

# Ainsley J Newson

## List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

108  
papers

1,759  
citations

21  
h-index

38  
g-index

126  
ext. papers

2,188  
ext. citations

3.4  
avg, IF

5.15  
L-index

#	Paper	IF	Citations
108	Intertwined Interests in Expanded Prenatal Genetic Testing: The State's Role in Facilitating Equitable Access.. <i>American Journal of Bioethics</i> , <b>2022</b> , 22, 45-47	1.1	2
107	Human Genetics Society of Australasia Position Statement: Use of Human Genetic and Genomic Information in Healthcare Settings.. <i>Twin Research and Human Genetics</i> , <b>2022</b> , 1-8	2.2	
106	Correspondence on "Screening for autosomal recessive and X-linked conditions during pregnancy and preconception: a practice resource of the American College of Medical Genetics and Genomics (ACMG)" by Gregg et al.. <i>Genetics in Medicine</i> , <b>2022</b> ,	8.1	1
105	Ethical aspects of the changing landscape for spinal muscular atrophy management in Australia.. <i>Australian Journal of General Practice</i> , <b>2022</b> , 51, 131-135	1.5	0
104	The perils of a broad approach to public interest in health data research: a response to Ballantyne and Schaefer. <i>Journal of Medical Ethics</i> , <b>2021</b> , 47, 580-582	2.5	2
103	The promise of public health ethics for precision medicine: the case of newborn preventive genomic sequencing. <i>Human Genetics</i> , <b>2021</b> , 1	6.3	1
102	"I wish that there was more info": characterizing the uncertainty experienced by carriers of pathogenic ATM and/or CHEK2 variants. <i>Familial Cancer</i> , <b>2021</b> , 1	3	2
101	Acceptability of risk-stratified population screening across cancer types: Qualitative interviews with the Australian public. <i>Health Expectations</i> , <b>2021</b> , 24, 1326-1336	3.7	3
100	Ethics of Reproductive Genetic Carrier Screening: From the Clinic to the Population. <i>Public Health Ethics</i> , <b>2021</b> , 14, 202-217	1.8	3
99	Knowledge, views and expectations for cancer polygenic risk testing in clinical practice: A cross-sectional survey of health professionals. <i>Clinical Genetics</i> , <b>2021</b> , 100, 430-439	4	1
98	Reproductive carrier screening: responding to the eugenics critique. <i>Journal of Medical Ethics</i> , <b>2021</b> ,	2.5	1
97	Gene selection for the Australian Reproductive Genetic Carrier Screening Project ("Mackenzie Mission"). <i>European Journal of Human Genetics</i> , <b>2021</b> , 29, 79-87	5.3	22
96	Ethical issues in reproductive genetic carrier screening. <i>Medical Journal of Australia</i> , <b>2021</b> , 214, 165-167.	4.1	6
95	Family communication about genomic sequencing: A qualitative study with cancer patients and relatives. <i>Patient Education and Counseling</i> , <b>2021</b> , 104, 944-952	3.1	2
94	The expectations and realities of nutrigenomic testing in australia: A qualitative study. <i>Health Expectations</i> , <b>2021</b> , 24, 670-686	3.7	2
93	The Emergence and Global Spread of Noninvasive Prenatal Testing. <i>Annual Review of Genomics and Human Genetics</i> , <b>2021</b> , 22, 309-338	9.7	6
92	Impact of personal genomic risk information on melanoma prevention behaviors and psychological outcomes: a randomized controlled trial. <i>Genetics in Medicine</i> , <b>2021</b> , 23, 2394-2403	8.1	2

91	To offer or request? Disclosing variants of uncertain significance in prenatal testing. <i>Bioethics</i> , <b>2021</b> , 35, 900-909	2	2
90	Ethical considerations in gene selection for reproductive carrier screening. <i>Human Genetics</i> , <b>2021</b> , 1	6.3	2
89	Human Genetics Society of Australasia Position Statement: Predictive and Presymptomatic Genetic Testing in Adults and Children. <i>Twin Research and Human Genetics</i> , <b>2020</b> , 23, 184-189	2.2	5
88	Implementation considerations for offering personal genomic risk information to the public: a qualitative study. <i>BMC Public Health</i> , <b>2020</b> , 20, 1028	4.1	5
87	Cancer patients' views and understanding of genome sequencing: a qualitative study. <i>Journal of Medical Genetics</i> , <b>2020</b> , 57, 671-676	5.8	7
86	Disclosure to genetic relatives without consent - Australian genetic professionals' awareness of the health privacy law. <i>BMC Medical Ethics</i> , <b>2020</b> , 21, 13	2.9	4
85	There is a lot of good in knowing, but there is also a lot of downsides: public views on ethical considerations in population genomic screening. <i>Journal of Medical Ethics</i> , <b>2020</b> ,	2.5	1
84	Obligations and preferences in knowing and not knowing: the importance of context. <i>Journal of Medical Ethics</i> , <b>2020</b> , 46, 306-307	2.5	2
83	Who should access germline genome sequencing? A mixed methods study of patient views. <i>Clinical Genetics</i> , <b>2020</b> , 97, 329-337	4	1
82	From Expectations to Experiences: Consumer Autonomy and Choice in Personal Genomic Testing. <i>AJOB Empirical Bioethics</i> , <b>2020</b> , 11, 63-76	3	6
81	Dynamic Consent: An Evaluation and Reporting Framework. <i>Journal of Empirical Research on Human Research Ethics</i> , <b>2020</b> , 15, 175-186	1.6	12
80	Advanced cancer patient preferences for receiving molecular profiling results. <i>Psycho-Oncology</i> , <b>2020</b> , 29, 1533-1539	3.9	1
79	"Who is watching the watchdog?": ethical perspectives of sharing health-related data for precision medicine in Singapore. <i>BMC Medical Ethics</i> , <b>2020</b> , 21, 118	2.9	5
78	Technical Categories and Ethical Justifications: Why the Wik Approach is the Wrong Way Around for Categorizing Germ-Line Gene Editing. <i>American Journal of Bioethics</i> , <b>2020</b> , 20, 27-29	1.1	1
77	Australians' views and experience of personal genomic testing: survey findings from the Genioz study. <i>European Journal of Human Genetics</i> , <b>2019</b> , 27, 711-720	5.3	8
76	Public attitudes towards novel reproductive technologies: a citizens' survey on mitochondrial donation. <i>Human Reproduction</i> , <b>2019</b> , 34, 751-757	5.7	4
75	Genetic counselors' perceptions of uncertainty in pretest counseling for genomic sequencing: A qualitative study. <i>Journal of Genetic Counseling</i> , <b>2019</b> , 28, 292-303	2.5	8
74	Patient perspectives on molecular tumor profiling: "Why wouldn't you?". <i>BMC Cancer</i> , <b>2019</b> , 19, 753	4.8	12

73	Australian Genomics: A Federated Model for Integrating Genomics into Healthcare. <i>American Journal of Human Genetics</i> , <b>2019</b> , 105, 7-14	11	39
72	GP attitudes to and expectations for providing personal genomic risk information to the public: a qualitative study. <i>BJGP Open</i> , <b>2019</b> , 3, bjgpopen18X101633	3.1	6
71	Rapid Challenges: Ethics and Genomic Neonatal Intensive Care. <i>Pediatrics</i> , <b>2019</b> , 143, S14-S21	7.4	20
70	Australians' perspectives on support around use of personal genomic testing: Findings from the Genioz study. <i>European Journal of Medical Genetics</i> , <b>2019</b> , 62, 290-299	2.6	10
69	The PiGeOn project: protocol for a longitudinal study examining psychosocial, behavioural and ethical issues and outcomes in cancer tumour genomic profiling. <i>BMC Cancer</i> , <b>2018</b> , 18, 389	4.8	7
68	The PiGeOn project: protocol of a longitudinal study examining psychosocial and ethical issues and outcomes in germline genomic sequencing for cancer. <i>BMC Cancer</i> , <b>2018</b> , 18, 454	4.8	10
67	Distress, uncertainty, and positive experiences associated with receiving information on personal genomic risk of melanoma. <i>European Journal of Human Genetics</i> , <b>2018</b> , 26, 1094-1100	5.3	17
66	Australians' views on personal genomic testing: focus group findings from the Genioz study. <i>European Journal of Human Genetics</i> , <b>2018</b> , 26, 1101-1112	5.3	8
65	Reconceptualizing Autonomy for Bioethics. <i>Kennedy Institute of Ethics Journal</i> , <b>2018</b> , 28, 171-203	1.1	13
64	Human Genetics Society of Australasia Position Statement: Genetic Testing and Personal Insurance Products in Australia. <i>Twin Research and Human Genetics</i> , <b>2018</b> , 21, 533-537	2.2	6
63	The melanoma genomics managing your risk study: A protocol for a randomized controlled trial evaluating the impact of personal genomic risk information on skin cancer prevention behaviors. <i>Contemporary Clinical Trials</i> , <b>2018</b> , 70, 106-116	2.3	13
62	Whole genome sequencing in children: ethics, choice and deliberation. <i>Journal of Medical Ethics</i> , <b>2017</b> , 43, 540-542	2.5	10
61	Does personalized melanoma genomic risk information trigger conversations about skin cancer prevention and skin examination with family, friends and health professionals?. <i>British Journal of Dermatology</i> , <b>2017</b> , 177, 779-790	4	11
60	Is Mitochondrial Donation Germ-Line Gene Therapy? Classifications and Ethical Implications. <i>Bioethics</i> , <b>2017</b> , 31, 55-67	2	19
59	Scanning the body, sequencing the genome: Dealing with unsolicited findings. <i>Bioethics</i> , <b>2017</b> , 31, 648-656		16
58	Key challenges in bringing CRISPR-mediated somatic cell therapy into the clinic. <i>Genome Medicine</i> , <b>2017</b> , 9, 85	14.4	10
57	Genetics and Insurance in Australia: Concerns around a Self-Regulated Industry. <i>Public Health Genomics</i> , <b>2017</b> , 20, 247-256	1.9	11
56	A Pilot Randomized Controlled Trial of the Feasibility, Acceptability, and Impact of Giving Information on Personalized Genomic Risk of Melanoma to the Public. <i>Cancer Epidemiology Biomarkers and Prevention</i> , <b>2017</b> , 26, 212-221	4	33

55	Genomic Testing in The Paediatric Population: Ethical Considerations in Light of Recent Policy Statements. <i>Molecular Diagnosis and Therapy</i> , <b>2016</b> , 20, 407-14	4.5	10
54	The need for ethics as well as evidence in evidence-based medicine. <i>Journal of Clinical Epidemiology</i> , <b>2016</b> , 77, 7-10	5.7	4
53	Regulating Risk and the Boundaries of State Conduct: A Relational Perspective on Home Birth in Australia. <i>American Journal of Bioethics</i> , <b>2016</b> , 16, 19-21	1.1	7
52	Genomic intensive care: should we perform genome testing in critically ill newborns?. <i>Archives of Disease in Childhood: Fetal and Neonatal Edition</i> , <b>2016</b> , 101, F94-8	4.7	18
51	A pilot randomised controlled trial examining the feasibility, acceptability and impact of giving information on personalised genomic risk of melanoma to the public, for motivating preventive behaviours.. <i>Journal of Clinical Oncology</i> , <b>2016</b> , 34, 1556-1556	2.2	
50	Compensated transnational surrogacy in Australia: time for a comprehensive review. <i>Medical Journal of Australia</i> , <b>2016</b> , 204, 33-5	4	2
49	Known unknowns: building an ethics of uncertainty into genomic medicine. <i>BMC Medical Genomics</i> , <b>2016</b> , 9, 57	3.7	45
48	Ethical and legal issues in mitochondrial transfer. <i>EMBO Molecular Medicine</i> , <b>2016</b> , 8, 589-91	12	26
47	Do We Need Ethical Theory to Achieve Quality Critical Engagement in Clinical Ethics?. <i>American Journal of Bioethics</i> , <b>2016</b> , 16, 43-5	1.1	1
46	Public preferences for communicating personal genomic risk information: a focus group study. <i>Health Expectations</i> , <b>2016</b> , 19, 1203-1214	3.7	24
45	Rethinking pediatric ethics consultations. <i>American Journal of Bioethics</i> , <b>2015</b> , 15, 26-8	1.1	5
44	Exploring the Potential Emotional and Behavioural Impact of Providing Personalised Genomic Risk Information to the Public: A Focus Group Study. <i>Public Health Genomics</i> , <b>2015</b> , 18, 309-17	1.9	12
43	Why should ethics approval be required prior to publication of health promotion research?. <i>Health Promotion Journal of Australia</i> , <b>2015</b> , 26, 170-175	1.7	9
42	For your interest? The ethical acceptability of using non-invasive prenatal testing to test purely for information. <i>Bioethics</i> , <b>2015</b> , 29, 19-25	2	39
41	The value of clinical ethics support in Australian health care. <i>Medical Journal of Australia</i> , <b>2015</b> , 202, 568-9		10
40	"What should happen before asymptomatic men decide whether or not to have a PSA test?" A report on three community juries. <i>Medical Journal of Australia</i> , <b>2015</b> , 203, 335	4	10
39	Australians' knowledge and perceptions of direct-to-consumer personal genome testing. <i>Internal Medicine Journal</i> , <b>2014</b> , 44, 27-31	1.6	9
38	Dynamics and ethics of comprehensive preimplantation genetic testing: a review of the challenges. <i>Human Reproduction Update</i> , <b>2013</b> , 19, 366-75	15.8	54

37	Ethical considerations for choosing between possible models for using NIPD for aneuploidy detection. <i>Journal of Medical Ethics</i> , <b>2012</b> , 38, 614-8	2.5	34
36	Clinical Ethics Committee Case 17: a paramedic sustains a bite while attending a callout and the assailant refuses testing for HIV or hepatitis C: what should we do?. <i>Clinical Ethics</i> , <b>2012</b> , 7, 1-6	1	
35	Should non-invasiveness change informed consent procedures for prenatal diagnosis?. <i>Health Care Analysis</i> , <b>2011</b> , 19, 122-32	2.3	58
34	Current ethical issues in synthetic biology: where should we go from here?. <i>Accountability in Research</i> , <b>2011</b> , 18, 181-93	1.9	5
33	Clinical Ethics Committee Case 16: A request from an accident and emergency department should we give our patient a blood transfusion?. <i>Clinical Ethics</i> , <b>2011</b> , 6, 154-158	1	
32	Clinical Ethics Committee Case 10: For the record: Should our patient's relatives be able to record her treatment?. <i>Clinical Ethics</i> , <b>2010</b> , 5, 57-62	1	
31	Clinical Ethics Committee Case 9: Should we inform our patient about animal products in his medicine?. <i>Clinical Ethics</i> , <b>2010</b> , 5, 7-12	1	3
30	Childhood genetic testing for familial cancer: should adoption make a difference?. <i>Familial Cancer</i> , <b>2010</b> , 9, 37-42	3	9
29	Will the introduction of non-invasive prenatal diagnostic testing erode informed choices? An experimental study of health care professionals. <i>Patient Education and Counseling</i> , <b>2010</b> , 78, 24-8	3.1	101
28	Prenatal diagnosis and abortion for congenital abnormalities: is it ethical to provide one without the other?. <i>American Journal of Bioethics</i> , <b>2009</b> , 9, 48-56	1.1	27
27	Clinical Ethics Committee case 6: Our patient wishes to take an unlisted drug even though we're not sure of his diagnosis. <i>Clinical Ethics</i> , <b>2009</b> , 4, 59-63	1	
26	Clinical ethics committee case 7: our young patient is in heart failure but has multiple co-morbidities. How can we best care for him and his family?. <i>Clinical Ethics</i> , <b>2009</b> , 4, 111-115	1	
25	Personal genomics as an interactive web broadcast. <i>American Journal of Bioethics</i> , <b>2009</b> , 9, 27-9	1.1	1
24	Clinical Ethics Committee Case 8: Should we carry out a predictive genetic test in our young patient?. <i>Clinical Ethics</i> , <b>2009</b> , 4, 169-172	1	
23	Response to open peer commentaries on "Prenatal diagnosis and abortion for congenital abnormalities: is it ethical to provide one without the other?". <i>American Journal of Bioethics</i> , <b>2009</b> , 9, W6-7	1.1	
22	Clinical Ethics Committee case 5: Should we discharge our vulnerable patient to a family who seem unable to look after her?. <i>Clinical Ethics</i> , <b>2009</b> , 4, 6-11	1	
21	Is informed choice in prenatal testing universally valued? A population-based survey in Europe and Asia. <i>BJOG: an International Journal of Obstetrics and Gynaecology</i> , <b>2009</b> , 116, 880-5	3.7	23
20	Ethical aspects arising from non-invasive fetal diagnosis. <i>Seminars in Fetal and Neonatal Medicine</i> , <b>2008</b> , 13, 103-8	3.7	71

19	Informed choice in prenatal testing: a survey among obstetricians and gynaecologists in Europe and Asia. <i>Prenatal Diagnosis</i> , <b>2008</b> , 28, 1238-44	3.2	5
18	Commentary: Consent and confidentiality in publishing--the view of the BMJQ ethics committee. <i>BMJ, The</i> , <b>2008</b> , 337, a1232	5.9	3
17	Communication of Genetic Information within Families: The Case for Familial Comity. <i>Journal of Bioethical Inquiry</i> , <b>2006</b> , 3, 161-166	1.9	10
16	Clinical genetics and the problem with unqualified confidentiality. <i>American Journal of Bioethics</i> , <b>2006</b> , 6, 41-3; discussion W32-4	1.1	2
15	Should parental refusals of newborn screening be respected?. <i>Cambridge Quarterly of Healthcare Ethics</i> , <b>2006</b> , 15, 135-46	0.9	16
14	Behavioural genetics: why eugenic selection is preferable to enhancement. <i>Journal of Applied Philosophy</i> , <b>2006</b> , 23, 157-71	0.7	16
13	Whither authenticity?. <i>American Journal of Bioethics</i> , <b>2005</b> , 5, 53-5; discussion W10-2	1.1	10
12	Cascade testing in familial hypercholesterolaemia: how should family members be contacted?. <i>European Journal of Human Genetics</i> , <b>2005</b> , 13, 401-8	5.3	79
11	Artificial gametes: new paths to parenthood?. <i>Journal of Medical Ethics</i> , <b>2005</b> , 31, 184-6	2.5	33
10	The nature and significance of behavioural genetic information. <i>Theoretical Medicine and Bioethics</i> , <b>2004</b> , 25, 89-111	0.9	3
9	Consent to the publication of patient information: incompetent patients may pose a problem. <i>BMJ, The</i> , <b>2004</b> , 329, 916	5.9	0
8	Partially functional Cenpa-GFP fusion protein causes increased chromosome missegregation and apoptosis during mouse embryogenesis. <i>Chromosome Research</i> , <b>2003</b> , 11, 345-57	4.4	25
7	White coat ceremonies: a second opinion. <i>Journal of Medical Ethics</i> , <b>2002</b> , 28, 60-60	2.5	78
6	Early disruption of centromeric chromatin organization in centromere protein A (Cenpa) null mice. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , <b>2000</b> , 97, 1148-53	11.5	342
5	Should we undertake genetic research on intelligence?. <i>Bioethics</i> , <b>1999</b> , 13, 327-42	2	13
4	Gene structure and sequence analysis of mouse centromere proteins A and C. <i>Genomics</i> , <b>1998</b> , 47, 108-14	4.3	22
3	Chromosomal localization of mouse Cenpa gene. <i>Cytogenetic and Genome Research</i> , <b>1997</b> , 79, 298-301	1.9	
2	Population screening		4

1 Personhood and Moral Status 277-283