

Christopher Morris

List of Publications by Year in descending order

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Version: 2024-02-01

87
papers

3,423
citations

136740

32
h-index

155451

55
g-index

90
all docs

90
docs citations

90
times ranked

4027
citing authors

#	ARTICLE	IF	CITATIONS
1	Mini-EDACS: Development of the Eating and Drinking Ability Classification System for young children with cerebral palsy. <i>Developmental Medicine and Child Neurology</i> , 2022, 64, 897-906.	1.1	5
2	Effect of different durations of using a standing frame on the rate of hip migration in children with moderate to severe cerebral palsy: a feasibility study for a randomised controlled trial. <i>Physiotherapy</i> , 2022, 116, 42-49.	0.2	1
3	Developing the FEEDS toolkit of parent-delivered interventions for eating, drinking and swallowing difficulties in young children with neurodisability: findings from a Delphi survey and stakeholder consultation workshops. <i>BMJ Paediatrics Open</i> , 2022, 6, e001425.	0.6	3
4	Feasibility of a randomised controlled trial to evaluate home-based virtual reality therapy in children with cerebral palsy. <i>Disability and Rehabilitation</i> , 2021, 43, 85-97.	0.9	27
5	Ankle foot orthoses for young children with cerebral palsy: a scoping review. <i>Disability and Rehabilitation</i> , 2021, 43, 726-738.	0.9	14
6	Parent-delivered interventions used at home to improve eating, drinking and swallowing in children with neurodisability: the FEEDS mixed-methods study. <i>Health Technology Assessment</i> , 2021, 25, 1-208.	1.3	14
7	Protocol for The Toxin Study: Understanding clinical and patient reported response of children and young people with cerebral palsy to intramuscular lower limb Botulinum neurotoxin-A injections, exploring all domains of the ICF. A pragmatic longitudinal observational study using a prospective one-group repeated measures design. <i>BMJ Open</i> , 2021, 11, e049542.	0.8	1
8	Implementation of training to improve communication with disabled children on the ward: A feasibility study. <i>Health Expectations</i> , 2021, 24, 1433-1442.	1.1	5
9	Children with neurodisability and feeding difficulties: a UK survey of parent-delivered interventions. <i>BMJ Paediatrics Open</i> , 2021, 5, e001095.	0.6	5
10	Healthy Parent Carers: feasibility randomised controlled trial of a peer-led group-based health promotion intervention for parent carers of disabled children. <i>Pilot and Feasibility Studies</i> , 2021, 7, 144.	0.5	9
11	Healthy Parent Carers programme: mixed methods process evaluation and refinement of a health promotion intervention. <i>BMJ Open</i> , 2021, 11, e045570.	0.8	5
12	Development of a core outcome set for evaluative research into paediatric cerebral visual impairment (CVI), in the UK and Ireland. <i>BMJ Open</i> , 2021, 11, e051014.	0.8	3
13	Online community engagement in response to COVID-19 pandemic. <i>Health Expectations</i> , 2021, 24, 728-730.	1.1	10
14	Improving continence in children and young people with neurodisability: a systematic review and survey. <i>Health Technology Assessment</i> , 2021, 25, 1-258.	1.3	1
15	Mapping epilepsy-specific patient-reported outcome measures for children to a proposed core outcome set for childhood epilepsy. <i>Epilepsy and Behavior</i> , 2020, 112, 107372.	0.9	11
16	Systematic review: measurement properties of patient-reported outcome measures evaluated with childhood brain tumor survivors or other acquired brain injury. <i>Neuro-Oncology Practice</i> , 2020, 7, 277-287.	1.0	7
17	Community-based family and carer-support programmes for children with disabilities. <i>Paediatrics and Child Health (United Kingdom)</i> , 2020, 30, 180-185.	0.2	2
18	Resources for parents raising a disabled child in the UK. <i>Paediatrics and Child Health (United Kingdom)</i> , 2020, 30, 180-185.	0.2	1

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19	Epilepsy-specific patient-reported outcome measures of children's health-related quality of life: A systematic review of measurement properties. <i>Epilepsia</i> , 2020, 61, 230-248.	2.6	19
20	Public involvement in health research: what does "good" look like in practice?. <i>Research Involvement and Engagement</i> , 2020, 6, 11.	1.1	46
21	Sleep positioning systems for children and adults with a neurodisability: A systematic review. <i>British Journal of Occupational Therapy</i> , 2019, 82, 5-14.	0.5	10
22	The Eating and Drinking Ability Classification System for cerebral palsy: A study of reliability and stability over time. <i>Journal of Pediatric Rehabilitation Medicine</i> , 2019, 12, 123-131.	0.3	8
23	Working with Patients and Members of the Public: Informing Health Economics in Child Health Research. <i>PharmacoEconomics - Open</i> , 2019, 3, 133-141.	0.9	9
24	"They've walked the walk" A systematic review of quantitative and qualitative evidence for parent-to-parent support for parents of babies in neonatal care. <i>Journal of Neonatal Nursing</i> , 2019, 25, 166-176.	0.3	6
25	Core Health Outcomes in Childhood Epilepsy (CHOICE): Development of a core outcome set using systematic review methods and a Delphi survey consensus. <i>Epilepsia</i> , 2019, 60, 857-871.	2.6	32
26	Healthy Parent Carers peer-led group-based health promotion intervention for parent carers of disabled children: protocol for a feasibility study using a parallel group randomised controlled trial design. <i>Pilot and Feasibility Studies</i> , 2019, 5, 137.	0.5	2
27	Autism and primary care dentistry: parents' experiences of taking children with autism or working diagnosis of autism for dental examinations. <i>International Journal of Paediatric Dentistry</i> , 2018, 28, 226-238.	1.0	46
28	Parents Suggest Which Indicators of Progress and Outcomes Should be Measured in Young Children with Autism Spectrum Disorder. <i>Journal of Autism and Developmental Disorders</i> , 2018, 48, 1041-1051.	1.7	54
29	Parent-to-parent support interventions for parents of babies cared for in a neonatal unit" protocol of a systematic review of qualitative and quantitative evidence. <i>Systematic Reviews</i> , 2018, 7, 179.	2.5	12
30	Healthy Parent Carers programme: development and feasibility of a novel group-based health-promotion intervention. <i>BMC Public Health</i> , 2018, 18, 270.	1.2	27
31	Therapeutic potential and ownership of commercially available consoles in children with cerebral palsy. <i>British Journal of Occupational Therapy</i> , 2017, 80, 108-116.	0.5	5
32	Interventions utilising contact with people with disabilities to improve children's attitudes towards disability: A systematic review and meta-analysis. <i>Disability and Health Journal</i> , 2017, 10, 11-22.	1.6	68
33	Improving communication between staff and disabled children in hospital wards: testing the feasibility of a training intervention developed through intervention mapping. <i>BMJ Paediatrics Open</i> , 2017, 1, e000103.	0.6	6
34	Reaching consensus on reporting patient and public involvement (PPI) in research: methods and lessons learned from the development of reporting guidelines. <i>BMJ Open</i> , 2017, 7, e016948.	0.8	35
35	Rasch analysis of the Chedoke-McMaster Attitudes towards Children with Handicaps scale. <i>Disability and Rehabilitation</i> , 2017, 39, 281-290.	0.9	13
36	Core Health Outcomes In Childhood Epilepsy (CHOICE): protocol for the selection of a core outcome set. <i>Trials</i> , 2017, 18, 572.	0.7	13

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37	Towards development and validation of an intraoperative assessment tool for robot-assisted radical prostatectomy training: results of a Delphi study. <i>International Braz J Urol: Official Journal of the Brazilian Society of Urology</i> , 2017, 43, 661-670.	0.7	5
38	Resources for parents raising a disabled child in the UK. <i>Paediatrics and Child Health (United Kingdom)</i> , 2017, 102, 702-707.	0.2	2
39	Measurement properties of multidimensional patient-reported outcome measures in neurodisability: a systematic review of evaluation studies. <i>Developmental Medicine and Child Neurology</i> , 2016, 58, 437-451.	1.1	30
40	Communicating with disabled children when inpatients: barriers and facilitators identified by parents and professionals in a qualitative study. <i>Health Expectations</i> , 2016, 19, 738-750.	1.1	26
41	Children's contact with people with disabilities and their attitudes towards disability: a cross-sectional study. <i>Disability and Rehabilitation</i> , 2016, 38, 879-888.	0.9	52
42	Re: The Incidence and Relative Risk of Cardiovascular Toxicity in Patients Treated with New Hormonal Agents for Castration-resistant Prostate Cancer. <i>European Urology</i> , 2016, 69, 177-178.	0.9	0
43	End-user involvement in a systematic review of quantitative and qualitative research of non-pharmacological interventions for attention deficit hyperactivity disorder delivered in school settings: reflections on the impacts and challenges. <i>Health Expectations</i> , 2016, 19, 1084-1097.	1.1	17
44	Sleep positioning systems for children with cerebral palsy. <i>The Cochrane Library</i> , 2015, 2015, CD009257.	1.5	13
45	Involving disabled children and young people as partners in research: a systematic review. <i>Child: Care, Health and Development</i> , 2015, 41, 505-514.	0.8	114
46	A Systematic Review of Generic Multidimensional Patient-Reported Outcome Measures for Children, Part I: Descriptive Characteristics. <i>Value in Health</i> , 2015, 18, 315-333.	0.1	56
47	A Systematic Review of Generic Multidimensional Patient-Reported Outcome Measures for Children, Part II: Evaluation of Psychometric Performance of English-Language Versions in a General Population. <i>Value in Health</i> , 2015, 18, 334-345.	0.1	37
48	Setting research priorities to improve the health of children and young people with neurodisability: a British Academy of Childhood Disability-James Lind Alliance Research Priority Setting Partnership. <i>BMJ Open</i> , 2015, 5, e006233-e006233.	0.8	90
49	Meaningful health outcomes for paediatric neurodisability: Stakeholder prioritisation and appropriateness of patient reported outcome measures. <i>Health and Quality of Life Outcomes</i> , 2015, 13, 87.	1.0	53
50	Engaging stakeholders in rehabilitation research: a scoping review of strategies used in partnerships and evaluation of impacts. <i>Disability and Rehabilitation</i> , 2015, 37, 1390-1400.	0.9	147
51	Systematic review of tools to measure outcomes for young children with autism spectrum disorder. <i>Health Technology Assessment</i> , 2015, 19, 1-506.	1.3	203
52	Non-pharmacological interventions for attention-deficit/hyperactivity disorder (ADHD) delivered in school settings: systematic reviews of quantitative and qualitative research. <i>Health Technology Assessment</i> , 2015, 19, 1-470.	1.3	55
53	The association between children's contact with people with disabilities and their attitudes towards disability: a systematic review. <i>Developmental Medicine and Child Neurology</i> , 2014, 56, 529-546.	1.1	69
54	Key health outcomes for children and young people with neurodisability: qualitative research with young people and parents. <i>BMJ Open</i> , 2014, 4, e004611.	0.8	63

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55	Development and reliability of a system to classify the eating and drinking ability of people with cerebral palsy. <i>Developmental Medicine and Child Neurology</i> , 2014, 56, 245-251.	1.1	187
56	Health outcomes for children with neurodisability: what do professionals regard as primary targets?. <i>Archives of Disease in Childhood</i> , 2014, 99, 927-932.	1.0	20
57	A systematic review of ordinal scales used to classify the eating and drinking abilities of individuals with cerebral palsy. <i>Developmental Medicine and Child Neurology</i> , 2014, 56, 313-322.	1.1	18
58	Informing the NHS Outcomes Framework: evaluating meaningful health outcomes for children with neurodisability using multiple methods including systematic review, qualitative research, Delphi survey and consensus meeting. <i>Health Services and Delivery Research</i> , 2014, 2, 1-224.	1.4	40
59	Peer support for parents of children with chronic disabling conditions: a systematic review of quantitative and qualitative studies. <i>Developmental Medicine and Child Neurology</i> , 2013, 55, 602-609.	1.1	176
60	Meeting the Information Needs of Families of Children with Chronic Health Conditions. <i>Physical and Occupational Therapy in Pediatrics</i> , 2013, 33, 265-270.	0.8	8
61	Towards a definition of neurodisability: a Delphi survey. <i>Developmental Medicine and Child Neurology</i> , 2013, 55, 1103-1108.	1.1	113
62	Why it is crucial to involve families in all stages of childhood disability research. <i>Developmental Medicine and Child Neurology</i> , 2011, 53, 769-771.	1.1	41
63	Orthotic management of cerebral palsy: Recommendations from a consensus conference. <i>NeuroRehabilitation</i> , 2011, 28, 37-46.	0.5	80
64	â€˜The relationship of cerebral palsy subtype and functional motor impairment: a populationâ€˜based studyâ€™. <i>Developmental Medicine and Child Neurology</i> , 2010, 52, 682-683.	1.1	7
65	The Oxford Ankle Foot Questionnaire for Children. <i>Prosthetics and Orthotics International</i> , 2010, 34, 238-244.	0.5	20
66	ISPO Cerebral Palsy Consensus Conference Report (available free at www.ispoweb.org). <i>Prosthetics and Orthotics International</i> , 2009, 33, 401-402.	0.5	6
67	The Oxford Ankle Foot Questionnaire for children: responsiveness and longitudinal validity. <i>Quality of Life Research</i> , 2009, 18, 1367-1376.	1.5	51
68	Measuring participation in childhood disability: how does the capability approach improve our understanding?. <i>Developmental Medicine and Child Neurology</i> , 2009, 51, 92-94.	1.1	59
69	Aiming to improve the health care of people with cerebral palsy worldwide: a report of an International Society for Prosthetics and Orthotics conference. <i>Developmental Medicine and Child Neurology</i> , 2009, 51, 689-689.	1.1	1
70	Current and future uses of the Gross Motor Function Classification System: the need to take account of other factors to explain functional outcomes. <i>Developmental Medicine and Child Neurology</i> , 2009, 51, 1003-1003.	1.1	2
71	Development of the Gross Motor Function Classification System (1997). <i>Developmental Medicine and Child Neurology</i> , 2008, 50, 5-5.	1.1	22
72	A year of foot and ankle orthotic provision for adults: Prospective consultations data, with patient satisfaction survey. <i>Foot</i> , 2008, 18, 75-83.	0.4	12

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73	Definition and classification of cerebral palsy: a historical perspective. <i>Developmental Medicine and Child Neurology</i> , 2007, 49, 3-7.	1.1	163
74	Do the abilities of children with cerebral palsy explain their activities and participation?. <i>Developmental Medicine and Child Neurology</i> , 2006, 48, 954.	1.1	96
75	Reliability of the Manual Ability Classification System for children with cerebral palsy. <i>Developmental Medicine and Child Neurology</i> , 2006, 48, 950.	1.1	105
76	Reliability of the Manual Ability Classification System for children with cerebral palsy. <i>Developmental Medicine and Child Neurology</i> , 2006, 48, 950-953.	1.1	80
77	Do the abilities of children with cerebral palsy explain their activities and participation?. <i>Developmental Medicine and Child Neurology</i> , 2006, 48, 954-961.	1.1	5
78	Editorial. <i>Prosthetics and Orthotics International</i> , 2005, 29, 115-117.	0.5	0
79	Reliability of family report for the Gross Motor Function Classification System. <i>Developmental Medicine and Child Neurology</i> , 2004, 46, 455-60.	1.1	55
80	Gross Motor Function Classification System: impact and utility. <i>Developmental Medicine and Child Neurology</i> , 2004, 46, 60-5.	1.1	83
81	Gross Motor Function Classification System: impact and utility. <i>Developmental Medicine and Child Neurology</i> , 2004, 46, 60-65.	1.1	155
82	Reliability of family report for the Gross Motor Function Classification System. <i>Developmental Medicine and Child Neurology</i> , 2004, 46, 455-460.	1.1	105
83	Orthotic Management of Children with Cerebral Palsy. <i>Journal of Prosthetics and Orthotics</i> , 2002, 14, 150-158.	0.2	24
84	A review of the efficacy of lower-limb orthoses used for cerebral palsy. <i>Developmental Medicine and Child Neurology</i> , 2002, 44, 205.	1.1	94
85	Variations in the orthotic management of cerebral palsy. <i>Child: Care, Health and Development</i> , 2002, 28, 139-147.	0.8	22
86	A review of the efficacy of lower-limb orthoses used for cerebral palsy. <i>Developmental Medicine and Child Neurology</i> , 2002, 44, 205-211.	1.1	4
87	Aspects of methodology relevant to patient-reported outcome measures (PROMs). , 0, , 1-9.		1