Amy L Mcguire

List of Publications by Year
in descending order

182	14,376	38720 50	21521 114
papers	citations	h-index	g-index

187	187	187	16195
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Researcher Views on Changes in Personality, Mood, and Behavior in Next-Generation Deep Brain Stimulation. AJOB Neuroscience, 2023, 14, 287-299.	0.6	13
2	Core values of genomic citizen science: results from a qualitative interview study. BioSocieties, 2022, 17, 203-228.	0.8	9
3	Effects of participation in a U.S. trial of newborn genomic sequencing on parents at risk for depression. Journal of Genetic Counseling, 2022, 31, 218-229.	0.9	5
4	Perceived Utility of Genomic Sequencing: Qualitative Analysis and Synthesis of a Conceptual Model to Inform Patient-Centered Instrument Development. Patient, 2022, 15, 317-328.	1.1	21
5	How NFTs could transform health information exchange. Science, 2022, 375, 500-502.	6.0	32
6	Family secrets: Experiences and outcomes of participating in direct-to-consumer genetic relative-finder services. American Journal of Human Genetics, 2022, 109, 486-497.	2.6	12
7	Should Pediatricians Dismiss Families Who Refuse a COVID-19 Vaccine?. Clinical Pediatrics, 2022, 61, 99-103.	0.4	0
8	Researchersâ€™ Ethical Concerns About Using Adaptive Deep Brain Stimulation for Enhancement. Frontiers in Human Neuroscience, 2022, 16, 813922.	1.0	10
9	Parental Attitudes Toward Standard Newborn Screening and Newborn Genomic Sequencing: Findings From the BabySeq Study. Frontiers in Genetics, 2022, 13, 867371.	1.1	19
10	Direct-to-Consumer Genetic Testing: Value and Risk. Annual Review of Medicine, 2021, 72, 151-166.	5.0	44
11	Direct-to-Consumer Drug Advertisement and Prescribing Practices: Evidence Review and Practical Guidance for Clinicians. Journal of General Internal Medicine, 2021, 36, 1390-1394.	1.3	7
12	Four misconceptions about investigative genetic genealogy. Journal of Law and the Biosciences, 2021, 8, lsab001.	0.8	20
13	Family-level impact of genetic testing: integrating health economics and ethical, legal, and social implications. Personalized Medicine, 2021, 18, 209-212.	0.8	6
14	Pediatric Oncologistsâ€™ Experiences Returning and Incorporating Genomic Sequencing Results into Cancer Care. Journal of Personalized Medicine, 2021, 11, 570.	1.1	2
15	Psychological Distress Among the U.S. General Population During the COVID-19 Pandemic. Frontiers in Psychiatry, 2021, 12, 642918.	1.3	22
16	Challenges to Building a Gene Variant Commons to Assess Hereditary Cancer Risk: Results of a Modified Policy Delphi Panel Deliberation. Journal of Personalized Medicine, 2021, 11, 646.	1.1	3
17	Psychosocial Effect of Newborn Genomic Sequencing on Families in the BabySeq Project. JAMA Pediatrics, 2021, 175, 1132.	3.3	35
18	Genetic testing in ambulatory cardiology clinics reveals high rate of findings with clinical management implications. Genetics in Medicine, 2021, 23, 2404-2414.	1.1	14

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19	Toward better governance of human genomic data. Nature Genetics, 2021, 53, 2-8.	9.4	31
20	Patient, Caregiver, and Decliner Perspectives on Whether to Enroll in Adaptive Deep Brain Stimulation Research. Frontiers in Neuroscience, 2021, 15, 734182.	1.4	4
21	Conceptualization of utility in translational clinical genomics research. American Journal of Human Genetics, 2021, 108, 2027-2036.	2.6	11
22	Whoâ€™s on third? Regulation of third-party genetic interpretation services. Genetics in Medicine, 2020, 22, 4-11.	1.1	39
23	The case for implementing sustainable routine, population-level genomic reanalysis. Genetics in Medicine, 2020, 22, 815-816.	1.1	11
24	Essential, not peripheral: Addressing health care workersâ€™ mental health concerns during the COVID-19 pandemic. Journal of Occupational Health, 2020, 62, e12169.	1.0	5
25	Researcher Perspectives on Ethical Considerations in Adaptive Deep Brain Stimulation Trials. Frontiers in Human Neuroscience, 2020, 14, 578695.	1.0	21
26	Ventilator Triage Policies During the COVID-19 Pandemic at U.S. Hospitals Associated With Members of the Association of Bioethics Program Directors. Annals of Internal Medicine, 2020, 173, 188-194.	2.0	137
27	Airmen and health-care providersâ€™ attitudes toward the use of genomic sequencing in the US Air Force: findings from the MilSeq Project. Genetics in Medicine, 2020, 22, 2003-2010.	1.1	2
28	The road ahead in genetics and genomics. Nature Reviews Genetics, 2020, 21, 581-596.	7.7	118
29	Ethical Challenges Arising in the COVID-19 Pandemic: An Overview from the Association of Bioethics Program Directors (ABPD) Task Force. American Journal of Bioethics, 2020, 20, 15-27.	0.5	102
30	Quantifying Downstream Healthcare Utilization in Studies of Genomic Testing. Value in Health, 2020, 23, 559-565.	0.1	6
31	Ethics in Genetic and Genomic Research. , 2020, , 91-110.		4
32	Researcher Perspectives on Data Sharing in Deep Brain Stimulation. Frontiers in Human Neuroscience, 2020, 14, 578687.	1.0	11
33	Ethical, Legal, and Social Implications. , 2020, , 431-442.		2
34	Biomedical Citizen Science or Something Else? Reflections on Terms and Definitions. American Journal of Bioethics, 2019, 19, 17-19.	0.5	11
35	Introduction: Sharing Data in a Medical Information Commons. Journal of Law, Medicine and Ethics, 2019, 47, 7-11.	0.4	11
36	Clarify the HIPAA right of access to individualsâ€™ research data. Nature Biotechnology, 2019, 37, 850-852.	9.4	28

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37	Neuroethics at 15: Keep the Kant but Add More Bacon. AJOB Neuroscience, 2019, 10, 97-100.	0.6	7
38	Returning a Genomic Result for an Adult-Onset Condition to the Parents of a Newborn: Insights From the BabySeq Project. Pediatrics, 2019, 143, S37-S43.	1.0	45
39	Characterizing the Biomedical Data-Sharing Landscape. Journal of Law, Medicine and Ethics, 2019, 47, 21-30.	0.4	18
40	Hopeful and Concerned: Public Input on Building a Trustworthy Medical Information Commons. Journal of Law, Medicine and Ethics, 2019, 47, 70-87.	0.4	15
41	What is a Medical Information Commons?. Journal of Law, Medicine and Ethics, 2019, 47, 41-50.	0.4	13
42	Device Removal Following Brain Implant Research. Neuron, 2019, 103, 759-761.	3.8	20
43	Genomic Data-Sharing Practices. Journal of Law, Medicine and Ethics, 2019, 47, 31-40.	0.4	12
44	Improving recommendations for genomic medicine: building an evolutionary process from clinical practice advisory documents to guidelines. Genetics in Medicine, 2019, 21, 2431-2438.	1.1	13
45	In support of mitochondrial replacement therapy. Nature Medicine, 2019, 25, 870-871.	15.2	10
46	Responsibility, culpability, and parental views on genomic testing for seriously ill children. Genetics in Medicine, 2019, 21, 2791-2797.	1.1	20
47	Psychological outcomes related to exome and genome sequencing result disclosure: a meta-analysis of seven Clinical Sequencing Exploratory Research (CSER) Consortium studies. Genetics in Medicine, 2019, 21, 2781-2790.	1.1	55
48	Who Owns the Data in a Medical Information Commons?. Journal of Law, Medicine and Ethics, 2019, 47, 62-69.	0.4	18
49	The Genomic Medicine Integrative Research Framework: A Conceptual Framework for Conducting Genomic Medicine Research. American Journal of Human Genetics, 2019, 104, 1088-1096.	2.6	35
50	Importance of Participant-Centricity and Trust for a Sustainable Medical Information Commons. Journal of Law, Medicine and Ethics, 2019, 47, 12-20.	0.4	20
51	Predispositional genome sequencing in healthy adults: design, participant characteristics, and early outcomes of the PeopleSeq Consortium. Genome Medicine, 2019, 11, 10.	3.6	41
52	Challenging the Current Recommendations for Carrier Testing in Children. Pediatrics, 2019, 143, S27-S32.	1.0	13
53	Perceived Benefits, Risks, and Utility of Newborn Genomic Sequencing in the BabySeq Project. Pediatrics, 2019, 143, S6-S13.	1.0	47
54	Donors, authors, and owners: how is genomic citizen science addressing interests in research outputs?. BMC Medical Ethics, 2019, 20, 84.	1.0	9

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55	Parental interest in genomic sequencing of newborns: enrollment experience from the BabySeq Project. Genetics in Medicine, 2019, 21, 622-630.	1.1	61
56	Agents of empathy: How medical interpreters bridge sociocultural gaps in genomic sequencing disclosures with Spanish-speaking families. Patient Education and Counseling, 2019, 102, 895-901.	1.0	14
57	Interpretation of Genomic Sequencing Results in Healthy and Ill Newborns: Results from the BabySeq Project. American Journal of Human Genetics, 2019, 104, 76-93.	2.6	176
58	Exome sequencing disclosures in pediatric cancer care: Patterns of communication among oncologists, genetic counselors, and parents. Patient Education and Counseling, 2019, 102, 680-686.	1.0	12
59	Secondary findings from clinical genomic sequencing: prevalence, patient perspectives, family history assessment, and health-care costs from a multisite study. Genetics in Medicine, 2019, 21, 1100-1110.	1.1	111
60	Medical information commons. , 2019, , 281-293.		1
61	Continued access to investigational brain implants. Nature Reviews Neuroscience, 2018, 19, 317-318.	4.9	38
62	How Primary Care Providers Talk to Patients about Genome Sequencing Results: Risk, Rationale, and Recommendation. Journal of General Internal Medicine, 2018, 33, 877-885.	1.3	16
63	Patient understanding of, satisfaction with, and perceived utility of whole-genome sequencing: findings from the MedSeq Project. Genetics in Medicine, 2018, 20, 1069-1076.	1.1	58
64	HEADS4: Social Media Screening in Adolescent Primary Care. Pediatrics, 2018, 141, .	1.0	19
65	The phenotypic spectrum of Xiaâ€Gibbs syndrome. American Journal of Medical Genetics, Part A, 2018, 176, 1315-1326.	0.7	34
66	Short-term costs of integrating whole-genome sequencing into primary care and cardiology settings: a pilot randomized trial. Genetics in Medicine, 2018, 20, 1544-1553.	1.1	25
67	Navigating the researchâ€“clinical interface in genomic medicine: analysis from the CSER Consortium. Genetics in Medicine, 2018, 20, 545-553.	1.1	34
68	Alienation, Quality of Life, and DBS for Depression. AJOB Neuroscience, 2018, 9, 223-225.	0.6	2
69	Standardizing return of participant results. Science, 2018, 362, 759-760.	6.0	3
70	Should police have access to genetic genealogy databases? Capturing the Golden State Killer and other criminals using a controversial new forensic technique. PLoS Biology, 2018, 16, e2006906.	2.6	97
71	The Clinical Sequencing Evidence-Generating Research Consortium: Integrating Genomic Sequencing in Diverse and Medically Underserved Populations. American Journal of Human Genetics, 2018, 103, 319-327.	2.6	122
72	Citizen science, public policy. Science, 2018, 361, 134-136.	6.0	120

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73	The BabySeq project: implementing genomic sequencing in newborns. BMC Pediatrics, 2018, 18, 225.	0.7	115
74	Reconciling newborn screening and a novel splice variant in BTD associated with partial biotinidase deficiency: a BabySeq Project case report. Journal of Physical Education and Sports Management, 2018, 4, a002873.	0.5	7
75	Genealogy databases and the future of criminal investigation. Science, 2018, 360, 1078-1079.	6.0	71
76	Newborn Sequencing in Genomic Medicine and Public Health. Pediatrics, 2017, 139, .	1.0	174
77	Should you profit from your genome?. Nature Biotechnology, 2017, 35, 18-20.	9.4	17
78	A curated gene list for reporting results of newborn genomic sequencing. Genetics in Medicine, 2017, 19, 809-818.	1.1	79
79	When bins blur: Patient perspectives on categories of results from clinical whole genome sequencing. AJOB Empirical Bioethics, 2017, 8, 82-88.	0.8	34
80	Myriad take two: Can genomic databases remain secret?. Science, 2017, 356, 586-587.	6.0	9
81	Sharing data under the 21st Century Cures Act. Genetics in Medicine, 2017, 19, 1289-1294.	1.1	20
82	Consumer Perspectives on Access to Directâ€toâ€Consumer Genetic Testing: Role of Demographic Factors and the Testing Experience. Milbank Quarterly, 2017, 95, 291-318.	2.1	22
83	Moving beyond Bermuda: sharing data to build a medical information commons. Genome Research, 2017, 27, 897-901.	2.4	27
84	The price of whole-genome sequencing may be decreasing, but who will be sequenced?. Personalized Medicine, 2017, 14, 203-211.	0.8	7
85	Communication challenges for nongeneticist physicians relaying clinical genomic results. Personalized Medicine, 2017, 14, 423-431.	0.8	36
86	Should We Be Concerned About Preserving Agency and Personal Identity in Patients With Adaptive Deep Brain Stimulation Systems?. AJOB Neuroscience, 2017, 8, 73-75.	0.6	11
87	Barriers to clinical adoption of next-generation sequencing: a policy Delphi panel's solutions. Personalized Medicine, 2017, 14, 339-354.	0.8	15
88	The Impact of Whole-Genome Sequencing on the Primary Care and Outcomes of Healthy Adult Patients. Annals of Internal Medicine, 2017, 167, 159.	2.0	145
89	Creating a data resource: what will it take to build a medical information commons?. Genome Medicine, 2017, 9, 84.	3.6	36
90	Portero versus portador: Spanish interpretation of genomic terminology during whole exome sequencing results disclosure. Personalized Medicine, 2017, 14, 503-514.	0.8	17

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91	Parental Perspectives on Whole-Exome Sequencing in Pediatric Cancer: A Typology of Perceived Utility. JCO Precision Oncology, 2017, 1, 1-10.	1.5	26
92	Do privacy and security regulations need a status update? Perspectives from an intergenerational survey. PLoS ONE, 2017, 12, e0184525.	1.1	8
93	Constraints on gene patent protection fuel secrecy concerns: a qualitative study. Journal of Law and the Biosciences, 2017, 4, 542-564.	0.8	2
94	Beyond Our Borders? Public Resistance to Global Genomic Data Sharing. PLoS Biology, 2016, 14, e2000206.	2.6	33
95	Developing context-specific next-generation sequencing policy. Nature Biotechnology, 2016, 34, 466-470.	9.4	5
96	Clinical Sequencing Exploratory Research Consortium: Accelerating Evidence-Based Practice of Genomic Medicine. American Journal of Human Genetics, 2016, 98, 1051-1066.	2.6	137
97	Return of individual genomic research results: what do consent forms tell participants?. European Journal of Human Genetics, 2016, 24, 1524-1529.	1.4	14
98	Ethical and Legal Challenges Associated with Public Molecular Autopsies. Journal of Law, Medicine and Ethics, 2016, 44, 309-318.	0.4	4
99	The ethics of conducting molecular autopsies in cases of sudden death in the young. Genome Research, 2016, 26, 1165-1169.	2.4	20
100	An open access pilot freely sharing cancer genomic data from participants in Texas. Scientific Data, 2016, 3, 160010.	2.4	19
101	Is Whole-Exome Sequencing an Ethically Disruptive Technology? Perspectives of Pediatric Oncologists and Parents of Pediatric Patients With Solid Tumors. Pediatric Blood and Cancer, 2016, 63, 511-515.	0.8	39
102	Legal Barriers to Adolescent Participation in Research About HIV and Other Sexually Transmitted Infections. American Journal of Public Health, 2016, 106, 40-44.	1.5	24
103	Barriers to clinical adoption of next generation sequencing: Perspectives of a policy Delphi panel. Applied & Translational Genomics, 2016, 10, 19-24.	2.1	30
104	Patientsâ€™ perceived utility of whole-genome sequencing for their healthcare: findings from the MedSeq project. Personalized Medicine, 2016, 13, 13-20.	0.8	31
105	Diagnostic Yield of Clinical Tumor and Germline Whole-Exome Sequencing for Children With Solid Tumors. JAMA Oncology, 2016, 2, 616.	3.4	378
106	Potential Psychosocial Risks of Sequencing Newborns. Pediatrics, 2016, 137, S24-S29.	1.0	47
107	Persistent confusion and controversy surrounding gene patents. Nature Biotechnology, 2016, 34, 145-147.	9.4	8
108	Community crystal gazing. Nature Biotechnology, 2016, 34, 276-283.	9.4	1

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109	Participants and Study Declinersâ€™ Perspectives About the Risks of Participating in a Clinical Trial of Whole Genome Sequencing. Journal of Empirical Research on Human Research Ethics, 2016, 11, 21-30.	0.6	41
110	Cultivating Administrative Support for a Clinical Ethics Consultation Service. Journal of Clinical Ethics, 2016, 27, 341-351.	0.1	1
111	Secondary findings and carrier test frequencies in a large multiethnic sample. Genome Medicine, 2015, 7, 54.	3.6	47
112	Marginally scientific? Genetic testing of children and adolescents for lifestyle and health promotion. Journal of Law and the Biosciences, 2015, 2, lsv038.	0.8	12
113	Pregnant patients' risk perception of prenatal test results with uncertain fetal clinical significance: ultrasound versus advanced genetic testing. Prenatal Diagnosis, 2015, 35, 1213-1217.	1.1	11
114	It depends whose data are being shared: considerations for genomic data sharing policies. Journal of Law and the Biosciences, 2015, 2, lsv030.	0.8	4
115	â€˜Someday it will be the normâ€™: physician perspectives on the utility of genome sequencing for patient care in the MedSeqProject. Personalized Medicine, 2015, 12, 23-32.	0.8	40
116	How behavioral economics can help to avoid â€˜The last mile problemâ€™ in whole genome sequencing. Genome Medicine, 2015, 7, 3.	3.6	14
117	GINA, Genetic Discrimination, and Genomic Medicine. New England Journal of Medicine, 2015, 372, 397-399.	13.9	141
118	Open Access Data Sharing in Genomic Research. Genes, 2014, 5, 739-747.	1.0	26
119	Obtaining informed consent for clinical tumor and germline exome sequencing of newly diagnosed childhood cancer patients. Genome Medicine, 2014, 6, 69.	3.6	60
120	Guidelines for return of research results from pediatric genomic studies: deliberations of the Boston Childrenâ€™s Hospital Gene Partnership Informed Cohort Oversight Board. Genetics in Medicine, 2014, 16, 547-552.	1.1	49
121	Social and behavioral research in genomic sequencing: approaches from the Clinical Sequencing Exploratory Research Consortium Outcomes and Measures Working Group. Genetics in Medicine, 2014, 16, 727-735.	1.1	60
122	Development of the clinical next-generation sequencing industry in a shifting policy climate. Nature Biotechnology, 2014, 32, 980-982.	9.4	25
123	Overcoming the Reimbursement Barriers for Clinical Sequencing. JAMA - Journal of the American Medical Association, 2014, 312, 1857.	3.8	35
124	Can I be sued for that? Liability risk and the disclosure of clinically significant genetic research findings. Genome Research, 2014, 24, 719-723.	2.4	31
125	Pediatric Data Sharing in Genomic Research: Attitudes and Preferences of Parents. Pediatrics, 2014, 133, 690-697.	1.0	36
126	The MedSeq Project: a randomized trial of integrating whole genome sequencing into clinical medicine. Trials, 2014, 15, 85.	0.7	122

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127	ACMG recommendations for reporting of incidental findings in clinical exome and genome sequencing. Genetics in Medicine, 2013, 15, 565-574.	1.1	2,186
128	Identifying Personal Genomes by Surname Inference. Science, 2013, 339, 321-324.	6.0	936
129	The Indispensable Role of Professional Judgment in Genomic Medicine. JAMA - Journal of the American Medical Association, 2013, 309, 1465.	3.8	32
130	Participants' Recall and Understanding of Genomic Research and Large-Scale Data Sharing. Journal of Empirical Research on Human Research Ethics, 2013, 8, 42-52.	0.6	42
131	Personalized genomic disease risk of volunteers. Proceedings of the National Academy of Sciences of the United States of America, 2013, 110, 16957-16962.	3.3	44
132	Policy Uncertainty, Sequencing, and Cell Lines. G3: Genes, Genomes, Genetics, 2013, 3, 1205-1207.	0.8	6
133	Ethics and Genomic Incidental Findings. Science, 2013, 340, 1047-1048.	6.0	160
134	Returning genetic research results: study type matters. Personalized Medicine, 2013, 10, 27-34.	0.8	20
135	Disclosing pathogenic genetic variants to research participants: Quantifying an emerging ethical responsibility. Genome Research, 2012, 22, 421-428.	2.4	79
136	The legal risks of returning results of genomics research. Genetics in Medicine, 2012, 14, 473-477.	1.1	110
137	Return of individual research results from genome-wide association studies: experience of the Electronic Medical Records and Genomics (eMERGE) Network. Genetics in Medicine, 2012, 14, 424-431.	1.1	94
138	Currents in Contemporary Bioethics. Journal of Law, Medicine and Ethics, 2012, 40, 1040-1046.	0.4	7
139	â€œSnake-oil,â€•â€œquack medicine,â€•and â€œindustrially cultured organisms:â€•biovalue and the commercialization of human microbiome research. BMC Medical Ethics, 2012, 13, 28.	1.0	39
140	Perspectives on Human Microbiome Research Ethics. Journal of Empirical Research on Human Research Ethics, 2012, 7, 1-14.	0.6	46
141	Direct-to-Consumer Genetic Testing: Perceptions, Problems, and Policy Responses. Annual Review of Medicine, 2012, 63, 23-33.	5.0	109
142	The Human Microbiome Project: A Community Resource for the Healthy Human Microbiome. PLoS Biology, 2012, 10, e1001377.	2.6	369
143	Exploring the ELSI universe: critical issues in the evolution of human genomic research. Genome Medicine, 2011, 3, 38.	3.6	14
144	Shaping Patients' Decisions. Chest, 2011, 139, 424-429.	0.4	25

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145	Ethical and practical challenges of sharing data from genome-wide association studies: The eMERGE Consortium experience. Genome Research, 2011, 21, 1001-1007.	2.4	68
146	To share or not to share: A randomized trial of consent for data sharing in genome research. Genetics in Medicine, 2011, 13, 948-955.	1.1	96
147	Health System Implications of Direct-to-Consumer Personal Genome Testing. Public Health Genomics, 2011, 14, 53-58.	0.6	37
148	Personal genome research : what should the participant be told?. Trends in Genetics, 2010, 26, 199-201.	2.9	68
149	Personalized genomic information: preparing for the future of genetic medicine. Nature Reviews Genetics, 2010, 11, 161-165.	7.7	106
150	Taking DNA from the dead. Nature Reviews Genetics, 2010, 11, 318-318.	7.7	8
151	Regulating Direct-to-Consumer Personal Genome Testing. Science, 2010, 330, 181-182.	6.0	43
152	Responseâ€"Regulating Genetic Tests: Who Owns the Data?. Science, 2010, 330, 1626-1627.	6.0	0
153	Confronting real time ethical, legal, and social issues in the Electronic Medical Records and Genomics (eMERGE) Consortium. Genetics in Medicine, 2010, 12, 616-620.	1.1	55
154	Whole-Genome Sequencing in a Patient with Charcotâ€"Marieâ€"Tooth Neuropathy. New England Journal of Medicine, 2010, 362, 1181-1191.	13.9	698
155	Informed Consent in Genomics and Genetic Research. Annual Review of Genomics and Human Genetics, 2010, 11, 361-381.	2.5	163
156	Ethical and Practical Guidelines for Reporting Genetic Research Results to Study Participants. Circulation: Cardiovascular Genetics, 2010, 3, 574-580.	5.1	328
157	The ethical use of existing samples for genome research. Genetics in Medicine, 2009, 11, 712-715.	1.1	52
158	Social Networkers' Attitudes Toward Direct-to-Consumer Personal Genome Testing. American Journal of Bioethics, 2009, 9, 3-10.	0.5	189
159	Developing a Tissue Resource to Characterize the Genome of Pancreatic Cancer. World Journal of Surgery, 2009, 33, 723-731.	0.8	9
160	Currents in Contemporary Ethics. Journal of Law, Medicine and Ethics, 2009, 37, 369-374.	0.4	35
161	The futility of genomic counseling: essential role of electronic health records. Genome Medicine, 2009, 1, 48.	3.6	21
162	Two cheers for GINA?. Genome Medicine, 2009, 1, 6.	3.6	34

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163	Paving the Way to Personalized Genomic Medicine: Steps to Successful Implementation. Current Pharmacogenomics and Personalized Medicine, 2009, 7, 125-132.	0.2	30
164	The complete genome of an individual by massively parallel DNA sequencing. Nature, 2008, 452, 872-876.	13.7	1,635
165	Research ethics and the challenge of whole-genome sequencing. Nature Reviews Genetics, 2008, 9, 152-156.	7.7	201
166	An Unwelcome Side Effect of Direct-to-Consumer Personal Genome Testing. JAMA - Journal of the American Medical Association, 2008, 300, 2669.	3.8	232
167	Ethical, legal, and social considerations in conducting the Human Microbiome Project. Genome Research, 2008, 18, 1861-1864.	2.4	68
168	Research Ethics Recommendations for Whole-Genome Research: Consensus Statement. PLoS Biology, 2008, 6, e73.	2.6	212
169	Identifiability of DNA Data: The Need for Consistent Federal Policy. American Journal of Bioethics, 2008, 8, 75-76.	0.5	21
170	Donâ€™t throw the baby out with the bathwater: Enabling a bottom-up approach in genome-wide association studies: Figure 1.. Genome Research, 2008, 18, 1683-1685.	2.4	49
171	1000 Genomes: on the road to personalized medicine. Personalized Medicine, 2008, 5, 195-197.	0.8	9
172	DNA data sharing: research participants' perspectives. Genetics in Medicine, 2008, 10, 46-53.	1.1	98
173	Confidentiality, privacy, and security of genetic and genomic test information in electronic health records: points to consider. Genetics in Medicine, 2008, 10, 495-499.	1.1	111
174	The Future of Personal Genomics. Science, 2007, 317, 1687-1687.	6.0	53
175	Currents in Contemporary Ethics. Journal of Law, Medicine and Ethics, 2006, 34, 809-812.	0.4	13
176	GENETICS: No Longer De-Identified. Science, 2006, 312, 370-371.	6.0	164
177	Missed Expectations?. Medical Care, 2005, 43, 466-470.	1.1	54
178	The Ethical Health Lawyer. Journal of Law, Medicine and Ethics, 2005, 33, 603-607.	0.4	2
179	Respect as an Organizing Normative Category for Research Ethics. American Journal of Bioethics, 2005, 5, W1-W2.	0.5	3
180	To the Editor. American Journal of Bioethics, 2005, 5, W1-W1.	0.5	9

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181	A Typology of Shared Decision Making, Informed Consent, and Simple Consent. Annals of Internal Medicine, 2004, 140, 54.	2.0	447
182	â€œIdealists and capitalistsâ€: ownership attitudes and preferences in genomic citizen science. New Genetics and Society, 0, , 1-22.	0.7	1