

John Massie Fracp

List of Publications by Year in descending order

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Version: 2024-02-01

108
papers

3,658
citations

201575

27
h-index

138417

58
g-index

110
all docs

110
docs citations

110
times ranked

3625
citing authors

#	ARTICLE	IF	CITATIONS
1	Vaccination of young people from 12 years of age for COVID-19 against parents' wishes. Medical Journal of Australia, 2022, , .	0.8	3
2	Factors in childhood associated with lung function decline to adolescence in cystic fibrosis. Journal of Cystic Fibrosis, 2022, 21, 977-983.	0.3	4
3	Lessons from Frankenstein 200 years on: brain organoids, chimaeras and other "monsters". Journal of Medical Ethics, 2021, 47, 567-571.	1.0	11
4	Exercise-induced laryngeal obstruction in children and adolescents: are we listening?. Archives of Disease in Childhood: Education and Practice Edition, 2021, 106, 66-70.	0.3	3
5	A narrative review of the experience and decision-making for children on home mechanical ventilation. Journal of Paediatrics and Child Health, 2021, 57, 791-796.	0.4	15
6	Lives changed: A new era for people with cystic fibrosis. Journal of Paediatrics and Child Health, 2021, 57, 968-970.	0.4	1
7	Suffering and the end of life. Journal of Paediatrics and Child Health, 2021, 57, 979-980.	0.4	1
8	A practical and ethical toolkit for last-minute refusal of anesthetic in children. Paediatric Anaesthesia, 2021, 31, 834-838.	0.6	7
9	Miracles in my time: Reflections of a pediatric respiratory physician. Pediatric Pulmonology, 2021, 56, 3586-3591.	1.0	1
10	Handshake or high-five: Greeting children. Journal of Paediatrics and Child Health, 2021, , .	0.4	0
11	Deciding with children. Archives of Disease in Childhood, 2021, , archdischild-2021-323048.	1.0	1
12	Investigating transmission of Mycobacterium abscessus amongst children in an Australian cystic fibrosis centre. Journal of Cystic Fibrosis, 2020, 19, 219-224.	0.3	47
13	How to interpret polysomnography. Archives of Disease in Childhood: Education and Practice Edition, 2020, 105, 130-135.	0.3	2
14	International perspectives on the implementation of reproductive carrier screening. Prenatal Diagnosis, 2020, 40, 301-310.	1.1	60
15	Which types of conditions should be included in reproductive genetic carrier screening? Views of parents of children with a genetic condition. European Journal of Medical Genetics, 2020, 63, 104075.	0.7	14
16	Seven Sins of Medicine: Reconsideration 70 years on. Journal of Paediatrics and Child Health, 2020, 56, 1846-1847.	0.4	0
17	Letter to the editor. Journal of Paediatrics and Child Health, 2020, 56, 660-661.	0.4	0
18	RISK AND JETSKIS. Journal of Paediatrics and Child Health, 2020, 56, 1323-1323.	0.4	0

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19	My love affair with the pleural space. Medical Journal of Australia, 2020, 213, 526.	0.8	1
20	Respiratory management of infants with chronic neonatal lung disease beyond the <scp>NICU</scp>: A position statement from the Thoracic Society of Australia and New Zealand*. Respirology, 2020, 25, 880-888.	1.3	15
21	Early markers of cystic fibrosis structural lung disease: follow-up of the ACFBAL cohort. European Respiratory Journal, 2020, 55, 1901694.	3.1	14
22	Letter to the editor. Journal of Paediatrics and Child Health, 2020, 56, 661-661.	0.4	0
23	Utility of Endobronchial Ultrasound in Assessment of Intrathoracic Lesions in Paediatric Patients. Respiration, 2019, 98, 340-346.	1.2	8
24	The Gutmacherâ€™ Lancet Commission on sexual and reproductive health and rights: how does Australia measure up?. Medical Journal of Australia, 2019, 211, 381.	0.8	0
25	Medical conditions revealed in fairy tales, folklore and literature. Journal of Paediatrics and Child Health, 2019, 55, 1295-1298.	0.4	1
26	Viral induced hypersecretion of mucous (Vlper). Pediatric Pulmonology, 2019, 54, 683-683.	1.0	0
27	Genetic neologisms: Collective nouns for the geneticist. Journal of Paediatrics and Child Health, 2019, 55, 1506-1507.	0.4	0
28	As You Take It. The Seven Ages of Coffee (with thanks to William Shakespeare, â€™As You Like Itâ€™™). Journal of Paediatrics and Child Health, 2019, 55, 1503-1503.	0.4	0
29	Repurposing medical equipment. Medical Journal of Australia, 2019, 211, 527-528.	0.8	1
30	Preconception and antenatal carrier screening for genetic conditions: The critical role of general practitioners. Australian Journal of General Practice, 2019, 48, 106-110.	0.3	23
31	Reproductive genetic carrier screening for cystic fibrosis, fragile X syndrome, and spinal muscular atrophy in Australia: outcomes of 12,000 tests. Genetics in Medicine, 2018, 20, 513-523.	1.1	80
32	Laboratory performance of sweat conductivity for the screening of cystic fibrosis. Clinical Chemistry and Laboratory Medicine, 2018, 56, 554-559.	1.4	7
33	Elimination of Australian epidemic strain (AES1) <i>pseudomonas aeruginosa</i> in a pediatric cystic fibrosis center. Pediatric Pulmonology, 2018, 53, 1498-1503.	1.0	7
34	MRÎ€: Inside the meat pie. Journal of Medical Imaging and Radiation Oncology, 2018, 62, 361-363.	0.9	1
35	Cystic Fibrosis Transmembrane Conductance Regulator-Related Metabolic Syndrome and Cystic Fibrosis Screen Positive, Inconclusive Diagnosis. Journal of Pediatrics, 2017, 181, S45-S51.e1.	0.9	95
36	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.	0.3	6

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37	A Review of Treatments That Improve Cystic Fibrosis Transmembrane Conductance Regulator Function. <i>Clinical Medicine Insights Therapeutics</i> , 2017, 9, 1179559X1771912.	0.4	2
38	Australasian Guideline (2nd Edition): an Annex to the CLSI and UK Guidelines for the Performance of the Sweat Test for the Diagnosis of Cystic Fibrosis. <i>Clinical Biochemist Reviews</i> , 2017, 38, 115-130.	3.3	11
39	Community-acquired pneumonia in children: what to do when there is no response to standard empirical treatment?. <i>Thorax</i> , 2016, 71, 957-959.	2.7	0
40	Newborn screening for cystic fibrosis. <i>Lancet Respiratory Medicine</i> , 2016, 4, 653-661.	5.2	118
41	Parents'™ experiences with requesting carrier testing for their unaffected children. <i>Genetics in Medicine</i> , 2016, 18, 1199-1205.	1.1	8
42	The story of cystic fibrosis 1965–2015. <i>Journal of Paediatrics and Child Health</i> , 2016, 52, 991-994.	0.4	4
43	Psychosocial characteristics and predictors of health care use in families of young children with cystic fibrosis in Western Australia. <i>Journal of Paediatrics and Child Health</i> , 2016, 52, 34-39.	0.4	7
44	Big drain, little drain, soft drain: Ouch! A few questions about the management of empyema thoracis (in the style of Dr Seuss). <i>Journal of Paediatrics and Child Health</i> , 2016, 52, 847-847.	0.4	0
45	Newborn screening and population carrier screening for cystic fibrosis: Two ends of the same rope. <i>Journal of Cystic Fibrosis</i> , 2016, 15, 407-408.	0.3	0
46	The authors reply. <i>Pediatric Pulmonology</i> , 2015, 50, 211-211.	1.0	0
47	“Suddenly Having two Positive People who are Carriers is a Whole New Thing”–Experiences of Couples Both Identified as Carriers of Cystic Fibrosis Through a Population-Based Carrier Screening Program in Australia. <i>Journal of Genetic Counseling</i> , 2015, 24, 987-1000.	0.9	25
48	Spirometry and regular follow-up do not improve quality of life in children or adolescents with asthma: Cluster randomized controlled trials. <i>Pediatric Pulmonology</i> , 2015, 50, 947-954.	1.0	15
49	Ethical considerations with the management of congenital central hypoventilation syndrome. <i>Pediatric Pulmonology</i> , 2015, 50, 503-510.	1.0	6
50	Population-based genetic carrier screening for cystic fibrosis in Victoria. <i>Medical Journal of Australia</i> , 2014, 200, 205-206.	0.8	5
51	Attitudes and opinions of pregnant women who are not offered cystic fibrosis carrier screening. <i>European Journal of Human Genetics</i> , 2014, 22, 859-865.	1.4	6
52	Population-based carrier screening for cystic fibrosis: a systematic review of 23 years of research. <i>Genetics in Medicine</i> , 2014, 16, 207-216.	1.1	72
53	Uncertain diagnosis after newborn screening for cystic fibrosis: An ethics-based approach to a clinical dilemma. <i>Pediatric Pulmonology</i> , 2014, 49, 1-7.	1.0	20
54	Human Genetics Society of Australasia Position Statement: Population-Based Carrier Screening for Cystic Fibrosis. <i>Twin Research and Human Genetics</i> , 2014, 17, 578-583.	0.3	15

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55	Newborn screening and carrier screening for cystic fibrosis: alternative or complementary?. European Respiratory Journal, 2014, 43, 20-23.	3.1	20
56	Prenatal and preconception population carrier screening for cystic fibrosis in Australia: Where are we up to?. Australian and New Zealand Journal of Obstetrics and Gynaecology, 2014, 54, 503-509.	0.4	9
57	Respiratory physicians and clinic coordinators' attitudes to population-based cystic fibrosis carrier screening. Journal of Cystic Fibrosis, 2014, 13, 99-105.	0.3	4
58	Sixty-five years since the New York heat wave: Advances in sweat testing for cystic fibrosis. Pediatric Pulmonology, 2014, 49, 106-117.	1.0	56
59	Carrier screening for cystic fibrosis in the new era of medications that restore CFTR function. Lancet, The, 2014, 383, 923-925.	6.3	21
60	Cystic Fibrosis Carrier Screening. Paediatric Respiratory Reviews, 2013, 14, 270-275.	1.2	16
61	Understanding the Costs of Care for Cystic Fibrosis: An Analysis by Age and Health State. Value in Health, 2013, 16, 345-355.	0.1	71
62	Cascade carrier testing after a child is diagnosed with cystic fibrosis through newborn screening: investigating why most relatives do not have testing. Genetics in Medicine, 2013, 15, 533-540.	1.1	22
63	Current Practice and Attitudes of Australian Obstetricians Toward Population-Based Carrier Screening for Inherited Conditions. Twin Research and Human Genetics, 2013, 16, 601-607.	0.3	22
64	The natural history and predictors of persistent problem behaviours in cystic fibrosis: a multicentre, prospective study. Archives of Disease in Childhood, 2012, 97, 625-631.	1.0	21
65	Australian epidemic strain pseudomonas (AES-1) declines further in a cohort segregated cystic fibrosis clinic. Journal of Cystic Fibrosis, 2012, 11, 49-52.	0.3	27
66	Cost-effectiveness of carrier screening for cystic fibrosis in Australia. Journal of Cystic Fibrosis, 2012, 11, 281-287.	0.3	22
67	Cochrane Review: Antibiotics for whooping cough (pertussis). Evidence-Based Child Health: A Cochrane Review Journal, 2012, 7, 893-956.	2.0	7
68	Infection, Inflammation, and Lung Function Decline in Infants with Cystic Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2011, 184, 75-81.	2.5	256
69	Spontaneous chylothorax in a 2-year-old child. Medical Journal of Australia, 2011, 195, 385-385.	0.8	4
70	A case for cystic fibrosis carrier testing in the general population. Medical Journal of Australia, 2011, 194, 208-209.	0.8	17
71	Emerging issues in cystic fibrosis newborn screening. Current Opinion in Pulmonary Medicine, 2010, 16, 584-590.	1.2	19
72	Uptake of carrier testing in families after cystic fibrosis diagnosis through newborn screening. European Journal of Human Genetics, 2010, 18, 1084-1089.	1.4	36

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73	A home respiratory support programme for children by parents and layperson carers. Journal of Paediatrics and Child Health, 2010, 46, 57-62.	0.4	53
74	Letters to the Editor. Journal of Paediatrics and Child Health, 2010, 46, 210-210.	0.4	0
75	Ethical considerations in choosing a model for population-based cystic fibrosis carrier screening. Medical Journal of Australia, 2010, 193, 157-160.	0.8	17
76	Reducing the burden of inherited disease: the Human Variome Project. Medical Journal of Australia, 2010, 193, 430-431.	0.8	2
77	Declining prevalence of cystic fibrosis since the introduction of newborn screening. Archives of Disease in Childhood, 2010, 95, 531-533.	1.0	33
78	Benchmarks for Cystic Fibrosis carrier screening: A European consensus document. Journal of Cystic Fibrosis, 2010, 9, 165-178.	0.3	75
79	Spontaneous chylothorax in a 2-year-old child. Medical Journal of Australia, 2009, 190, 262-264.	0.8	10
80	Population-based carrier screening for cystic fibrosis in Victoria: The first three years experience. Australian and New Zealand Journal of Obstetrics and Gynaecology, 2009, 49, 484-489.	0.4	42
81	Chylothorax: Diagnosis and Management in Children. Paediatric Respiratory Reviews, 2009, 10, 199-207.	1.2	139
82	Carrier screening for cystic fibrosis. Lancet, The, 2009, 374, 978.	6.3	3
83	Safety of bronchoalveolar lavage in young children with cystic fibrosis. Pediatric Pulmonology, 2008, 43, 965-972.	1.0	48
84	Guidelines for Diagnosis of Cystic Fibrosis in Newborns through Older Adults: Cystic Fibrosis Foundation Consensus Report. Journal of Pediatrics, 2008, 153, S4-S14.	0.9	904
85	Re: Acrodermatitis enteropathica-like eruption and failure to thrive as presenting signs of cystic fibrosis. Journal of the American Academy of Dermatology, 2008, 59, 720-721.	0.6	3
86	Economic evaluation of cystic fibrosis screening: A review of the literature. Health Policy, 2008, 85, 133-147.	1.4	27
87	Guidelines for Implementation of Cystic Fibrosis Newborn Screening Programs: Cystic Fibrosis Foundation Workshop Report. Pediatrics, 2007, 119, e495-e518.	1.0	139
88	Estimating inspired oxygen concentration delivered by nasal prongs in children with bronchiolitis. Journal of Paediatrics and Child Health, 2007, 44, 070719023208002-???.	0.4	7
89	Community-wide screening for cystic fibrosis carriers could replace newborn screening for the diagnosis of cystic fibrosis. Journal of Paediatrics and Child Health, 2007, 43, 721-723.	0.4	16
90	Cough in children: when does it matter?. Paediatric Respiratory Reviews, 2006, 7, 9-14.	1.2	8

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91	Sweat testing for cystic fibrosis: How good is your laboratory?. Journal of Paediatrics and Child Health, 2006, 42, 153-154.	0.4	7
92	Parental attitudes to the identification of their infants as carriers of cystic fibrosis by newborn screening. Journal of Paediatrics and Child Health, 2006, 42, 533-537.	0.4	61
93	Pharmacokinetic profile of once daily intravenous tobramycin in children with cystic fibrosis. Journal of Paediatrics and Child Health, 2006, 42, 601-605.	0.4	34
94	Changing Their Minds With Time: A Comparison of Hypothetical and Actual Reproductive Behaviors in Parents of Children With Cystic Fibrosis. Pediatrics, 2006, 118, e649-e656.	1.0	69
95	Diagnosis of cystic fibrosis after newborn screening: The Australasian experience?twenty years and five million babies later: A consensus statement from the Australasian paediatric respiratory group. Pediatric Pulmonology, 2005, 39, 440-446.	1.0	79
96	Screening couples for cystic fibrosis carrier status: why are we waiting?. Medical Journal of Australia, 2005, 183, 501-502.	0.8	25
97	Effects of Segregation on an Epidemic Pseudomonas aeruginosa Strain in a Cystic Fibrosis Clinic. American Journal of Respiratory and Critical Care Medicine, 2005, 171, 1020-1025.	2.5	77
98	The relevance of sweat testing for the diagnosis of cystic fibrosis in the genomic era. Clinical Biochemist Reviews, 2005, 26, 135-53.	3.3	79
99	Internal mandibular distraction to relieve airway obstruction in infants and young children with micrognathia. Pediatric Pulmonology, 2004, 37, 230-235.	1.0	31
100	National study of infants hospitalized with pertussis in the acellular vaccine era. Pediatric Infectious Disease Journal, 2004, 23, 246-252.	1.1	105
101	Exercise-Induced Asthma in Children. Paediatric Drugs, 2002, 4, 267-278.	1.3	10
102	Sweat testing following newborn screening for cystic fibrosis. , 2000, 29, 452-456.		48
103	Long-term outcome of surgically treated acquired subglottic stenosis in infancy. Pediatric Pulmonology, 2000, 30, 125-130.	1.0	12
104	Newborn screening for cystic fibrosis in Victoria: 10 years' experience (1989-1998). Medical Journal of Australia, 2000, 172, 584-587.	0.8	86
105	Bronchial granuloma "where's the foreign body?. International Journal of Pediatric Otorhinolaryngology, 2000, 53, 215-219.	0.4	9
106	Long-term outcome of surgically treated acquired subglottic stenosis in infancy. , 2000, 30, 125.		1
107	Cystic fibrosis: The twilight zone. , 1999, 28, 222-224.		1
108	The Economic Aspects of Drug Delivery in Asthma. Pharmacoeconomics, 1997, 11, 398-407.	1.7	5