Law, Medicine and Ethics</i> , 2019, 47, 12-20.	0.4	20
58	Health professionals' perspectives on breast cancer risk stratification: understanding evaluation of risk versus screening for disease. <i>Public Health Reviews</i> , 2019, 40, 2.	1.3	27
59	Letter: Relearning the 3 R's? Reinterpretation, recontact, and return of genetic variants. <i>Genetics in Medicine</i> , 2019, 21, 2401-2402.	1.1	9
60	'CRISPR babies': What does this mean for science and Canada?. <i>Cmaj</i> , 2019, 191, E91-E92.	0.9	7
61	A response to 'Personalised medicine and population health: breast and ovarian cancer'. <i>Human Genetics</i> , 2019, 138, 287-289.	1.8	14
62	The 'serious' factor in germline modification. <i>Journal of Medical Ethics</i> , 2019, 45, 508-513.	1.0	15
63	Biotechnologies nibbling at the legal 'human'. <i>Science</i> , 2019, 366, 1455-1457.	6.0	11
64	Pre-implantation Genetic Diagnosis: The Road Forward in Canada. <i>Journal of Obstetrics and Gynaecology Canada</i> , 2019, 41, 68-71.	0.3	7
65	Organizational challenges to equity in the delivery of services within a new personalized risk-based approach to breast cancer screening. <i>New Genetics and Society</i> , 2019, 38, 38-59.	0.7	12
66	Envisioning Implementation of a Personalized Approach in Breast Cancer Screening Programs: Stakeholder Perspectives. <i>Healthcare Policy</i> , 2019, 15, 39-54.	0.3	12
67	The Genomic Commons. <i>Annual Review of Genomics and Human Genetics</i> , 2018, 19, 429-453.	2.5	26
68	Key Implications of Data Sharing in Pediatric Genomics. <i>JAMA Pediatrics</i> , 2018, 172, 476.	3.3	29
69	Responsible sharing of biomedical data and biospecimens via the 'Automatable Discovery and Access Matrix' (ADA-M). <i>Npj Genomic Medicine</i> , 2018, 3, 17.	1.7	38
70	Overcoming barriers to facilitate the regulation of multi-centre regenerative medicine clinical trials. <i>Stem Cell Research and Therapy</i> , 2018, 9, 307.	2.4	25
71	Bridging stem cell research and medicine: a learning health system. <i>Regenerative Medicine</i> , 2018, 13, 741-752.	0.8	11
72	Research on Human Embryos and Reproductive Materials: Revisiting Canadian Law and Policy. <i>Healthcare Policy</i> , 2018, 13, 10-19.	0.3	7

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73	BRCA Challenge: BRCA Exchange as a global resource for variants in BRCA1 and BRCA2. PLoS Genetics, 2018, 14, e1007752.	1.5	148
74	Genetic database software as medical devices. Human Mutation, 2018, 39, 1702-1712.	1.1	10
75	Ethical, Legal, and Regulatory Issues for the Implementation of Omics-Based Risk Prediction of Women's Cancer: Points to Consider. Public Health Genomics, 2018, 21, 37-44.	0.6	13
76	Genetically Enhanced Minors: Whose Responsibility?. American Journal of Bioethics, 2018, 18, 1-3.	0.5	2
77	Registered access: authorizing data access. European Journal of Human Genetics, 2018, 26, 1721-1731.	1.4	33
78	Ethics, big data and computing in epidemiology and public health. Annals of Epidemiology, 2017, 27, 297-301.	0.9	61
79	International Cooperation to Enable the Diagnosis of All Rare Genetic Diseases. American Journal of Human Genetics, 2017, 100, 695-705.	2.6	305
80	“Matching” consent to purpose: The example of the Matchmaker Exchange. Human Mutation, 2017, 38, 1281-1285.	1.1	13
81	Human gene editing: revisiting Canadian policy. Npj Regenerative Medicine, 2017, 2, 3.	2.5	14
82	Reply to C Harling. European Journal of Human Genetics, 2017, 25, 1030-1030.	1.4	0
83	Comparative Approaches to Genetic Discrimination: Chasing Shadows?. Trends in Genetics, 2017, 33, 299-302.	2.9	63
84	Criminal Prohibition of Wrongful Re-identification: Legal Solution or Minefield for Big Data?. Journal of Bioethical Inquiry, 2017, 14, 527-539.	0.9	17
85	Oversight of Genomic Data Sharing: What Roles for Ethics and Data Access Committees?. Biopreservation and Biobanking, 2017, 15, 469-474.	0.5	25
86	Mitochondrial Replacement Therapy: The Road to the Clinic in Canada. Journal of Obstetrics and Gynaecology Canada, 2017, 39, 916-918.	0.3	10
87	Ethics and Big Data in health. Current Opinion in Systems Biology, 2017, 4, 53-57.	1.3	84
88	Data Sharing and Privacy. , 2017, , 143-160.		2
89	The Gatekeeping Function in Personalized Medicine Initiatives. Current Pharmacogenomics and Personalized Medicine, 2017, 14, 36-49.	0.2	0
90	Population Biobanks and the Principle of Reciprocity. , 2017, , 99-109.		1

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91	Ethics approval in applications for open-access clinical trial data: An analysis of researcher statements to <a href="http://clinicalstudydatarequest.com">clinicalstudydatarequest.com</a> . <i>PLoS ONE</i> , 2017, 12, e0184491.	1.1	7
92	P3G: Towards an International Policy Platform for Population Genomics. , 2017, , 155-167.		0
93	Consent Codes: Upholding Standard Data Use Conditions. <i>PLoS Genetics</i> , 2016, 12, e1005772.	1.5	65
94	Precision medicine: a matter of regulation or collaboration?. <i>Journal of Law and the Biosciences</i> , 2016, 3, 687-690.	0.8	2
95	Don't Take It Personal: European Union Legal Aspects of Procuring and Protecting Environmental Exposure Data in Population Biobanks Through the Use of a Geo-Information-Systems Toolkit. <i>Biopreservation and Biobanking</i> , 2016, 14, 217-223.	0.5	3
96	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
97	Registered access: a "Triple-A"™ approach. <i>European Journal of Human Genetics</i> , 2016, 24, 1676-1680.	1.4	33
98	A step forward for data protection and biomedical research. <i>Lancet, The</i> , 2016, 387, 1374-1375.	6.3	11
99	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
100	Facilitating a culture of responsible and effective sharing of cancer genome data. <i>Nature Medicine</i> , 2016, 22, 464-471.	15.2	83
101	Do It Yourself Newborn Screening. <i>JAMA Pediatrics</i> , 2016, 170, 523.	3.3	7
102	Clinical Sequencing Exploratory Research Consortium: Accelerating Evidence-Based Practice of Genomic Medicine. <i>American Journal of Human Genetics</i> , 2016, 98, 1051-1066.	2.6	137
103	Next-Generation Sequencing and the Return of Results. <i>Cold Spring Harbor Perspectives in Medicine</i> , 2016, 6, a026724.	2.9	8
104	Streamlining ethical review of data intensive research. <i>BMJ, The</i> , 2016, 354, i4181.	3.0	9
105	Legal approaches regarding health-care decisions involving minors: implications for next-generation sequencing. <i>European Journal of Human Genetics</i> , 2016, 24, 1559-1564.	1.4	12
106	Are Data Sharing and Privacy Protection Mutually Exclusive?. <i>Cell</i> , 2016, 167, 1150-1154.	18.5	50
107	Data Sharing "Is the Juice Worth the Squeeze?". <i>New England Journal of Medicine</i> , 2016, 375, 1608-1609.	13.9	49
108	The discombobulation of de-identification. <i>Nature Biotechnology</i> , 2016, 34, 1102-1103.	9.4	20

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109	Sharing health-related data: a privacy test?. Npj Genomic Medicine, 2016, 1, 160241-160246.	1.7	34
110	Access Governance for Biobanks: The Case of the BioSHaRE-EU Cohorts. Biopreservation and Biobanking, 2016, 14, 201-206.	0.5	20
111	A decision tool to guide the ethics review of a challenging breed of emerging genomic projects. European Journal of Human Genetics, 2016, 24, 1099-1103.	1.4	0
112	Ethics review for international data-intensive research. Science, 2016, 351, 1399-1400.	6.0	44
113	Population genetic testing for cancer susceptibility: founder mutations to genomes. Nature Reviews Clinical Oncology, 2016, 13, 41-54.	12.5	86
114	Exposing participants? Population biobanks go geo. European Journal of Human Genetics, 2016, 24, 155-156.	1.4	5
115	The Adoption of Cloud Computing in the Field of Genomics Research: The Influence of Ethical and Legal Issues. PLoS ONE, 2016, 11, e0164347.	1.1	22
116	Influence des facteurs organisationnels sur lâ€™implantation dâ€™une approche personnalisÃ©e de dÃ©pistage du cancer du sein. Sante Publique, 2016, Vol. 28, 353-361.	0.0	4
117	Data Safe Havens in health research and healthcare. Bioinformatics, 2015, 31, 3241-3248.	1.8	58
118	The ethics weathervane. BMC Medical Ethics, 2015, 16, 58.	1.0	8
119	From the principles of genomic data sharing to the practices of data access committees. EMBO Molecular Medicine, 2015, 7, 507-509.	3.3	51
120	Ethical challenges and innovations in the dissemination of genomic data: the experience of the PERSPECTIVE project. Advances in Genomics and Genetics, 2015, , 283.	0.8	1
121	Harmonizing Privacy Laws to Enable International Biobank Research. Journal of Law, Medicine and Ethics, 2015, 43, 673-674.	0.4	12
122	Data analysis: Create a cloud commons. Nature, 2015, 523, 149-151.	13.7	89
123	Streamlining review of research involving humans: Canadian models: TableÂ¹. Journal of Medical Genetics, 2015, 52, 566-569.	1.5	7
124	Statement of principles on the return of research results and incidental findings in paediatric research: a multi-site consultative process. Genome, 2015, 58, 541-548.	0.9	25
125	Whole-genome sequencing in newborn screening? A statement on the continued importance of targeted approaches in newborn screening programmes. European Journal of Human Genetics, 2015, 23, 1593-1600.	1.4	87
126	Return of genetic testing results in the era of whole-genome sequencing. Nature Reviews Genetics, 2015, 16, 553-559.	7.7	127



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127	International Charter of principles for sharing bio-specimens and data. <i>European Journal of Human Genetics</i> , 2015, 23, 721-728.	1.4	112
128	CRISPR germline engineering—the community speaks. <i>Nature Biotechnology</i> , 2015, 33, 478-486.	9.4	110
129	Oversight of human inheritable genome modification. <i>Nature Biotechnology</i> , 2015, 33, 454-455.	9.4	17
130	Genomic cloud computing: legal and ethical points to consider. <i>European Journal of Human Genetics</i> , 2015, 23, 1271-1278.	1.4	80
131	Return of whole-genome sequencing results in paediatric research: a statement of the P3G international paediatrics platform. <i>European Journal of Human Genetics</i> , 2014, 22, 3-5.	1.4	61
132	Does policy grow on trees?. <i>BMC Medical Ethics</i> , 2014, 15, 87.	1.0	4
133	Framework for responsible sharing of genomic and health-related data. <i>The HUGO Journal</i> , 2014, 8, 3.	4.1	185
134	The best interests of the child and the return of results in genetic research: international comparative perspectives. <i>BMC Medical Ethics</i> , 2014, 15, 72.	1.0	22
135	Towards an ethics safe harbor for global biomedical research. <i>Journal of Law and the Biosciences</i> , 2014, 1, 3-51.	0.8	39
136	Data sharing in large research consortia: experiences and recommendations from ENGAGE. <i>European Journal of Human Genetics</i> , 2014, 22, 317-321.	1.4	54
137	Beyond public health genomics: proposals from an international working group. <i>European Journal of Public Health</i> , 2014, 24, 877-879.	0.1	22
138	Data Sharing, Year 1 — Access to Data from Industry-Sponsored Clinical Trials. <i>New England Journal of Medicine</i> , 2014, 371, 2052-2054.	13.9	101
139	Attitudes of parents toward the return of targeted and incidental genomic research findings in children. <i>Genetics in Medicine</i> , 2014, 16, 633-640.	1.1	82
140	The Challenge of Informed Consent and Return of Results in Translational Genomics: Empirical Analysis and Recommendations. <i>Journal of Law, Medicine and Ethics</i> , 2014, 42, 344-355.	0.4	63
141	From the Right to Know to the Right Not to Know. <i>Journal of Law, Medicine and Ethics</i> , 2014, 42, 6-10.	0.4	23
142	Management Strategies for Ethics in International Research. <i>Current Genetic Medicine Reports</i> , 2014, 2, 255-260.	1.9	1
143	An implementation framework for the feedback of individual research results and incidental findings in research. <i>BMC Medical Ethics</i> , 2014, 15, 88.	1.0	39
144	Whole-Genome Sequencing in Newborn Screening Programs. <i>Science Translational Medicine</i> , 2014, 6, 229cm2.	5.8	59

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145	The ethical framing of personalized medicine. <i>Current Opinion in Allergy and Clinical Immunology</i> , 2014, 14, 404-408.	1.1	26
146	DataSHIELD: taking the analysis to the data, not the data to the analysis. <i>International Journal of Epidemiology</i> , 2014, 43, 1929-1944.	0.9	188
147	A human rights approach to an international code of conduct for genomic and clinical data sharing. <i>Human Genetics</i> , 2014, 133, 895-903.	1.8	104
148	Reporting results from whole-genome and whole-exome sequencing in clinical practice: a proposal for Canada?. <i>Journal of Medical Genetics</i> , 2014, 51, 68-70.	1.5	27
149	International ethics harmonization and the global alliance for genomics and health. <i>Genome Medicine</i> , 2014, 6, 13.	3.6	38
150	Building a data sharing model for global genomic research. <i>Genome Biology</i> , 2014, 15, 430.	3.8	37
151	Data protection and consent to biomedical research: a step forward?. <i>Lancet, The</i> , 2014, 384, 855.	6.3	11
152	Stem Cell Research Funding Policies and Dynamic Innovation: A Survey of Open Access and Commercialization Requirements. <i>Stem Cell Reviews and Reports</i> , 2014, 10, 455-471.	5.6	5
153	Special Issue "From Biobanks to the Clinic". <i>Applied &amp; Translational Genomics</i> , 2014, 3, 21-22.	2.1	1
154	Identifiability and Privacy in Pluripotent Stem Cell Research. <i>Cell Stem Cell</i> , 2014, 14, 427-430.	5.2	35
155	Anonymity 2.0: direct-to-consumer genetic testing and donor conception. <i>Fertility and Sterility</i> , 2014, 101, 630-632.	0.5	21
156	FORGE Canada Consortium: Outcomes of a 2-Year National Rare-Disease Gene-Discovery Project. <i>American Journal of Human Genetics</i> , 2014, 94, 809-817.	2.6	219
157	Population Biobanking and International Collaboration. <i>Pathobiology</i> , 2014, 81, 276-285.	1.9	13
158	Genotype-driven recruitment: a strategy whose time has come?. <i>BMC Medical Genomics</i> , 2013, 6, 19.	0.7	14
159	Recommendations for returning genomic incidental findings? We need to talk!. <i>Genetics in Medicine</i> , 2013, 15, 854-859.	1.1	272
160	An ethics safe harbor for international genomics research?. <i>Genome Medicine</i> , 2013, 5, 99.	3.6	23
161	Genomics: from persons to populations and back again. <i>Genome</i> , 2013, 56, 537-539.	0.9	4
162	Rare diseases and now rare data?. <i>Nature Reviews Genetics</i> , 2013, 14, 372-372.	7.7	19

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163	Whole-genome sequencing in health care. <i>European Journal of Human Genetics</i> , 2013, 21, 580-584.	1.4	330
164	Cohort profile of the CARTaGENE study: Quebec's population-based biobank for public health and personalized genomics. <i>International Journal of Epidemiology</i> , 2013, 42, 1285-1299.	0.9	172
165	From Tissues to Genomes. <i>G3: Genes, Genomes, Genetics</i> , 2013, 3, 1203-1204.	0.8	2
166	A P3G generic access agreement for population genomic studies. <i>Nature Biotechnology</i> , 2013, 31, 384-385.	9.4	24
167	Demystifying Biobanks. <i>Hastings Center Report</i> , 2013, 43, 4-5.	0.7	10
168	Personalized medicine and access to health care: potential for inequitable access?. <i>European Journal of Human Genetics</i> , 2013, 21, 143-147.	1.4	45
169	Population studies: return of research results and incidental findings Policy Statement. <i>European Journal of Human Genetics</i> , 2013, 21, 245-247.	1.4	94
170	Pediatric research "personalized"? International perspectives on the return of results. <i>Personalized Medicine</i> , 2013, 10, 89-95.	0.8	7
171	A new twist on an old problem: primary care physicians and results from direct-to-consumer genetic testing. <i>Personalized Medicine</i> , 2013, 10, 827-833.	0.8	5
172	Beyond ELSIs. , 2013, , 405-428.		0
173	Data Sharing in the Post-Genomic World: The Experience of the International Cancer Genome Consortium (ICGC) Data Access Compliance Office (DACO). <i>PLoS Computational Biology</i> , 2012, 8, e1002549.	1.5	100
174	International normative perspectives on the return of individual research results and incidental findings in genomic biobanks. <i>Genetics in Medicine</i> , 2012, 14, 484-489.	1.1	51
175	ELSI 2.0 for Genomics and Society. <i>Science</i> , 2012, 336, 673-674.	6.0	39
176	Power to the people: a wiki-governance model for biobanks. <i>Genome Biology</i> , 2012, 13, 158.	13.9	36
177	Managing incidental findings and research results in genomic research involving biobanks and archived data sets. <i>Genetics in Medicine</i> , 2012, 14, 361-384.	1.1	418
178	Questioning the Limits of Genomic Privacy. <i>American Journal of Human Genetics</i> , 2012, 91, 577-578.	2.6	16
179	Sampling Populations of Humans Across the World: ELSI Issues. <i>Annual Review of Genomics and Human Genetics</i> , 2012, 13, 395-413.	2.5	57
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