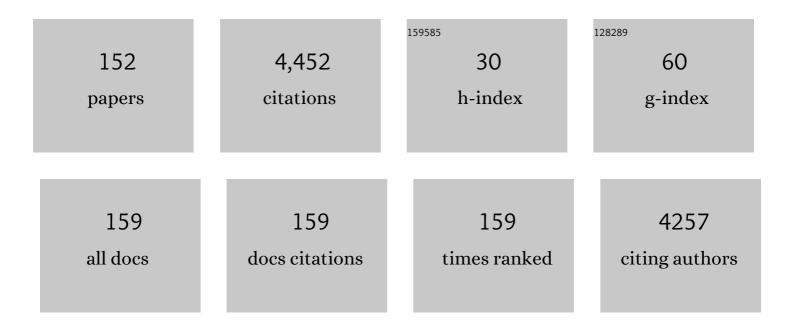
List of Publications by Year in descending order

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IVNIN CILLAM

#	Article	IF	CITATIONS
1	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
2	Mild matters: parental insights into the conundrums of managing mild congenital hearing loss. International Journal of Audiology, 2022, 61, 500-506.	1.7	6
3	Whom are we seeking to protect? Extremely preterm babies and moral distress. Seminars in Perinatology, 2022, 46, 151549.	2.5	5
4	How can the education sector support children's mental health? Views of Australian healthcare clinicians. PLoS ONE, 2022, 17, e0261827.	2.5	2
5	Decision-making approaches for children with life-limiting conditions: results from a qualitative phenomenological study. BMC Medical Ethics, 2022, 23, 52.	2.4	10
6	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
7	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
8	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
9	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
10	An institutional audit of the use of novel drugs in pediatric oncology. Cancer Reports, 2021, , e1404.	1.4	1
11	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
12	Clinical Utility of Precision Medicine in Pediatric Oncology: A Systematic Review. JCO Precision Oncology, 2021, 5, 1088-1102.	3.0	14
13	A practical and ethical toolkit for lastâ€minute refusal of anesthetic in children. Paediatric Anaesthesia, 2021, 31, 834-838.	1.1	7
14	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
15	Moral Distress in Neonatology. Pediatrics, 2021, 148, .	2.1	19
16	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
17	Treatment limitation and advance planning: Hospitalâ€wide audit of paediatric death. Journal of Paediatrics and Child Health, 2020, 56, 893-899.	0.8	6
18	The practice of balancing in clinical ethics case consultation. Clinical Ethics, 2020, 15, 49-55.	0.7	4

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19	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
20	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
21	Telling the Truth to Child Cancer Patients in COVID-19 Times. Journal of Bioethical Inquiry, 2020, 17, 797-801.	1.5	6
22	Forever young? The ethics of ongoing puberty suppression for non-binary adults. Journal of Medical Ethics, 2020, 46, 743-752.	1.8	17
23	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
24	Simulation-Based Research: A Scoping Review. Qualitative Health Research, 2020, 30, 2351-2360.	2.1	13
25	Managing aggression in hospitals: A role for clinical ethicists. Clinical Ethics, 2020, , 147775092097180.	0.7	2
26	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
27	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
28	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
29	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
30	Long-term Puberty Suppression for a Nonbinary Teenager. Pediatrics, 2020, 145, .	2.1	14
31	Ethical complexities in child co-research. Research Ethics, 2019, 15, 1-16.	1.7	21
32	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
33	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
34	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
35	Public attitudes towards novel reproductive technologies: a citizens' jury on mitochondrial donation. Human Reproduction, 2019, 34, 751-757.	0.9	8
36	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

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37	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
38	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
39	The enactment stage of end-of-life decision-making for children. Palliative and Supportive Care, 2019, 17, 165-171.	1.0	2
40	Providing clarity around ethical discussion: development of a neonatal intervention score. Acta Paediatrica, International Journal of Paediatrics, 2019, 108, 1453-1459.	1.5	4
41	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
42	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
43	Can the Ethical Best Practice of Shared Decision-Making lead to Moral Distress?. Journal of Bioethical Inquiry, 2018, 15, 259-268.	1.5	28
44	Do Research Participants Trust Researchers or Their Institution?. Journal of Empirical Research on Human Research Ethics, 2018, 13, 285-294.	1.3	28
45	The use and misuse of moral distress in neonatology. Seminars in Fetal and Neonatal Medicine, 2018, 23, 39-43.	2.3	49
46	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
47	"l Don't See That as a Medical Problem― Clinicians' Attitudes and Responses to Requests for Cosmet Genital Surgery by Adolescents. Journal of Bioethical Inquiry, 2018, 15, 535-548.	^{:ic} 1.5	8
48	Facial Feminization Surgery: Privacy, Personal Identity, Compensatory Justice, and Resource Allocation. American Journal of Bioethics, 2018, 18, 12-15.	0.9	4
49	The zone of parental discretion and the complexity of paediatrics: A response to Alderson. Clinical Ethics, 2018, 13, 172-174.	0.7	7
50	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
51	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
52	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
53	Cosmetic Labiaplasty: Defining "Normality―Is Not the Issue. Journal of Pediatric and Adolescent Gynecology, 2017, 30, 438.	0.7	4
54	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

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55	"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
56	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
57	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
58	The Concept of "Continuing Creation―Will Not Save Us From Difficult Decisions. American Journal of Bioethics, 2017, 17, 23-25.	0.9	1
59	Primary care physicians' views about gatekeeping in clinical research recruitment: A qualitative study. AJOB Empirical Bioethics, 2017, 8, 99-105.	1.6	16
60	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
61	Therapeutic appropriation: a new concept in the ethics of clinical research. Journal of Medical Ethics, 2016, 42, 805-808.	1.8	7
62	Parents' experiences with requesting carrier testing for their unaffected children. Genetics in Medicine, 2016, 18, 1199-1205.	2.4	8
63	Body Dysmorphic Disorder: Contraindication or Ethical Justification for Female Genital Cosmetic Surgery in Adolescents. Bioethics, 2016, 30, 706-713.	1.4	19
64	"Doing Trust― Journal of Empirical Research on Human Research Ethics, 2016, 11, 370-381.	1.3	9
65	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
66	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
67	Extraordinary Survival: What Does It Take?. Pediatric Neurology, 2016, 64, 8-9.	2.1	0
68	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
69	"We're checking them out†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
70	Moral distress within neonatal and paediatric intensive care units: a systematic review. Archives of Disease in Childhood, 2016, 101, 701-708.	1.9	165
71	<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
72	"l'm scared of being like mumâ€: TheÂExperience of Adolescents Living inÂFamilies with Huntington Disease. Journal of Huntington's Disease, 2015, 4, 209-217.	1.9	13

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73	The authors reply. Pediatric Pulmonology, 2015, 50, 211-211.	2.0	0
74	Emotions, Narratives, and Ethical Mindfulness. Academic Medicine, 2015, 90, 726-731.	1.6	37
75	Deception of children in research. Journal of Medical Ethics, 2015, 41, 179-182.	1.8	4
76	Balancing obligations: should written information about life-sustaining treatment be neutral?. Journal of Medical Ethics, 2015, 41, 234-239.	1.8	5
77	Carrier testing in children: exploration of genetic health professionals' practices in Australia. Genetics in Medicine, 2015, 17, 380-385.	2.4	9
78	A Cautionary Note on Anonymous Referrals for Clinical Ethics Case Consultations. American Journal of Bioethics, 2015, 15, 32-33.	0.9	0
79	Making Meaning From Experience: A Working Typology for Pediatrics Ethics Consultations. American Journal of Bioethics, 2015, 15, 24-26.	0.9	7
80	Ethical considerations with the management of congenital central hypoventilation syndrome. Pediatric Pulmonology, 2015, 50, 503-510.	2.0	6
81	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
82	Fifty years of paediatric ethics. Journal of Paediatrics and Child Health, 2015, 51, 8-11.	0.8	9
83	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
84	Doctors' "judgements―and parents' "wishes― ethical implications in conflict situations. Medical Journal of Australia, 2014, 200, 372-372.	1.7	6
85	Collaboration in Clinical Ethics Consultation: A Method for Achieving "Balanced Accountabilityâ€. American Journal of Bioethics, 2014, 14, 47-48.	0.9	6
86	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
87	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
88	The role of emotions in health professional ethics teaching. Journal of Medical Ethics, 2014, 40, 331-335.	1.8	26
89	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
90	Fertility in <scp>T</scp> urner syndrome. Clinical Endocrinology, 2013, 79, 606-614.	2.4	39

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91	"lt was the missing pieceâ€: adolescent experiences of predictive genetic testing for adult-onset conditions. Genetics in Medicine, 2013, 15, 643-649.	2.4	39
92	Ethical controversies in prenatal microarray. Current Opinion in Obstetrics and Gynecology, 2013, 25, 133-137.	2.0	23
93	Should longâ€ŧerm ventilation be offered in severe spinal muscular atrophy. Journal of Paediatrics and Child Health, 2013, 49, 813-814.	0.8	5
94	Predictive testing in minors: the need for empirical evidence. Journal of Medical Ethics, 2012, 38, 533-534.	1.8	1
95	Human Research Ethics Committees: Examining Their Roles and Practices. Journal of Empirical Research on Human Research Ethics, 2012, 7, 38-49.	1.3	84
96	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
97	Genetic counselling and ethical issues with chromosome microarray analysis in prenatal testing. Prenatal Diagnosis, 2012, 32, 389-395.	2.3	82
98	Ethical Principles for the Management of Children with Disorders of Sex Development: A Systematic Approach for Individual Cases. , 2012, , 147-157.		1
99	The psychosocial impact of Klinefelter syndrome and factors influencing quality of life. Genetics in Medicine, 2011, 13, 632-642.	2.4	58
100	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
101	Ethics at the end of life: Who should make decisions about treatment limitation for young children with lifeâ€threatening or lifeâ€limiting conditions?. Journal of Paediatrics and Child Health, 2011, 47, 594-598.	0.8	49
102	Postnatal screening for Klinefelter syndrome: is there a rationale?. Acta Paediatrica, International Journal of Paediatrics, 2011, 100, 923-933.	1.5	24
103	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
104	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
105	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
106	"Testing Times, Challenging Choices― An Australian Study of Prenatal Genetic Counseling. Journal of Genetic Counseling, 2010, 19, 22-37.	1.6	32
107	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
108	Democratizing "Psychotropic Neuroenhancement― AJOB Neuroscience, 2010, 1, 19-20.	1.1	2

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109	Ethical Principles for the Management of Infants with Disorders of Sex Development. Hormone Research in Paediatrics, 2010, 74, 412-418.	1.8	72
110	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
111	Ethical Issues in Fetal Management: A Cardiac Perspective. International Journal of Pediatrics (United) Tj ETQq1 1	0.784314 0.8	l rgBT /Over
112	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
113	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
114	Children's bioethics and the zone of parental discretion. Monash Bioethics Review, 2010, 20, 09.1-3.	0.8	7
115	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
116	Medical and ethical considerations in twin pregnancies discordant for serious cardiac disease. Journal of Perinatology, 2009, 29, 662-667.	2.0	11
117	Genetic selection for deafness: the views of hearing children of deaf adults. Journal of Medical Ethics, 2009, 35, 722-728.	1.8	24
118	Human Research Ethics in Practice. Monash Bioethics Review, 2009, 28, 34-50.	0.8	13
119	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
120	"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
121	End of life decisionâ€making in paediatrics. Journal of Paediatrics and Child Health, 2008, 44, 389-391.	0.8	9
122	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
123	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
124	How to facilitate decisions about surplus embryos: patients' views. Human Reproduction, 2007, 22, 3129-3138.	0.9	65
125	Fetal Diagnosis – Obligations of the Clinician. Fetal Diagnosis and Therapy, 2007, 22, 233-237.	1.4	10
126	Articulation and transparency of decision-making by human research ethics committees. Monash Bioethics Review, 2007, 26, 46-56.	0.8	0

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127	Ethical Mindfulness: Narrative Analysis and Everyday Ethics in Health Care. Advances in Medical Sociology, 2007, , 157-178.	0.1	5
128	"Holding your breathâ€i 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
129	â€~Obstructive and power hungry'?: the Australian human research ethics process. Monash Bioethics Review, 2006, 25, S30-S38.	0.8	5
130	Attitudes to Genetic Testing for Deafness: The Importance of Informed Choice. Journal of Genetic Counseling, 2006, 15, 51-59.	1.6	23
131	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
132	Enhancing the ethical conduct of genetic research: investigating views of parents on including their healthy children in a study on mild hearing loss. Journal of Medical Ethics, 2006, 32, 537-541.	1.8	6
133	Slight/Mild Sensorineural Hearing Loss in Children. Pediatrics, 2006, 118, 1842-1851.	2.1	130
134	Protecting researchers is not the role of HRECs. Australian and New Zealand Journal of Public Health, 2005, 29, 580-581.	1.8	4
135	An international survey of predictive genetic testing in children for adult onset conditions. Genetics in Medicine, 2005, 7, 390-396.	2.4	75
136	Medical research and involuntary mental health patients: implications of proposed changes to legislation in Victoria. Monash Bioethics Review, 2005, 24, S45-S49.	0.8	0
137	Expertise in research ethics: Is there any such thing?. Monash Bioethics Review, 2004, 23, S58-S64.	0.8	3
138	Secret ethics business?. Monash Bioethics Review, 2003, 22, S52-S62.	0.8	2
139	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
140	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
141	An ethics core curriculum for Australasian medical schools. Medical Journal of Australia, 2001, 175, 205-210.	1.7	59
142	Confidentiality. Medical Journal of Australia, 2001, 174, 296-297.	1.7	3
143	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

144 Cloning to avoid genetic disease. , 2001, , 37-50.

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145	The Declaration of Helsinki, CIOMS and the ethics of research on vulnerable populations. Nature Medicine, 2000, 6, 615-617.	30.7	89
146	The new National Statement on Ethical Conduct in Research Involving Humans: A commentary. Monash Bioethics Review, 2000, 19, 1-13.	0.8	3
147	Prenatal diagnosis and discrimination against the disabled Journal of Medical Ethics, 1999, 25, 163-171.	1.8	42
148	The 'More-Abortions' Objection to Fetal Tissue Transplantation. Journal of Medicine and Philosophy, 1998, 23, 411-427.	0.8	3
149	Arguing by Analogy in the Fetal Tissue Debate. Bioethics, 1997, 11, 397-412.	1.4	12
150	Bioethics and Public Policy in Australia. Politics and the Life Sciences, 1994, 13, 87-88.	0.7	2
151	Ethical Issues in Transplantation. Monash Bioethics Review, 1992, 11, 28-39.	0.8	3
152	The Abortion Debate. Monash Bioethics Review, 1991, 11, 39-44.	0.8	1