Kirstie Haywood

List of Publications by Year in descending order

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61 4,507 28 61 papers citations h-index g-index

63 63 63 63 7009

times ranked

citing authors

docs citations

all docs

#	Article	IF	CITATIONS
1	Qualitative interview study exploring the patient experience of living with axial spondyloarthritis and fatigue: difficult, demanding and draining. BMJ Open, 2022, 12, e053958.	0.8	7
2	European Resuscitation Council and European Society of Intensive Care Medicine guidelines 2021: post-resuscitation care. Intensive Care Medicine, 2021, 47, 369-421.	3.9	450
3	European Resuscitation Council and European Society of Intensive Care Medicine Guidelines 2021: Post-resuscitation care. Resuscitation, 2021, 161, 220-269.	1.3	358
4	Measuring health-related quality of life in chronic headache: A comparative evaluation of the Chronic Headache Quality of Life Questionnaire and Headache Impact Test (HIT-6). Cephalalgia, 2021, 41, 1100-1123.	1.8	13
5	SPIRIT-PRO Extension explanation and elaboration: guidelines for inclusion of patient-reported outcomes in protocols of clinical trials. BMJ Open, 2021, 11, e045105.	0.8	65
6	Moving beyond survival and navigating survivorship: Mind the gaps!. Resuscitation, 2021, 167, 395-397.	1.3	2
7	Brain injury after cardiac arrest. Lancet, The, 2021, 398, 1269-1278.	6.3	86
8	Core outcome set for preventive intervention trials in chronic and episodic migraine (COSMIG): an international, consensus-derived and multistakeholder initiative. BMJ Open, 2021, 11, e043242.	0.8	18
9	Patient and public involvement in a UK National Institute for Health Research Programme Grant for Applied Research: experiences from the Chronic Headache Education and Self-management Study (CHESS). Primary Health Care Research and Development, 2021, 22, e72.	0.5	5
10	Trapped in a disrupted normality: Survivors' and partners' experiences of life after a sudden cardiac arrest. Resuscitation, 2020, 147, 81-87.	1.3	29
11	P71â€fWorking collaboratively with patients as research partners in the co-production of the WASTEd: a measure of fatigue and energy in axial spondyloarthritis (axSpA). Rheumatology, 2020, 59, .	0.9	0
12	Use of the nominal group technique to identify UK stakeholder views of the measures and domains used in the assessment of therapeutic exercise adherence for patients with musculoskeletal disorders. BMJ Open, 2020, 10, e031591.	0.8	9
13	Development and feasibility of an intervention featuring individual supported work placements to aid return to work for unemployed people living with chronic pain. Pilot and Feasibility Studies, 2020, 6, 49.	0.5	4
14	Usual care and a self-management support programme versus usual care and a relaxation programme for people living with chronic headache disorders: a randomised controlled trial protocol (CHESS). BMJ Open, 2020, 10, e033520.	0.8	12
15	Survival and quality of life following a cardiac arrest: Capturing what really matters to survivors. Resuscitation, 2019, 135, 221-223.	1.3	3
16	Implementing patient-reported outcome measures in clinical practice: a companion guide to the ISOQOL user's guide. Quality of Life Research, 2019, 28, 621-627.	1.5	67
17	Testing a support programme for opioid reduction for people with chronic non-malignant pain: the I-WOTCH randomised controlled trial protocol. BMJ Open, 2019, 9, e028937.	0.8	6
18	Guidelines for Inclusion of Patient-Reported Outcomes in Clinical Trial Protocols. JAMA - Journal of the American Medical Association, 2018, 319, 483.	3.8	507

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19	Life after cardiac arrest: The importance of engaging with the †forgotten patient'. Resuscitation, 2018, 128, A1-A2.	1.3	22
20	COSCA (Core Outcome Set for Cardiac Arrest) in Adults: An Advisory Statement From the International Liaison Committee on Resuscitation. Resuscitation, 2018, 127, 147-163.	1.3	141
21	COSCA (Core Outcome Set for Cardiac Arrest) in Adults: An Advisory Statement From the International Liaison Committee on Resuscitation. Circulation, 2018, 137, e783-e801.	1.6	171
22	Assessing the impact of headaches and the outcomes of treatment: A systematic review of patient-reported outcome measures (PROMs). Cephalalgia, 2018, 38, 1374-1386.	1.8	71
23	Development and prospective external validation of a tool to predict poor recovery at 9 months after acute ankle sprain in UK emergency departments: the SPRAINED prognostic model. BMJ Open, 2018, 8, e022802.	0.8	5
24	Influence of co-morbid fibromyalgia on disease activity measures and response to tumour necrosis factor inhibitors in axial spondyloarthritis: results from a UK national register. Rheumatology, 2018, 57, 1982-1990.	0.9	26
25	Prognostic models for identifying risk of poor outcome in people with acute ankle sprains: the SPRAINED development and external validation study. Health Technology Assessment, 2018, 22, 1-112.	1.3	124
26	Quality and acceptability of measures of exercise adherence in musculoskeletal settings: a systematic review. Rheumatology, 2017, 56, kew422.	0.9	21
27	Children's experiences of chronic fatigue syndrome/myalgic encephalomyelitis (CFS/ME): a systematic review and meta-ethnography of qualitative studies. BMJ Open, 2017, 7, e012633.	0.8	33
28	Establishing the values for patient engagement (PE) in health-related quality of life (HRQoL) research: an international, multiple-stakeholder perspective. Quality of Life Research, 2017, 26, 1393-1404.	1.5	27
29	Patient-reported outcome measures in older people with hip fracture: a systematic review of quality and acceptability. Quality of Life Research, 2017, 26, 799-812.	1.5	38
30	UNDERSTANDING DISEASE FLARES IN INFLAMMATORY ARTHRITISI96. $\hat{a} \in f$ THE PATIENT EXPERIENCE OF FLARES: WHAT DO PATIENTS MEAN WHEN THEY SAY THEY HAVE A FLARE?. Rheumatology, 2017, 56, .	0.9	0
31	Facet joint injections for people with persistent non-specific low back pain (Facet Injection Study): a feasibility study for a randomised controlled trial. Health Technology Assessment, 2017, 21, 1-184.	1.3	9
32	Evolution of medical ethics in resuscitation and end of life. Trends in Anaesthesia and Critical Care, 2016, 10, 7-14.	0.4	14
33	Using PROMs in Healthcare: Who Should Be in the Driving Seat—Policy Makers, Health Professionals, Methodologists or Patients?. Patient, 2016, 9, 495-498.	1.1	19
34	A survey of key opinion leaders on ethical resuscitation practices in 31 European Countries. Resuscitation, 2016, 100, 11-17.	1.3	40
35	International Society for Quality of Life Research commentary on the draft European Medicines Agency reflection paper on the use of patient-reported outcome (PRO) measures in oncology studies. Quality of Life Research, 2016, 25, 359-362.	1.5	19
36	Facet-joint injections for people with persistent non-specific low back pain (FIS): study protocol for a randomised controlled feasibility trial. Trials, 2015, 16, 588.	0.7	5

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37	Patient and public engagement in health-related quality of life and patient-reported outcomes research: what is important and why should we care? Findings from the first ISOQOL patient engagement symposium. Quality of Life Research, 2015, 24, 1069-1076.	1.5	65
38	Evaluating recovery following hip fracture: a qualitative interview study of what is important to patients. BMJ Open, 2015, 5, e005406-e005406.	0.8	69
39	A systematic review of the outcomes reported in cardiac arrest clinical trials: The need for a core outcome set. Resuscitation, 2015, 88, 150-157.	1.3	83
40	What matters to children with CFS/ME? A conceptual model as the first stage in developing a PROM. Archives of Disease in Childhood, 2015, 100, 1141-1147.	1.0	33
41	Intraarticular Facet Injections for Low Back Pain: Design Considerations, Consensus Methodology to Develop the Protocol for a Randomized Controlled Trial. Pain Physician, 2015, 18, 473-93.	0.3	9
42	Assessing severity of illness and outcomes of treatment in children with <scp>C</scp> hronic <scp>F</scp> atigue <scp>S</scp> yndrome/ <scp>M</scp> yalgic <scp>E</scp> ncephalomyelitis (<scp>CFS</scp> / <scp>ME</scp>): a systematic review of patientâ€reported outcome measures (<scp>PROMs</scp>). Child: Care, Health and Development, 2014, 40, 806-824.	0.8	25
43	Developing a core outcome set for hip fracture trials. Bone and Joint Journal, 2014, 96-B, 1016-1023.	1.9	127
44	Assessing fatigue in ankylosing spondylitis: the importance of frequency and severity. Rheumatology, 2014, 53, 552-556.	0.9	30
45	The psychosocial outcomes of cardiac arrest: Relevant and robust patient-centred assessment is essential. Resuscitation, 2014, 85, 718-719.	1.3	10
46	Patients with well-established ankylosing spondylitis show limited deterioration in a ten-year prospective cohort study. Clinical Rheumatology, 2013, 32, 67-72.	1.0	11
47	Evaluation of Ankylosing Spondylitis Quality of Life questionnaire: responsiveness of a new patient-reported outcome measure. Rheumatology, 2012, 51, 707-714.	0.9	22
48	Quality and acceptability of patient-reported outcome measures used in chronic fatigue syndrome/myalgic encephalomyelitis (CFS/ME): a systematic review. Quality of Life Research, 2012, 21, 35-52.	1.5	90
49	Impact of ankylosing spondylitis on work in patients across the UK. Scandinavian Journal of Rheumatology, 2011, 40, 34-40.	0.6	63
50	Evaluation of Ankylosing Spondylitis Quality of Life (EASi-QoL): Reliability and Validity of a New Patient-reported Outcome Measure. Journal of Rheumatology, 2010, 37, 2100-2109.	1.0	27
51	Continence specialists use of quality of life information in routine practice: a national survey of practitioners. Quality of Life Research, 2009, 18, 423-433.	1.5	12
52	The direct healthcare costs associated with ankylosing spondylitis patients attending a UK secondary care rheumatology unit. Rheumatology, 2008, 47, 68-71.	0.9	44
53	Impact of patient-reported outcome measures on routine practice: a structured review. Journal of Evaluation in Clinical Practice, 2006, 12, 559-568.	0.9	509
54	Patient participation in the consultation process: A structured review of intervention strategies. Patient Education and Counseling, 2006, 63, 12-23.	1.0	203

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55	Quality of life in older people: A structured review of generic self-assessed health instruments. Quality of Life Research, 2005, 14, 1651-1668.	1.5	314
56	Patient-assessed health in ankylosing spondylitis: a structured review. Rheumatology, 2005, 44, 577-586.	0.9	56
57	Spinal mobility in ankylosing spondylitis: reliability, validity and responsiveness. Rheumatology, 2004, 43, 750-757.	0.9	76
58	Reviewing measures of outcome: reliability of data extraction. Journal of Evaluation in Clinical Practice, 2004, 10, 329-337.	0.9	18
59	Multi-item outcome measures for lateral ligament injury of the ankle: a structured review. Journal of Evaluation in Clinical Practice, 2004, 10, 339-352.	0.9	26
60	Generic measures of health-related quality of life in ankylosing spondylitis: reliability, validity and responsiveness. British Journal of Rheumatology, 2002, 41, 1380-1387.	2.5	61
61	Disease-specific, patient-assessed measures of health outcome in ankylosing spondylitis: reliability, validity and responsiveness. British Journal of Rheumatology, 2002, 41, 1295-1302.	2.5	71