

Literature Reviews on Palliative Care Family Carer Research

Note:

This list of literature reviews in palliative care family carer research is compiled by the IPCFRC Administrator via [PubMed](#) search using the search terms: 'caregiver', 'carer', 'hospice', 'family', 'research' and 'palliative'; and spanning the period of **1 Jan 2016 – 17 Mar 2017**. The search yielded a total number of **29** publications which are listed below. This list will be reviewed and updated annually and serves as a basic resource to aid those who are looking for publications related to family carer research. Please note that this is not a comprehensive list of publications in the area of family carer research.

1. Sarmiento VP, Gysels M, Higginson IJ, Gomes B. (2017). "Home palliative care works: but how? A meta-ethnography of the experiences of patients and family caregivers." *BMJ Support Palliat Care*. doi: 10.1136/bmjspcare-2016-001141. [Epub ahead of print]
2. Cagle JG, Bunting M, Kelemen A, Lee J, Terry D, Harris R. (2017). "Psychosocial needs and interventions for heart failure patients and families receiving palliative care support: a systematic review." *Heart Fail Rev*. doi: 10.1007/s10741-017-9596-5. [Epub ahead of print]
3. Ankuda CK, Kersting K, Guetterman TC, Haefner J, Fonger E, Paletta M, Hopp F. (2017). "What Matters Most? A Mixed Methods Study of Critical Aspects of a Home-Based Palliative Program". *Am J Hosp Palliat Care*. doi: 10.1177/1049909117691929. [Epub ahead of print]
4. Epstein RM, Duberstein PR, Fenton JJ, Fiscella K, Hoerger M, Tancredi DJ, Xing G, Gramling R, Mohile S, Franks P, Kaesberg P, Plumb S, Cipri CS, Street RL Jr, Shields CG, Back AL, Butow P, Walczak A, Tattersall M, Venuti A, Sullivan P, Robinson M, Hoh B, Lewis L, Kravitz RL. (2017). "Effect of a Patient-Centered Communication Intervention on Oncologist-Patient Communication, Quality of Life, and Health Care Utilization in Advanced Cancer: The VOICE Randomized Clinical Trial." *JAMA Oncol*. 3(1):92-100.
5. Spatuzzi R, Giulietti MV, Ricciuti M, Merico F, Meloni C, Fabbietti P, Ottaviani M, Violani C, Cormio C, Vespa A. (2016). "Quality of life and burden in family caregivers of patients with advanced cancer in active treatment settings and hospice care: A comparative study." *Death Stud*. doi: 10.1080/07481187.2016.1273277. [Epub ahead of print]
6. Woodman C, Baillie J, Sivell S. (2016). "The preferences and perspectives of family caregivers towards place of care for their relatives at the end-of-life. A systematic review and thematic synthesis of the qualitative evidence." *BMJ Support Palliat Care* 6(4):418-429.
7. Collins A, Hennessy-Anderson N, Hosking S, Hynson J, Remedios C, Thomas K. (2016). "Lived experiences of parents caring for a child with a life-limiting condition in Australia: A qualitative study." *Palliat Med*. 30(10):950-959.
8. Harrop E, Morgan F, Byrne A, Nelson A. (2016). ""It still haunts me whether we did the right thing": a qualitative analysis of free text survey data on the bereavement experiences and support needs of family caregivers." *BMC Palliat Care*. 2016 Nov 8;15(1):92.



9. Harano K, Yonemori K, Hirakawa A, Shimizu C, Katsumata N, Gemma A, Fujiwara Y, Tamura K. (2016). "The influence of familial factors on the choice of the place of death for terminally ill breast cancer patients: a retrospective single-center study." *Breast Cancer* 23(5):797-806.
10. Jack BA, Mitchell TK, Cope LC, O'Brien MR. (2016). "Supporting older people with cancer and life-limiting conditions dying at home: a qualitative study of patient and family caregiver experiences of Hospice at Home care." *J Adv Nurs.* 72(9):2162-72.
11. Lee YJ, Kim JE, Choi YS, Hwang IC, Hwang SW, Kim YS, Kim HM, Ahn HY, Kim SJ. (2016) "Quality of life discordance between terminal cancer patients and family caregivers: a multicenter study." *Support Care Cancer* 24(7):2853-60.
12. Baird-Bower D, Roach J, Andrews M, Onslow F, Curnin E. (2016). "Help is just a phone call away: after-hours support for palliative care patients wishing to die at home." *Int J Palliat Nurs.* 22(6):286-91.
13. DiBiasio EL, Teno JM, Clark MA, Spence C, Casarett D. (2016). "Development of an Assessment to Examine Training of the Hospice Primary Caregiver." *J Palliat Med.* 19(6):639-45.
14. Kobayakawa M, Okamura H, Yamagishi A, Morita T, Kawagoe S, Shimizu M, Ozawa T, An E, Tsuneto S, Shima Y, Miyashita M. (2016). "Family caregivers require mental health specialists for end-of-life psychosocial problems at home: a nationwide survey in Japan." *Psychooncology* 25(6):641-7.
15. Sutherland N, Ward-Griffin C, McWilliam C, Stajduhar K. (2016). "Gendered Processes in Hospice Palliative Home Care for Seniors With Cancer and Their Family Caregivers." *Qual Health Res.* 26(7):907-20.
16. Meier EA, Gallegos JV, Thomas LP, Depp CA, Irwin SA, Jeste DV. (2016). "Defining a Good Death (Successful Dying): Literature Review and a Call for Research and Public Dialogue." *Am J Geriatr Psychiatry* 24(4):261-71.
17. Holtslander L, Duggleby W, Teucher U, Cooper D, Bally JM, Solar J, Steeves M. (2016) "Developing and pilot-testing a Finding Balance Intervention for older adult bereaved family caregivers: A randomized feasibility trial." *Eur J Oncol Nurs.* 21:66-74.
18. Gleeson A, Larkin P, O'Sullivan N. (2016). "The impact of meticillin-resistant *Staphylococcus aureus* on patients with advanced cancer and their family members: A qualitative study." *Palliat Med.* 30(4):382-91.
19. Ewing G, Austin L, Grande G. (2016) "The role of the Carer Support Needs Assessment Tool in palliative home care: A qualitative study of practitioners' perspectives of its impact and mechanisms of action." *Palliat Med.* 30(4):392-400.
20. Hudson P, Collins A, Bostanci A, Willenberg L, Stepanov N, Philip J. (2016) "Toward a systematic approach to assessment and care planning in palliative care: A practical review of clinical tools." *Palliat Support Care.* 14(2):161-73.
21. Washington KT, Oliver DP, Gage LA, Albright DL, Demiris G.(2016) "A multimethod analysis of shared decision-making in hospice interdisciplinary team meetings including family caregivers." *Palliat Med.* 30(3):270-8.



22. Johnson MJ, Allgar V, Macleod U, Jones A, Oliver S, Currow D. (2016). "Family Caregivers Who Would Be Unwilling to Provide Care at the End of Life Again: Findings from the Health Survey for England Population Survey." *PLoS One*. 11(1):e0146960.
23. Schnakenberg R, Goeldlin A, Boehm-Stiel C, Bleckwenn M, Weckbecker K, Radbruch L. (2016). "Written survey on recently deceased patients in Germany and Switzerland: how do general practitioners see their role?" *BMC Health Serv Res*. 16:22.
24. Kataoka-Yahiro MR, McFarlane S, Kojane J, Li D. (2016). "Culturally Competent Palliative and Hospice Care Training for Ethnically Diverse Staff in Long-Term Care Facilities." *Am J Hosp Palliat Care*. doi: 10.1177/1049909116638347. [Epub ahead of print]
25. Chi NC, Demiris G. (2016) "Family Caregivers' Pain Management in End-of-Life Care." *Am J Hosp Palliat Care*. doi: 10.1177/1049909116637359. [Epub ahead of print]
26. O'Callaghan CC, McDermott F, Reid P, Michael N, Hudson P, Zalcborg JR, Edwards J. (2016). "Music's Relevance for People Affected by Cancer: A Meta-Ethnography and Implications for Music Therapists." *J Music Ther*. 53(4):398-429.
27. MacKenzie MA, Buck HG, Meghani SH, Riegel B. (2016). "Unique Correlates of Heart Failure and Cancer Caregiver Satisfaction With Hospice Care." *J Pain Symptom Manage* 51(1):71-8.
28. Thienprayoon R, Marks E, Funes M, Martinez-Puente LM, Winick N, Lee SC. (2016) "Perceptions of the Pediatric Hospice Experience among English- and Spanish-Speaking Families." *J Palliat Med*. 19(1):30-41.
29. Akiyama M, Hirai K, Takebayashi T, Morita T, Miyashita M, Takeuchi A, Yamagishi A, Kinoshita H, Shirahige Y, Eguchi K. (2016) "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." *Support Care Cancer* 24(1):347-56.