

# 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

31  
papers

958  
citations

516710

16  
h-index

501196

28  
g-index

32  
all docs

32  
docs citations

32  
times ranked

1181  
citing authors

#	ARTICLE	IF	CITATIONS
1	“I am in other people's hands as regards my health” A sociological critique of health care encounters of people with cirrhosis. A secondary analysis. <i>Chronic Illness</i> , 2023, 19, 102-117.	1.5	2
2	Do screening tools assess palliative care needs and 12-month mortality in patients admitted to hepatology in-patient wards?. <i>Frontline Gastroenterology</i> , 2022, 13, 211-217.	1.8	5
3	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. <i>Journal of Liver Transplantation</i> , 2022, , 100103.	0.4	0
4	Variations in provision of psychological care to hematopoietic cell transplant recipients: results of a national survey of UK transplant centers. <i>Bone Marrow Transplantation</i> , 2022, 57, 1586-1588.	2.4	1
5	Transitioning out of prognostic talk in discussions with families of hospice patients at the end of life: A conversation analytic study. <i>Patient Education and Counseling</i> , 2021, 104, 1075-1085.	2.2	5
6	Palliative Care for Patients with End-Stage Liver Disease on the Liver Transplant Waiting List: An International Systematic Review. <i>Digestive Diseases and Sciences</i> , 2021, 66, 4072-4089.	2.3	15
7	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. <i>International Journal of Environmental Research and Public Health</i> , 2021, 18, 8194.	2.6	1
8	Improving Communication in Outpatient Consultations in People With Cirrhosis. <i>Journal of Clinical Gastroenterology</i> , 2020, 54, 898-904.	2.2	8
9	Managing uncertainty and references to time in prognostic conversations with family members at the end of life: A conversation analytic study. <i>Palliative Medicine</i> , 2020, 34, 896-905.	3.1	25
10	Communication between healthcare professionals and relatives of patients approaching the end-of-life: A systematic review of qualitative evidence. <i>Palliative Medicine</i> , 2019, 33, 926-941.	3.1	121
11	Acceptance and commitment therapy for adults with advanced cancer (CanACT): A feasibility randomised controlled trial. <i>Psycho-Oncology</i> , 2019, 28, 488-496.	2.3	35
12	Social and structural conditions for the avoidance of advance care planning in neuro-oncology: a qualitative study. <i>BMJ Open</i> , 2018, 8, e019057.	1.9	11
13	Why do we offer palliative care so late to patients with liver disease?. <i>The Lancet Gastroenterology and Hepatology</i> , 2018, 3, 225-226.	8.1	0
14	End-of-life care for homeless people: A qualitative analysis exploring the challenges to access and provision of palliative care. <i>Palliative Medicine</i> , 2018, 32, 36-45.	3.1	66
15	Supportive and palliative care in people with cirrhosis: International systematic review of the perspective of patients, family members and health professionals. <i>Journal of Hepatology</i> , 2018, 69, 1260-1273.	3.7	66
16	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. <i>BMJ Open</i> , 2017, 7, e016887.	1.9	46
17	Challenges to discussing palliative care with people experiencing homelessness: a qualitative study. <i>BMJ Open</i> , 2017, 7, e017502.	1.9	24
18	What support do people with cirrhosis need from liver healthcare professionals in a liver transplant centre?. <i>BMJ Supportive and Palliative Care</i> , 2016, 6, 389.3-390.	1.6	0

#	ARTICLE	IF	CITATIONS
19	Acceptance and commitment therapy for adults with advanced cancer (CanACT): study protocol for a feasibility randomised controlled trial. <i>Trials</i> , 2016, 17, 77.	1.6	26
20	Palliative care for cirrhosis: a UK survey of health professionals' perceptions, current practice and future needs. <i>Frontline Gastroenterology</i> , 2016, 7, 4-9.	1.8	32
21	SP707 VALIDATION OF THE DISTRESS THERMOMETER IN A UK RENAL POPULATION. <i>Nephrology Dialysis Transplantation</i> , 2015, 30, iii612-iii612.	0.7	6
22	SP714 HAEMODIALYSIS PATIENTS EXPERIENCE HIGHER LEVELS OF PSYCHOSOCIAL DISTRESS THAN EQUIVALENT CKD PATIENTS. <i>Nephrology Dialysis Transplantation</i> , 2015, 30, iii614-iii615.	0.7	1
23	The experiences of close persons caring for people with chronic kidney disease stage 5 on conservative kidney management: Contested discourses of ageing. <i>Health (United Kingdom)</i> , 2014, 18, 613-630.	1.5	19
24	Narratives of continuity among older people with late stage chronic kidney disease who decline dialysis. <i>Social Science and Medicine</i> , 2014, 114, 49-56.	3.8	30
25	The Role of Acceptance in Rehabilitation in Life-Threatening Illness. <i>Journal of Pain and Symptom Management</i> , 2012, 43, 20-28.	1.2	23
26	The Subjective Index for Physical and Social Outcome (SIPSO) in Stroke: investigation of its subscale structure. <i>BMC Neurology</i> , 2010, 10, 26.	1.8	14
27	Screening for Distress and Depression in Cancer Patients: Is Ultrashort Depression Screening a Valid Measure in the UK? A Prospective Validation Study. <i>Journal of Pain and Symptom Management</i> , 2009, 38, 234-243.	1.2	12
28	The impact of end-stage kidney disease (ESKD) on close persons: a literature review. <i>CKJ: Clinical Kidney Journal</i> , 2008, 1, 67-79.	2.9	38
29	Measuring spiritual belief: development and standardization of a Beliefs and Values Scale. <i>Psychological Medicine</i> , 2006, 36, 417-425.	4.5	76
30	The Subjective Index of Physical and Social Outcome: its usefulness in a younger stroke population. <i>International Journal of Rehabilitation Research</i> , 2004, 27, 59-63.	1.3	17
31	The impact of stroke on informal carers: a literature review. <i>Social Science and Medicine</i> , 1999, 49, 711-725.	3.8	233