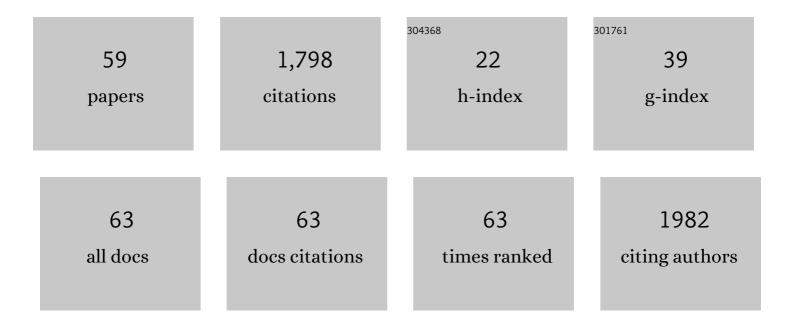
Ian Rees Jones

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/3201112/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	â€`We're happy as we are': the experience of living with possible undiagnosed dementia. Ageing and Society, 2023, 43, 2041-2066.	1.2	4
2	Psychological processes in adapting to dementia: Illness representations among the IDEAL cohort Psychology and Aging, 2022, 37, 524-541.	1.4	5
3	Longitudinal Trajectories of Quality of Life Among People With Mild-to-Moderate Dementia: A Latent Growth Model Approach With IDEAL Cohort Study Data. Journals of Gerontology - Series B Psychological Sciences and Social Sciences, 2022, 77, 1037-1050.	2.4	9
4	The Use and Costs of Paid and Unpaid Care for People with Dementia: Longitudinal Findings from the IDEAL Cohort. Journal of Alzheimer's Disease, 2022, 86, 135-153.	1.2	10
5	Characteristics of people living with undiagnosed dementia: findings from the CFAS Wales study. BMC Geriatrics, 2022, 22, 409.	1.1	7
6	Explaining health system responses to public reporting of cardiac surgery mortality in England and the USA. Health Economics, Policy and Law, 2021, 16, 183-200.	1.1	1
7	Promotion of Healthy Aging Within a Community Center Through Behavior Change: Health and Fitness Findings From the AgeWell Pilot Randomized Controlled Trial. Journal of Aging and Physical Activity, 2021, 29, 80-88.	0.5	1
8	Perceived and objective availability of green and blue spaces and quality of life in people with dementia: results from the IDEAL programme. Social Psychiatry and Psychiatric Epidemiology, 2021, 56, 1601-1610.	1.6	8
9	The impact of relationship quality on life satisfaction and well-being in dementia caregiving dyads: findings from the IDEAL study. Aging and Mental Health, 2020, 24, 1411-1420.	1.5	35
10	â€~All the world's a stage': Accounting for the dementia experience – insights from the IDEAL study. Qualitative Research, 2020, 20, 703-720.	2.2	3
11	Factors associated with self- and informant ratings of quality of life, well-being and life satisfaction in people with mild-to-moderate dementia: results from the Improving the experience of Dementia and Enhancing Active Life programme. Age and Ageing, 2020, 49, 446-452.	0.7	20
12	Professional autonomy and surveillance: the case of public reporting in cardiac surgery. Sociology of Health and Illness, 2019, 41, 1040-1055.	1.1	9
13	Caregivers' beliefs about dementia: findings from the IDEAL study. Psychology and Health, 2019, 34, 1214-1230.	1.2	10
14	Use and costs of services and unpaid care for people with mildâ€toâ€moderate dementia: Baseline results from the IDEAL cohort study. Alzheimer's and Dementia: Translational Research and Clinical Interventions, 2019, 5, 685-696.	1.8	18
15	A Comprehensive Model of Factors Associated With Capability to "Live Well―for Family Caregivers of People Living With Mild-to-Moderate Dementia. Alzheimer Disease and Associated Disorders, 2019, 33, 29-35.	0.6	35
16	A Comprehensive Model of Factors Associated With Subjective Perceptions of "Living Well―With Dementia. Alzheimer Disease and Associated Disorders, 2019, 33, 36-41.	0.6	50
17	Goal-setting to Promote a Healthier Lifestyle in Later Life: Qualitative Evaluation of the AgeWell Trial. Clinical Gerontologist, 2018, 41, 335-345.	1.2	16
18	Elite city-deals for economic growth? Problematizing the complexities of devolution, city-region building, and the (re)positioning of civil society. Space and Polity, 2018, 22, 307-327.	0.8	23

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19	Protocol for the IDEAL-2 longitudinal study: following the experiences of people with dementia and their primary carers to understand what contributes to living well with dementia and enhances active life. BMC Public Health, 2018, 18, 1214.	1.2	35
20	Inequalities in living well with dementia—The impact of deprivation on wellâ€being, quality of life and life satisfaction: Results from the improving the experience of dementia and enhancing active life study. International Journal of Geriatric Psychiatry, 2018, 33, 1736-1742.	1.3	24
21	Dualities of dementia illness narratives and their role in a narrative economy. Sociology of Health and Illness, 2018, 40, 874-891.	1.1	22
22	Living well with dementia: a systematic review and correlational meta-analysis of factors associated with quality of life, well-being and life satisfaction in people with dementia. Psychological Medicine, 2018, 48, 2130-2139.	2.7	181
23	Illness representations in caregivers of people with dementia. Aging and Mental Health, 2017, 21, 553-561.	1.5	15
24	Editorial: Bourdieu, capitals and health. Sociology of Health and Illness, 2017, 39, 3-4.	1.1	6
25	Social class, dementia and the fourth age. Sociology of Health and Illness, 2017, 39, 303-317.	1.1	16
26	Making sense: further studies of living with chronic illness. Sociology of Health and Illness, 2017, 39, 653-658.	1.1	3
27	Connected growth: Developing a framework to drive inclusive growth across a city-region. Local Economy, 2017, 32, 565-575.	0.8	11
28	Editorial: Childbirth and Reproduction. Sociology of Health and Illness, 2016, 38, 687-688.	1.1	0
29	"l Don't Think Of It As An Illnessâ€i Illness Representations in Mild to Moderate Dementia. Journal of Alzheimer's Disease, 2016, 51, 139-150.	1.2	35
30	Regulation, governance and agglomeration: making links in city-region research. Regional Studies, Regional Science, 2016, 3, 509-530.	0.7	22
31	Defining continuity of care from the perspectives of mental health service users and professionals: an exploratory, comparative study. Health Expectations, 2016, 19, 973-987.	1.1	28
32	The Agewell trial: a pilot randomised controlled trial of a behaviour change intervention to promote healthy ageing and reduce risk of dementia in later life. BMC Psychiatry, 2015, 15, 25.	1.1	43
33	Improving the experience of dementia and enhancing active life - living well with dementia: study protocol for the IDEAL study. Health and Quality of Life Outcomes, 2014, 12, 164.	1.0	97
34	Continuity of care for carers of people with severe mental illness: Results of a longitudinal study. International Journal of Social Psychiatry, 2013, 59, 663-670.	1.6	5
35	Continuity of care for people with psychotic illness: Its relationship to clinical and social functioning. International Journal of Social Psychiatry, 2013, 59, 5-17.	1.6	31
36	Choice, consumerism and devolution: growing old in the welfare state(s) of Scotland, Wales and England. Ageing and Society, 2012, 32, 725-746.	1.2	34

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37	Understanding service user-defined continuity of care and its relationship to health and social measures: a cross-sectional study. BMC Health Services Research, 2012, 12, 145.	0.9	39
38	The AgeWell study of behavior change to promote health and wellbeing in later life: study protocol for a randomized controlled trial. Trials, 2012, 13, 115.	0.7	33
39	Towards a Sociology of Disclosure: The Case of Surgical Performance. Sociology Compass, 2012, 6, 908-922.	1.4	8
40	Class and lifestyle †lockâ€in' among middleâ€aged and older men: a Multiple Correspondence Analysis of the British Regional Heart Study. Sociology of Health and Illness, 2011, 33, 399-419.	1.1	23
41	Achieving Continuity of Care: Facilitators and Barriers in Community Mental Health Teams. Implementation Science, 2011, 6, 23.	2.5	73
42	The natural, the normal and the normative: Contested terrains in ageing and old age. Social Science and Medicine, 2010, 71, 1513-1519.	1.8	82
43	Review: Taking care of yourself in later life: A qualitative study into the use of non-prescription medicines by people aged 60+. Health (United Kingdom), 2010, 14, 213-231.	0.9	8
44	The Experience of Retirement in Second Modernity. Sociology, 2010, 44, 103-120.	1.7	30
45	Disclosing clinical performance: the case of cardiac surgery. Journal of Health Organization and Management, 2010, 24, 571-583.	0.6	11
46	Not just old and sick – the â€~will to health' in later life. Ageing and Society, 2009, 29, 687-707.	1.2	126
47	Illness careers and continuity of care in mental health services: A qualitative study of service users and carers. Social Science and Medicine, 2009, 69, 632-639.	1.8	72
48	Dignity in the care of older people $\hat{a} \in $ a review of the theoretical and empirical literature. BMC Nursing, 2008, 7, 11.	0.9	142
49	With an attack I associate it more with going into hospital: Understandings of asthma and psychosocial stressors; are they related to use of services?. Social Science and Medicine, 2008, 66, 765-775.	1.8	20
50	`Habitus and Bureaucratic Routines', Cultural and Structural Factors in the Experience of Informal Care. Current Sociology, 2008, 56, 57-76.	0.8	21
51	The long shadow of workdoes time since labour market exit affect the association between socioeconomic position and health in a post-working population. Journal of Epidemiology and Community Health, 2007, 61, 533-539.	2.0	24
52	Ethical principles and the rationing of health care: a qualitative study in general practice. British Journal of General Practice, 2005, 55, 620-5.	0.7	18
53	Is patient involvement possible when decisions involve scarce resources? A qualitative study of decision-making in primary care. Social Science and Medicine, 2004, 59, 93-102.	1.8	73
54	Power, Present and Past: For a Historical Sociology of Health and Illness. Social Theory and Health, 2003, 1, 130-148.	1.0	4

#	Article	IF	CITATIONS
55	Social identities and the 'new genetics': Scientific and social consequences. Critical Public Health, 2002, 12, 265-282.	1.4	32
56	Evolutionary Psychology and Health: Confronting an Evolving Paradigm. Journal of Health Services Research and Policy, 1999, 4, 187-190.	0.8	1
57	Oregon, public health and social control. Critical Public Health, 1992, 3, 12-16.	1.4	1
58	Correspondence Analysis: A Case for Methodological Pluralism?. , 0, , 139-149.		1
59	Medical Sociology and Old Age. , 0, , .		38