

# 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

471509  
17  
h-index

414414  
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. BMC Geriatrics, 2022, 22, .	2.7	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. Quality of Life Research, 2021, 30, 1641-1652.	3.1	2
3	Using rapid cycle tests of change to develop the Carers Assistive Technology Experience Questionnaire: a cognitive interview study in the UK. BMJ Open, 2021, 11, e042361.	1.9	10
4	Impact of COVID-19 restrictions on carers of persons with dementia in the UK: a qualitative study. Age and Ageing, 2021, 50, 1876-1885.	1.6	19
5	Carers' experiences of assistive technology use in dementia care: a cross sectional survey. BMC Geriatrics, 2021, 21, 471.	2.7	7
6	Enhancing primary care support for informal carers: A scoping study with professional stakeholders. Health and Social Care in the Community, 2020, 28, 642-650.	1.6	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. Journal of Human Nutrition and Dietetics, 2020, 33, 453-464.	2.5	7
8	Awareness of Appropriate Antibiotic Use in Primary Care for Influenza-Like Illness: Evidence of Improvement from UK Population-Based Surveys. Antibiotics, 2020, 9, 690.	3.7	3
9	Rasch analysis of the long-term conditions questionnaire (LTCQ) and development of a short-form (LTCQ-8). Health and Quality of Life Outcomes, 2020, 18, 375.	2.4	7
10	Healthcare experiences and quality of life of adults with coeliac disease: a cross-sectional study. Journal of Human Nutrition and Dietetics, 2020, 33, 741-751.	2.5	5
11	Carers' experience of using assistive technology for dementia care at home: a qualitative study. BMJ Open, 2020, 10, e034460.	1.9	16
12	Reducing expectations for antibiotics in primary care: a randomised experiment to test the response to fear-based messages about antimicrobial resistance. BMC Medicine, 2020, 18, 110.	5.5	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. BMJ Open, 2020, 10, e041709.	1.9	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. BMJ Open, 2020, 10, e041709.	1.9	1
15	Informal carers' experience of assistive technology use in dementia care at home: a systematic review. BMC Geriatrics, 2019, 19, 160.	2.7	93
16	Aids to management of headache disorders in primary care (2nd edition). Journal of Headache and Pain, 2019, 20, 57.	6.0	195
17	Self-efficacy and health-related quality of life: a cross-sectional study of primary care patients with multi-morbidity. Health and Quality of Life Outcomes, 2019, 17, 37.	2.4	119
18	The context of coping: a qualitative exploration of underlying inequalities that influence health services support for people living with long-term conditions. Sociology of Health and Illness, 2018, 40, 130-145.	2.1	16

#	ARTICLE	IF	CITATIONS
19	Quality of life in coeliac disease: qualitative interviews to develop candidate items for the Coeliac Disease Assessment Questionnaire. Patient Related Outcome Measures, 2018, Volume 9, 211-220.	1.2	9
20	Quality of life in coeliac disease: item reduction, scale development and psychometric evaluation of the Coeliac Disease Assessment Questionnaire (<scp>CDAQ</scp>). Alimentary Pharmacology and Therapeutics, 2018, 48, 852-862.	3.7	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. Eurosurveillance, 2018, 23, .	7.0	33
22	The role of life context and self-defined well-being in the outcomes that matter to people with a diagnosis of schizophrenia. Health Expectations, 2017, 20, 1061-1072.	2.6	17
23	Long-Term Conditions Questionnaire (LTCQ): initial validation survey among primary care patients and social care recipients in England. BMJ Open, 2017, 7, e019235.	1.9	15
24	Treatment outcomes in schizophrenia: qualitative study of the views of family carers. BMC Psychiatry, 2017, 17, 266.	2.6	13
25	The Long-Term Conditions Questionnaire: conceptual framework and item development. Patient Related Outcome Measures, 2016, Volume 7, 109-125.	1.2	14
26	Refinement of the Long-Term Conditions Questionnaire (LTCQ): patient and expert stakeholder opinion. Patient Related Outcome Measures, 2016, Volume 7, 183-193.	1.2	8
27	Psychometric properties of carer-reported outcome measures in palliative care: A systematic review. Palliative Medicine, 2016, 30, 23-44.	3.1	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. BMJ Open, 2015, 5, e006986-e006986.	1.9	17
29	The routine collection of patient-reported outcome measures (PROMs) for long-term conditions in primary care: a cohort survey. BMJ Open, 2014, 4, e003968.	1.9	72
30	Carer quality of life and experiences of health services: a cross-sectional survey across three neurological conditions. Health and Quality of Life Outcomes, 2013, 11, 103.	2.4	38
31	Patients' experiences of health and social care in long-term neurological conditions in England: A cross-sectional survey. Journal of Health Services Research and Policy, 2013, 18, 28-33.	1.7	22
32	The impact of perceived lack of support provided by health and social care services to caregivers of people with motor neuron disease. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2012, 13, 223-228.	2.1	19
33	Quality in the provision of headache care. 2: defining quality and its indicators. Journal of Headache and Pain, 2012, 13, 449-457.	6.0	22
34	Quality in the provision of headache care. 1: systematic review of the literature and commentary. Journal of Headache and Pain, 2012, 13, 437-447.	6.0	19
35	The development and validation of a quality of life measure for the carers of people with Parkinson's disease (the PDQ-Carer). Parkinsonism and Related Disorders, 2012, 18, 483-487.	2.2	55
36	Factors Influencing Quality of Life in Caregivers of People with Parkinson's Disease and Implications for Clinical Guidelines. Parkinson's Disease, 2012, 2012, 1-6.	1.1	37

#	ARTICLE	IF	CITATIONS
37	Does self-reported well-being of patients with Parkinson's disease influence caregiver strain and quality of life?. Parkinsonism and Related Disorders, 2011, 17, 348-352.	2.2	59
38	Translating Instruments for Cross-Cultural Studies in Headache Research. Headache, 2006, 46, 82-91.	3.9	58