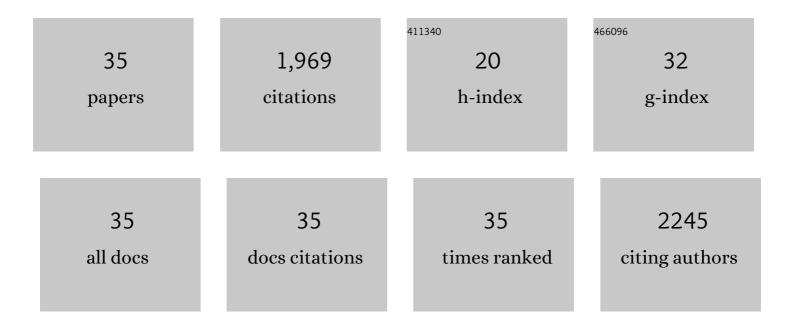
Christine J Mcpherson

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Involving family in discharge education: A nurse-led intervention to reduce hospital readmissions for older adults with heart failure and cognitive impairment. Evidence-based Nursing, 2021, 24, 126-126.	0.1	0
2	Without adequate nursing support for families, dying at home threatens the values of a good death. Evidence-based Nursing, 2021, 24, 45-45.	0.1	0
3	Qualitative synthesis of research on healthcare providers' perceptions of the transition from in-patient to community palliative care. Evidence-based Nursing, 2021, , ebnurs-2020-103352.	0.1	0
4	Visitor Restrictions, Palliative Care, and Epistemic Agency: A Qualitative Study of Nurses' Relational Practice During the Coronavirus Pandemic. Global Qualitative Nursing Research, 2021, 8, 233339362110517.	0.7	12
5	Public health nurses' experiences during the H1N1/09 response. Public Health Nursing, 2020, 37, 533-540.	0.7	9
6	Exploring the Experiences of Parent Caregivers of Adult Children With Schizophrenia: A Systematic Review. Archives of Psychiatric Nursing, 2019, 33, 93-103.	0.7	33
7	Unregulated care providers' engagement in palliative care to older clients and their families in the home setting: a mixed methods study. BMC Palliative Care, 2019, 18, 52.	0.8	6
8	Being Parent Caregivers for Adult Children with Schizophrenia. Issues in Mental Health Nursing, 2019, 40, 297-303.	0.6	10
9	Medical Assistance in Dying. Journal of Hospice and Palliative Nursing, 2019, 21, 46-53.	0.5	16
10	The meaning of being an oncology nurse: Investing to make a difference. Canadian Oncology Nursing Journal = Revue Canadienne De Nursing Oncologique, 2017, 27, 9-14.	0.1	4
11	Battling a Tangled Web: The Lived Experience of Nurses Providing End-of-Life Care on an Acute Medical Unit. Research and Theory for Nursing Practice, 2016, 30, 353-378.	0.2	5
12	Experiences of Family Caregivers Making the Transition From Home to the Palliative Care Unit. Journal of Hospice and Palliative Nursing, 2015, 17, 404-412.	0.5	2
13	A qualitative investigation of the roles and perspectives of older patients with advanced cancer and their family caregivers in managing pain in the home. BMC Palliative Care, 2014, 13, 39.	0.8	25
14	Cancer-Related Pain in Older Adults Receiving Palliative Care: Patient and Family Caregiver Perspectives on the Experience of Pain. Pain Research and Management, 2013, 18, 293-300.	0.7	27
15	Impact of Patient Smoking Behavior on Empathic Helping by Family Caregivers in Lung Cancer. Oncology Nursing Forum, 2012, 39, E112-E121.	0.5	10
16	A comparison of patient and family caregiver prospective control over lung cancer. Journal of Advanced Nursing, 2012, 68, 1122-1133.	1.5	4
17	Assessing Agreement Between Terminally Ill Cancer Patients' Reports of Their Quality of Life and Family Caregiver and Palliative Care Physician Proxy Ratings. Journal of Pain and Symptom Management, 2011, 42, 354-365.	0.6	80
18	The caregiving relationship and quality of life among partners of stroke survivors: A cross-sectional study. Health and Quality of Life Outcomes, 2011, 9, 29.	1.0	63

#	Article	IF	CITATIONS
19	The balance of give and take in caregiver–partner relationships: An examination of self-perceived burden, relationship equity, and quality of life from the perspective of care recipients following stroke Rehabilitation Psychology, 2010, 55, 194-203.	0.7	104
20	Prognostic Acceptance and the Well-Being of Patients Receiving Palliative Care for Cancer. Journal of Clinical Oncology, 2009, 27, 5757-5762.	0.8	89
21	Family Caregivers' Assessment of Symptoms in Patients with Advanced Cancer: Concordance with Patients and Factors Affecting Accuracy. Journal of Pain and Symptom Management, 2008, 35, 70-82.	0.6	87
22	Does Blaming the Patient With Lung Cancer Affect the Helping Behavior of Primary Caregivers?. Oncology Nursing Forum, 2008, 35, 681-689.	0.5	25
23	A Dyadic Affair. Cancer Nursing, 2008, 31, 435-443.	0.7	20
24	Suffering With Advanced Cancer. Journal of Clinical Oncology, 2007, 25, 1691-1697.	0.8	135
25	Desire for euthanasia or physician-assisted suicide in palliative cancer care Health Psychology, 2007, 26, 314-323.	1.3	136
26	Feeling like a burden to others: a systematic review focusing on the end of life. Palliative Medicine, 2007, 21, 115-128.	1.3	179
27	Self-Perceived Burden to Others: Patient and Family Caregiver Correlates. Journal of Palliative Care, 2007, 23, 135-142.	0.4	54
28	Feeling like a burden: Exploring the perspectives of patients at the end of life. Social Science and Medicine, 2007, 64, 417-427.	1.8	172
29	Self-perceived burden to others: patient and family caregiver correlates. Journal of Palliative Care, 2007, 23, 135-42.	0.4	9
30	Caring for patients with terminal delirium: palliative care unit and home care nurses' experiences. International Journal of Palliative Nursing, 2006, 12, 150-156.	0.2	38
31	A Burden to Others: A Common Source of Distress for the Terminally Ill. Cognitive Behaviour Therapy, 2005, 34, 115-123.	1.9	147
32	How do Proxies' Perceptions of Patients' Pain, Anxiety, and Depression Change during the Bereavement Period?. Journal of Palliative Care, 2004, 20, 12-19.	0.4	30
33	Evaluating palliative care: bereaved family members' evaluations of patients' pain, anxiety and depression. Journal of Pain and Symptom Management, 2004, 28, 104-114.	0.6	43
34	After-Death Interviews with Surrogates/Bereaved Family Members. Journal of Pain and Symptom Management, 2001, 22, 784-790.	0.6	171
35	Effective methods of giving information in cancer: a systematic literature review of randomized controlled trials. Journal of Public Health, 2001, 23, 227-234.	1.0	224