

# Barbara Prainsack

## List of Publications by Year in descending order

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Version: 2024-02-01

161  
papers

4,643  
citations

94433

37  
h-index

175258

52  
g-index

177  
all docs

177  
docs citations

177  
times ranked

4102  
citing authors

#	ARTICLE	IF	CITATIONS
1	Rethinking value construction in biomedicine and healthcare. <i>BioSocieties</i> , 2022, 17, 391-414.	1.3	15
2	Normative positions towards COVID-19 contact-tracing apps: findings from a large-scale qualitative study in nine European countries. <i>Critical Public Health</i> , 2022, 32, 5-18.	2.4	22
3	The second pandemic: Examining structural inequality through reverberations of COVID-19 in Europe. <i>Social Science and Medicine</i> , 2022, 292, 114634.	3.8	32
4	The benefits, costs and feasibility of a low incidence COVID-19 strategy. <i>Lancet Regional Health - Europe</i> , The, 2022, 13, 100294.	5.6	17
5	Beyond Vaccination Mandates: Solidarity and Freedom During COVID-19. <i>American Journal of Public Health</i> , 2022, 112, 232-233.	2.7	3
6	Return of genomic results does not motivate intent to participate in research for all: Perspectives across 22 countries. <i>Genetics in Medicine</i> , 2022, 24, 1120-1129.	2.4	8
7	Anticipating hopes, fears and expectations towards COVID-19 vaccines: A qualitative interview study in seven European countries. <i>SSM Qualitative Research in Health</i> , 2022, 2, 100035.	1.5	26
8	Value-creation in the health data domain: a typology of what health data help us do. <i>BioSocieties</i> , 2022, , 1-25.	1.3	4
9	Public involvement in the governance of population-level biomedical research: unresolved questions and future directions. <i>Journal of Medical Ethics</i> , 2021, 47, 522-525.	1.8	20
10	Antimicrobial resistance as a problem of values? Views from three continents. <i>Critical Public Health</i> , 2021, 31, 451-463.	2.4	53
11	The meaning and enactment of openness in Personalised and Precision Medicine. <i>Science and Public Policy</i> , 2021, 47, 647-654.	2.4	2
12	The Austrian Corona Panel Project: monitoring individual and societal dynamics amidst the COVID-19 crisis. <i>European Political Science</i> , 2021, 20, 318-344.	1.2	54
13	Not all biases are bad: equitable and inequitable biases in machine learning and radiology. <i>Insights Into Imaging</i> , 2021, 12, 13.	3.4	27
14	An action plan for pan-European defence against new SARS-CoV-2 variants. <i>Lancet</i> , The, 2021, 397, 469-470.	13.7	101
15	Early Perceptions of COVID-19 Contact Tracing Apps in German-Speaking Countries: Comparative Mixed Methods Study. <i>Journal of Medical Internet Research</i> , 2021, 23, e25525.	4.3	86
16	Demonstrating trustworthiness when collecting and sharing genomic data: public views across 22 countries. <i>Genome Medicine</i> , 2021, 13, 92.	8.2	39
17	Constructing the "Future of Work": An analysis of the policy discourse. <i>New Technology, Work and Employment</i> , 2021, 36, 307-326.	4.0	20
18	A look into the future of the COVID-19 pandemic in Europe: an expert consultation. <i>Lancet Regional Health - Europe</i> , The, 2021, 8, 100185.	5.6	72

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19	Toward better governance of human genomic data. <i>Nature Genetics</i> , 2021, 53, 2-8.	21.4	31
20	Solidarit�t in Zeiten einer Pandemie: Alltagspraktiken und Priorisierungentscheidungen im Lichte des Solidarit�tskonzeptes. , 2021, , 29-43.		1
21	Reply to Letter to the Editor on "Not all biases are bad: equitable and inequitable biases in machine learning and radiology". <i>Insights Into Imaging</i> , 2021, 12, 157.	3.4	0
22	Open science, data sharing and solidarity: who benefits?. <i>History and Philosophy of the Life Sciences</i> , 2021, 43, 115.	1.1	19
23	A sociology of precision" in practice: The affective and temporal complexities of everyday clinical care. <i>Sociology of Health and Illness</i> , 2021, 43, 2178-2195.	2.1	14
24	Assessing policies for the implementation of new technological interventions to combat COVID-19. <i>European Journal of Public Health</i> , 2021, 31, .	0.3	1
25	Big Data Governance Needs More Collective Responsibility: The Role of Harm Mitigation in the Governance of Data Use in Medicine and Beyond. <i>Medical Law Review</i> , 2020, 28, 155-182.	0.5	27
26	Quadruple Decision Making for Parkinson's Disease Patients: Combining Expert Opinion, Patient Preferences, Scientific Evidence, and Big Data Approaches to Reach Precision Medicine. <i>Journal of Parkinson's Disease</i> , 2020, 10, 223-231.	2.8	24
27	Members of the public in the USA, UK, Canada and Australia expressing genetic exceptionalism say they are more willing to donate genomic data. <i>European Journal of Human Genetics</i> , 2020, 28, 424-434.	2.8	29
28	The double-edged sword of digital self-care: Physician perspectives from Northern Germany. <i>Social Science and Medicine</i> , 2020, 260, 113174.	3.8	31
29	Personalized medicine in Austria: expectations and limitations. <i>Personalized Medicine</i> , 2020, 17, 423-428.	1.5	3
30	COVID-19 and Contact Tracing Apps: Ethical Challenges for a Social Experiment on a Global Scale. <i>Journal of Bioethical Inquiry</i> , 2020, 17, 835-839.	1.5	66
31	Solidarity in Times of Pandemics. <i>Democratic Theory</i> , 2020, 7, 124-133.	0.7	41
32	Global Public Perceptions of Genomic Data Sharing: What Shapes the Willingness to Donate DNA and Health Data?. <i>American Journal of Human Genetics</i> , 2020, 107, 743-752.	6.2	76
33	Data Mining in Systems Medicine and the Project of Solidarity: The Interface of Genomics and Society Revisited. , 2020, , 97-117.		2
34	Genomic Sequencing Capacity, Data Retention, and Personal Access to Raw Data in Europe. <i>Frontiers in Genetics</i> , 2020, 11, 303.	2.3	9
35	Transparency, consent and trust in the use of customers' data by an online genetic testing company: an Exploratory survey among 23andMe users. <i>New Genetics and Society</i> , 2020, 39, 459-482.	1.2	10
36	Oil crisis: the political economy of digital data. Conclusion of the special issue. <i>Policy Studies</i> , 2020, 41, 563-566.	1.6	2

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37	The political economy of digital data: introduction to the special issue. <i>Policy Studies</i> , 2020, 41, 439-446.	1.6	43
38	Integrating artificial intelligence into the clinical practice of radiology: challenges and recommendations. <i>European Radiology</i> , 2020, 30, 3576-3584.	4.5	113
39	The value of healthcare data: to nudge, or not?. <i>Policy Studies</i> , 2020, 41, 547-562.	1.6	20
40	Willingness to donate genomic and other medical data: results from Germany. <i>European Journal of Human Genetics</i> , 2020, 28, 1000-1009.	2.8	28
41	Face Masks During the COVID-19 Pandemic: A Simple Protection Tool With Many Meanings. <i>Frontiers in Public Health</i> , 2020, 8, 606635.	2.7	100
42	Shifting Solidarities: Personalisation in Insurance and Medicine. , 2020, , 127-151.		12
43	Caring for data: Value creation in a data-intensive research laboratory. <i>Social Studies of Science</i> , 2020, 50, 175-197.	2.5	49
44	Citizen Science in Health Domain. , 2020, , 274-279.		1
45	Health Information Counselors: A New Profession for the Age of Big Data. <i>Academic Medicine</i> , 2019, 94, 37-41.	1.6	35
46	Trust in genomic data sharing among members of the general public in the UK, USA, Canada and Australia. <i>Human Genetics</i> , 2019, 138, 1237-1246.	3.8	69
47	Civil society stakeholder views on forensic DNA phenotyping: Balancing risks and benefits. <i>Forensic Science International: Genetics</i> , 2019, 43, 102157.	3.1	20
48	Response to Dr Uluçanlar. <i>Journal of Evaluation in Clinical Practice</i> , 2019, 25, 728-728.	1.8	0
49	Muireann Quigley, Self-Ownership, Property Rights, and the Human Body: A Legal and Philosophical Analysis. <i>Medical Law Review</i> , 2019, 27, 358-362.	0.5	2
50	Consent insufficient for data release. <i>Science</i> , 2019, 364, 445-446.	12.6	9
51	Moral entrepreneurship, the power-knowledge nexus, and the Cochrane "crisis". <i>Journal of Evaluation in Clinical Practice</i> , 2019, 25, 717-725.	1.8	15
52	Logged out: Ownership, exclusion and public value in the digital data and information commons. <i>Big Data and Society</i> , 2019, 6, 205395171982977.	4.5	67
53	Precision Medicine Needs a Cure for Inequality. <i>Current History</i> , 2019, 118, 11-15.	0.7	7
54	Meeting the needs of underserved populations: setting the agenda for more inclusive citizen science of medicine. <i>Journal of Medical Ethics</i> , 2019, 45, 617-622.	1.8	37

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55	In Reply to Kringos et al. Academic Medicine, 2019, 94, 748-749.	1.6	0
56	Attitudes of publics who are unwilling to donate DNA data for research. European Journal of Medical Genetics, 2019, 62, 316-323.	1.3	53
57	Forensic DNA phenotyping in Europe: views on the ground from those who have a professional stake in the technology. New Genetics and Society, 2019, 38, 119-141.	1.2	39
58	Data Donation: How to Resist the iLeviathan. Philosophical Studies Series, 2019, , 9-22.	1.9	24
59	Conceptual and Ethical Considerations for Citizen Science in Biomedicine. Öffentliche Wissenschaft Und Gesellschaftlicher Wandel, 2019, , 195-217.	0.1	22
60	Data Work: Meaning-Making in the Era of Data-Rich Medicine. Journal of Medical Internet Research, 2019, 21, e11672.	4.3	30
61	The Use of Forensic DNA Phenotyping in Predicting Appearance and Biogeographic Ancestry. Deutsches A&#x0308;rzteblatt International, 2019, 51-52, 873-880.	0.9	47
62	The Gender of Biomedical Data: Challenges for Personalised and Precision Medicine. Somatechnics, 2019, 9, 170-187.	0.7	3
63	The "We" in the "Me" Science Technology and Human Values, 2018, 43, 21-44.	3.1	88
64	Markers as mediators: A review and synthesis of epigenetics literature. BioSocieties, 2018, 13, 276-303.	1.3	30
65	Solidarity can make a difference: Addressing transformations in healthcare, demographics and technological replacement. Bioethics, 2018, 32, 537-540.	1.4	6
66	The value of work: Addressing the future of work through the lens of solidarity. Bioethics, 2018, 32, 585-592.	1.4	20
67	The Genomic Medicine Alliance: A Global Effort to Facilitate the Introduction of Genomics into Healthcare in Developing Nations. , 2018, , 173-188.		1
68	A response to the forensic genetics policy initiative's report "Establishing Best Practice for Forensic DNA Databases". Forensic Science International: Genetics, 2018, 36, e19-e21.	3.1	11
69	Is there a duty to participate in digital epidemiology?. Life Sciences, Society and Policy, 2018, 14, 9.	3.2	19
70	APPLaUD: access for patients and participants to individual level uninterpreted genomic data. Human Genomics, 2018, 12, 7.	2.9	45
71	"Your DNA, Your Say": global survey gathering attitudes toward genomics: design, delivery and methods. Personalized Medicine, 2018, 15, 311-318.	1.5	26
72	Motivations of participants in the citizen science of microbiomics: data from the British Gut Project. Genetics in Medicine, 2017, 19, 959-961.	2.4	19

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73	Every participant is a PI. Citizen science and participatory governance in population studies. <i>International Journal of Epidemiology</i> , 2017, 46, dyw204.	1.9	33
74	Beyond individualism: Is there a place for relational autonomy in clinical practice and research?. <i>Clinical Ethics</i> , 2017, 12, 150-165.	0.7	148
75	Citizen Science in Health Domain. , 2017, , 1-6.		0
76	Emerging ethical issues regarding digital health data. On the World Medical Association Draft Declaration on Ethical Considerations Regarding Health Databases and Biobanks. <i>Croatian Medical Journal</i> , 2016, 57, 207-213.	0.7	29
77	Test Pricing and Reimbursement in Genomic Medicine: Towards a General Strategy. <i>Public Health Genomics</i> , 2016, 19, 352-363.	1.0	37
78	Thinking ethical and regulatory frameworks in medicine from the perspective of solidarity on both sides of the Atlantic. <i>Theoretical Medicine and Bioethics</i> , 2016, 37, 489-501.	0.8	31
79	Beyond clinical utility: The multiple values of DTC genetics. <i>Applied &amp; Translational Genomics</i> , 2016, 8, 4-8.	2.1	32
80	Approaching ethical, legal and social issues of emerging forensic DNA phenotyping (FDP) technologies comprehensively: Reply to "Forensic DNA phenotyping: Predicting human appearance from crime scene material for investigative purposes" by Manfred Kayser. <i>Forensic Science International: Genetics</i> , 2016, 22, e1-e4.	3.1	44
81	Crowdsourcing the Human Gut. Is crowdsourcing also 'citizen science'?. <i>Journal of Science Communication</i> , 2016, 15, A03.	0.8	34
82	Sticks and carrots: encouraging open science at its source. <i>Geo: Geography and Environment</i> , 2015, 2, 12-16.	0.8	43
83	Genetics and Forensics. , 2015, , 955-961.		0
84	Evaluating the Consent Preferences of UK Research Volunteers for Genetic and Clinical Studies. <i>PLoS ONE</i> , 2015, 10, e0118027.	2.5	36
85	Forensic Genetic Databases: Ethical and Social Dimensions. , 2015, , 339-345.		4
86	Ethical Standards for Research Biobank Donation. <i>JAMA - Journal of the American Medical Association</i> , 2015, 313, 1573.	7.4	2
87	The lifestylisation of healthcare? "Consumer genomics" and mobile health as technologies for healthy lifestyle. <i>Applied &amp; Translational Genomics</i> , 2015, 4, 44-49.	2.1	44
88	Is personalized medicine different? (Reinscription: the sequel) A response to Roy Duster. <i>British Journal of Sociology</i> , 2015, 66, 28-35.	1.5	21
89	Ethics of Healthcare Policy and the Concept of Solidarity. , 2015, , 649-664.		6
90	Bioethics in the Post-genomic Era. , 2015, , 625-630.		0

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91	DIY genetics: the right to know your own genome. , 2014, , 100-115.		5
92	The Powers of Participatory Medicine. PLoS Biology, 2014, 12, e1001837.	5.6	54
93	Ready to Put Metadata on the Post-2015 Development Agenda? Linking Data Publications to Responsible Innovation and Science Diplomacy. OMICS A Journal of Integrative Biology, 2014, 18, 1-9.	2.0	31
94	Raw Personal Data: Providing Access. Science, 2014, 343, 373-374.	12.6	57
95	Personhood and solidarity: what kind of personalized medicine do we want?. Personalized Medicine, 2014, 11, 651-657.	1.5	43
96	Valediction for Herbert Gottweis. BioSocieties, 2014, 9, 457-480.	1.3	0
97	Genetic Data and the Law: A Critical Perspective on Privacy Protection. Medical Law Review, 2014, 22, 291-295.	0.5	0
98	Raw Data: Access to Inaccuracyâ€™Response. Science, 2014, 343, 969-969.	12.6	2
99	Bridging genomics research between developed and developing countries: the Genomic Medicine Alliance. Personalized Medicine, 2014, 11, 615-623.	1.5	22
100	Obituary for Herbert Gottweis, Professor of Political Science, University of Vienna: Born 8 February 1958 in Vienna, died 31 March 2014 in Vienna. Life Sciences, Society and Policy, 2014, 10, 12.	3.2	1
101	Working towards personalization of Medicine: Genomics in 2014. Personalized Medicine, 2014, 11, 611-613.	1.5	4
102	â€žRoteâ€™œ Biowissenschaften, Biotechnologie und Biomedizin. , 2014, , 331-340.		1
103	Performing the Union: The PrÃ¼m Decision and the European dream. Studies in History and Philosophy of Science Part C:Studies in History and Philosophy of Biological and Biomedical Sciences, 2013, 44, 71-79.	1.3	35
104	The challenge of personal genomics in Germany. Nature Biotechnology, 2013, 31, 16-17.	17.5	11
105	Beyond the clinic: â€™direct-to-consumerâ€™™ genomic profiling services and pharmacogenomics. Pharmacogenomics, 2013, 14, 403-412.	1.3	30
106	Opening Pandoraâ€™™s box in the UK: a hypothetical pharmacogenetic test for clozapine. Pharmacogenomics, 2013, 14, 1907-1914.	1.3	11
107	Regulating Genomics: Time for a Broader Vision. Science Translational Medicine, 2013, 5, 198ed12.	12.4	12
108	A SOLIDARITY-BASED APPROACH TO THE GOVERNANCE OF RESEARCH BIOBANKS. Medical Law Review, 2013, 21, 71-91.	0.5	89

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109	Situated bio-regulation: Ethnographic sensibility at the interface of STS, policy studies and the social studies of medicine. <i>BioSocieties</i> , 2013, 8, 336-359.	1.3	16
110	Let's get real about virtual: online health is here to stay. <i>Genetical Research</i> , 2013, 95, 111-113.	0.9	5
111	Risky profiles: societal dimensions of forensic uses of DNA profiling technologies. <i>New Genetics and Society</i> , 2012, 31, 249-258.	1.2	13
112	Understanding Solidarity (With a Little Help from Your Friends): Response to Dawson and Verweij. <i>Public Health Ethics</i> , 2012, 5, 206-210.	1.0	16
113	Experiences of Early Users of Direct-to-Consumer Genomics in Switzerland: An Exploratory Study. <i>Public Health Genomics</i> , 2012, 15, 352-362.	1.0	53
114	Solidarity. Reflections on an Emerging Concept in Bioethics. Summary. <i>Jahrbuch Fur Wissenschaft Und Ethik</i> , 2012, 17, 331-344.	0.2	28
115	Lifestyle-related diseases and individual responsibility through the prism of solidarity. <i>Clinical Ethics</i> , 2012, 7, 79-85.	0.7	29
116	Bracketing off population does not advance ethical reflection on EVCs: A reply to Kayser and Schneider. <i>Forensic Science International: Genetics</i> , 2012, 6, e16-e17.	3.1	31
117	The future of technologies for personalised medicine. <i>New Biotechnology</i> , 2012, 29, 625-633.	4.4	87
118	SOLIDARITY IN CONTEMPORARY BIOETHICS – TOWARDS A NEW APPROACH. <i>Bioethics</i> , 2012, 26, 343-350.	1.4	100
119	Enterprising or altruistic selves? Making up research subjects in genetics research. <i>Sociology of Health and Illness</i> , 2011, 33, 1081-1095.	2.1	71
120	Voting with their Mice: Personal Genome Testing and the ‘Participatory Turn’ in Disease Research. <i>Accountability in Research</i> , 2011, 18, 132-147.	2.4	42
121	Overcoming embryonic exceptionalism? Lessons from analyzing human stem cell research regulation in Israel. <i>New Genetics and Society</i> , 2011, 30, 267-277.	1.2	5
122	Genomic stuff: Governing the (im)matter of life. <i>International Journal of the Commons</i> , 2011, 5, 259.	1.4	10
123	DNA profiling versus fingerprint evidence: more of the same?. , 2010, , 105-128.		2
124	A Survey of UK Public Interest in Internet-Based Personal Genome Testing. <i>PLoS ONE</i> , 2010, 5, e13473.	2.5	93
125	The Prum Regime: Situated Dis/Empowerment in Transnational DNA Profile Exchange. <i>British Journal of Criminology</i> , 2010, 50, 1117-1135.	2.1	47
126	Attitudes of social science students in Israel and Austria towards the Belated Twins scenario – an exploratory study. <i>Public Understanding of Science</i> , 2010, 19, 435-451.	2.8	3



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127	How do we collaborate? Social science researchersâ€™ experience of multidisciplinary in biomedical settings. <i>BioSocieties</i> , 2010, 5, 278-286.	1.3	15
128	Direct-to-consumer genome testing: opportunities for pharmacogenomics research?. <i>Pharmacogenomics</i> , 2010, 11, 651-655.	1.3	35
129	Die Verflüssigung der Norm: Selbstregierung und personalisierte Gesundheit. , 2010, , 39-53.		1
130	Reply: Attitudes towards human reproductive cloning, ART and gene selection. <i>Human Reproduction</i> , 2009, 24, 2674-2674.	0.9	0
131	DNA Behind Bars. <i>Social Studies of Science</i> , 2009, 39, 51-79.	2.5	46
132	Separating The Social From the Natural. <i>Metascience</i> , 2009, 18, 475-479.	0.3	0
133	Being a member of the club: the transnational (self-)governance of networks of biobanks. <i>International Journal of Risk Assessment and Management</i> , 2009, 12, 64.	0.1	34
134	Misdirected precaution. <i>Nature</i> , 2008, 456, 34-35.	27.8	95
135	The Politics of Life Itself: Biomedicine, Power and Subjectivity in the Twenty-First Century - By N. Rose. <i>British Journal of Sociology</i> , 2008, 59, 818-819.	1.5	0
136	Clean Soil and Common Ground: The Biopolitics of Human Embryonic Stem Cell Research in Austria. <i>Science As Culture</i> , 2008, 17, 377-395.	3.2	12
137	Stem Cell Technologies 1998â€“2008: Controversies and Silences. <i>Science As Culture</i> , 2008, 17, 351-362.	3.2	27
138	Stem Cell Stories 1998â€“2008. <i>Science As Culture</i> , 2008, 17, 1-11.	3.2	33
139	What are the stakes? Genetic nondiscrimination legislation and personal genomics. <i>Personalized Medicine</i> , 2008, 5, 415-418.	1.5	7
140	Attitudes towards human reproductive cloning, assisted reproduction and gene selection: a survey of 4600 British twins. <i>Human Reproduction</i> , 2007, 22, 2302-2308.	0.9	10
141	Research populations: biobanks in Israel. <i>New Genetics and Society</i> , 2007, 26, 85-103.	1.2	21
142	â€˜Private fears in public places?â€™ Ethical and regulatory concerns regarding human genomic databases. <i>Personalized Medicine</i> , 2007, 4, 447-452.	1.5	7
143	â€˜Negotiating Lifeâ€™. <i>Social Studies of Science</i> , 2006, 36, 173-205.	2.5	108
144	â€˜Science for survivalâ€™: biotechnology regulation in Israel. <i>Science and Public Policy</i> , 2006, 33, 33-46.	2.4	34

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145	Relocating health governance: personalized medicine in times of 'global genes. <i>Personalized Medicine</i> , 2006, 3, 349-355.	1.5	4
146	Twins: A cloning experience. <i>Social Science and Medicine</i> , 2006, 63, 2739-2752.	3.8	7
147	The Rise of Genetic Couplehood? A Comparative View of Premarital Genetic Testing. <i>BioSocieties</i> , 2006, 1, 17-36.	1.3	47
148	Emotion in political discourse: contrasting approaches to stem cell governance in the USA, UK, Israel and Germany. <i>Regenerative Medicine</i> , 2006, 1, 823-829.	1.7	37
149	Genetically modified survival: Red and green biotechnology in Israel. <i>Science As Culture</i> , 2005, 14, 355-372.	3.2	7
150	Key issues in DNA profiling and databasing: implications for governance. , 0, , 15-39.		7
151	Partners in crime: the use of forensic DNA technologies in Austria. , 0, , 153-174.		5
152	Forensic utilization of voluntarily collected DNA samples: law enforcement versus human rights. , 0, , 40-62.		5
153	Base assumptions? Racial aspects of US DNA forensics. , 0, , 63-84.		4
154	Portuguese forensic DNA database: political enthusiasm, public trust and probable issues in future practice. , 0, , 218-239.		3
155	Beyond borders: trends and challenges in global forensic profiling and databasing. , 0, , 333-341.		0
156	DNA databases and the forensic imaginary. , 0, , 131-152.		14
157	DNA the Nor-way: black-boxing the evidence and monopolising the key. , 0, , 197-217.		3
158	Blue Chips and White Collars: Whose Data Science Is It?. , 0, , .		2
159	The Austrian Corona Panel Project: Monitoring Individual and Societal Dynamics amidst the COVID-19 Crisis. <i>SSRN Electronic Journal</i> , 0, , .	0.4	9
160	Labouring towards self-fulfilment: personal genomics and the "participatory turn". <i>Critical Public Health</i> , 0, , 1-13.	2.4	0
161	Ethics of Healthcare Policy and the Concept of Solidarity. , 0, , .		0