Christopher Morris

List of Publications by Year in descending order

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87 papers 3,423 citations

32 h-index 55 g-index

90 all docs 90 docs citations

90 times ranked 4027 citing authors

#	Article	IF	CITATIONS
1	Systematic review of tools to measure outcomes for young children with autism spectrum disorder. Health Technology Assessment, 2015, 19, 1-506.	1.3	203
2	Development and reliability of a system to classify the eating and drinking ability of people with cerebral palsy. Developmental Medicine and Child Neurology, 2014, 56, 245-251.	1.1	187
3	Peer support for parents of children with chronic disabling conditions: a systematic review of quantitative and qualitative studies. Developmental Medicine and Child Neurology, 2013, 55, 602-609.	1.1	176
4	Definition and classification of cerebral palsy: a historical perspective. Developmental Medicine and Child Neurology, 2007, 49, 3-7.	1.1	163
5	Gross Motor Function Classification System: impact and utility. Developmental Medicine and Child Neurology, 2004, 46, 60-65.	1.1	155
6	Engaging stakeholders in rehabilitation research: a scoping review of strategies used in partnerships and evaluation of impacts. Disability and Rehabilitation, 2015, 37, 1390-1400.	0.9	147
7	Involving disabled children and young people as partners in research: a systematic review. Child: Care, Health and Development, 2015, 41, 505-514.	0.8	114
8	Towards a definition of neurodisability: a Delphi survey. Developmental Medicine and Child Neurology, 2013, 55, 1103-1108.	1.1	113
9	Reliability of the Manual Ability Classification System for children with cerebral palsy. Developmental Medicine and Child Neurology, 2006, 48, 950.	1.1	105
10	Reliability of family report for the Gross Motor Function Classification System. Developmental Medicine and Child Neurology, 2004, 46, 455-460.	1.1	105
11	Do the abilities of children with cerebral palsy explain their activities and participation?. Developmental Medicine and Child Neurology, 2006, 48, 954.	1.1	96
12	A review of the efficacy of lower-limb orthoses used for cerebral palsy. Developmental Medicine and Child Neurology, 2002, 44, 205.	1.1	94
13	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. BMJ Open, 2015, 5, e006233-e006233.	0.8	90
14	Gross Motor Function Classification System: impact and utility. Developmental Medicine and Child Neurology, 2004, 46, 60-5.	1.1	83
15	Reliability of the Manual Ability Classification System for children with cerebral palsy. Developmental Medicine and Child Neurology, 2006, 48, 950-953.	1.1	80
16	Orthotic management of cerebral palsy: Recommendations from a consensus conference. NeuroRehabilitation, 2011, 28, 37-46.	0.5	80
17	The association between children's contact with people with disabilities and their attitudes towards disability: a systematic review. Developmental Medicine and Child Neurology, 2014, 56, 529-546.	1.1	69
18	Interventions utilising contact with people with disabilities to improve children's attitudes towards disability: A systematic review and meta-analysis. Disability and Health Journal, 2017, 10, 11-22.	1.6	68

#	Article	IF	Citations
19	Key health outcomes for children and young people with neurodisability: qualitative research with young people and parents. BMJ Open, 2014, 4, e004611.	0.8	63
20	Measuring participation in childhood disability: how does the capability approach improve our understanding?. Developmental Medicine and Child Neurology, 2009, 51, 92-94.	1.1	59
21	A Systematic Review of Generic Multidimensional Patient-Reported Outcome Measures for Children, Part I: Descriptive Characteristics. Value in Health, 2015, 18, 315-333.	0.1	56
22	Reliability of family report for the Gross Motor Function Classification System. Developmental Medicine and Child Neurology, 2004, 46, 455-60.	1.1	55
23	Non-pharmacological interventions for attention-deficit/hyperactivity disorder (ADHD) delivered in school settings: systematic reviews of quantitative and qualitative research. Health Technology Assessment, 2015, 19, 1-470.	1.3	55
24	Parents Suggest Which Indicators of Progress and Outcomes Should be Measured in Young Children with Autism Spectrum Disorder. Journal of Autism and Developmental Disorders, 2018, 48, 1041-1051.	1.7	54
25	Meaningful health outcomes for paediatric neurodisability: Stakeholder prioritisation and appropriateness of patient reported outcome measures. Health and Quality of Life Outcomes, 2015, 13, 87.	1.0	53
26	Children's contact with people with disabilities and their attitudes towards disability: a cross-sectional study. Disability and Rehabilitation, 2016, 38, 879-888.	0.9	52
27	The Oxford Ankle Foot Questionnaire for children: responsiveness and longitudinal validity. Quality of Life Research, 2009, 18, 1367-1376.	1.5	51
28	Autism and primary care dentistry: parents' experiences of taking children with autism or working diagnosis of autism for dental examinations. International Journal of Paediatric Dentistry, 2018, 28, 226-238.	1.0	46
29	Public involvement in health research: what does â€~good' look like in practice?. Research Involvement and Engagement, 2020, 6, 11.	1.1	46
30	Why it is crucial to involve families in all stages of childhood disability research. Developmental Medicine and Child Neurology, 2011, 53, 769-771.	1.1	41
31	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. Health Services and Delivery Research, 2014, 2, 1-224.	1.4	40
32	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. Value in Health, 2015, 18, 334-345.	0.1	37
33	Reaching consensus on reporting patient and public involvement (PPI) in research: methods and lessons learned from the development of reporting guidelines. BMJ Open, 2017, 7, e016948.	0.8	35
34	Core Health Outcomes in Childhood Epilepsy (CHOICE): Development of a core outcome set using systematic review methods and a Delphi survey consensus. Epilepsia, 2019, 60, 857-871.	2.6	32
35	Measurement properties of multidimensional patientâ€reported outcome measures in neurodisability: a systematic review of evaluation studies. Developmental Medicine and Child Neurology, 2016, 58, 437-451.	1.1	30
36	Healthy Parent Carers programme: development and feasibility of a novel group-based health-promotion intervention. BMC Public Health, 2018, 18, 270.	1.2	27

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37	Feasibility of a randomised controlled trial to evaluate home-based virtual reality therapy in children with cerebral palsy. Disability and Rehabilitation, 2021, 43, 85-97.	0.9	27
38	Communicating with disabled children when inpatients: barriers and facilitators identified by parents and professionals in a qualitative study. Health Expectations, 2016, 19, 738-750.	1.1	26
39	Orthotic Management of Children with Cerebral Palsy. Journal of Prosthetics and Orthotics, 2002, 14, 150-158.	0.2	24
40	Variations in the orthotic management of cerebral palsy. Child: Care, Health and Development, 2002, 28, 139-147.	0.8	22
41	Development of the Gross Motor Function Classification System (1997). Developmental Medicine and Child Neurology, 2008, 50, 5-5.	1.1	22
42	The Oxford Ankle Foot Questionnaire for Children. Prosthetics and Orthotics International, 2010, 34, 238-244.	0.5	20
43	Health outcomes for children with neurodisability: what do professionals regard as primary targets?. Archives of Disease in Childhood, 2014, 99, 927-932.	1.0	20
44	Epilepsyâ€specific patientâ€reported outcome measures of children's healthâ€related quality of life: A systematic review of measurement properties. Epilepsia, 2020, 61, 230-248.	2.6	19
45	A systematic review of ordinal scales used to classify the eating and drinking abilities of individuals with cerebral palsy. Developmental Medicine and Child Neurology, 2014, 56, 313-322.	1.1	18
46	Endâ€user involvement in a systematic review of quantitative and qualitative research of nonâ€pharmacological interventions for attention deficit hyperactivity disorder delivered in school settings: reflections on the impacts and challenges. Health Expectations, 2016, 19, 1084-1097.	1,1	17
47	Ankle foot orthoses for young children with cerebral palsy: a scoping review. Disability and Rehabilitation, 2021, 43, 726-738.	0.9	14
48	Parent-delivered interventions used at home to improve eating, drinking and swallowing in children with neurodisability: the FEEDS mixed-methods study. Health Technology Assessment, 2021, 25, 1-208.	1.3	14
49	Sleep positioning systems for children with cerebral palsy. The Cochrane Library, 2015, 2015, CD009257.	1.5	13
50	Rasch analysis of the Chedoke–McMaster Attitudes towards Children with Handicaps scale. Disability and Rehabilitation, 2017, 39, 281-290.	0.9	13
51	Core Health Outcomes In Childhood Epilepsy (CHOICE): protocol for the selection of a core outcome set. Trials, 2017, 18, 572.	0.7	13
52	A year of foot and ankle orthotic provision for adults: Prospective consultations data, with patient satisfaction survey. Foot, 2008, 18, 75-83.	0.4	12
53	Parent-to-parent support interventions for parents of babies cared for in a neonatal unitâ€"protocol of a systematic review of qualitative and quantitative evidence. Systematic Reviews, 2018, 7, 179.	2.5	12
54	Mapping epilepsy-specific patient-reported outcome measures for children to a proposed core outcome set for childhood epilepsy. Epilepsy and Behavior, 2020, 112, 107372.	0.9	11

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55	Sleep positioning systems for children and adults with a neurodisability: A systematic review. British Journal of Occupational Therapy, 2019, 82, 5-14.	0.5	10
56	Online community engagement in response to COVIDâ€19 pandemic. Health Expectations, 2021, 24, 728-730.	1.1	10
57	Working with Patients and Members of the Public: Informing Health Economics in Child Health Research. PharmacoEconomics - Open, 2019, 3, 133-141.	0.9	9
58	Healthy Parent Carers: feasibility randomised controlled trial of a peer-led group-based health promotion intervention for parent carers of disabled children. Pilot and Feasibility Studies, 2021, 7, 144.	0.5	9
59	Meeting the Information Needs of Families of Children with Chronic Health Conditions. Physical and Occupational Therapy in Pediatrics, 2013, 33, 265-270.	0.8	8
60	The Eating and Drinking Ability Classification System for cerebral palsy: A study of reliability and stability over time. Journal of Pediatric Rehabilitation Medicine, 2019, 12, 123-131.	0.3	8
61	†The relationship of cerebral palsy subtype and functional motor impairment: a populationâ€based study'. Developmental Medicine and Child Neurology, 2010, 52, 682-683.	1.1	7
62	Systematic review: measurement properties of patient-reported outcome measures evaluated with childhood brain tumor survivors or other acquired brain injury. Neuro-Oncology Practice, 2020, 7, 277-287.	1.0	7
63	ISPO Cerebral Palsy Consensus Conference Report (available free at www.ispoweb.org). Prosthetics and Orthotics International, 2009, 33, 401-402.	0.5	6
64	Improving communication between staff and disabled children in hospital wards: testing the feasibility of a training intervention developed through intervention mapping. BMJ Paediatrics Open, 2017, 1, e000103.	0.6	6
65	"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. Journal of Neonatal Nursing, 2019, 25, 166-176.	0.3	6
66	Do the abilities of children with cerebral palsy explain their activities and participation?. Developmental Medicine and Child Neurology, 2006, 48, 954-961.	1.1	5
67	Therapeutic potential and ownership of commercially available consoles in children with cerebral palsy. British Journal of Occupational Therapy, 2017, 80, 108-116.	0.5	5
68	Towards development and validation of an intraoperative assessment tool for robot-assisted radical prostatectomy training: results of a Delphi study. International Braz J Urol: Official Journal of the Brazilian Society of Urology, 2017, 43, 661-670.	0.7	5
69	Implementation of training to improve communication with disabled children on the ward: A feasibility study. Health Expectations, 2021, 24, 1433-1442.	1.1	5
70	Children with neurodisability and feeding difficulties: a UK survey of parent-delivered interventions. BMJ Paediatrics Open, 2021, 5, e001095.	0.6	5
71	Healthy Parent Carers programme: mixed methods process evaluation and refinement of a health promotion intervention. BMJ Open, 2021, 11, e045570.	0.8	5
72	Miniâ€EDACS: Development of the Eating and Drinking Ability Classification System for young children with cerebral palsy. Developmental Medicine and Child Neurology, 2022, 64, 897-906.	1.1	5

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73	A review of the efficacy of lowerâ€limb orthoses used for cerebral palsy. Developmental Medicine and Child Neurology, 2002, 44, 205-211.	1.1	4
74	Development of a core outcome set for evaluative research into paediatric cerebral visual impairment (CVI), in the UK and Eire. BMJ Open, 2021, 11, e051014.	0.8	3
75	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. BMJ Paediatrics Open, 2022, 6, e001425.	0.6	3
76	Current and future uses of the Gross Motor Function Classification System: the need to take account of other factors to explain functional outcomes. Developmental Medicine and Child Neurology, 2009, 51, 1003-1003.	1.1	2
77	Resources for parents raising a disabled child in the UK. Paediatrics and Child Health (United) Tj ETQq1 1 0.7843	14,rgBT /	Overlock 10 Tf
78	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. Pilot and Feasibility Studies, 2019, 5, 137.	0.5	2
79	Community-based family and carer-support programmes for children with disabilities. Paediatrics and Child Health (United Kingdom), 2020, 30, 180-185.	0.2	2
80	Aiming to improve the health care of people with cerebral palsy worldwide: a report of an International Society for Prosthetics and Orthotics conference. Developmental Medicine and Child Neurology, 2009, 51, 689-689.	1.1	1
81	Resources for parents raising a disabled child in the UK. Paediatrics and Child Health (United) Tj ETQq1 1 0.7843	14,rgBT /	Overlock 10 Tf
82	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. BMJ Open, 2021, 11, e049542.	0.8	1
83	Aspects of methodology relevant to patient-reported outcome measures (PROMs)., 0,, 1-9.		1
84	Improving continence in children and young people with neurodisability: a systematic review and survey. Health Technology Assessment, 2021, 25, 1-258.	1.3	1
85	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. Physiotherapy, 2022, 116, 42-49.	0.2	1
86	Editorial. Prosthetics and Orthotics International, 2005, 29, 115-117.	0.5	0
87	Re: The Incidence and Relative Risk of Cardiovascular Toxicity in Patients Treated with New Hormonal Agents for Castration-resistant Prostate Cancer. European Urology, 2016, 69, 177-178.	0.9	O