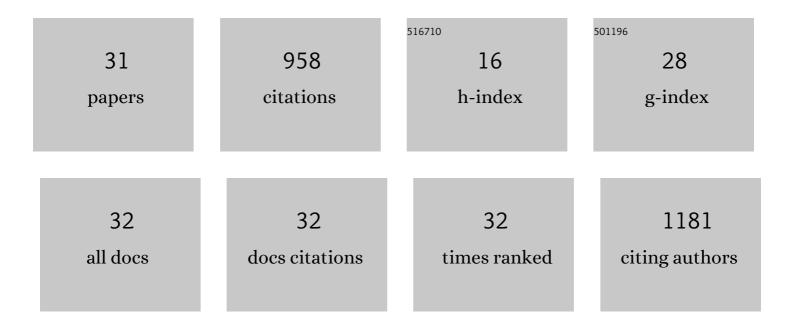
## Joseph T S Low

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/7898778/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	The impact of stroke on informal carers: a literature review. Social Science and Medicine, 1999, 49, 711-725.	3.8	233
2	Communication between healthcare professionals and relatives of patients approaching the end-of-life: A systematic review of qualitative evidence. Palliative Medicine, 2019, 33, 926-941.	3.1	121
3	Measuring spiritual belief: development and standardization of a Beliefs and Values Scale. Psychological Medicine, 2006, 36, 417-425.	4.5	76
4	End-of-life care for homeless people: A qualitative analysis exploring the challenges to access and provision of palliative care. Palliative Medicine, 2018, 32, 36-45.	3.1	66
5	Supportive and palliative care in people with cirrhosis: International systematic review of the perspective of patients, family members and health professionals. Journal of Hepatology, 2018, 69, 1260-1273.	3.7	66
6	Advanced chronic liver disease in the last year of life: a mixed methods study to understand how care in a specialist liver unit could be improved. BMJ Open, 2017, 7, e016887.	1.9	46
7	The impact of end-stage kidney disease (ESKD) on close persons: a literature review. CKJ: Clinical Kidney Journal, 2008, 1, 67-79.	2.9	38
8	Acceptance and commitment therapy for adults with advanced cancer (CanACT): A feasibility randomised controlled trial. Psycho-Oncology, 2019, 28, 488-496.	2.3	35
9	Palliative care for cirrhosis: a UK survey of health professionals' perceptions, current practice and future needs. Frontline Gastroenterology, 2016, 7, 4-9.	1.8	32
10	Narratives of continuity among older people with late stage chronic kidney disease who decline dialysis. Social Science and Medicine, 2014, 114, 49-56.	3.8	30
11	Acceptance and commitment therapy for adults with advanced cancer (CanACT): study protocol for a feasibility randomised controlled trial. Trials, 2016, 17, 77.	1.6	26
12	Managing uncertainty and references to time in prognostic conversations with family members at the end of life: A conversation analytic study. Palliative Medicine, 2020, 34, 896-905.	3.1	25
13	Challenges to discussing palliative care with people experiencing homelessness: a qualitative study. BMJ Open, 2017, 7, e017502.	1.9	24
14	The Role of Acceptance in Rehabilitation in Life-Threatening Illness. Journal of Pain and Symptom Management, 2012, 43, 20-28.	1.2	23
15	The experiences of close persons caring for people with chronic kidney disease stage 5 on conservative kidney management: Contested discourses of ageing. Health (United Kingdom), 2014, 18, 613-630.	1.5	19
16	The Subjective Index of Physical and Social Outcome: its usefulness in a younger stroke population. International Journal of Rehabilitation Research, 2004, 27, 59-63.	1.3	17
17	Palliative Care for Patients with End-Stage Liver Disease on the Liver Transplant Waiting List: An International Systematic Review. Digestive Diseases and Sciences, 2021, 66, 4072-4089.	2.3	15
18	The Subjective Index for Physical and Social Outcome (SIPSO) in Stroke: investigation of its subscale structure. BMC Neurology, 2010, 10, 26.	1.8	14

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19	Screening for Distress and Depression inÂCancer Patients: Is Ultrashort Depression Screening a Valid Measure in the UK? AÂProspective Validation Study. Journal of Pain and Symptom Management, 2009, 38, 234-243.	1.2	12
20	Social and structural conditions for the avoidance of advance care planning in neuro-oncology: a qualitative study. BMJ Open, 2018, 8, e019057.	1.9	11
21	Improving Communication in Outpatient Consultations in People With Cirrhosis. Journal of Clinical Gastroenterology, 2020, 54, 898-904.	2.2	8
22	SP707VALIDATION OF THE DISTRESS THERMOMETER IN A UK RENAL POPULATION. Nephrology Dialysis Transplantation, 2015, 30, iii612-iii612.	0.7	6
23	Transitioning out of prognostic talk in discussions with families of hospice patients at the end of life: A conversation analytic study. Patient Education and Counseling, 2021, 104, 1075-1085.	2.2	5
24	Do screening tools assess palliative care needs and 12-month mortality in patients admitted to hepatology in-patient wards?. Frontline Gastroenterology, 2022, 13, 211-217.	1.8	5
25	"l am in other people's hands as regards my health―A sociological critique of health care encounters of people with cirrhosis. A secondary analysis. Chronic Illness, 2023, 19, 102-117.	1.5	2
26	SP714HAEMODIALYSIS PATIENTS EXPERIENCE HIGHER LEVELS OF PSYCHOSOCIAL DISTRESS THAN EQUIVALENT CKD PATIENTS. Nephrology Dialysis Transplantation, 2015, 30, iii614-iii615.	0.7	1
27	Exploring the Impact of Different Types of Do-Not-Resuscitate Consent on End-of-Life Treatments among Patients with Advanced Kidney Disease: An Observational Study. International Journal of Environmental Research and Public Health, 2021, 18, 8194.	2.6	1
28	Variations in provision of psychological care to hematopoietic cell transplant recipients: results of a national survey of UK transplant centers. Bone Marrow Transplantation, 2022, 57, 1586-1588.	2.4	1
29	What support do people with cirrhosis need from liver healthcare professionals in a liver transplant centre?. BMJ Supportive and Palliative Care, 2016, 6, 389.3-390.	1.6	0
30	Why do we offer palliative care so late to patients with liver disease?. The Lancet Gastroenterology and Hepatology, 2018, 3, 225-226.	8.1	0
31	Understanding why Advance Care Plans and Goals of Care Discussions are so difficult to perform in Liver Transplantation – a new methodological approach from the Social Sciences. Journal of Liver Transplantation, 2022, , 100103.	0.4	0