Mitsunori Miyashita

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

4,801 285 37 57 h-index g-index citations papers 2.6 5.31 5,590 304 L-index ext. citations avg, IF ext. papers

#	Paper	IF	Citations
285	Family experience of palliative sedation therapy: proportional vs. continuous deep sedation <i>Supportive Care in Cancer</i> , 2022 , 30, 3903	3.9	O
284	Quality Indicators in Palliative Radiation Oncology: Development and Pilot Testing <i>Advances in Radiation Oncology</i> , 2022 , 7, 100856	3.3	
283	Palliative care physicians' recognition of patients after immune checkpoint inhibitors and immune-related adverse events. <i>Supportive Care in Cancer</i> , 2022 , 30, 775-784	3.9	
282	Appropriate referral timing to specialized palliative care service: survey of bereaved families of cancer patients who died in palliative care units. <i>Supportive Care in Cancer</i> , 2022 , 30, 931-940	3.9	1
281	Association between temporary discharge from the inpatient palliative care unit and achievement of good death in end-of-life cancer patients: A nationwide survey of bereaved family members Japan Journal of Nursing Science, 2022, e12474	1.7	
280	Development of a list of competencies and entrustable professional activities for resident physicians during death pronouncement: a modified Delphi study <i>BMC Medical Education</i> , 2022 , 22, 119	3.3	1
279	Private or shared room? A nationwide questionnaire survey on bereaved family members' perceptions of where patients spend their last days <i>BMJ Open</i> , 2022 , 12, e055942	3	
278	Influence of financial burden on withdrawal or change of cancer treatment in Japan: results of a bereavement survey <i>Supportive Care in Cancer</i> , 2022 , 1	3.9	0
277	Development and Validation of the Death Pronouncement Burden Scale for Oncology Practice <i>Palliative Medicine Reports</i> , 2022 , 3, 39-46	1	1
276	Comparison of the quality of death between primary malignant brain tumor patients and other cancer patients: results from a nationwide bereavement survey in Japan <i>Journal of Neuro-Oncology</i> , 2022 , 1	4.8	О
275	Bathing in Terminal Care of Cancer Patients and Its Relation to Perceptions of a "Good Death": A Nationwide Bereavement Survey in Japan <i>Palliative Medicine Reports</i> , 2022 , 3, 55-64	1	
274	Nationwide survey on family caregiver-perceived experiences of patients with cancer of unknown primary site <i>Supportive Care in Cancer</i> , 2022 , 1	3.9	
273	Depression and complicated grief in bereaved caregivers in cardiovascular diseases: prevalence and determinants. <i>BMJ Supportive and Palliative Care</i> , 2021 ,	2.2	2
272	The prevalence of artificially administered nutrition and hydration in different age groups among patients with advanced cancer admitted to palliative care units. <i>Clinical Nutrition Open Science</i> , 2021 , 40, 69-78		
271	Bereaved Family Members' Perceived Care at the End of Life for Patients with Noncancerous Respiratory Diseases <i>Palliative Medicine Reports</i> , 2021 , 2, 265-271	1	O
270	How Successful Is Parenteral Oxycodone for Relieving Terminal Cancer Dyspnea Compared With Morphine? A Multicenter Prospective Observational Study. <i>Journal of Pain and Symptom Management</i> , 2021 , 62, 336-345	4.8	1
269	Patients' understanding of communication about palliative care and health condition in Japanese patients with unresectable or recurrent cancer: a cross-sectional survey. <i>Annals of Palliative Medicine</i> , 2021 , 10, 2650-2661	1.7	1

(2021-2021)

268	C-Reactive Protein and Its Relationship with Pain in Patients with Advanced Cancer Cachexia: Secondary Cross-Sectional Analysis of a Multicenter Prospective Cohort Study. <i>Palliative Medicine Reports</i> , 2021 , 2, 122-131	1	O
267	Are cancer patients living alone more or less likely to achieve a good death? Two cross-sectional surveys of bereaved families. <i>Journal of Advanced Nursing</i> , 2021 , 77, 3745-3758	3.1	O
266	Unmet need for palliative rehabilitation in inpatient hospices/palliative care units: a nationwide post-bereavement survey. <i>Japanese Journal of Clinical Oncology</i> , 2021 , 51, 1334-1338	2.8	
265	The Association of Family Functioning With Possible Major Depressive Disorders and Complicated Grief Among Bereaved Family Members of Patients With Cancer: Results From the J-HOPE4 Study, a Nationwide Cross-Sectional Follow-Up Survey in Japan. <i>Journal of Pain and Symptom Management</i> ,	4.8	O
264	Growth and Challenges in Hospital Palliative Cancer Care Services: An Analysis of Nationwide Surveys Over a Decade in Japan. <i>Journal of Pain and Symptom Management</i> , 2021 , 61, 1155-1164	4.8	1
263	Factors related to specialized palliative care use and aggressive care at end of life in Japanese patients with advanced solid cancers: a cohort study. <i>Supportive Care in Cancer</i> , 2021 , 29, 7805-7813	3.9	
262	Effects of enteral nutrition and parenteral nutrition on survival in patients with advanced cancer cachexia: Analysis of a multicenter prospective cohort study. <i>Clinical Nutrition</i> , 2021 , 40, 1168-1175	5.9	10
261	A Population-Based Mortality Follow-Back Survey Evaluating Good Death for Cancer and Noncancer Patients: A Randomized Feasibility Study. <i>Journal of Pain and Symptom Management</i> , 2021 , 61, 42-53.e.	2 ^{4.8}	O
260	Comparison of two measures for Complicated Grief: Brief Grief Questionnaire (BGQ) and Inventory of Complicated Grief (ICG). <i>Japanese Journal of Clinical Oncology</i> , 2021 , 51, 252-257	2.8	2
259	Cancer Care Evaluation Scale (CCES): measuring the quality of the structure and process of cancer care from the perspective of patients with cancer. <i>Japanese Journal of Clinical Oncology</i> , 2021 , 51, 92-9	9 ^{2.8}	
258	The association of family functioning and psychological distress in the bereaved families of patients with advanced cancer: a nationwide survey of bereaved family members. <i>Psycho-Oncology</i> , 2021 , 30, 74-83	3.9	3
257	Differences in aggressive treatments during the actively dying phase in patients with cancer and heart disease: an exploratory study using the sampling dataset of the National Database of Health Insurance Claims. <i>Heart and Vessels</i> , 2021 , 36, 724-730	2.1	1
256	High Feasibility and Safety, but Negligible Efficacy of Acupressure for Treating Nausea in Cancer Patients Admitted to the Palliative Care Unit: A Pilot Study. <i>Tohoku Journal of Experimental Medicine</i> , 2021 , 254, 155-161	2.4	
255	Development and validation of the Terminal Delirium-Related Distress Scale to assess irreversible terminal delirium. <i>Palliative and Supportive Care</i> , 2021 , 19, 287-293	2.5	O
254	Conditions, possibility and priority for admission into inpatient hospice/palliative care units in Japan: a nationwide survey. <i>Japanese Journal of Clinical Oncology</i> , 2021 , 51, 1437-1443	2.8	
253	Effects of financial status on major depressive disorder and complicated grief among bereaved family members of patients with cancer. <i>Psycho-Oncology</i> , 2021 , 30, 844-852	3.9	1
252	Predicting models of depression or complicated grief among bereaved family members of patients with cancer. <i>Psycho-Oncology</i> , 2021 , 30, 1151-1159	3.9	О
251	Comparison between patient-reported and clinician-reported outcomes: Validation of the Japanese version of the Integrated Palliative care Outcome Scale for staff. <i>Palliative and Supportive Care</i> , 2021 , 1-7	2.5	2

250	Physician's Communication in Code Status Discussions for Terminally Ill Cancer Patients in Inpatient Hospice/Palliative Care Units in Japan: A Nationwide Post-Bereavement Survey. <i>Journal of Pain and Symptom Management</i> , 2021 , 62, e120-e129	4.8	1
249	Effects of Bathing in a Tub on Physical and Psychological Symptoms of End-of-Life Cancer Patients: An Observational, Controlled Study. <i>Journal of Hospice and Palliative Nursing</i> , 2021 ,	2.2	1
248	Care needs level in long-term care insurance system and family caregivers' self-perceived time-dependent burden in patients with home palliative care for cancer: a cross-sectional study. <i>Supportive Care in Cancer</i> , 2021 , 1	3.9	0
247	Care Associated With Satisfaction of Bereaved Family Members of Terminally Ill Cancer Patients With Dyspnea: A Cross-sectional Nationwide Survey. <i>Journal of Pain and Symptom Management</i> , 2021 , 62, 796-804	4.8	O
246	Defining a good death for people with dementia: A scoping review. <i>Japan Journal of Nursing Science</i> , 2021 , 18, e12402	1.7	5
245	Spiritual Care Training Program Effects on Physicians Confidence, Self-reported Practice, and Attitudes in Caring for Terminally Ill Cancer Patients Who Express Meaninglessness. <i>Palliative Care Research</i> , 2021 , 16, 45-54	0.1	
244	Palliative care knowledge test for nurses and physicians: validation and cross-cultural adaptation. <i>BMJ Supportive and Palliative Care</i> , 2020 ,	2.2	4
243	Quality-of-life of Pediatric Cancer Patients Receiving End-of-life Care and Related Factors: Using a Proxy QOL Rating Scale from the Nursell Perspective. <i>Palliative Care Research</i> , 2020 , 15, 53-64	0.1	
242	Insomnia and changes in alcohol consumption: Relation between possible complicated grief and depression among bereaved family caregivers. <i>Journal of Affective Disorders</i> , 2020 , 275, 1-6	6.6	3
241	Perceptions of physicians and nurses concerning advanced care planning for patients with heart failure in Japan. <i>Annals of Palliative Medicine</i> , 2020 , 9, 1718-1731	1.7	7
240	Advance care planning for adults with heart failure. <i>The Cochrane Library</i> , 2020 , 2, CD013022	5.2	11
239	Impressions of Interfaith Chaplain's Activities among Patients in a Palliative Care Unit: A Semi-Structured Interview-Based Qualitative Study. <i>Tohoku Journal of Experimental Medicine</i> , 2020 , 251, 91-96	2.4	1
238	A Nationwide Survey of Bereaved Family Members' Perception of the Place Patients Spent Their Final Days: Is the Inpatient Hospice Like or Unlike a Home? Why?. <i>Palliative Medicine Reports</i> , 2020 , 1, 174-178	1	1
237	Prevalence and associated factors of perceived cancer-related stigma in Japanese cancer survivors. Japanese Journal of Clinical Oncology, 2020 , 50, 1325-1329	2.8	4
236	Rehabilitation for Cancer Patients in Inpatient Hospices/Palliative Care Units and Achievement of a Good Death: Analyses of Combined Data From Nationwide Surveys Among Bereaved Family Members. <i>Journal of Pain and Symptom Management</i> , 2020 , 60, 1163-1169	4.8	3
235	The Bereaved Families' Preferences for Individualized Goals of Care for Terminal Dyspnea: What Is an Acceptable Balance between Dyspnea Intensity and Communication Capacity?. <i>Palliative Medicine Reports</i> , 2020 , 1, 42-49	1	3
234	Potential Measurement Properties of a Questionnaire for Eating-Related Distress Among Advanced Cancer Patients With Cachexia: Preliminary Findings of Reliability and Validity Analysis. <i>Journal of Palliative Care</i> , 2020 , 825859720951356	1.8	3
233	Investigating associations between pain and complicated grief symptoms in bereaved Japanese older adults. <i>Aging and Mental Health</i> , 2020 , 24, 1472-1478	3.5	1

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232	Development of a Proxy Quality-of-Life Rating Scale for the End-of-Life Care of Pediatric Cancer Patients Evaluated from a Nurse's Perspective. <i>Journal of Palliative Medicine</i> , 2020 , 23, 82-89	2.2	2
231	Beliefs and Perceptions About Parenteral Nutrition and Hydration by Family Members of Patients With Advanced Cancer Admitted to Palliative Care Units: A Nationwide Survey of Bereaved Family Members in Japan. <i>Journal of Pain and Symptom Management</i> , 2020 , 60, 355-361	4.8	11
230	Quality indicators of palliative care for acute cardiovascular diseases. <i>Journal of Cardiology</i> , 2020 , 76, 177-183	3	6
229	Effectiveness of spiritual care using spiritual pain assessment sheet for advanced cancer patients: A pilot non-randomized controlled trial. <i>Palliative and Supportive Care</i> , 2019 , 17, 46-53	2.5	4
228	Validation of the Integrated Palliative care Outcome Scale (IPOS) - Japanese Version. <i>Japanese Journal of Clinical Oncology</i> , 2019 , 49, 257-262	2.8	40
227	Development and Validity of the Nursing Care Scale and Nurse's Difficulty Scale in Caring for Dying Patients With Cancer and Their Families in General Hospitals in Japan. <i>Journal of Hospice and Palliative Nursing</i> , 2019 , 21, 174-182	2.2	2
226	Validity and Reliability of the Dying Care Process and Outcome Scales Before and After Death From the Bereaved Family Members' Perspective. <i>American Journal of Hospice and Palliative Medicine</i> , 2019 , 36, 130-137	2.6	2
225	End-of-life Care and Good Death of Dying Non-cancer Patients from the Perspective of Bereaved Family Members. <i>Palliative Care Research</i> , 2019 , 14, 177-185	0.1	
224	Changes in Difficulty with Cancer Care among Nurses at Tohoku University Hospital between 2010 and 2016. <i>Palliative Care Research</i> , 2019 , 14, 259-267	0.1	O
223	Achievement of a good death among young adult patients with cancer: analyses of combined data from three nationwide surveys among bereaved family members. <i>Supportive Care in Cancer</i> , 2019 , 27, 1519-1527	3.9	3
222	Difference in Opinions About Continuous Deep Sedation Among Cancer Patients, Bereaved Families, and Physicians. <i>Journal of Pain and Symptom Management</i> , 2019 , 57, e5-e9	4.8	
221	Development and validation of the Comprehensive Quality of Life Outcome (CoQoLo) inventory for patients with advanced cancer. <i>BMJ Supportive and Palliative Care</i> , 2019 , 9, 75-83	2.2	3
220	Bereaved Family Members' Perceptions of the Distressing Symptoms of Terminal Patients With Cancer. <i>American Journal of Hospice and Palliative Medicine</i> , 2018 , 35, 972-977	2.6	1
219	Development of an Instrument for Evaluating Nurses' Knowledge and Attitude Toward End-of-Life Care: End-of-Life Nursing Education Consortium-Japan Core Quiz. <i>Journal of Hospice and Palliative Nursing</i> , 2018 , 20, 55-62	2.2	5
218	Communication about the impending death of patients with cancer to the family: a nationwide survey. <i>BMJ Supportive and Palliative Care</i> , 2018 , 8, 221-228	2.2	13
217	The distress and benefit to bereaved family members of participating in a post-bereavement survey. <i>Japanese Journal of Clinical Oncology</i> , 2018 , 48, 135-143	2.8	3
216	Factors associated with possible complicated grief and major depressive disorders. <i>Psycho-Oncology</i> , 2018 , 27, 915-921	3.9	25
215	Communication Disparity Between the Bereaved and Others: What Hurts Them and What Is Unhelpful? A Nationwide Study of the Cancer Bereaved. <i>Journal of Pain and Symptom Management</i> , 2018 , 55, 1061-1067.e1	4.8	4

214	The Japan hospice and palliative evaluation study 4: a cross-sectional questionnaire survey. <i>BMC Palliative Care</i> , 2018 , 17, 66	3	17
213	Families' Sense of Abandonment When Patients Are Referred to Hospice. <i>Oncologist</i> , 2018 , 23, 1109-11	15 7	5
212	Prevalence and predictors of conflict in the families of patients with advanced cancer: A nationwide survey of bereaved family members. <i>Psycho-Oncology</i> , 2018 , 27, 302-308	3.9	15
211	End-of-life experiences of family caregivers of deceased patients with cancer: A nation-wide survey. <i>Psycho-Oncology</i> , 2018 , 27, 272-278	3.9	4
210	Which Research Questions Are Important for the Bereaved Families of Palliative Care Cancer Patients? A Nationwide Survey. <i>Journal of Pain and Symptom Management</i> , 2018 , 55, 379-386	4.8	3
209	Improved knowledge of and difficulties in palliative care among physicians during 2008 and 2015 in Japan: Association with a nationwide palliative care education program. <i>Cancer</i> , 2018 , 124, 626-635	6.4	25
208	Physician Behavior toward Death Pronouncement in Palliative Care Units. <i>Journal of Palliative Medicine</i> , 2018 , 21, 368-372	2.2	2
207	Are Bereaved Family Members Satisfied With Information Provision About Palliative Care Units in Japan?. <i>American Journal of Hospice and Palliative Medicine</i> , 2018 , 35, 275-283	2.6	2
206	Changes in Nurses' Knowledge, Difficulties, and Self-reported Practices Toward Palliative Care for Cancer Patients in Japan: An Analysis of Two Nationwide Representative Surveys in 2008 and 2015. Journal of Pain and Symptom Management, 2018, 55, 402-412	4.8	8
205	Acculturation and Perceptions of a Good Death Among Japanese Americans and Japanese Living in the U.S. <i>Journal of Pain and Symptom Management</i> , 2018 , 55, 31-38	4.8	15
204	Advance care planning for heart failure. The Cochrane Library, 2018,	5.2	1
203	Patient perceptions of curability and physician-reported disclosures of incurability in Japanese patients with unresectable/recurrent cancer: a cross-sectional survey. <i>Japanese Journal of Clinical Oncology</i> , 2018 , 48, 913-919	2.8	1
202	End-of-life care for cancer patients in Japanese acute care hospitals: A nationwide retrospective administrative database survey. <i>Japanese Journal of Clinical Oncology</i> , 2018 , 48, 877-883	2.8	5
201	Development of a Scale for the Japanese Version of the Quality of Dying and Death in the Intensive Care Unit Instrument for Use by Nurses. <i>Palliative Care Research</i> , 2018 , 13, 121-128	0.1	1
200	Cancer Survivors Experiences of Changes in Employment and Income, and the Factors Affecting Their Experience, Quality of Life and Mental Health. <i>Palliative Care Research</i> , 2018 , 13, 209-218	0.1	2
199	Intervention and Implementation Studies on Integrated Care Pathway for End-of-Life Care in Long-term Care Facilities: A Scoping Review. <i>Palliative Care Research</i> , 2018 , 13, 313-327	0.1	O
198	The Job Types and Related Factors That Physicians and Nurses Feel Easy to Consult about Pediatric Cancer Patients and Their Families End of Life Care. <i>Palliative Care Research</i> , 2018 , 13, 89-98	0.1	
197	"What I Did for My Loved One Is More Important than Whether We Talked About Death": A Nationwide Survey of Bereaved Family Members. <i>Journal of Palliative Medicine</i> , 2018 , 21, 335-341	2.2	8

196	Desirable Information of Opioids for Families of Patients With Terminal Cancer. <i>American Journal of Hospice and Palliative Medicine</i> , 2017 , 34, 248-253	2.6	O
195	The Japan HOspice and Palliative Care Evaluation Study 3: Study Design, Characteristics of Participants and Participating Institutions, and Response Rates. <i>American Journal of Hospice and Palliative Medicine</i> , 2017 , 34, 654-664	2.6	34
194	Development the Care Evaluation Scale Version 2.0: a modified version of a measure for bereaved family members to evaluate the structure and process of palliative care for cancer patient. <i>BMC Palliative Care</i> , 2017 , 16, 8	3	18
193	Decision Making Regarding the Place of End-of-Life Cancer Care: The Burden on Bereaved Families and Related Factors. <i>Journal of Pain and Symptom Management</i> , 2017 , 53, 862-870	4.8	29
192	Potential palliative care quality indicators in heart disease patients: A review of the literature. Journal of Cardiology, 2017 , 70, 335-341	3	13
191	Effects of End-of-Life Discussions on the Mental Health of Bereaved Family Members and Quality of Patient Death and Care. <i>Journal of Pain and Symptom Management</i> , 2017 , 54, 17-26.e1	4.8	44
190	Collaborative care intervention for the perceived care needs of women with breast cancer undergoing adjuvant therapy after surgery: a feasibility study. <i>Japanese Journal of Clinical Oncology</i> , 2017 , 47, 213-220	2.8	7
189	The relationship between cancer patients' place of death and bereaved caregivers' mental health status. <i>Psycho-Oncology</i> , 2017 , 26, 1959-1964	3.9	14
188	Validation Study for the Brief Measure of Quality of Life and Quality of Care: A Questionnaire for the National Random Sampling Hospital Survey. <i>American Journal of Hospice and Palliative Medicine</i> , 2017 , 34, 622-631	2.6	1
187	Pediatric Cancer Patients' Important End-of-Life Issues, Including Quality of Life: A Survey of Pediatric Oncologists and Nurses in Japan. <i>Journal of Palliative Medicine</i> , 2017 , 20, 487-493	2.2	6
186	Unfinished Business in Families of Terminally Ill With Cancer Patients. <i>Journal of Pain and Symptom Management</i> , 2017 , 54, 861-869	4.8	9
185	Variations in Denominators and Cut-off Points of Pain Intensity in the Pain Management Index: A Methodological Systematic Review. <i>Journal of Pain and Symptom Management</i> , 2017 , 54, e1-e4	4.8	
184	Meaningful Communication Before Death, but Not Present at the Time of Death Itself, Is Associated With Better Outcomes on Measures of Depression and Complicated Grief Among Bereaved Family Members of Cancer Patients. <i>Journal of Pain and Symptom Management</i> , 2017 , 54, 273	4.8 - 279	49
183	Trust in Physicians, Continuity and Coordination of Care, and Quality of Death in Patients with Advanced Cancer. <i>Journal of Palliative Medicine</i> , 2017 , 20, 1252-1259	2.2	5
182	Talking About Death With Terminally-Ill Cancer Patients: What Contributes to the Regret of Bereaved Family Members?. <i>Journal of Pain and Symptom Management</i> , 2017 , 54, 853-860.e1	4.8	12
181	Psychological and psychiatric symptoms of terminally ill patients with cancer and their family caregivers in the home-care setting: A nation-wide survey from the perspective of bereaved family members in Japan. <i>Journal of Psychosomatic Research</i> , 2017 , 103, 127-132	4.1	9
180	Empowering Nurses Through Translating the End-of-Life Nursing Education Consortium. <i>Journal of Hospice and Palliative Nursing</i> , 2017 , 19, 539-549	2.2	7
179	Caregiving Consequence for Dying Elderly Patients with and without Comorbid Dementia from the Perspective of Bereaved Family Members. <i>Palliative Care Research</i> , 2017 , 12, 159-168	0.1	1

178	Views on life and death of physicians, nurses, cancer patients and general population in Japan. <i>PLoS ONE</i> , 2017 , 12, e0176648	3.7	3
177	Exploratory Qualitative Study of Regret Stemming from Ending Terminal Treatment and Psychological Coping among the Bereaved Family Members of Cancer Patients: What Does the Family Regret about Terminal Treatment Choices and Why?. <i>Palliative Care Research</i> , 2017 , 12, 753-760	0.1	O
176	Practical Use of Feedback Report of Nationwide Bereavement Survey in Participated Institutions. <i>Palliative Care Research</i> , 2017 , 12, 131-139	0.1	
175	Dementia Provision for Elderly Cancer Patients in Designated Regional Cancer Centers. <i>Palliative Care Research</i> , 2017 , 12, 116-124	0.1	1
174	Why Patients in Institutions with Religious Background Achieve Good Death ?: Findings from National Bereavement Survey of 127 Hospice and Palliative Care Units. <i>Palliative Care Research</i> , 2017, 12, 211-220	0.1	2
173	Guidelines for parenteral fluid management for terminal cancer patients. <i>Japanese Journal of Clinical Oncology</i> , 2016 , 46, 986-992	2.8	12
172	A simple home-based self-monitoring tool for early detection of hand-foot syndrome in cancer patients. <i>Japanese Journal of Clinical Oncology</i> , 2016 , 46, 979-985	2.8	2
171	The effects of community-wide dissemination of information on perceptions of palliative care, knowledge about opioids, and sense of security among cancer patients, their families, and the general public. <i>Supportive Care in Cancer</i> , 2016 , 24, 347-356	3.9	25
170	Japanese Bereaved Family Members' Perspectives of Palliative Care Units and Palliative Care: J-HOPE Study Results. <i>American Journal of Hospice and Palliative Medicine</i> , 2016 , 33, 425-30	2.6	4
169	Changes in Perceptions of Opioids Before and After Admission to Palliative Care Units in Japan: Results of a Nationwide Bereaved Family Member Survey. <i>American Journal of Hospice and Palliative Medicine</i> , 2016 , 33, 431-8	2.6	1
168	The Association Between Pain and Quality of Life for Patients With Cancer in an Outpatient Clinic, an Inpatient Oncology Ward, and Inpatient Palliative Care Units. <i>American Journal of Hospice and Palliative Medicine</i> , 2016 , 33, 782-90	2.6	11
167	Population-Based Quality Indicators for Palliative Care Programs for Cancer Patients in Japan: A Delphi Study. <i>Journal of Pain and Symptom Management</i> , 2016 , 51, 652-661	4.8	12
166	Association Between Bereaved Families' Sense of Security and Their Experience of Death in Cancer Patients: Cross-Sectional Population-Based Study. <i>Journal of Pain and Symptom Management</i> , 2016 , 51, 926-32	4.8	5
165	A Preliminary Survey to Measure the Quality Indicators of End-of-life Cancer Care Using the Japanese National Database. <i>Palliative Care Research</i> , 2016 , 11, 156-165	0.1	3
164	Factors that Influence the Decision Maker regarding End-of-life Care. <i>Palliative Care Research</i> , 2016 , 11, 189-200	0.1	4
163	Quality Evaluation and Improvement for End-of-life Care toward Residents in Long Term Care Facilities in Japan: A Literature Review. <i>Palliative Care Research</i> , 2016 , 11, 401-412	0.1	3
162	A Second Time Nationwide Survey of Quality of End-of-Life Cancer Care in General Hospitals, Inpatient Palliative Care Units, and Clinics in Japan: The J-HOPE 2 Study. <i>Palliative Care Research</i> , 2016 , 11, 254-264	0.1	6
161	Family caregivers require mental health specialists for end-of-life psychosocial problems at home: a nationwide survey in Japan. <i>Psycho-Oncology</i> , 2016 , 25, 641-7	3.9	5

160	Anxiety and depression in patients after surgery for head and neck cancer in Japan. <i>Palliative and Supportive Care</i> , 2016 , 14, 269-77	2.5	19
159	Evaluation of care for leukemia and lymphoma patients during their last hospitalization from the perspective of the bereaved family. <i>Leukemia Research</i> , 2016 , 47, 93-9	2.7	4
158	Development and validation of scales for attitudes, self-reported practices, difficulties and knowledge among home care nurses providing palliative care. <i>European Journal of Oncology Nursing</i> , 2016 , 22, 8-22	2.8	17
157	Changes in Relatives' Perspectives on Quality of Death, Quality of Care, Pain Relief, and Caregiving Burden Before and After a Region-Based Palliative Care Intervention. <i>Journal of Pain and Symptom Management</i> , 2016 , 52, 637-645	4.8	10
156	Nationwide Japanese Survey About Deathbed Visions: "My Deceased Mother Took Me to Heaven". Journal of Pain and Symptom Management, 2016 , 52, 646-654.e5	4.8	15
155	End-of-Life Medical Treatments in the Last Two Weeks of Life in Palliative Care Units in Japan, 2005-2006: A Nationwide Retrospective Cohort Survey. <i>Journal of Palliative Medicine</i> , 2016 , 19, 1188-11	36 ²	8
154	Eating-related distress and need for nutritional support of families of advanced cancer patients: a nationwide survey of bereaved family members. <i>Journal of Cachexia, Sarcopenia and Muscle</i> , 2016 , 7, 527-534	10.3	57
153	The importance of good death components among cancer patients, the general population, oncologists, and oncology nurses in Japan: patients prefer "fighting against cancer". <i>Supportive Care in Cancer</i> , 2015 , 23, 103-10	3.9	16
152	Prevalence, associated factors and source of support concerning supportive care needs among Japanese cancer survivors. <i>Psycho-Oncology</i> , 2015 , 24, 635-42	3.9	21
151	A Nationwide Survey of Quality of End-of-Life Cancer Care in Designated Cancer Centers, Inpatient Palliative Care Units, and Home Hospices in Japan: The J-HOPE Study. <i>Journal of Pain and Symptom Management</i> , 2015 , 50, 38-47.e3	4.8	54
150	Independent Validation of the Japanese Version of the EORTC QLQ-C15-PAL for Patients With Advanced Cancer. <i>Journal of Pain and Symptom Management</i> , 2015 , 49, 953-9	4.8	8
149	Outcome evaluation of the Palliative care Emphasis program on symptom management and Assessment for Continuous Medical Education: nationwide physician education project for primary palliative care in Japan. <i>Journal of Palliative Medicine</i> , 2015 , 18, 45-9	2.2	25
148	Association between depressive symptoms and changes in sleep condition in the grieving process. Supportive Care in Cancer, 2015 , 23, 1925-31	3.9	12
147	Validation of the Japanese version of HFS-14, a disease-specific quality of life scale for patients suffering from hand-foot syndrome. <i>Supportive Care in Cancer</i> , 2015 , 23, 2739-45	3.9	8
146	Place of death and the differences in patient quality of death and dying and caregiver burden. Journal of Clinical Oncology, 2015 , 33, 357-63	2.2	104
145	Anxiety in disease-free breast cancer patients might be alleviated by provision of psychological support, not of information. <i>Japanese Journal of Clinical Oncology</i> , 2015 , 45, 929-33	2.8	12
144	Improvements in Physicians' Knowledge, Difficulties, and Self-Reported Practice After a Regional Palliative Care Program. <i>Journal of Pain and Symptom Management</i> , 2015 , 50, 232-40	4.8	15
143	Views on death with regard to end-of-life care preferences among cancer patients at a Japanese university hospital. <i>Palliative and Supportive Care</i> , 2015 , 13, 969-79	2.5	1

142	Cross-sectional online survey of research productivity in young Japanese nursing faculty. <i>Japan Journal of Nursing Science</i> , 2015 , 12, 198-207	1.7	5
141	Length of home hospice care, family-perceived timing of referrals, perceived quality of care, and quality of death and dying in terminally ill cancer patients who died at home. <i>Supportive Care in Cancer</i> , 2015 , 23, 491-9	3.9	19
140	Strategies for Development of Palliative Care From the Perspectives of General Population and Health Care Professionals: A Japanese Outreach Palliative Care Trial of Integrated Regional Model Study. <i>American Journal of Hospice and Palliative Medicine</i> , 2015 , 32, 604-10	2.6	2
139	Changes in and Associations Among Functional Status and Perceived Quality of Life of Patients With Metastatic/Locally Advanced Cancer Receiving Rehabilitation for General Disability. <i>American Journal of Hospice and Palliative Medicine</i> , 2015 , 32, 695-702	2.6	15
138	Why people accept opioids: role of general attitudes toward drugs, experience as a bereaved family, information from medical professionals, and personal beliefs regarding a good death. <i>Journal of Pain and Symptom Management</i> , 2015 , 49, 45-54	4.8	6
137	Factors influencing death or the cessation of palliative care in home-based setting among patients with cancer. <i>Palliative Care Research</i> , 2015 , 10, 116-123	0.1	3
136	Analysis of administrative data to investigate end-of-life cancer care in a Japanese university hospital: development of methodology. <i>Palliative Care Research</i> , 2015 , 10, 177-185	0.1	2
135	Family experience and evaluation about caring the body of deceased patient in Japanese inpatient hospices: a content analysis of free comments in questionnaire survey. <i>Palliative Care Research</i> , 2015 , 10, 209-216	0.1	
134	Current home palliative care for terminally ill cancer patients in Japan. <i>Palliative Care Research</i> , 2015 , 10, 153-161	0.1	5
133	Progressive development and enhancement of palliative care services in Japan: nationwide surveys of designated cancer care hospitals for three consecutive years. <i>Journal of Pain and Symptom Management</i> , 2014 , 48, 364-73	4.8	9
132	Does a regional comprehensive palliative care program improve pain in outpatient cancer patients?. <i>Supportive Care in Cancer</i> , 2014 , 22, 2445-55	3.9	3
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128	Care evaluation scale-patient version: measuring the quality of the structure and process of palliative care from the patient's perspective. <i>Journal of Pain and Symptom Management</i> , 2014 , 48, 110-8	₄ .8	10
127	A pilot study of adaptation of the transtheoretical model to narratives of bereaved family members in the bereavement life review. <i>American Journal of Hospice and Palliative Medicine</i> , 2014 , 31, 422-7	2.6	1
126	Symptom burden and achievement of good death of elderly cancer patients. <i>Journal of Palliative Medicine</i> , 2014 , 17, 887-93	2.2	13
125	Nurse education program on meaninglessness in terminally ill cancer patients: a randomized controlled study of a novel two-day workshop. <i>Journal of Palliative Medicine</i> , 2014 , 17, 1298-305	2.2	14

124	Association between quality of end-of-life care and possible complicated grief among bereaved family members. <i>Journal of Palliative Medicine</i> , 2014 , 17, 1025-31	2.2	36
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121	One-year follow-up of an educational intervention for palliative care consultation teams. <i>Japanese Journal of Clinical Oncology</i> , 2014 , 44, 172-9	2.8	6
120	A population-based survey on perceptions of opioid treatment and palliative care units: OPTIM study. <i>American Journal of Hospice and Palliative Medicine</i> , 2014 , 31, 155-60	2.6	6
119	What bereavement follow-up does family members request in Japanese palliative care units? A qualitative study. <i>American Journal of Hospice and Palliative Medicine</i> , 2014 , 31, 485-94	2.6	7
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109	Depressive symptoms after treatment in hepatocellular carcinoma survivors: prevalence, determinants, and impact on health-related quality of life. <i>Psycho-Oncology</i> , 2013 , 22, 2347-53	3.9	15
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94	Making the decision for home hospice: perspectives of bereaved Japanese families who had loved ones in home hospice. <i>Japanese Journal of Clinical Oncology</i> , 2012 , 42, 498-505	2.8	4
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90	Leaflet for health professionals based on patient and family surveys: ^ ^ldquo;Voices of cancer patients and families^ ^rdquo;. <i>Palliative Care Research</i> , 2012 , 7, 342-347	0.1	2
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82	Cross-cultural application of the Korean version of the European Organization for Research and Treatment of Cancer Quality of Life Questionnaire-Core 15-Palliative Care. <i>Journal of Pain and Symptom Management</i> , 2011 , 41, 478-84	4.8	29
81	Experience with prognostic disclosure of families of Japanese patients with cancer. <i>Journal of Pain and Symptom Management</i> , 2011 , 41, 594-603	4.8	11
80	Measuring comprehensive outcomes in palliative care: validation of the Korean version of the Good Death Inventory. <i>Journal of Pain and Symptom Management</i> , 2011 , 42, 632-42	4.8	21
79	The relationships between complicated grief, depression, and alexithymia according to the seriousness of complicated grief in the Japanese general population. <i>Journal of Affective Disorders</i> , 2011 , 135, 122-7	6.6	12
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69	Multiple evaluation of a hospital-based palliative care consultation team in a university hospital: activities, patient outcome, and referring staff's view. <i>Palliative and Supportive Care</i> , 2010 , 8, 49-57	2.5	18
68	The palliative care self-reported practices scale and the palliative care difficulties scale: reliability and validity of two scales evaluating self-reported practices and difficulties experienced in palliative care by health professionals. <i>Journal of Palliative Medicine</i> , 2010 , 13, 427-37	2.2	49
67	Preference of place for end-of-life cancer care and death among bereaved Japanese families who experienced home hospice care and death of a loved one. <i>Supportive Care in Cancer</i> , 2010 , 18, 1445-53	3.9	37
66	Changes in medical and nursing care after admission to palliative care units: a potential method for improving regional palliative care. <i>Supportive Care in Cancer</i> , 2010 , 18, 1107-13	3.9	3
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61	Value of religious care for relief of psycho-existential suffering in Japanese terminally ill cancer patients: the perspective of bereaved family members. <i>Psycho-Oncology</i> , 2010 , 19, 750-5	3.9	12
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54	Meaninglessness in terminally ill cancer patients: a randomized controlled study. <i>Journal of Pain and Symptom Management</i> , 2009 , 37, 649-58	4.8	24
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51	Administrators' perspectives on end-of-life care for cancer patients in Japanese long-term care facilities. <i>Supportive Care in Cancer</i> , 2009 , 17, 1247-54	3.9	4
50	Caregiving Consequences Inventory: a measure for evaluating caregiving consequences from the bereaved family member's perspective. <i>Psycho-Oncology</i> , 2009 , 18, 657-66	3.9	35
49	Care burden and depression in caregivers caring for patients with intractable neurological diseases at home in Japan. <i>Journal of the Neurological Sciences</i> , 2009 , 276, 148-52	3.2	51
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47	The Long-Term effect of A Population-Based Educational Intervention Focusing on End-Of-Life Home Care, Life-Prolongation Treatment, and Knowledge about Palliative Care. <i>Journal of Palliative Care</i> , 2009 , 25, 206-212	1.8	7
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40	Effect of a population-based educational intervention focusing on end-of-life home care, life-prolonging treatment and knowledge about palliative care. <i>Palliative Medicine</i> , 2008 , 22, 376-82	5.5	20
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37	Evaluation of end-of-life cancer care from the perspective of bereaved family members: the Japanese experience. <i>Journal of Clinical Oncology</i> , 2008 , 26, 3845-52	2.2	69
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35	Quality of life, day hospice needs, and satisfaction of community-dwelling patients with advanced	2.2	21

34	Reliability assessment and findings of a newly developed quality measurement instrument: quality indicators of end-of-life cancer care from medical chart review at a Japanese regional cancer center. <i>Journal of Palliative Medicine</i> , 2008 , 11, 729-37	2.2	16
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28	The current status of palliative care teams in Japanese university hospitals: a nationwide questionnaire survey. <i>Supportive Care in Cancer</i> , 2007 , 15, 801-6	3.9	7
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25	Good death in cancer care: a nationwide quantitative study. <i>Annals of Oncology</i> , 2007 , 18, 1090-7	10.3	228
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19	Good death in Japanese cancer care: a qualitative study. <i>Journal of Pain and Symptom Management</i> , 2006 , 31, 140-7	4.8	145
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17	Knowledge and beliefs about end-of-life care and the effects of specialized palliative care: a	4.8	83

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10	Association between hydration volume and symptoms in terminally ill cancer patients with abdominal malignancies. <i>Annals of Oncology</i> , 2005 , 16, 640-7	10.3	91
9	Emotional burden of nurses in palliative sedation therapy. <i>Palliative Medicine</i> , 2004 , 18, 550-7	5.5	69
8	Development of a diabetes diet-related quality-of-life scale. <i>Diabetes Care</i> , 2004 , 27, 1271-5	14.6	22
7	Physician- and nurse-reported effects of intravenous hydration therapy on symptoms of terminally ill patients with cancer. <i>Journal of Palliative Medicine</i> , 2004 , 7, 683-93	2.2	30
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4	Effects of an off-site walking program on energy expenditure, serum lipids, and glucose metabolism in middle-aged women. <i>Biological Research for Nursing</i> , 2003 , 4, 181-92	2.6	26
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1	Phase I/II study of vinorelbine, mitomycin, and cisplatin for stage IIIB or IV non-small-cell lung cancer. <i>Journal of Clinical Oncology</i> , 1999 , 17, 3195-200	2.2	8