



## End-of-Life Care Bibliography

- 2006 Monographs Task Force of the American Society for Healthcare Risk Management. (2006, August). *Enterprise risk management: Perspectives on advance directives*. Chicago: American Society for Healthcare Risk Management. Available from <http://www.ashrm.org>
- Achtenberg, B., & Mitchell, C. (2001). *Caring at the end of life* [Videotape]. Boston, MA: Fanlight Productions.
- Arnold, R. L., & Egan, K. (2004). Breaking the “bad” news to patients and families: Preparing to have the conversation about end-of-life and hospice care. *American Journal of Geriatric Cardiology*, 13(6), 307-312.
- Astrow, A. B., Wexler, A., Teixeira, K., He, M. K., & Sulmasy, D. P. (2007). Is failure to meet spiritual needs associated with cancer patients’ perceptions of quality of care and their satisfaction with care? *Journal of Clinical Oncology*, 25(36), 5753-5757.
- Back, A. L., Arnold, R. M., Baile, W. F., Fryer-Edwards, K. A., & Tulsky, J. M. (2005). On saying goodbye: Acknowledging the end of the patient–physician relationship with patients who are near death. *Annals of Internal Medicine*, 142(8), 683-685.
- Block, S. D., & Billings, J. A. (2005). Learning from the dying. *The New England Journal of Medicine*, 353(13), 1313-1315.
- Bronner, E. (2003). The foundation’s end-of-life programs: Changing the American way of death. In *The Robert Wood Johnson Foundation anthology: To improve health and health care, Volume VI*. Retrieved January 2, 2009, from [http://www.rwjf.org/files/publications/books/2003/chapter\\_04.html](http://www.rwjf.org/files/publications/books/2003/chapter_04.html)
- Casarett, D., Karlawish, J., Morales, K., Crowley, R., Mirsch, T., & Asch, D. A. (2005). Improving the use of hospice services in nursing homes: A randomized controlled trial. *Journal of the American Medical Association*, 294(2), 211-217.
- Clarke, E. B., Curtis, J. R., Luse, J. M., Levy, M., Danis, M., Nelson, J., et al. (2003). Quality indicators for end-of-life care in the intensive care unit. *Critical Care Medicine*, 31(5 Sup), S367-S372.
- Curtis, R. J., Engelberg, R. A., Wenrich, M. D., Shannon, S. E., Treece, P. D., & Rubenfeld, G. D. (2005). Missed opportunities during family conferences about end-of-life care in the intensive care unit. *American Journal of Respiratory and Critical Care Medicine*, 272, 844-849.
- Farber, S. J., Egnaw, T. R., Herman-Bertsch, J. L., Taylor, T. R., & Guildin, G. E. (2003). Issues in end-of-life care: Patient, caregiver, and clinician perspectives. *Journal of Palliative Medicine*, 6(1), 19-31.
- Field, M. J., & Behrman, R. E. (2003). *When children die: Improving palliative and end-of-life care for children and their families*. Washington, DC: National Academies Press. Available from [http://www.nap.edu/catalog.php?record\\_id=10390](http://www.nap.edu/catalog.php?record_id=10390)
- Hallenbeck, J. (2005). Palliative care in the final days of life: They were expecting it at any time. *Journal of the American Medical Association*, 293(18), 2265-2271.
- Innovations in End-of-Life Care*, an international journal of leaders in end-of-life care, published bimonthly (January 1999 through September 2003). All 28 past issues are archived and available from <http://www2.edc.org/lastacts>
- Last Acts (2002). *Means to a better end: A report on dying in America today*. Retrieved January 5, 2009, from <http://www.rwjf.org/files/publications/other/meansbetterend.pdf>

- Levy, M. M. (2007). A view from the other side. *Critical Care Medicine*, 35(2), 603-604.
- Longden, J. V., & Mayer, A. P. T. (2007). Family involvement in end-of-life care in a paediatric intensive care unit. *Nursing in Critical Care*, 12(4), 181-187.
- Meyer, E. C., Ritholz, M. D., Burns, J. P., & Truog, R. D. (2006). Improving the quality of end-of-life care in the pediatric intensive care unit: Parents' priorities and recommendations. *Pediatrics*, 117(3), 649-657.
- Meyer, E. C., Sellers, D. E., Browning, D. M., McGuffie, K., Solomon, M. Z., & Truog, R. D. (2009). Difficult conversations: Improving communication skills and relational abilities in health care. *Pediatric Critical Care Medicine*, 10(3), 352-359.
- Monterosso, L., Kristjanson, L., Aoun, S., & Phillips, M. B. (2007). Supportive and palliative care needs of families of children with life-threatening illnesses in Western Australia: Evidence to guide the development of a palliative care service. *Palliative Medicine*, 21(8), 689-696.
- National Consensus Project for Quality Palliative Care. (2004). *Clinical practice guidelines for quality palliative care*. Brooklyn, NY: Author.
- Ngo-Metzger, Q., August, K. J., Srinivasan, M., Liao, S., & Meyskens, F. L. (2008). End-of-life care: Guidelines for patient-centered communication. *American Family Physician*, 77(2), 167-174. Available from <http://www.aafp.org/afp>
- Perkins, P., Barclay, S., & Booth, S. (2007). What are patients' priorities for palliative care research? Focus group study. *Palliative Medicine*, 21(3), 219-225.
- Pollak, K. I., Arnold, R. M., Jeffrey, A. S., Alexander, S. C., Olsen, M. K., Abernethy, A. P., et al. (2007). Oncologist communication about emotion during visits with patients with advanced cancer. *Journal of Clinical Oncology*, 25(36), 5748-5752.
- Public Affairs Television, Inc., Owen, E. (Producer). (2000). *On our own terms: Moyers on dying*. Available from <http://www.pbs.org/wnet/onourown/terms>
- Ruder, D. B. (2006). Life lessons: Gravely ill patients teach medical students about listening and compassion. *Harvard Magazine*, 108(3), 44-51, 90-91.
- Sanders, C., Rogers, A., Gately, C., & Kennedy, A. (2008). Planning for end of life care within lay-led chronic illness self-management training: The significance of 'death awareness' and biographical context in participant accounts. *Social Science & Medicine*, 66(4), 892-893.
- Sullivan, A. M., Lakoma, M. J., Billings, A., Peters, A. S., Block, S., & PCEP Core Faculty. (2005). Teaching and learning end-of-life care: Evaluation of a faculty development program in palliative care. *Academic Medicine*, 80(77), 657-668. Retrieved January 2, 2009, from <http://www.hms.harvard.edu/cdi/pallcare/AcadMedJuly05.pdf>
- Sweeney, L., Halpert, A., & Waranoff, J. (2007). Patient-centered management of complex patients can reduce costs without shortening life. *The American Journal of Managed Care*, 13(2), 84-92.
- Take Ten Inc. Productions (Producer). Emmot, H., & Russell, J. (Writers). (2004). *Ethics thru drama* [Videotape]. Boston, MA: Fanlight Productions.
- Takayesu, J. K., & Hutson, H. R. (2004). Communicating life-threatening diagnoses to patients in the emergency department. *Annals of Emergency Medicine*, 43(6), 749-755.
- Teno, J. M., Clarridge, B. R., Casey, V., Welch, L. C., Wetle, T., Shield, R., et al. (2004). Family perspectives on end-of-life care at the last place of care. *Journal of the American Medical Association*, 291(1), 88-93.
- Truog, R. D., Meyer, E. C., & Burns, J. P. (2006). Toward interventions to improve end-of-life care in the pediatric intensive care unit. *Critical Care Medicine*, 34(11), S373-S379.
- Wall, R. J., Curtis, J. R., Cooke, C. R., & Engelberg, R. A. (2007). Family satisfaction in the ICU. Differences between families of survivors and nonsurvivors. *Chest*, 132(5), 1425-1433.
- Wetle, T., Shield, R., Teno, J., Miller, S. C., & Welch, L. (2005). Family perspectives on end-of-life care experiences in nursing homes. *Gerontologist*, 45(5), 642-650.