Joanna Murray

List of Publications by Year in descending order

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43 papers

1,839 citations

304743

22

h-index

265206 42 g-index

45 all docs

45 docs citations

45 times ranked

2586 citing authors

#	Article	IF	CITATIONS
1	Psychological and demographic characteristics of 368 patients with dissociative seizures: data from the CODES cohort. Psychological Medicine, 2021, 51, 2433-2445.	4.5	24
2	Cognitive–behavioural therapy compared with standardised medical care for adults with dissociative non-epileptic seizures: the CODES RCT. Health Technology Assessment, 2021, 25, 1-144.	2.8	8
3	"We should see her like part of the team― an investigation into care home staff's experiences of being part of an RCT of a complex psychosocial intervention. Aging and Mental Health, 2020, 24, 178-185.	2.8	12
4	The experience of trial participation, treatment approaches and perceptions of change among participants with dissociative seizures within the CODES randomized controlled trial: A qualitative study. Epilepsy and Behavior, 2020, 111, 107230.	1.7	7
5	The experiences of therapists providing cognitive behavioral therapy (CBT) for dissociative seizures in the CODES randomized controlled trial: A qualitative study. Epilepsy and Behavior, 2020, 105, 106943.	1.7	4
6	Cognitive behavioural therapy for adults with dissociative seizures (CODES): a pragmatic, multicentre, randomised controlled trial. Lancet Psychiatry,the, 2020, 7, 491-505.	7.4	175
7	Improving mental health and reducing antipsychotic use in people with dementia in care homes: the WHELD research programme including two RCTs. Programme Grants for Applied Research, 2020, 8, 1-98.	1.0	15
8	Characteristics of 698 patients with dissociative seizures: A <scp>UK</scp> multicenter study. Epilepsia, 2019, 60, 2182-2193.	5.1	51
9	Exploring psychiatrists' perspectives of working with patients with dissociative seizures in the UK healthcare system as part of the CODES trial: a qualitative study. BMJ Open, 2019, 9, e026493.	1.9	4
10	Measuring the quality of life of family carers of people with dementia: development and validation of C-DEMQOL. Quality of Life Research, 2019, 28, 2299-2310.	3.1	24
11	Understanding the quality of life of family carers of people with dementia: Development of a new conceptual framework. International Journal of Geriatric Psychiatry, 2019, 34, 79-86.	2.7	31
12	Visual hallucinations in dementia and Parkinson's disease: A qualitative exploration of patient and caregiver experiences. International Journal of Geriatric Psychiatry, 2018, 33, 1327-1334.	2.7	17
13	Factors associated with the quality of life of family carers of people with dementia: A systematic review. Alzheimer's and Dementia, 2017, 13, 572-581.	0.8	228
14	Experience of stigma in the public life of relatives of people diagnosed with schizophrenia in the Republic of Belarus. Social Psychiatry and Psychiatric Epidemiology, 2017, 52, 493-501.	3.1	10
15	Instruments measuring the disease-specific quality of life of family carers of people with neurodegenerative diseases: a systematic review. BMJ Open, 2017, 7, e013611.	1.9	28
16	Epidemiology of Pain in People With Dementia Living in Care Homes: Longitudinal Course, Prevalence, and Treatment Implications. Journal of the American Medical Directors Association, 2017, 18, 453.e1-453.e6.	2.5	70
17	Systematic review of the effective components of psychosocial interventions delivered by care home staff to people with dementia. BMJ Open, 2017, 7, e014177.	1.9	52
18	COgnitive behavioural therapy versus standardised medical care for adults with Dissociative non-Epileptic Seizures (CODES): statistical and economic analysis plan for a randomised controlled trial. Trials, 2017, 18, 258.	1.6	13

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19	Helping staff to implement psychosocial interventions in care homes: augmenting existing practices and meeting needs for support. International Journal of Geriatric Psychiatry, 2016, 31, 284-293.	2.7	30
20	UPBEAT-UK: a programme of research into the relationship between coronary heart disease and depression in primary care patients. Programme Grants for Applied Research, 2016, 4, 1-172.	1.0	4
21	Comparative Analysis of Informal Caregiver Burden in Advanced Cancer, Dementia, and Acquired Brain Injury. Journal of Pain and Symptom Management, 2015, 50, 445-452.	1.2	79
22	Improving the diagnosis of physical illness in patients with mental illness who present in Emergency Departments: Consensus study. Journal of Psychosomatic Research, 2015, 78, 346-351.	2.6	26
23	COgnitive behavioural therapy vs standardised medical care for adults with Dissociative non-Epileptic Seizures (CODES): a multicentre randomised controlled trial protocol. BMC Neurology, 2015, 15, 98.	1.8	77
24	User-generated quality standards for youth mental health in primary care: a participatory research design using mixed methods. BMJ Quality and Safety, 2014, 23, 857-866.	3.7	24
25	Improving Well-being and Health for People with Dementia (WHELD): study protocol for a randomised controlled trial. Trials, 2014, 15, 284.	1.6	24
26	Diagnostic Overshadowing and Other Challenges Involved in the Diagnostic Process of Patients with Mental Illness Who Present in Emergency Departments with Physical Symptoms $\hat{a} \in A$ Qualitative Study. PLoS ONE, 2014, 9, e111682.	2.5	173
27	Feasibility study of an optimised person-centred intervention to improve mental health and reduce antipsychotics amongst people with dementia in care homes: study protocol for a randomised controlled trial. Trials, 2013, 14, 13.	1.6	15
28	Cost-effectiveness analyses for mirtazapine and sertraline in dementia: randomised controlled trial. British Journal of Psychiatry, 2013, 202, 121-128.	2.8	43
29	Improving quality of life for people with dementia in care homes: making psychosocial interventions work. British Journal of Psychiatry, 2012, 201, 344-351.	2.8	147
30	People with dementia and their family carers' satisfaction with a memory service: A qualitative evaluation generating quality indicators for dementia care. Journal of Mental Health, 2009, 18, 26-37.	1.9	20
31	When to Delete Recorded Qualitative Research Data. Research Ethics, 2008, 4, 76-77.	1.7	2
32	Primary care professionals' perceptions of depression in older people: a qualitative study. Social Science and Medicine, 2006, 63, 1363-1373.	3.8	89
33	Personal Control in Social and Life-Course Contexts , Editors: S <scp>TEVEN</scp> H. Z <scp>ARIT</scp> , L <scp>EONARD</scp> I. P <scp>EARLIN</scp> <scp>AND</scp> W <scp>ARNER</scp> S <scp>CHAIE</scp> , Springer, New York, 2003, U.S. \$49.95. Hardback, pp. 328, ISBN 0 8261 2402 X. International Psychogeriatrics, 2005, 17, 713-714.	1.0	0
34	Prospects for mental health social work: A qualitative study of attitudes of service managers and academic staff. Journal of Mental Health, 2004, 13, 305-317.	1.9	16
35	Formal and informal care for people with dementia: variations in costs over time. Ageing and Society, 2003, 23, 303-326.	1.7	30
36	New approach to translating instruments for cross-cultural research: a combined qualitative and quantitative approach for translation and consensus generation. International Journal of Methods in Psychiatric Research, 2000, 9, 87-95.	2.1	65

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37	Qualitative methods. International Review of Psychiatry, 1998, 10, 312-316.	2.8	12
38	A qualitative study of adjustment to caring for an older spouse with psychiatric illness. Ageing and Society, 1998, 18, 659-671.	1.7	22
39	Prevention and the identification of high risk groups. International Review of Psychiatry, 1992, 4, 281-286.	2.8	14
40	The use of health diaries in the field of psychiatric illness in general practice. Psychological Medicine, 1985, 15, 827-840.	4.5	13
41	Self-assessment of health: an exploration of the effects of physical and psychological symptoms. Psychological Medicine, 1982, 12, 371-378.	4.5	53
42	Minor psychiatric morbidity and the threat of redundancy in a professional group. Psychological Medicine, 1982, 12, 799-807.	4.5	63
43	Long-term psychotropic drug-taking and the process of withdrawal. Psychological Medicine, 1981, 11, 853-858.	4.5	18