

# Bartha M Knoppers

## List of Publications by Year in descending order

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239  
papers

8,994  
citations

41344  
49  
h-index

58581  
82  
g-index

245  
all docs

245  
docs citations

245  
times ranked

12350  
citing authors

#	ARTICLE	IF	CITATIONS
1	Ethical challenges of precision cancer medicine. Seminars in Cancer Biology, 2022, 84, 263-270.	9.6	8
2	Genetic counselors and legal recognition: A madeâ€forâ€Canada approach. Journal of Genetic Counseling, 2022, 31, 49-58.	1.6	9
3	Artificial Intelligence in Cardiovascular Imaging: â€Unexplainableâ€Legal and Ethical Challenges?. Canadian Journal of Cardiology, 2022, 38, 225-233.	1.7	14
4	The international data governance landscape. Journal of Law and the Biosciences, 2022, 9, Isac005.	1.6	11
5	The Serious Factor in Expanded Prenatal Genetic Testing. American Journal of Bioethics, 2022, 22, 23-25.	0.9	7
6	Re-contact Following Withdrawal of Minors from Research. Canadian Journal of Bioethics, 2022, 5, 45.	0.1	1
7	Genomics4RD: An integrated platform to share Canadian deep-phenotype and multiomic data for international rare disease gene discovery.. Human Mutation, 2022, , .	2.5	4
8	Gynecologic Cancer Risk and Genetics: Informing an Ideal Model of Gynecologic Cancer Prevention. Current Oncology, 2022, 29, 4632-4646.	2.2	1
9	Raising standards for global data-sharingâ€Response. Science, 2021, 371, 134-135.	12.6	0
10	Frontline Ethico-Legal Issues in Childhood Cancer Genetics Research. , 2021, , 387-414.		1
11	Womenâ€™s Views on Multifactorial Breast Cancer Risk Assessment and Risk-Stratified Screening: A Population-Based Survey from Four Provinces in Canada. Journal of Personalized Medicine, 2021, 11, 95.	2.5	28
12	Parental Access to Children's Raw Genomic Data in Canada: Legal Rights and Professional Responsibility. Frontiers in Genetics, 2021, 12, 535340.	2.3	2
13	Coming Out to Play: Privacy, Data Protection, Childrenâ€™s Health, and COVID-19 Research. Frontiers in Genetics, 2021, 12, 659027.	2.3	8
14	The Equitable Implementation of Cystic Fibrosis Personalized Medicines in Canada. Journal of Personalized Medicine, 2021, 11, 382.	2.5	2
15	Three decades of genetic privacy: a metaphoric journey. Human Molecular Genetics, 2021, 30, R156-R160.	2.9	8
16	Personalized Risk Assessment for Prevention and Early Detection of Breast Cancer: Integration and Implementation (PERSPECTIVE I&I). Journal of Personalized Medicine, 2021, 11, 511.	2.5	59
17	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.	2.4	1
18	Longitudinal Health Studies: Secondary Uses Serving the Future. Biopreservation and Biobanking, 2021, 19, 404-413.	1.0	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. Journal of Personalized Medicine, 2021, 11, 726.	2.5	5
20	Of Screening, Stratification, and Scores. Journal of Personalized Medicine, 2021, 11, 736.	2.5	13
21	Ethical and Analytic Challenges With Genomic Sequencing of Relapsed Hematologic Malignancies Following Allogeneic Hematopoietic Stem-Cell Transplantation. JCO Precision Oncology, 2021, 5, 1339-1347.	3.0	2
22	A marathon, not a sprint – neuroimaging, Open Science and ethics. NeuroImage, 2021, 236, 118041.	4.2	14
23	Should Age-Dependent Absolute Risk Thresholds Be Used for Risk Stratification in Risk-Stratified Breast Cancer Screening?. Journal of Personalized Medicine, 2021, 11, 916.	2.5	8
24	Toward better governance of human genomic data. Nature Genetics, 2021, 53, 2-8.	21.4	31
25	GA4GH: International policies and standards for data sharing across genomic research and healthcare. Cell Genomics, 2021, 1, 100029.	6.5	94
26	Computational tools for genomic data de-identification: facilitating data protection law compliance. Nature Communications, 2021, 12, 6949.	12.8	3
27	A centralized rare disease database and whole-genome sequencing as a standard of care: two essential implementations for the future of health. Facets, 2021, 6, 1831-1834.	2.4	1
28	When information is the treatment? Precision medicine in healthcare. Healthcare Management Forum, 2020, 33, 120-125.	1.4	0
29	How to fix the GDPR's frustration of global biomedical research. Science, 2020, 370, 40-42.	12.6	29
30	The Right to Benefit from Science and Its Implications for Genomic Data Sharing. European Journal of International Law, 2020, 31, 665-691.	0.5	8
31	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.	1.6	7
32	Of the Rights and Best Interests of Future Generations. American Journal of Bioethics, 2020, 20, 38-40.	0.9	0
33	Modeling consent in the time of COVID-19. Journal of Law and the Biosciences, 2020, 7, 150-160.	1.6	9
34	The Genetic Family as Patient?. American Journal of Bioethics, 2020, 20, 77-80.	0.9	8
35	International mHealth Research: Old Tools and New Challenges. Journal of Law, Medicine and Ethics, 2020, 48, 178-186.	0.9	6
36	How Can We Not Waste Legacy Genomic Research Data?. Frontiers in Genetics, 2020, 11, 446.	2.3	11

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37	Doping controls and the “Mature Minor” elite athlete: towards clarification?. International Journal of Sport Policy and Politics, 2020, 12, 179-187.	1.6	9
38	The Provision of Genetic Testing and Related Services in Quebec, Canada. Frontiers in Genetics, 2020, 11, 127.	2.3	13
39	Pandemics, privacy, and public health research. Canadian Journal of Public Health, 2020, 111, 454-457.	2.3	9
40	Genomics: data sharing needs an international code of conduct. Nature, 2020, 578, 31-33.	27.8	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. Biopreservation and Biobanking, 2020, 18, 204-212.	1.0	15
43	“Serious” factor”a relevant starting point for further debate: a response. Journal of Medical Ethics, 2020, 46, 153-155.	1.8	4
44	Bringing Code to Data: Do Not Forget Governance. Journal of Medical Internet Research, 2020, 22, e18087.	4.3	11
45	Genome-wide sequencing in acutely ill infants: genomic medicine’s critical application?. Genetics in Medicine, 2019, 21, 498-504.	2.4	42
46	Development of a consent resource for genomic data sharing in the clinical setting. Genetics in Medicine, 2019, 21, 81-88.	2.4	20
47	Privacy-Preserving Linkage of Genomic and Clinical Data Sets. IEEE/ACM Transactions on Computational Biology and Bioinformatics, 2019, 16, 1342-1348.	3.0	18
48	Model consent clauses for rare disease research. BMC Medical Ethics, 2019, 20, 55.	2.4	16
49	Regulatory Landscape of International Direct-to-Participant (DTP) Genomic Research: Time to Untie the Gordian Knot?. Journal of Law, Medicine and Ethics, 2019, 47, 336-341.	0.9	2
50	The Human Right to Science and the Regulation of Human Germline Engineering. CRISPR Journal, 2019, 2, 134-142.	2.9	12
51	A Tale of Two Capacities: Including Children and Decisionally Vulnerable Adults in Biomedical Research. Frontiers in Genetics, 2019, 10, 289.	2.3	16
52	Canada's Assisted Human Reproduction Act: Pragmatic Reforms in Support of Research. Frontiers in Medicine, 2019, 6, 157.	2.6	7
53	Heritable Genome Editing: Who Speaks for “Future” Children?. CRISPR Journal, 2019, 2, 285-292.	2.9	14
54	Reciprocity and the Quest for Meaningful Disclosure. American Journal of Bioethics, 2019, 19, 36-38.	0.9	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.	2.8	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.	0.9	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.	0.9	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.	3.2	27
59	Letter: Relearning the 3 R's? Reinterpretation, recontact, and return of genetic variants. <i>Genetics in Medicine</i> , 2019, 21, 2401-2402.	2.4	9
60	“CRISPR babies”: What does this mean for science and Canada?. <i>Cmaj</i> , 2019, 191, E91-E92.	2.0	7
61	A response to “Personalised medicine and population health: breast and ovarian cancer”. <i>Human Genetics</i> , 2019, 138, 287-289.	3.8	14
62	The “serious” factor in germline modification. <i>Journal of Medical Ethics</i> , 2019, 45, 508-513.	1.8	15
63	Biotechnologies nibbling at the legal “human”. <i>Science</i> , 2019, 366, 1455-1457.	12.6	11
64	Pre-implantation Genetic Diagnosis: The Road Forward in Canada. <i>Journal of Obstetrics and Gynaecology Canada</i> , 2019, 41, 68-71.	0.7	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.	1.2	12
66	Envisioning Implementation of a Personalized Approach in Breast Cancer Screening Programs: Stakeholder Perspectives. <i>Healthcare Policy</i> , 2019, 15, 39-54.	0.6	12
67	The Genomic Commons. <i>Annual Review of Genomics and Human Genetics</i> , 2018, 19, 429-453.	6.2	26
68	Key Implications of Data Sharing in Pediatric Genomics. <i>JAMA Pediatrics</i> , 2018, 172, 476.	6.2	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.	3.8	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.	5.5	25
71	Bridging stem cell research and medicine: a learning health system. <i>Regenerative Medicine</i> , 2018, 13, 741-752.	1.7	11
72	Research on Human Embryos and Reproductive Materials: Revisiting Canadian Law and Policy. <i>Healthcare Policy</i> , 2018, 13, 10-19.	0.6	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.	3.5	148
74	Genetic database software as medical devices. Human Mutation, 2018, 39, 1702-1712.	2.5	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.	1.0	13
76	Genetically Enhanced Minors: Whose Responsibility?. American Journal of Bioethics, 2018, 18, 1-3.	0.9	2
77	Registered access: authorizing data access. European Journal of Human Genetics, 2018, 26, 1721-1731.	2.8	33
78	Ethics, big data and computing in epidemiology and public health. Annals of Epidemiology, 2017, 27, 297-301.	1.9	61
79	International Cooperation to Enable the Diagnosis of All Rare Genetic Diseases. American Journal of Human Genetics, 2017, 100, 695-705.	6.2	305
80	“Matching” consent to purpose: The example of the Matchmaker Exchange. Human Mutation, 2017, 38, 1281-1285.	2.5	13
81	Human gene editing: revisiting Canadian policy. Npj Regenerative Medicine, 2017, 2, 3.	5.2	14
82	Reply to C Harling. European Journal of Human Genetics, 2017, 25, 1030-1030.	2.8	0
83	Comparative Approaches to Genetic Discrimination: Chasing Shadows?. Trends in Genetics, 2017, 33, 299-302.	6.7	63
84	Criminal Prohibition of Wrongful Re-identification: Legal Solution or Minefield for Big Data?. Journal of Bioethical Inquiry, 2017, 14, 527-539.	1.5	17
85	Oversight of Genomic Data Sharing: What Roles for Ethics and Data Access Committees?. Biopreservation and Biobanking, 2017, 15, 469-474.	1.0	25
86	Mitochondrial Replacement Therapy: The Road to the Clinic in Canada. Journal of Obstetrics and Gynaecology Canada, 2017, 39, 916-918.	0.7	10
87	Ethics and Big Data in health. Current Opinion in Systems Biology, 2017, 4, 53-57.	2.6	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="https://clinicalstudydatarequest.com">clinicalstudydatarequest.com</a> . <i>PLoS ONE</i> , 2017, 12, e0184491.	2.5	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.	3.5	65
94	Precision medicine: a matter of regulation or collaboration?. <i>Journal of Law and the Biosciences</i> , 2016, 3, 687-690.	1.6	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.	1.0	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.	1.5	5
97	Registered access: a "Triple-A" approach. <i>European Journal of Human Genetics</i> , 2016, 24, 1676-1680.	2.8	33
98	A step forward for data protection and biomedical research. <i>Lancet, The</i> , 2016, 387, 1374-1375.	13.7	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.	1.0	35
100	Facilitating a culture of responsible and effective sharing of cancer genome data. <i>Nature Medicine</i> , 2016, 22, 464-471.	30.7	83
101	Do It Yourself Newborn Screening. <i>JAMA Pediatrics</i> , 2016, 170, 523.	6.2	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.	6.2	137
103	Next-Generation Sequencing and the Return of Results. <i>Cold Spring Harbor Perspectives in Medicine</i> , 2016, 6, a026724.	6.2	8
104	Streamlining ethical review of data intensive research. <i>BMJ, The</i> , 2016, 354, i4181.	6.0	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.	2.8	12
106	Are Data Sharing and Privacy Protection Mutually Exclusive?. <i>Cell</i> , 2016, 167, 1150-1154.	28.9	50
107	Data Sharing "Is the Juice Worth the Squeeze?. <i>New England Journal of Medicine</i> , 2016, 375, 1608-1609.	27.0	49
108	The discombobulation of de-identification. <i>Nature Biotechnology</i> , 2016, 34, 1102-1103.	17.5	20

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109	Sharing health-related data: a privacy test?. Npj Genomic Medicine, 2016, 1, 160241-160246.	3.8	34
110	Access Governance for Biobanks: The Case of the BioSHaRE-EU Cohorts. Biopreservation and Biobanking, 2016, 14, 201-206.	1.0	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.	2.8	0
112	Ethics review for international data-intensive research. Science, 2016, 351, 1399-1400.	12.6	44
113	Population genetic testing for cancer susceptibility: founder mutations to genomes. Nature Reviews Clinical Oncology, 2016, 13, 41-54.	27.6	86
114	Exposing participants? Population biobanks go geo. European Journal of Human Genetics, 2016, 24, 155-156.	2.8	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.	2.5	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.	0.1	4
117	Data Safe Havens in health research and healthcare. Bioinformatics, 2015, 31, 3241-3248.	4.1	58
118	The ethics weathervane. BMC Medical Ethics, 2015, 16, 58.	2.4	8
119	From the principles of genomic data sharing to the practices of data access committees. EMBO Molecular Medicine, 2015, 7, 507-509.	6.9	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.	0.9	12
122	Data analysis: Create a cloud commons. Nature, 2015, 523, 149-151.	27.8	89
123	Streamlining review of research involving humans: Canadian models: TableÂ1. Journal of Medical Genetics, 2015, 52, 566-569.	3.2	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.	2.0	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.	2.8	87
126	Return of genetic testing results in the era of whole-genome sequencing. Nature Reviews Genetics, 2015, 16, 553-559.	16.3	127



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127	International Charter of principles for sharing bio-specimens and data. European Journal of Human Genetics, 2015, 23, 721-728.	2.8	112
128	CRISPR germline engineeringâ€”the community speaks. Nature Biotechnology, 2015, 33, 478-486.	17.5	110
129	Oversight of human inheritable genome modification. Nature Biotechnology, 2015, 33, 454-455.	17.5	17
130	Genomic cloud computing: legal and ethical points to consider. European Journal of Human Genetics, 2015, 23, 1271-1278.	2.8	80
131	Return of whole-genome sequencing results in paediatric research: a statement of the P3C international paediatrics platform. European Journal of Human Genetics, 2014, 22, 3-5.	2.8	61
132	Does policy grow on trees?. BMC Medical Ethics, 2014, 15, 87.	2.4	4
133	Framework for responsible sharing of genomic and health-related data. The HUGO Journal, 2014, 8, 3.	4.1	185
134	The best interests of the child and the return of results in genetic research: international comparative perspectives. BMC Medical Ethics, 2014, 15, 72.	2.4	22
135	Towards an ethics safe harbor for global biomedical research. Journal of Law and the Biosciences, 2014, 1, 3-51.	1.6	39
136	Data sharing in large research consortia: experiences and recommendations from ENGAGE. European Journal of Human Genetics, 2014, 22, 317-321.	2.8	54
137	Beyond public health genomics: proposals from an international working group. European Journal of Public Health, 2014, 24, 877-879.	0.3	22
138	Data Sharing, Year 1 â€” Access to Data from Industry-Sponsored Clinical Trials. New England Journal of Medicine, 2014, 371, 2052-2054.	27.0	101
139	Attitudes of parents toward the return of targeted and incidental genomic research findings in children. Genetics in Medicine, 2014, 16, 633-640.	2.4	82
140	The Challenge of Informed Consent and Return of Results in Translational Genomics: Empirical Analysis and Recommendations. Journal of Law, Medicine and Ethics, 2014, 42, 344-355.	0.9	63
141	From the Right to Know to the Right Not to Know. Journal of Law, Medicine and Ethics, 2014, 42, 6-10.	0.9	23
142	Management Strategies for Ethics in International Research. Current Genetic Medicine Reports, 2014, 2, 255-260.	1.9	1
143	An implementation framework for the feedback of individual research results and incidental findings in research. BMC Medical Ethics, 2014, 15, 88.	2.4	39
144	Whole-Genome Sequencing in Newborn Screening Programs. Science Translational Medicine, 2014, 6, 229cm2.	12.4	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.	2.3	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.	1.9	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.	3.8	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.	3.2	27
149	International ethics harmonization and the global alliance for genomics and health. <i>Genome Medicine</i> , 2014, 6, 13.	8.2	38
150	Building a data sharing model for global genomic research. <i>Genome Biology</i> , 2014, 15, 430.	8.8	37
151	Data protection and consent to biomedical research: a step forward?. <i>Lancet, The</i> , 2014, 384, 855.	13.7	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.	11.1	35
155	Anonymity 2.0: direct-to-consumer genetic testing and donor conception. <i>Fertility and Sterility</i> , 2014, 101, 630-632.	1.0	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.	6.2	219
157	Population Biobanking and International Collaboration. <i>Pathobiology</i> , 2014, 81, 276-285.	3.8	13
158	Genotype-driven recruitment: a strategy whose time has come?. <i>BMC Medical Genomics</i> , 2013, 6, 19.	1.5	14
159	Recommendations for returning genomic incidental findings? We need to talk!. <i>Genetics in Medicine</i> , 2013, 15, 854-859.	2.4	272
160	An ethics safe harbor for international genomics research?. <i>Genome Medicine</i> , 2013, 5, 99.	8.2	23
161	Genomics: from persons to populations and back again. <i>Genome</i> , 2013, 56, 537-539.	2.0	4
162	Rare diseases and now rare data?. <i>Nature Reviews Genetics</i> , 2013, 14, 372-372.	16.3	19

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163	Whole-genome sequencing in health care. <i>European Journal of Human Genetics</i> , 2013, 21, 580-584.	2.8	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.	1.9	172
165	From Tissues to Genomes. <i>G3: Genes, Genomes, Genetics</i> , 2013, 3, 1203-1204.	1.8	2
166	A P3G generic access agreement for population genomic studies. <i>Nature Biotechnology</i> , 2013, 31, 384-385.	17.5	24
167	Demystifying Biobanks. <i>Hastings Center Report</i> , 2013, 43, 4-5.	1.0	10
168	Personalized medicine and access to health care: potential for inequitable access?. <i>European Journal of Human Genetics</i> , 2013, 21, 143-147.	2.8	45
169	Population studies: return of research results and incidental findings Policy Statement. <i>European Journal of Human Genetics</i> , 2013, 21, 245-247.	2.8	94
170	Pediatric research "personalized"? International perspectives on the return of results. <i>Personalized Medicine</i> , 2013, 10, 89-95.	1.5	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.	1.5	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.	3.2	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.	2.4	51
175	ELSI 2.0 for Genomics and Society. <i>Science</i> , 2012, 336, 673-674.	12.6	39
176	Power to the people: a wiki-governance model for biobanks. <i>Genome Biology</i> , 2012, 13, 158.	9.6	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.	2.4	418
178	Questioning the Limits of Genomic Privacy. <i>American Journal of Human Genetics</i> , 2012, 91, 577-578.	6.2	16
179	Sampling Populations of Humans Across the World: ELSI Issues. <i>Annual Review of Genomics and Human Genetics</i> , 2012, 13, 395-413.	6.2	57
180	Disclosure and management of research findings in stem cell research and banking: policy statement. <i>Regenerative Medicine</i> , 2012, 7, 439-448.	1.7	31

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181	Toward a roadmap in global biobanking for health. <i>European Journal of Human Genetics</i> , 2012, 20, 1105-1111.	2.8	139
182	Paediatric research and the communication of not-so incidental findings. <i>Paediatrics and Child Health</i> , 2012, 17, 190-192.	0.6	17
183	Editorial (An Idea Whose Time Has Come? An African Foresight Observatory on Genomics Medicine and) <i>Tj ETQq1 1 0,784314 rgBT /O</i>	0.2	11
184	Attitudes of Researchers to the Return of Incidental and Targeted Genomic Findings Obtained in a Research Setting to Participants. <i>Blood</i> , 2012, 120, 2069-2069.	1.4	0
185	Towards a data sharing Code of Conduct for international genomic research. <i>Genome Medicine</i> , 2011, 3, 46.	8.2	95
186	Policy and Data-Intensive Scientific Discovery in the Beginning of the 21st Century. <i>OMICS A Journal of Integrative Biology</i> , 2011, 15, 221-225.	2.0	21
187	From Banking to International Governance: Fostering Innovation in Stem Cell Research. <i>Stem Cells International</i> , 2011, 2011, 1-8.	2.5	21
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