

DAFTAR PUSTAKA

Aljawi, M. D., & Harfort, B. J. (2012). Palliative Care in the Muslim-Majority Countries: The Need for more and better care in Contemporary and innovative practice in palliative care, Prof. Esther Chang (Ed.), pp. 138-150, InTech, www.intechopen.com [accessed September 11, 2016].

Anon, N. D., et al. Modul 1: Qualitative Research Methods Overview pada Qualitative Research Methods: A Data Collector's Field Guide. www.ccs.neu.edu/course/is4800sp12/resources/qualmethods.pdf [accessed August 21, 2015].

Badr, H., Carmack, L. C., & Diefenbach, A. M. (2015). Psychosocial Interventions for Patients and Caregivers in the Age of New Communication Technologies: Opportunities and Challenges in Cancer Care, *J Health Commun.*, March, pp. 328-342.

Balboni, A. T., Paulk, E. M., & Balboni, J. M. (2010). Provision of spiritual care to patients with advanced cancer: associations with medical care and quality of life near death, *Journal of Clinical Oncology*, Vol. 28, No. 3, pp. 445-453.

Balboni, A. T., Balboni, M., Enzinger, C. A., et al. (2013). Provision of Spiritual Support to Patients With Advanced Cancer by Religious Communities and Associations with Medical Care at The End of Life, *JAMA Intern Med.*, June, pp. 1109-1117.

Balls, P. (2009). Phenomenology in nursing research: methodology, interviewing and transcribing. www.medscape.com [accessed August 18, 2015].

Boog, K., & Tester, C. (2008). *Palliative Care A Practical Guide for The Health Professional in Finding meaning and purpose in life and death*. Churchill Livingstone: Elsevier. www.sciencedirect.com [accessed June 08, 2015].

Bruera, E., & Hui, D. (2012). Conceptual Models for Integrating Palliative Care at Cancer Centers. *Journal of Palliative Medicine*, Vol 15, no. 11.

Bruera, E., & Yennurajalingam, S. (2012). Palliative Care in Advanced Cancer Patient: How and When, www.TheOncologist.com, pp 267-273.

California Health Care Foundation. (2007). *Palliative Care in California: an Overview of Hospital-based Programs*. www.chcf.org [accessed September 11, 2016].

Eerden, M., Csikos, A., Busa, C., et al. (2014). Experiences of patients, family and professional caregivers with Integrated Palliative Care in Europe: protocol for an international, multicenter, prospective, mixed method study, *BMC Palliative Care*. www.biomedcentral.com [accessed Juni 24, 2015].

Farooqui, M., Hassali, A. M., Shatar, K. A., et al. (2011). A qualitative exploration of Malaysian cancer patients perspective on cancer and its Treatment, *BMC Public Health*. www.biomedcentral.com [accessed November 09, 2016].

Gupta, D. (2012). Dilemma in Palliation in Contemporary and Innovative Practice in Palliative Care, Prof. Esther Chang (Ed.), pp. 105-114, InTech, www.intechopen.com [accessed September 11, 2016].

Hak, T., & Champion, P., (1999). Achieving a patient-centred consultation by giving feedback in its early phases, *Postgrad Med J*, pp. 405-409, www.researchgate.net [accessed July 05, 2015].

Hannon, B., et al., (2016). Experiences of patients and caregivers with early palliative care: a qualitative study. pmj.sagepub.com [accessed September 11, 2016].

Holm, M., Carlander, I., Furst, J. C., et al. (2015). Delivering and participating in a psycho-educational intervention for family caregivers during palliative home care: a qualitative study from the perspectives of health professionals and family caregivers. *BMC Palliative Care* [accessed November 11, 2016].

King, M., Llewellyn, H., Leurent, B., et al. (2013). Spiritual beliefs near the end of life: a prospective cohort study of people with cancer receiving palliative care. *Psycho-oncology*, pp. 2505-2512.

Maizes, V., Rakel, D., Niemiec, C., (2009). Integrative Medicine and Patient-Centered Care. www.ncbi.nlm.nih.gov [accessed Auguts 09, 2015].

Mazanec, Polly., Daly, J. B., Pitorak, F. E., et al. (2009). A New Model of Palliative Care for Oncology Patients with Advanced Disease, *Journal of hospice and palliative nursing*, Vol. 11, No. 6, pp. 324-331. www.medscape.com [accessed May 22, 2015].

McLeod, D. L., Tapp, D. M., Moules, N. J., et al. (2010). Knowing the family: interpretations of family nursing in oncology and palliative care. *European*

Journal of Oncology Nursing, Vol. 14, No. 2, pp. 93-100. www.elsevier.com [accessed August 09, 2015].

McPherson, C. J., Wilson, K. G., & Murray, M. A., (2006). Feeling like a burden: Exploring the perspectives of patients at the end of life, *Social science and medicine*, Vol. 64, No. 2, pp. 417-427. www.elsevier.com [accessed July 05, 2015].

Montgogery, K., & Liittle, M., (2011). Enriching Patient-Centered Care in Serious Illness: A Focus on Patients Experiences of Agency. *The Milbank Quarterly A Multidisciplinary Journal of Population Health and Health Policy*, Vol. 89, No. 3, pp. 381-398.

NCCN Guideline, (2015). Palliative Care versi 2. www.nccn.org [accessed September 11, 2016].

Pusat Data dan Informasi Kementerian Kesehatan RI, (2015). Stop Kanker. www.depkes.go.id [accessed September 11, 2016].

Sage, (2016). Ethical Issues in Qualitative Research. uk.sagepub.com [accessed August 21, 2015].

Slevin, M. L., Nichols, S. E., Downer, S. M., et al., (1996). Emotional support for cancer patient: what do patients really wants?, *British journal of cancer*, pp. 1275-1279. www.nature.com/bjc/journal [accessed August 09, 2015].

Terry, W., Olson, L. G., Wills, L., et al. (2006). Experience of dying: concerns of dying patients and of carers, *Internal medicine journal*, Vol. 36, pp. 338-346. www.ncbi.nlm.nih.gov [accessed July 05, 2015].

Tiby-Levy, Y., & Bungener, M., (2012). Information Needs in Palliative Care: Patient and Family Perspectives in Contemporary and Innovative Practice in Palliative Care, Prof. Esther Chang (Ed.), pp. 138-150, InTech, www.intechopen.com [accessed September 11, 2016].

Tong, A., & Sainsbury, P., (2007). Consolidated criteria for reporting qualitative research (COREQ): a 32-item checklist for interviews and focus group. mat.or.th/journal/download/COREQ [accessed August 18, 2016].

Tsianakas, V., Robert, G., Maben, J., et al, (2012). Implementing patient-centred cancer care: using experience-based co-design to improve patient experience in

breast and lung cancer services in Supportive care in cancer. link.springer.com/article [accessed August 09, 2015].

Vadivelu, N., Kaye, D. A. & Berger, M. J. (2013). *Essential of Palliative Care*. New York: Springer-Verlag New York. www.springer.com [accessed May 30, 2015].

Vilalta, A., Valls, J., Porta, J., et al. (2014). Evaluation of Spiritual Needs of Patients with Advanced Cancer in a Palliative Care Unit. *Journal of Palliative Medicine*, Volume 17, p. 5.

Witjaksono, M., Sutandiyo, N., Suardi, D. (2014). Regional support for palliative care in Indonesia. www.ehospice.com [accessed August 09, 2015].

Connor, R. S., et al. (2014). *Global Atlas of Palliative Care at the End of Life*. Diunduh dari www.who.int/nmh/GlobalAtlasofPalliativeCare pada tanggal 2 Juni 2015.

Keputusan Menteri Kesehatan nomor 812/Menkes/SK/VII tahun 2007 tentang *Kebijakan Perawatan Paliatif*. Diunduh dari <https://perpustakaan.depkes.go.id> pada tanggal 2 Juni 2015.

Chang, Esther dan Johnson, Amanda. 2012. *Contemporary and Innovative Practice in Palliative Care*. Diunduh dari <https://handle.uws.edu.au/> pada tanggal 30 Mei 2015.

Pusat Data dan Informasi Kementerian Kesehatan RI. *Stop Kanker*. 2015. Diunduh dari www.depkes.go.id pada tanggal 20 Agustus 2015.

Effendy, Christantie. 2014. *Pengembangan Manajemen Pelayanan Paliatif* pada *Jurnal Manajemen Pelayanan Kesehatan*. Diunduh dari jurnal.ugm.ac.id pada tanggal 09 Agustus 2015.

Dewi, et al. 2013. *Person-centered care in the Indonesian health-care system* pada *International Journal of Nursing Practice*. Diunduh dari www.researchgate.net pada tanggal 09 Agustus 2015.

Quebec End-Of-Life Palliative Care Policy. Diunduh dari www.chpca.net pada tanggal 18 Agustus 2015.

Palliative Care in California: an Overview of Hospital-based Programs. California Health Care Foundation. Diunduh dari www.chcf.org pada tanggal 18 Agustus 2015.