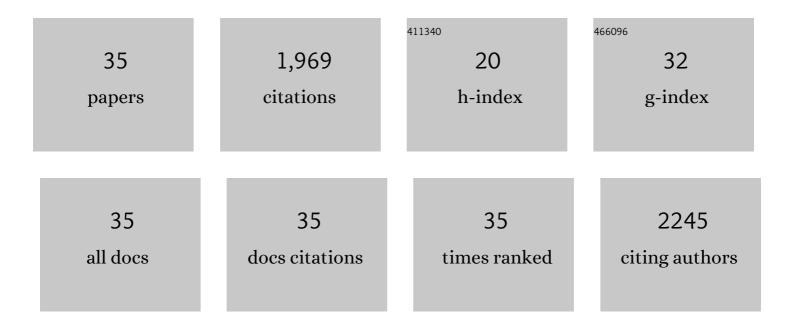
## **Christine J Mcpherson**

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/7824314/publications.pdf Version: 2024-02-01



| #  | 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.<br>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<br>Practice During the Coronavirus Pandemic. Global Qualitative Nursing Research, 2021, 8,<br>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<br>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,<br>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<br>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<br>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<br>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.<br>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<br>Family Caregiver and Palliative Care Physician Proxy Ratings. Journal of Pain and Symptom<br>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        |

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|----|---|-----|-----------|
| 19 | The balance of give and take in caregiver–partner relationships: An examination of self-perceived<br>burden, relationship equity, and quality of life from the perspective of care recipients following<br>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<br>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?.<br>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.<br>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<br>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<br>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       |