

Michele Peters

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/2431039/publications.pdf>

Version: 2024-02-01

38
papers

1,123
citations

471061

17
h-index

414034

32
g-index

41
all docs

41
docs citations

41
times ranked

1540
citing authors

#	ARTICLE	IF	CITATIONS
1	Carers using assistive technology in dementia care at home: a mixed methods study. <i>BMC Geriatrics</i> , 2022, 22, .	1.1	2
2	Use of the Long-Term Conditions Questionnaire (LTCQ) for monitoring health-related quality of life in people affected by cognitive impairment including dementia: pilot study in UK memory clinic services. <i>Quality of Life Research</i> , 2021, 30, 1641-1652.	1.5	2
3	Using rapid cycle tests of change to develop the Carers Assistive Technology Experience Questionnaire: a cognitive interview study in the UK. <i>BMJ Open</i> , 2021, 11, e042361.	0.8	10
4	Impact of COVID-19 restrictions on carers of persons with dementia in the UK: a qualitative study. <i>Age and Ageing</i> , 2021, 50, 1876-1885.	0.7	19
5	Carers'™ experiences of assistive technology use in dementia care: a cross sectional survey. <i>BMC Geriatrics</i> , 2021, 21, 471.	1.1	7
6	Enhancing primary care support for informal carers: A scoping study with professional stakeholders. <i>Health and Social Care in the Community</i> , 2020, 28, 642-650.	0.7	18
7	Withdrawing gluten-free food from prescriptions in England: a mixed-methods study to examine the impact of policy changes on quality of life. <i>Journal of Human Nutrition and Dietetics</i> , 2020, 33, 453-464.	1.3	7
8	Awareness of Appropriate Antibiotic Use in Primary Care for Influenza-Like Illness: Evidence of Improvement from UK Population-Based Surveys. <i>Antibiotics</i> , 2020, 9, 690.	1.5	3
9	Rasch analysis of the long-term conditions questionnaire (LTCQ) and development of a short-form (LTCQ-8). <i>Health and Quality of Life Outcomes</i> , 2020, 18, 375.	1.0	7
10	Healthcare experiences and quality of life of adults with coeliac disease: a cross-sectional study. <i>Journal of Human Nutrition and Dietetics</i> , 2020, 33, 741-751.	1.3	5
11	Carers'™ experience of using assistive technology for dementia care at home: a qualitative study. <i>BMJ Open</i> , 2020, 10, e034460.	0.8	16
12	Reducing expectations for antibiotics in primary care: a randomised experiment to test the response to fear-based messages about antimicrobial resistance. <i>BMC Medicine</i> , 2020, 18, 110.	2.3	24
13	How can patient experience scores be used to predict quality inspection ratings? A retrospective cross-sectional study of national primary care datasets in the UK. <i>BMJ Open</i> , 2020, 10, e041709.	0.8	0
14	How can patient experience scores be used to predict quality inspection ratings? A retrospective cross-sectional study of national primary care datasets in the UK. <i>BMJ Open</i> , 2020, 10, e041709.	0.8	1
15	Informal carers'™ experience of assistive technology use in dementia care at home: a systematic review. <i>BMC Geriatrics</i> , 2019, 19, 160.	1.1	93
16	Aids to management of headache disorders in primary care (2nd edition). <i>Journal of Headache and Pain</i> , 2019, 20, 57.	2.5	195
17	Self-efficacy and health-related quality of life: a cross-sectional study of primary care patients with multi-morbidity. <i>Health and Quality of Life Outcomes</i> , 2019, 17, 37.	1.0	119
18	The context of coping: a qualitative exploration of underlying inequalities that influence health services support for people living with long-term conditions. <i>Sociology of Health and Illness</i> , 2018, 40, 130-145.	1.1	16

#	ARTICLE	IF	CITATIONS
19	Quality of life in coeliac disease: qualitative interviews to develop candidate items for the Coeliac Disease Assessment Questionnaire. <i>Patient Related Outcome Measures</i> , 2018, Volume 9, 211-220.	0.7	9
20	Quality of life in coeliac disease: item reduction, scale development and psychometric evaluation of the Coeliac Disease Assessment Questionnaire (<scp>CDAQ</scp>). <i>Alimentary Pharmacology and Therapeutics</i> , 2018, 48, 852-862.	1.9	21
21	Reducing demand for antibiotic prescriptions: evidence from an online survey of the general public on the interaction between preferences, beliefs and information, United Kingdom, 2015. <i>Eurosurveillance</i> , 2018, 23, .	3.9	33
22	The role of life context and self-defined well-being in the outcomes that matter to people with a diagnosis of schizophrenia. <i>Health Expectations</i> , 2017, 20, 1061-1072.	1.1	17
23	Long-Term Conditions Questionnaire (LTCQ): initial validation survey among primary care patients and social care recipients in England. <i>BMJ Open</i> , 2017, 7, e019235.	0.8	15
24	Treatment outcomes in schizophrenia: qualitative study of the views of family carers. <i>BMC Psychiatry</i> , 2017, 17, 266.	1.1	13
25	The Long-Term Conditions Questionnaire: conceptual framework and item development. <i>Patient Related Outcome Measures</i> , 2016, Volume 7, 109-125.	0.7	14
26	Refinement of the Long-Term Conditions Questionnaire (LTCQ): patient and expert stakeholder opinion. <i>Patient Related Outcome Measures</i> , 2016, Volume 7, 183-193.	0.7	8
27	Psychometric properties of carer-reported outcome measures in palliative care: A systematic review. <i>Palliative Medicine</i> , 2016, 30, 23-44.	1.3	30
28	Perspectives from health, social care and policy stakeholders on the value of a single self-report outcome measure across long-term conditions: a qualitative study. <i>BMJ Open</i> , 2015, 5, e006986-e006986.	0.8	17
29	The routine collection of patient-reported outcome measures (PROMs) for long-term conditions in primary care: a cohort survey. <i>BMJ Open</i> , 2014, 4, e003968.	0.8	72
30	Carer quality of life and experiences of health services: a cross-sectional survey across three neurological conditions. <i>Health and Quality of Life Outcomes</i> , 2013, 11, 103.	1.0	38
31	Patients' experiences of health and social care in long-term neurological conditions in England: A cross-sectional survey. <i>Journal of Health Services Research and Policy</i> , 2013, 18, 28-33.	0.8	22
32	The impact of perceived lack of support provided by health and social care services to caregivers of people with motor neuron disease. <i>Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders</i> , 2012, 13, 223-228.	2.3	19
33	Quality in the provision of headache care. 2: defining quality and its indicators. <i>Journal of Headache and Pain</i> , 2012, 13, 449-457.	2.5	22
34	Quality in the provision of headache care. 1: systematic review of the literature and commentary. <i>Journal of Headache and Pain</i> , 2012, 13, 437-447.	2.5	19
35	The development and validation of a quality of life measure for the carers of people with Parkinson's disease (the PDQ-Carer). <i>Parkinsonism and Related Disorders</i> , 2012, 18, 483-487.	1.1	55
36	Factors Influencing Quality of Life in Caregivers of People with Parkinson's Disease and Implications for Clinical Guidelines. <i>Parkinson's Disease</i> , 2012, 2012, 1-6.	0.6	37

#	ARTICLE	IF	CITATIONS
37	Does self-reported well-being of patients with Parkinson's disease influence caregiver strain and quality of life?. <i>Parkinsonism and Related Disorders</i> , 2011, 17, 348-352.	1.1	59
38	Translating Instruments for Cross-Cultural Studies in Headache Research. <i>Headache</i> , 2006, 46, 82-91.	1.8	58