

# Carole Mockford

## List of Publications by Year in descending order

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Version: 2024-02-01

28  
papers

3,312  
citations

331259

21  
h-index

525886

27  
g-index

28  
all docs

28  
docs citations

28  
times ranked

4512  
citing authors

#	ARTICLE	IF	CITATIONS
1	Using patient experience data to support improvements in inpatient mental health care: the EURIPIDES multimethod study. <i>Health Services and Delivery Research</i> , 2020, 8, 1-338.	1.4	6
2	Experiences of in-patient mental health services: systematic review. <i>British Journal of Psychiatry</i> , 2019, 214, 329-338.	1.7	79
3	Facilitating Implementation of Research Evidence (FIRE): an international cluster randomised controlled trial to evaluate two models of facilitation informed by the Promoting Action on Research Implementation in Health Services (PARiHS) framework. <i>Implementation Science</i> , 2018, 13, 137.	2.5	50
4	A realist process evaluation within the Facilitating Implementation of Research Evidence (FIRE) cluster randomised controlled international trial: an exemplar. <i>Implementation Science</i> , 2018, 13, 138.	2.5	66
5	The development of service user-led recommendations for health and social care services on leaving hospital with memory loss or dementia - the SHARED study. <i>Health Expectations</i> , 2017, 20, 495-507.	1.1	8
6	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
7	Recommendations to support informal carers of people living with motor neurone disease. <i>British Journal of Community Nursing</i> , 2016, 21, 518-524.	0.2	17
8	A SHARED study-the benefits and costs of setting up a health research study involving lay co-researchers and how we overcame the challenges. <i>Research Involvement and Engagement</i> , 2016, 2, 8.	1.1	33
9	A Bridge Over Turbulent Waters: Illustrating the Interaction Between Managerial Leaders and Facilitators When Implementing Research Evidence. <i>Worldviews on Evidence-Based Nursing</i> , 2016, 13, 25-31.	1.2	56
10	Do-not-attempt-cardiopulmonary-resuscitation decisions: an evidence synthesis. <i>Health Services and Delivery Research</i> , 2016, 4, 1-154.	1.4	29
11	A review of family carers&#39; experiences of hospital discharge for people with dementia, and the rationale for involving service users in health research. <i>Journal of Healthcare Leadership</i> , 2015, 7, 21.	1.5	14
12	Do not attempt cardiopulmonary resuscitation (DNACPR) orders: A systematic review of the barriers and facilitators of decision-making and implementation. <i>Resuscitation</i> , 2015, 88, 99-113.	1.3	99
13	Mapping the impact of patient and public involvement on health and social care research: a systematic review. <i>Health Expectations</i> , 2014, 17, 637-650.	1.1	986
14	A Systematic Review of the Impact of Patient and Public Involvement on Service Users, Researchers and Communities. <i>Patient</i> , 2014, 7, 387-395.	1.1	471
15	A systematic review of do-not-attempt-cardiopulmonary-resuscitation (DNACPR) orders: Summarising the evidence around decision making and implementation. <i>Resuscitation</i> , 2014, 85, S85.	1.3	1
16	The impact of patient and public involvement on UK NHS health care: a systematic review. <i>International Journal for Quality in Health Care</i> , 2012, 24, 28-38.	0.9	407
17	FIRE (facilitating implementation of research evidence): a study protocol. <i>Implementation Science</i> , 2012, 7, 25.	2.5	68
18	Developing the evidence base of patient and public involvement in health and social care research: the case for measuring impact. <i>International Journal of Consumer Studies</i> , 2011, 35, 628-632.	7.2	118

#	ARTICLE	IF	CITATIONS
19	The GRIPP checklist: Strengthening the quality of patient and public involvement reporting in research. <i>International Journal of Technology Assessment in Health Care</i> , 2011, 27, 391-399.	0.2	253
20	Moving forward: understanding the negative experiences and impacts of patient and public involvement in health service planning, development and evaluation. , 2011, , 129-141.		3
21	Development of the Motor Neuron Disease Carer Questionnaire. <i>Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders</i> , 2009, 10, 463-469.	2.3	2
22	Measuring the impact of patient and public involvement: the need for an evidence base. <i>International Journal for Quality in Health Care</i> , 2008, 20, 373-374.	0.9	83
23	Role of home visiting in improving parenting and health in families at risk of abuse and neglect: results of a multicentre randomised controlled trial and economic evaluation. <i>Archives of Disease in Childhood</i> , 2007, 92, 229-233.	1.0	127
24	A Review: Carers, MND and service provision. <i>Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders</i> , 2006, 7, 132-141.	2.3	69
25	Parenting programmes: some unintended consequences. <i>Primary Health Care Research and Development</i> , 2004, 5, 219-227.	0.5	34
26	Impact of a general practice based group parenting programme: quantitative and qualitative results from a controlled trial at 12 months. <i>Archives of Disease in Childhood</i> , 2004, 89, 519-525.	1.0	69
27	Need and demand for parenting programmes in general practice. <i>Archives of Disease in Childhood</i> , 2002, 87, 468-471.	1.0	30
28	Improving mental health through parenting programmes: block randomised controlled trial. <i>Archives of Disease in Childhood</i> , 2002, 87, 472-477.	1.0	99