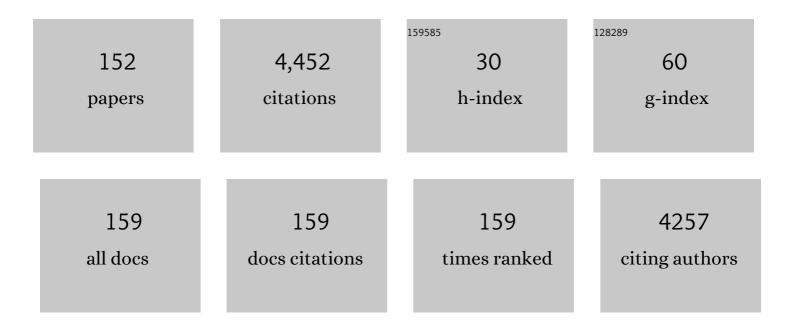
List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	The zone of parental discretion: An ethical tool for dealing with disagreement between parents and doctors about medical treatment for a child. Clinical Ethics, 2016, 11, 1-8.	0.7	174
2	Moral distress within neonatal and paediatric intensive care units: a systematic review. Archives of Disease in Childhood, 2016, 101, 701-708.	1.9	165
3	Slight/Mild Sensorineural Hearing Loss in Children. Pediatrics, 2006, 118, 1842-1851.	2.1	130
4	The Declaration of Helsinki, CIOMS and the ethics of research on vulnerable populations. Nature Medicine, 2000, 6, 615-617.	30.7	89
5	Human Research Ethics Committees: Examining Their Roles and Practices. Journal of Empirical Research on Human Research Ethics, 2012, 7, 38-49.	1.3	84
6	"You're one of us now― Young people describe their experiences of predictive genetic testing for Huntington disease (HD) and familial adenomatous polyposis (FAP). American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2008, 148C, 47-55.	1.6	82
7	Genetic counselling and ethical issues with chromosome microarray analysis in prenatal testing. Prenatal Diagnosis, 2012, 32, 389-395.	2.3	82
8	An international survey of predictive genetic testing in children for adult onset conditions. Genetics in Medicine, 2005, 7, 390-396.	2.4	75
9	Ethical Principles for the Management of Infants with Disorders of Sex Development. Hormone Research in Paediatrics, 2010, 74, 412-418.	1.8	72
10	How to facilitate decisions about surplus embryos: patients' views. Human Reproduction, 2007, 22, 3129-3138.	0.9	65
11	Should patients with self-inflicted illness receive lower priority in access to healthcare resources? Mapping out the debate. Journal of Medical Ethics, 2010, 36, 661-665.	1.8	60
12	An ethics core curriculum for Australasian medical schools. Medical Journal of Australia, 2001, 175, 205-210.	1.7	59
13	The psychosocial impact of Klinefelter syndrome and factors influencing quality of life. Genetics in Medicine, 2011, 13, 632-642.	2.4	58
14	Predictive genetic testing in minors for late-onset conditions: a chronological and analytical review of the ethical arguments: Figure 1. Journal of Medical Ethics, 2012, 38, 519-524.	1.8	55
15	What parents want from doctors in end-of-life decision-making for children. Archives of Disease in Childhood, 2014, 99, 216-220.	1.9	52
16	"Holding your breath― Interviews with young people who have undergone predictive genetic testing for Huntington disease. American Journal of Medical Genetics, Part A, 2007, 143A, 1984-1989.	1.2	51
17	Ethics at the end of life: Who should make decisions about treatment limitation for young children with lifeâ€threatening or life″imiting conditions?. Journal of Paediatrics and Child Health, 2011, 47, 594-598.	0.8	49
18	The use and misuse of moral distress in neonatology. Seminars in Fetal and Neonatal Medicine, 2018, 23, 39-43.	2.3	49

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19	Always a burden? Healthcare providers' perspectives on moral distress. Archives of Disease in Childhood: Fetal and Neonatal Edition, 2018, 103, F441-F445.	2.8	44
20	Ethics of fertility preservation for prepubertal children: should clinicians offer procedures where efficacy is largely unproven?. Journal of Medical Ethics, 2018, 44, 27-31.	1.8	43
21	Prenatal diagnosis and discrimination against the disabled Journal of Medical Ethics, 1999, 25, 163-171.	1.8	42
22	"We're checking them outâ€! Indigenous and non-Indigenous research participants' accounts of deciding to be involved in research. International Journal for Equity in Health, 2016, 15, 8.	3.5	40
23	Fertility in <scp>T</scp> urner syndrome. Clinical Endocrinology, 2013, 79, 606-614.	2.4	39
24	"lt was the missing piece― adolescent experiences of predictive genetic testing for adult-onset conditions. Genetics in Medicine, 2013, 15, 643-649.	2.4	39
25	The contribution of GJB2 mutations to slight or mild hearing loss in Australian elementary school children. Journal of Medical Genetics, 2006, 43, 850-855.	3.2	37
26	Emotions, Narratives, and Ethical Mindfulness. Academic Medicine, 2015, 90, 726-731.	1.6	37
27	Clinical, ethical and legal considerations in the treatment of newborns with non-ketotic hyperglycinaemia. Molecular Genetics and Metabolism, 2008, 94, 143-147.	1.1	34
28	Developing "Ethical Mindfulness―in Continuing Professional Development in Healthcare: Use of a Personal Narrative Approach. Cambridge Quarterly of Healthcare Ethics, 2009, 18, 197-208.	0.8	33
29	"Testing Times, Challenging Choices― An Australian Study of Prenatal Genetic Counseling. Journal of Genetic Counseling, 2010, 19, 22-37.	1.6	32
30	Role and function of a paediatric clinical ethics service: Experiences at the Royal Children's Hospital, Melbourne. Journal of Paediatrics and Child Health, 2011, 47, 632-636.	0.8	32
31	Conscientious objection to abortion, the law and its implementation in Victoria, Australia: perspectives of abortion service providers. BMC Medical Ethics, 2019, 20, 11.	2.4	31
32	Balancing health worker well-being and duty to care: an ethical approach to staff safety in COVID-19 and beyond. Journal of Medical Ethics, 2020, , medethics-2020-106557.	1.8	30
33	Can the Ethical Best Practice of Shared Decision-Making lead to Moral Distress?. Journal of Bioethical Inquiry, 2018, 15, 259-268.	1.5	28
34	Do Research Participants Trust Researchers or Their Institution?. Journal of Empirical Research on Human Research Ethics, 2018, 13, 285-294.	1.3	28
35	Fertility Preservation in Children and Adolescents With Cancer: Pilot of a Decision Aid for Parents of Children and Adolescents With Cancer. JMIR Pediatrics and Parenting, 2018, 1, e10463.	1.6	28
36	The challenge of developmentally appropriate care: predictive genetic testing in young people for familial adenomatous polyposis. Familial Cancer, 2010, 9, 27-35.	1.9	27

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37	The role of emotions in health professional ethics teaching. Journal of Medical Ethics, 2014, 40, 331-335.	1.8	26
38	A Study Protocol for the Australasian Oncofertility Registry: Monitoring Referral Patterns and the Uptake, Quality, and Complications of Fertility Preservation Strategies in Australia and New Zealand. Journal of Adolescent and Young Adult Oncology, 2016, 5, 215-225.	1.3	25
39	Genetic selection for deafness: the views of hearing children of deaf adults. Journal of Medical Ethics, 2009, 35, 722-728.	1.8	24
40	Postnatal screening for Klinefelter syndrome: is there a rationale?. Acta Paediatrica, International Journal of Paediatrics, 2011, 100, 923-933.	1.5	24
41	Attitudes to Genetic Testing for Deafness: The Importance of Informed Choice. Journal of Genetic Counseling, 2006, 15, 51-59.	1.6	23
42	Ethical controversies in prenatal microarray. Current Opinion in Obstetrics and Gynecology, 2013, 25, 133-137.	2.0	23
43	Decisionâ€making at the borderline of viability: Who should decide and on what basis?. Journal of Paediatrics and Child Health, 2017, 53, 105-111.	0.8	23
44	Satisfaction, disappointment and regret surrounding fertility preservation decisions in the paediatric and adolescent cancer population. Journal of Assisted Reproduction and Genetics, 2019, 36, 1805-1822.	2.5	23
45	Fertility Preservation Toolkit: A Clinician Resource to Assist Clinical Discussion and Decision Making in Pediatric and Adolescent Oncology. Journal of Pediatric Hematology/Oncology, 2018, 40, e133-e139.	0.6	22
46	Consent in paediatric research: an evaluation of the guidance provided in the 2007 NHMRC National statement on ethical conduct in human research. Medical Journal of Australia, 2008, 188, 360-362.	1.7	21
47	Parents and end-of-life decision-making for their child: roles and responsibilities. BMJ Supportive and Palliative Care, 2015, 5, 240-248.	1.6	21
48	Ethical complexities in child co-research. Research Ethics, 2019, 15, 1-16.	1.7	21
49	Uncertain diagnosis after newborn screening for cystic fibrosis: An ethicsâ€based approach to a clinical dilemma. Pediatric Pulmonology, 2014, 49, 1-7.	2.0	20
50	Telling the truth to young children: Ethical reasons for information disclosure in paediatrics. Journal of Paediatrics and Child Health, 2019, 55, 13-17.	0.8	20
51	<i>Caring Decisions:</i> The Development of a Written Resource for Parents Facing End-of-Life Decisions. Journal of Palliative Medicine, 2015, 18, 945-955.	1.1	19
52	Body Dysmorphic Disorder: Contraindication or Ethical Justification for Female Genital Cosmetic Surgery in Adolescents. Bioethics, 2016, 30, 706-713.	1.4	19
53	Moral Distress in Neonatology. Pediatrics, 2021, 148, .	2.1	19
54	Forever young? The ethics of ongoing puberty suppression for non-binary adults. Journal of Medical Ethics, 2020, 46, 743-752.	1.8	17

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55	Finding Out What Matters in Decision-Making Related to Genomics and Personalized Medicine in Pediatric Oncology: Developing Attributes to Include in a Discrete Choice Experiment. Patient, 2020, 13, 347-361.	2.7	17
56	Primary care physicians' views about gatekeeping in clinical research recruitment: A qualitative study. AJOB Empirical Bioethics, 2017, 8, 99-105.	1.6	16
57	Ethical considerations for paediatrics during the COVID â€19 pandemic: A discussion paper from the Australian Paediatric Clinical Ethics Collaboration. Journal of Paediatrics and Child Health, 2020, 56, 847-851.	0.8	15
58	Clinicians' perceptions of the Australian Paediatric Mental Health Service System: Problems and solutions. Australian and New Zealand Journal of Psychiatry, 2021, 55, 494-505.	2.3	15
59	Long-term Puberty Suppression for a Nonbinary Teenager. Pediatrics, 2020, 145, .	2.1	14
60	Understanding the Reasons Behind Healthcare Providers' Conscientious Objection to Voluntary Assisted Dying in Victoria, Australia. Journal of Bioethical Inquiry, 2021, 18, 277-289.	1.5	14
61	Clinical Utility of Precision Medicine in Pediatric Oncology: A Systematic Review. JCO Precision Oncology, 2021, 5, 1088-1102.	3.0	14
62	Human Research Ethics in Practice. Monash Bioethics Review, 2009, 28, 34-50.	0.8	13
63	Assessing the risks and benefits of diagnosing genetic conditions with variable phenotypes through population screening: Klinefelter syndrome as an example. Journal of Community Genetics, 2010, 1, 41-46.	1.2	13
64	"l'm scared of being like mumâ€i TheÂExperience of Adolescents Living inÂFamilies with Huntington Disease. Journal of Huntington's Disease, 2015, 4, 209-217.	1.9	13
65	Simulation-Based Research: A Scoping Review. Qualitative Health Research, 2020, 30, 2351-2360.	2.1	13
66	Arguing by Analogy in the Fetal Tissue Debate. Bioethics, 1997, 11, 397-412.	1.4	12
67	Resources Employed by Health Researchers to Ensure Ethical Research Practice. Journal of Empirical Research on Human Research Ethics, 2010, 5, 21-34.	1.3	12
68	Why Do Parents Want to Know their Child's Carrier Status? A Qualitative Study. Journal of Genetic Counseling, 2016, 25, 1257-1266.	1.6	12
69	Reflecting Before, During, and After the Heat of the Moment: A Review of Four Approaches for Supporting Health Staff to Manage Stressful Events. Journal of Bioethical Inquiry, 2021, 18, 573-587.	1.5	12
70	Medical and ethical considerations in twin pregnancies discordant for serious cardiac disease. Journal of Perinatology, 2009, 29, 662-667.	2.0	11
71	Ethical Issues in Fetal Management: A Cardiac Perspective. International Journal of Pediatrics (United) Tj ETQq1	1 0.784314	l rgBT /Over
72	Fetal Diagnosis – Obligations of the Clinician. Fetal Diagnosis and Therapy, 2007, 22, 233-237.	1.4	10

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73	The Unique Nature of Clinical Ethics in Allied Health Pediatrics: Implications for Ethics Education. Cambridge Quarterly of Healthcare Ethics, 2010, 19, 471-480.	0.8	10
74	Ethical Principles: An Essential Part of the Process in Disorders of Sex Development Care. Hormone Research in Paediatrics, 2011, 76, 367-368.	1.8	10
75	Decision-making approaches for children with life-limiting conditions: results from a qualitative phenomenological study. BMC Medical Ethics, 2022, 23, 52.	2.4	10
76	A commentary on the NH&MRC Draft Values and Ethics in Aboriginal and Torres Strait Islander Health Research. Monash Bioethics Review, 2003, 22, 8-19.	0.8	9
77	End of life decisionâ€making in paediatrics. Journal of Paediatrics and Child Health, 2008, 44, 389-391.	0.8	9
78	Carrier testing in children: exploration of genetic health professionals' practices in Australia. Genetics in Medicine, 2015, 17, 380-385.	2.4	9
79	Fifty years of paediatric ethics. Journal of Paediatrics and Child Health, 2015, 51, 8-11.	0.8	9
80	"Doing Trust― Journal of Empirical Research on Human Research Ethics, 2016, 11, 370-381.	1.3	9
81	How should neonatal clinicians act in the presence of moral distress?. Archives of Disease in Childhood: Fetal and Neonatal Edition, 2020, 105, 348-349.	2.8	9
82	Parents' experiences with requesting carrier testing for their unaffected children. Genetics in Medicine, 2016, 18, 1199-1205.	2.4	8
83	"l Don't See That as a Medical Problemâ€ŧ Clinicians' Attitudes and Responses to Requests for Cosme Genital Surgery by Adolescents. Journal of Bioethical Inquiry, 2018, 15, 535-548.	tic _{1.5}	8
84	A Clinical Decision Support System to Assist Pediatric Oncofertility: A Short Report. Journal of Adolescent and Young Adult Oncology, 2018, 7, 509-513.	1.3	8
85	Public attitudes towards novel reproductive technologies: a citizens' jury on mitochondrial donation. Human Reproduction, 2019, 34, 751-757.	0.9	8
86	Thinking outside the square: considering gender in Klinefelter syndrome and 47, XXY. Journal of Developmental and Physical Disabilities, 2011, 34, e348-e349.	3.6	7
87	Making Meaning From Experience: A Working Typology for Pediatrics Ethics Consultations. American Journal of Bioethics, 2015, 15, 24-26.	0.9	7
88	Therapeutic appropriation: a new concept in the ethics of clinical research. Journal of Medical Ethics, 2016, 42, 805-808.	1.8	7
89	The zone of parental discretion and the complexity of paediatrics: A response to Alderson. Clinical Ethics, 2018, 13, 172-174.	0.7	7
90	A practical and ethical toolkit for lastâ€minute refusal of anesthetic in children. Paediatric Anaesthesia, 2021, 31, 834-838.	1.1	7

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91	Children's bioethics and the zone of parental discretion. Monash Bioethics Review, 2010, 20, 09.1-3.	0.8	7
92	Enhancing the ethical conduct of genetic research: investigating views of parents on including their hearly children in a study on mild hearing loss. Journal of Medical Ethics, 2006, 32, 537-541.	1.8	6
93	Doctors' "judgements―and parents' "wishes― ethical implications in conflict situations. Medical Journal of Australia, 2014, 200, 372-372.	1.7	6
94	Collaboration in Clinical Ethics Consultation: A Method for Achieving "Balanced Accountability― American Journal of Bioethics, 2014, 14, 47-48.	0.9	6
95	Ethical considerations with the management of congenital central hypoventilation syndrome. Pediatric Pulmonology, 2015, 50, 503-510.	2.0	6
96	When is too little care, too much harm in cystic fibrosis? Psychological and ethical approaches to the problem. Journal of Cystic Fibrosis, 2017, 16, 299-303.	0.7	6
97	A good resource for parents, but will clinicians use it?: Evaluation of a resource for paediatric end-of-life decision making. BMC Palliative Care, 2017, 16, 12.	1.8	6
98	The Value of Open Deliberation in Clinical Ethics, and the Role of Parents' Reasons in the Zone of Parental Discretion. American Journal of Bioethics, 2018, 18, 47-49.	0.9	6
99	Treatment limitation and advance planning: Hospitalâ€wide audit of paediatric death. Journal of Paediatrics and Child Health, 2020, 56, 893-899.	0.8	6
100	Telling the Truth to Child Cancer Patients in COVID-19 Times. Journal of Bioethical Inquiry, 2020, 17, 797-801.	1.5	6
101	After an endâ€ofâ€life decision: Parents' reflections on living with an endâ€ofâ€life decision for their child. Journal of Paediatrics and Child Health, 2020, 56, 1060-1065.	0.8	6
102	Mild matters: parental insights into the conundrums of managing mild congenital hearing loss. International Journal of Audiology, 2022, 61, 500-506.	1.7	6
103	â€~Obstructive and power hungry'?: the Australian human research ethics process. Monash Bioethics Review, 2006, 25, S30-S38.	0.8	5
104	Ethical Mindfulness: Narrative Analysis and Everyday Ethics in Health Care. Advances in Medical Sociology, 2007, , 157-178.	0.1	5
105	Should longâ€ŧerm ventilation be offered in severe spinal muscular atrophy. Journal of Paediatrics and Child Health, 2013, 49, 813-814.	0.8	5
106	Paediatricians' decision making about prescribing stimulant medications for children with attentionâ€deficit/hyperactivity disorder. Child: Care, Health and Development, 2014, 40, 301-308.	1.7	5
107	Balancing obligations: should written information about life-sustaining treatment be neutral?. Journal of Medical Ethics, 2015, 41, 234-239.	1.8	5
108	Understanding decisions to participate in genomic medicine in children's cancer care: A comparison of what influences parents, health care providers, and the general community. Pediatric Blood and Cancer, 2021, 68, e29101.	1.5	5

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109	Whom are we seeking to protect? Extremely preterm babies and moral distress. Seminars in Perinatology, 2022, 46, 151549.	2.5	5
110	Conceptualising paediatric advance care planning: a qualitative phenomenological study of paediatricians caring for children with life-limiting conditions in Australia. BMJ Open, 2022, 12, e060077.	1.9	5
111	Telling the truth to seriously ill children: Considering children's interests when parents veto telling the truth. Bioethics, 2022, 36, 765-773.	1.4	5
112	Protecting researchers is not the role of HRECs. Australian and New Zealand Journal of Public Health, 2005, 29, 580-581.	1.8	4
113	Reply to Richards: "Predictive Genetic Testing of Adolescents for Huntington Disease: A Question of Autonomy and Harm― American Journal of Medical Genetics, Part A, 2008, 146A, 2447-2448.	1.2	4
114	Deception of children in research. Journal of Medical Ethics, 2015, 41, 179-182.	1.8	4
115	Predictive genetic testing for neurodegenerative conditions: how should conflicting interests within families be managed?. Journal of Medical Ethics, 2016, 42, 640-642.	1.8	4
116	Cosmetic Labiaplasty: Defining "Normality―Is Not the Issue. Journal of Pediatric and Adolescent Gynecology, 2017, 30, 438.	0.7	4
117	Facial Feminization Surgery: Privacy, Personal Identity, Compensatory Justice, and Resource Allocation. American Journal of Bioethics, 2018, 18, 12-15.	0.9	4
118	The practice of balancing in clinical ethics case consultation. Clinical Ethics, 2020, 15, 49-55.	0.7	4
119	Providing clarity around ethical discussion: development of a neonatal intervention score. Acta Paediatrica, International Journal of Paediatrics, 2019, 108, 1453-1459.	1.5	4
120	Ethical Issues in Transplantation. Monash Bioethics Review, 1992, 11, 28-39.	0.8	3
121	The 'More-Abortions' Objection to Fetal Tissue Transplantation. Journal of Medicine and Philosophy, 1998, 23, 411-427.	0.8	3
122	The new National Statement on Ethical Conduct in Research Involving Humans: A commentary. Monash Bioethics Review, 2000, 19, 1-13.	0.8	3
123	Confidentiality. Medical Journal of Australia, 2001, 174, 296-297.	1.7	3
124	Expertise in research ethics: Is there any such thing?. Monash Bioethics Review, 2004, 23, S58-S64.	0.8	3
125	Two's company, is three a crowd? Ethical cognition in decision making and the role of industry third parties in pediatric diabetes care. Pediatric Diabetes, 2019, 20, 15-22.	2.9	3
126	Bioethics and Public Policy in Australia. Politics and the Life Sciences, 1994, 13, 87-88.	0.7	2

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127	Secret ethics business?. Monash Bioethics Review, 2003, 22, S52-S62.	0.8	2
128	Democratizing "Psychotropic Neuroenhancement― AJOB Neuroscience, 2010, 1, 19-20.	1.1	2
129	Participants' Understanding of Informed Consent in a Randomized Controlled Trial for Chronic Knee Pain. Journal of Empirical Research on Human Research Ethics, 2015, 10, 435-443.	1.3	2
130	"They Just Want to Know――Genetic Health Professionals' Beliefs About Why Parents Want to Know their Child's Carrier Status. Journal of Genetic Counseling, 2017, 26, 1314-1323.	1.6	2
131	â€`Operating is the easy part': Surgeons' decision-making processes and responses to parental requests for elective paediatric appearance-altering facial surgery. Journal of Plastic, Reconstructive and Aesthetic Surgery, 2019, 72, 1379-1387.	1.0	2
132	The enactment stage of end-of-life decision-making for children. Palliative and Supportive Care, 2019, 17, 165-171.	1.0	2
133	Identity, well-being and autonomy in ongoing puberty suppression for non-binary adults: a response to the commentaries. Journal of Medical Ethics, 2020, 46, 761-762.	1.8	2
134	Managing aggression in hospitals: A role for clinical ethicists. Clinical Ethics, 2020, , 147775092097180.	0.7	2
135	Dissonance in views between parents and clinicians of children with serious illness: How can we bridge the gap?. Journal of Paediatrics and Child Health, 2021, 57, 1370-1375.	0.8	2
136	How can the education sector support children's mental health? Views of Australian healthcare clinicians. PLoS ONE, 2022, 17, e0261827.	2.5	2
137	Regulation, consultation and divergent community views: the case of access to ART by lesbian and single women. Journal of Law & Medicine, 2002, 9, 498-510.	0.0	2
138	Crossing the line? Ethics of parenteral nutrition in paediatric neurodisability complicated by intestinal failure. Archives of Disease in Childhood, 2023, 108, 11-14.	1.9	2
139	The Abortion Debate. Monash Bioethics Review, 1991, 11, 39-44.	0.8	1
140	Predictive testing in minors: the need for empirical evidence. Journal of Medical Ethics, 2012, 38, 533-534.	1.8	1
141	The Concept of "Continuing Creation―Will Not Save Us From Difficult Decisions. American Journal of Bioethics, 2017, 17, 23-25.	0.9	1
142	Ethical Considerations in Multiple Pregnancy: Preterm Delivery in the Setting of Discordant Fetal Anomaly. Twin Research and Human Genetics, 2019, 22, 120-123.	0.6	1
143	Protocol for a single patient therapy plan: A randomised, doubleâ€blind, placeboâ€controlled Nâ€ofâ€1 trial to assess the efficacy of cannabidiol in patients with intractable epilepsy. Journal of Paediatrics and Child Health, 2020, 56, 1918-1923.	0.8	1
144	An institutional audit of the use of novel drugs in pediatric oncology. Cancer Reports, 2021, , e1404.	1.4	1

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145	Ethical Principles for the Management of Children with Disorders of Sex Development: A Systematic Approach for Individual Cases. , 2012, , 147-157.		1
146	The Reproduction Revolution-A Christian Appraisal of Sexuality, Reproductive Technologies and the Family: Edited by John F Kilner, Paige C Cunningham and W David Hager, Grand Rapids Michigan, William B Eardmans Publishing Company, 2000, 290 pages, \$20, pound12.99 Journal of Medical Ethics, 2001, 27, 415-415.	1.8	0
147	Medical research and involuntary mental health patients: implications of proposed changes to legislation in Victoria. Monash Bioethics Review, 2005, 24, S45-S49.	0.8	Ο
148	Articulation and transparency of decision-making by human research ethics committees. Monash Bioethics Review, 2007, 26, 46-56.	0.8	0
149	The authors reply. Pediatric Pulmonology, 2015, 50, 211-211.	2.0	0
150	A Cautionary Note on Anonymous Referrals for Clinical Ethics Case Consultations. American Journal of Bioethics, 2015, 15, 32-33.	0.9	0
151	Extraordinary Survival: What Does It Take?. Pediatric Neurology, 2016, 64, 8-9.	2.1	0
152	Cloning to avoid genetic disease. , 2001, , 37-50.		0