

# Carole Mockford

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/4449179/publications.pdf>

Version: 2024-02-01

28  
papers

3,312  
citations

331538

21  
h-index

526166

27  
g-index

28  
all docs

28  
docs citations

28  
times ranked

4512  
citing authors

#	ARTICLE	IF	CITATIONS
1	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
2	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
3	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
4	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
5	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
6	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
7	Improving mental health through parenting programmes: block randomised controlled trial. <i>Archives of Disease in Childhood</i> , 2002, 87, 472-477.	1.0	99
8	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
9	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
10	Experiences of in-patient mental health services: systematic review. <i>British Journal of Psychiatry</i> , 2019, 214, 329-338.	1.7	79
11	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
12	A Review: Carers, MND and service provision. <i>Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders</i> , 2006, 7, 132-141.	2.3	69
13	FIRE (facilitating implementation of research evidence): a study protocol. <i>Implementation Science</i> , 2012, 7, 25.	2.5	68
14	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
15	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
16	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
17	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
18	Parenting programmes: some unintended consequences. <i>Primary Health Care Research and Development</i> , 2004, 5, 219-227.	0.5	34

#	ARTICLE	IF	CITATIONS
19	A SHARED study-the benefits and costs of setting up a health research study involving lay co-researchers and how we overcame the challenges. Research Involvement and Engagement, 2016, 2, 8.	1.1	33
20	Need and demand for parenting programmes in general practice. Archives of Disease in Childhood, 2002, 87, 468-471.	1.0	30
21	Do-not-attempt-cardiopulmonary-resuscitation decisions: an evidence synthesis. Health Services and Delivery Research, 2016, 4, 1-154.	1.4	29
22	Recommendations to support informal carers of people living with motor neurone disease. British Journal of Community Nursing, 2016, 21, 518-524.	0.2	17
23	A review of family carers&#39; experiences of hospital discharge for people with dementia, and the rationale for involving service users in health research. Journal of Healthcare Leadership, 2015, 7, 21.	1.5	14
24	The development of service user-led recommendations for health and social care services on leaving hospital with memory loss or dementia - the SHARED study. Health Expectations, 2017, 20, 495-507.	1.1	8
25	Using patient experience data to support improvements in inpatient mental health care: the EURIPIDES multimethod study. Health Services and Delivery Research, 2020, 8, 1-338.	1.4	6
26	Moving forward: understanding the negative experiences and impacts of patient and public involvement in health service planning, development and evaluation. , 2011, , 129-141.		3
27	Development of the Motor Neuron Disease Carer Questionnaire. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2009, 10, 463-469.	2.3	2
28	A systematic review of do-not-attempt-cardiopulmonary-resuscitation (DNACPR) orders: Summarising the evidence around decision making and implementation. Resuscitation, 2014, 85, S85.	1.3	1