

DAFTAR PUSTAKA

- Abraham, L., Hareendran, A., Mills, I. W., Martin, M. L., Abrams, P., Drake, M. J., MacDonagh, R. P., & Noble, J. G. (2004). Development and validation of a quality-of-life measure for men with nocturia. *Urology*, *63*(3), 481–486. <https://doi.org/10.1016/j.urology.2003.10.019>
- Adelman, R. D., Tmanova, L. L., Delgado, D., Dion, S., & Lachs, M. S. (2014). Caregiver burden: a clinical review. *JAMA*, *311*(10), 1052–1060. <https://doi.org/10.1001/jama.2014.304>
- Aggarwal, S., Chadha, P., Kalia, S., Richardson, S., Winterbottom, L., & Shields, L. (2009). Perceptions of family-centred care: A UK pilot study of the Shields and Tanner questionnaires. *Neonatal, Paediatric and Child Health Nursing*, *12*.
- Alderfer MA, et al. (2010) Psychosocial adjustment of siblings of children with cancer: a systematic review. *Psycho-oncology*, *19*(8):789/805.
- Ali, K., Sutaryo, S., Purwanto, I., Mulatsih, S., Supriyadi, E., Widjajanto, P. H., Sumadiono, S., & Nurse, J. (2010). Yogyakarta Pediatric Cancer Registry: an international collaborative project of University Gadjah Mada, University of Saskatchewan, and the Saskatchewan Cancer Agency. *Asian Pacific Journal of Cancer Prevention : APJCP*, *11*(1), 131–136.
- American Cancer Society. (2010). The Atlas of Canada Evapotranspiration.pdf. *American Cancer Society*.
- American Cancer Society. (2013). Cancer Facts & Figures 2013. *American Cancer Society*.
- Amery, J. (2012). *A Really Practical Handbook of Children's Palliative Care*. Lulu Publishing Services. <https://books.google.co.id/books?id=pAu3CwAAQBAJ>
- Anderson, C. P., Matthay, K. K., Perentesis, J. P., Neglia, J. P., Bailey, H. H., Villablanca, J. G., Groshen, S., Hasenauer, B., Maris, J. M., Seeger, R. C., & Reynolds, C. P. (2015). Pilot study of intravenous melphalan combined with continuous infusion L-S,R-buthionine sulfoximine for children with recurrent neuroblastoma. *Pediatric Blood & Cancer*, *62*(10), 1739–1746. <https://doi.org/10.1002/pbc.25594>
- Arslan, F. T., Basbakkal, Z., & Kantar, M. (2013). Quality of life and chemotherapy-related symptoms of Turkish cancer children undergoing chemotherapy. *Asian Pacific Journal of Cancer Prevention : APJCP*, *14*(3), 1761–1768. <https://doi.org/10.7314/apjcp.2013.14.3.1761>
- Association for Children's Palliative Care. (2009). A Guide to the Development of Children's Palliative Care Services. *A Guide to the Development of Children's Palliative Care Services*, *34*. http://www.togetherforshortlives.org.uk/assets/0001/1649/ACT_Guide_to_Developing_Services.pdf
- Azwar, S. (2015). *Reliabilitas dan Validitas* (4th ed.). Pustaka Pelajar.
- Azwar, S. (2016). *Penyusunan Skala Psikologis* (2nd ed.). Pustaka Pelajar.
- Ball, J. W., & Bindler, R. C. (2003). *Pediatric Nursing: Caring for Children, 3rd Edition*. Pearson.
- Ball, J. W., Bindler, R. C., & Cowen, K. (2015). *Principles of Pediatric Nursing: Caring for Children, 6th Edition*. Pearson. <https://www.pearson.com/us/higher-education/product/Ball-Principles-of-Pediatric-Nursing-Caring-for-Children-6th-Edition/9780133898064.html?tab=authors>
- Bessell, A. G. (2001). Children Surviving Cancer: Psychosocial Adjustment, Quality of Life, and School Experiences. *Exceptional Children*, *67*(3), 345–359. <https://doi.org/10.1177/001440290106700304>
- Blazin, L. J., Cecchini, C., Habashy, C., Kaye, E. C., & Baker, J. N. (2018). Communicating Effectively in Pediatric Cancer Care: Translating Evidence into Practice. *Children (Basel, Switzerland)*, *5*(3), 40. <https://doi.org/10.3390/children5030040>
- Brien, T. O. (2017). Report of the National Advisory Committee on Palliative Care.

Departement of Health and Children.

- Brown, S. M., Rozenblum, R., Aboumatar, H., Fagan, M. B., Milic, M., Lee, B. S., Turner, K., & Frosch, D. L. (2015). Defining patient and family engagement in the intensive care unit'. *American Journal of Respiratory and Critical Care Medicine*, 191(3), 358-360. <https://doi.org/10.1164/rccm.201410-1936LE>
- Bruce, B., & Ritchie, J. (1997). Nurses' practices and perceptions of family-centered care. *Journal of Pediatric Nursing*, 12(4), 214-222. [https://doi.org/10.1016/S0882-5963\(97\)80004-8](https://doi.org/10.1016/S0882-5963(97)80004-8)
- Campbell, R. J., Kagan, N., & Krathwohl, D. R. (1971). The development and validation of a scale to measure affective sensitivity (empathy). *Journal of Counseling Psychology*, 18(5), 407-412. <https://doi.org/10.1037/h0031492>
- Campbell, S. M., Braspenning, J., Hutchinson, A., & Marshall, M. (2003). Research methods used in developing and applying quality indicators in primary care. *Quality & Safety in Health Care*, 326, 816-819. <https://doi.org/10.1136/qhc.11.4.358>
- Catlin, A., Ford, M., & Maloney, C. (2015). Determining Family Needs on an Oncology Hospital Unit Using Interview, Art, and Survey. *Clinical Nursing Research*, 25(2), 209-231. <https://doi.org/10.1177/1054773815578806>
- Chang, A. M., Gardner, G. E., Duffield, C., & Ramis, M.-A. (2010). A Delphy study to validate an advanced practice nursing tool. *Journal of Advanced Nursing*, 66(10), 2320-2330. <https://doi.org/10.1111/j.1365-2648.2010.05367.x>
- Cheng, B. T., & Wangmo, T. (2020). Palliative care utilization in hospitalized children with cancer. *Pediatric blood & cancer*, 67(1), e28013. <https://doi.org/10.1002/pbc.28013>
- Coats, H., Bourget, E., Starks, H., Lindhorst, T., Saiki-Craighill, S., Curtis, J. R., Hays, R., & Doorenbos, A. (2018). Nurses' Reflections on Benefits and Challenges of Implementing Family-Centered Care in Pediatric Intensive Care Units. *American journal of critical care : an official publication, American Association of Critical-Care Nurses*, 27(1), 52-58. <https://doi.org/10.4037/ajcc2018353>
- Corden, A., Sloper, P., & Sainsbury, R. (2002). Financial effects for families after the death of a disabled or chronically ill child: a neglected dimension of bereavement. *Child: Care, Health and Development*, 28(3), 199-204. <https://doi.org/10.1046/j.1365-2214.2002.00267.x>
- Coyne, I., Murphy, M., Costello, T., O'Neill, C., & Donnellan, C. (2013). A survey of nurses' practices and perceptions of family-centered care in Ireland. *Journal of Family Nursing*, 19(4), 469-488. <https://doi.org/10.1177/1074840713508224>
- Crespo, C., Santos, S., Tavares, A., & Salvador, Á. (2016). "Care that matters": Family-centered care, caregiving burden, and adaptation in parents of children with cancer. *Families, Systems & Health : The Journal of Collaborative Family Healthcare*, 34(1), 31-40. <https://doi.org/10.1037/fsh0000166>
- Creswell, J. W., & Clark, V. L. P. (2011). *Designing and Conducting Mixed Methods Research* (2nd ed.). SAGE Publication.
- Cypress, B. S. (2012). Family Presence on Rounds: A Systematic Review of Literature. *Dimensions of Critical Care Nursing*, 31(1). https://journals.lww.com/dccjournal/Fulltext/2012/01000/Family_Presence_on_Rounds_A_Systematic_Review_of.15.aspx
- da Silva, F. M., Jacob, E., & Nascimento, L. C. (2010). Impact of childhood cancer on parents' relationships: an integrative review. *Journal of Nursing Scholarship : An Official Publication of Sigma Theta Tau International Honor Society of Nursing*, 42(3), 250-261. <https://doi.org/10.1111/j.1547-5069.2010.01360.x>
- Davidson, J. E., Aslakson, R. A., Long, A. C., Puntillo, K. A., Kross, E. K., Hart, J., Cox, C. E., Wunsch, H., Wickline, M. A., Nunnally, M. E., Netzer, G., Kentish-Barnes, N., Sprung, C. L., Hartog, C. S., Coombs, M., Gerritsen, R. T., Hopkins, R. O., Franck, L. S., Skrobik, Y., ... Curtis, J. R. (2017). Guidelines for Family-Centered Care in the Neonatal,

- Pediatric, and Adult ICU. *Critical Care Medicine*, 45(1), 103–128. <https://doi.org/10.1097/CCM.0000000000002169>
- Davidson, J. E., Powers, K., Hedayat, K. M., Tieszen, M., Kon, A. A., Shepard, E., Spuhler, V., Todres, I. D., Levy, M., Barr, J., Ghandi, R., Hirsch, G., & Armstrong, D. (2007). Clinical practice guidelines for support of the family in the patient-centered intensive care unit: American College of Critical Care Medicine Task Force 2004-2005. *Critical Care Medicine*, 35(2), 605–622. <https://doi.org/10.1097/01.CCM.0000254067.14607.EB>
- Dellve, L., Samuelsson, L., Tallborn, A., Fasth, A., & Hallberg, L. R.-M. (2006). Stress and well-being among parents of children with rare diseases: a prospective intervention study. *Journal of Advanced Nursing*, 53(4), 392–402. <https://doi.org/10.1111/j.1365-2648.2006.03736.x>
- DeVellis, R. F. (2012). *Scale Development: Theory and Applications* (3rd ed.). SAGE Publication.
- Dorothee, A. (1998). *Spiritual Care and Palliative Care: Opportunities and Challenges for Pastoral Care*. www.who.int/cancer/palliative/definition/en/
- Downing, J., Jassal, S. S., Mathews, L., Brits, H., & Friedrichsdorf, S. J. (2015). Pediatric pain management in palliative care. *Pain Management*, 5(1), 23–35. <https://doi.org/10.2217/pmt.14.45>
- Doyle, D., Hanks, G., Cherny, N. I., & Calman, S. K. (2013). *Oxford Textbook of Palliative Medicine* (3rd Editio). Oxford Medical Publications (OUP).
- Effendy, C., Vissers, K., Tejawinata, S., Vernooij-Dassen, M., & Engels, Y. (2015). Dealing with Symptoms and Issues of Hospitalized Patients with Cancer in Indonesia: The Role of Families, Nurses, and Physicians. *Pain Practice*, 15(5), 441–446. <https://doi.org/10.1111/papr.12203>
- Ekstedt, M., Stenberg, U., Olsson, M., & Ruland, C. M. (2014). Health care professionals' perspectives of the experiences of family caregivers during in-patient cancer care. *Journal of Family Nursing*, 20(4), 462–486. <https://doi.org/10.1177/1074840714556179>
- Finley, G. A., Kristjánssdóttir, O., & Forgeron, P. A. (2009). Cultural influences on the assessment of children's pain. *Pain Research & Management*, 14(1), 33–37. <https://doi.org/10.1155/2009/763031>
- Fitch, K., Bernstein, S. J. J., Aguilar, M. D., Burnand, B., LaCalle, J. R., Lazaro, P., Van Het Loo, M., McDonnell, J., Vader, J. P., & Kahan, J. P. (2001). The RAND / UCLA Appropriateness Method User's Manual. In *Transformation*.
- Fochtman, D. (2006). The Concept of Suffering in Children and Adolescents With Cancer. *Journal of Pediatric Oncology Nursing*, 23(2), 92–102. <https://doi.org/10.1177/1043454205285870>
- Foster, M., Whitehead, L., & Maybee, P. (2016). The Parents', Hospitalized Child's, and Health Care Providers' Perceptions and Experiences of Family-Centered Care Within a Pediatric Critical Care Setting: A Synthesis of Quantitative Research. *Journal of Family Nursing*, 22(1), 6–73. <https://doi.org/10.1177/1074840715618193>
- França, J. R. F. de S., da Costa, S. F. G., Lopes, M. E. L., da Nóbrega, M. M. L., & de França, I. S. X. (2013). The importance of communication in pediatric oncology palliative care: focus on Humanistic Nursing Theory. *Revista Latino-Americana de Enfermagem*, 21(3), 780–786. <https://doi.org/10.1590/S0104-11692013000300018>
- Friedman. (1998). *Family Nursing : Research, Theory and Practice* (4th Ed). Alpleton & Lange.
- Friedman, M. ., Bowden, V. R., & Jones, E. G. (2003). *Family Nursing: Research, Theory, & Practice* (5th ed.). Pearson Education Inc.
- Ganz, F. D., & Yoffe, F. (2012). Intensive care nurses' perspectives of family-centered care and their attitudes toward family presence during resuscitation. *The Journal of*

- Cardiovascular Nursing*, 27(3), 220–227. <https://doi.org/10.1097/JCN.0b013e31821888b4>
- Gill, F. J., Pascoe, E., Monterosso, L., Young, J., Charlotte Burr, Tanner, A., & Shields, L. (2014). *Parent and staff perceptions of family-centered care in two Australian children's hospitals*. 1(2), 317–325. http://researchonline.nd.edu.au/cgi/viewcontent.cgi?article=1077&context=nursing_article
- Glaser, B. G., & Strauss, A. L. (2006). *The discovery of grounded theory: strategies for qualitative research*. Aldine transaction.
- Grinyer, A. (2006). Caring for a young adult with cancer: the impact on mothers' health. *Health & Social Care in the Community*, 14(4), 311–318. <https://doi.org/10.1111/j.1365-2524.2006.00622.x>
- Hallström, I. (2008). Commentary on Wong MY-F & Chan SW-C (2006) The qualitative experience of Chinese parents with children diagnosed of cancer. *Journal of Clinical Nursing* 15, 710-717. *Journal of Clinical Nursing*, 17(2), 282–283; author reply 283. <https://doi.org/10.1111/j.1365-2702.2006.01678.x>
- Harding, R., Powell, R. A., Downing, J., Connor, S. R., Mwangi-Powell, F., Defilippi, K., Cameron, S., Garanganga, E., Kikule, E., & Alexander, C. (2008). Generating an African palliative care evidence base: the context, need, challenges, and strategies. *Journal of Pain and Symptom Management*, 36(3), 304–309. <https://doi.org/10.1016/j.jpainsymman.2008.04.008>
- Hendryadi. (2014). *Content validity (validitas isi) (Issue 01)*.
- Hetland, B., Hickman, R., McAndrew, N., & Daly, B. (2017). Factors Influencing Active Family Engagement in Care Among Critical Care Nurses. *AACN Advanced Critical Care*, 28(2), 160–170. <https://doi.org/10.4037/aacnacc2017118>
- Hidayat, A. A. A. (2007). *Metode Penelitian Keperawatan dan Teknik Analisis Data*. Salemba Medika.
- Hockenberry, M. J., & Wilson, D. (2008). *Wong's Clinical Manual of Pediatric Nursing, 8th Edition*. Elsevier Ltd.
- Holm, K., Patterson, J., & Gurney, J. (2003). Parental Involvement and Family-Centered Care in the Diagnostic and Treatment Phases of Childhood Cancer: Results from a Qualitative Study. *Journal of Pediatric Oncology Nursing: Official Journal of the Association of Pediatric Oncology Nurses*, 20, 301–313. <https://doi.org/10.1177/1043454203254984>
- Hsu, C.-C., & Sandford, B. A. (2007). The Delphi Technique: Mkaing Sense of Consensus. *Practical Assessment, Research and Evaluation*, 12(10), 1–8.
- Institute For Family Centered Care. (2003). Family-Centered Care and the Pediatrician's Role. *Pediatrics*, 112(3), 691 LP – 696. <https://doi.org/10.1542/peds.112.3.691>
- Institute for Patient and Family Centered Care. (2011). *Advancing the practice of patient- and family-centered care in hospitals - how to get started*. 20814(301), 1–29. www.ipfcc.org
- Irwanto, I. (2012). PENILAIAN KUALITAS HIDUP ANAK PENDERITA KANKER (Assessment of Quality of Life in Cancer Children). *Media Jurnal Ners*.
- Ji, Y., & Lacey, N. (2011). Experimental Transplantation and Immunotherapy Branch. *National Institutes of Health*.
- Johnson, Bev, Abraham, M., Conway, J., Simmons, L., Edgman-Levitan, S., Sodomka, P., Schlucter, J., & Ford, D. (2008). Partnering with Patients and Families to design a Patient- and Family-Centered health Care system. *Institute for Family-Centered Care*, 34. <http://www.ipfcc.org/pdf/Roadmap.pdf>
- Johnson, Beverley. (2008). Institute for Family-Centered Care. *HERD*, 1, 137–138.
- Kamus Besar Bahasa Indonesia*. (2016).
- Keeney, S., Hasson, F., & Mckenna, H. (2001). Keeney S, Hasson F, McKenna HP. A critical review of the Delphi technique as a research methodology for nursing. *Int J Nurs Stud*. 38: 195-200. *International Journal of Nursing Studies*, 38, 195–200. <https://doi.org/>

10.1016/ S0020-7489(00)00044-4

- Kementrian Kesehatan Republik Indonesia. (2016). *Profil Kesehatan Indonesia Tahun 2015*. Kementrian Kesehatan Republik Indonesia.
- Kinahan KE, et al. Scarring, disfigurement and quality of life in long-term survivors of childhood cancer: a report from the childhood cancer survivor study. *JCO*. 2012. 30:2466-2474.
- Khajeh, M., Dehghan Nayeri, N., Bahramnezhad, F., & Sadat Hoseini, A. S. (2017). Family centered care of hospitalized children: A hybrid concept analysis in Iran. *Health Promotion Perspectives*, 7(4), 210–215. <https://doi.org/10.15171/hpp.2017.37>
- Klassen, A., Gulati, S., & Dix, D. (2012). Health care providers' perspectives about working with parents of children with cancer: a qualitative study. *Journal of Pediatric Oncology Nursing : Official Journal of the Association of Pediatric Oncology Nurses*, 29(2), 92–97. <https://doi.org/10.1177/1043454212438405>
- Klick, J. C., & Ballantine, A. (2007). Providing care in chronic disease: the ever-changing balance of integrating palliative and restorative medicine. *Pediatric Clinics of North America*, 54(5), 799–812, xii. <https://doi.org/10.1016/j.pcl.2007.07.003>
- Klick, J. C., & Hauer, J. (2010). Pediatric Palliative Care. *Current Problems in Pediatric and Adolescent Health Care*, 40(6), 120–151. <https://doi.org/10.1016/j.cppeds.2010.05.001>
- Kokorelias, K., Gignac, M., Naglie, G., & Cameron, J. (2019). Towards a universal model of family centered care: A scoping review. *BMC Health Services Research*, 19. <https://doi.org/10.1186/s12913-019-4394-5>
- Labmandat Balitbangkes. (2014). Studi Diet Total (SDT) 2014-2015. In *Labmandat Balitbangkes*. <http://labdata.litbang.kemkes.go.id/riset-badan-litbangkes/menu-risikesnas/menu-rikus/392-rikus-sdt-2014>
- Lawoko, S., & Soares, J. J. F. (2003). Quality of life among parents of children with congenital heart disease, parents of children with other diseases and parents of healthy children. *Quality of Life Research : An International Journal of Quality of Life Aspects of Treatment, Care and Rehabilitation*, 12(6), 655–666. <https://doi.org/10.1023/a:1025114331419>
- Liben, S., Papadatou, D., & Wolfe, J. (2008). Paediatric palliative care: challenges and emerging ideas. *Lancet (London, England)*, 371(9615), 852–864. [https://doi.org/10.1016/S0140-6736\(07\)61203-3](https://doi.org/10.1016/S0140-6736(07)61203-3)
- Linstone, H. A., & Turoff, M. (2002). *Introduction*. <https://web.njit.edu/~turoff/pubs/delphybook/ch1.pdf>
- Liput, S. A., Kane-Gill, S. L., Seybert, A. L., & Smithburger, P. L. (2016). A Review of the Perceptions of Healthcare Providers and Family Members Toward Family Involvement in Active Adult Patient Care in the ICU. *Critical Care Medicine*, 44(6), 1191–1197. <https://doi.org/10.1097/CCM.0000000000001641>
- Loonen, J. J., Blijlevens, N. M., Prins, J., Dona, D. J., Den Hartogh, J., Senden, T., van Dulmen-Den Broeder, E., van der Velden, K., & Hermens, R. P. (2018). Cancer Survivorship Care: Person Centered Care in a Multidisciplinary Shared Care Model. *International journal of integrated care*, 18(1), 4. <https://doi.org/10.5334/ijic.3046>
- MacKay, L., & Gregory, D. (2011). Exploring Family-Centered Care Among Pediatric Oncology Nurses. *Journal of Pediatric Oncology Nursing : Official Journal of the Association of Pediatric Oncology Nurses*, 28, 43–52. <https://doi.org/10.1177/1043454210377179>
- Mainz, J. (2003). Developing evidence-based clinical indicators: a state of the art methods primer. *International Journal for Quality in Health Care : Journal of the International Society for Quality in Health Care / ISQua*, 15(1), i5–i11. <https://doi.org/10.1093/intqhc/mzg084>

- Makworo, Bwibo, & Omoni. (2016). Implementation of Family Centered Care in Child Health Nursing: Kenya Paediatric Nurses' Experiences. *Nursing & Care Open Access Journal*, 2(3), 49–51. <https://doi.org/10.15406/ncoaj.2016.01.00015>
- McConnell, B., & Moroney, T. (2015). Involving relatives in ICU patient care: critical care nursing challenges. *Journal of Clinical Nursing*, 24(7–8), 991–998. <https://doi.org/10.1111/jocn.12755>
- McGrath, P. (2002). Beginning treatment for childhood acute lymphoblastic leukemia: insights from the parents' perspective. *Oncology Nursing Forum*, 29(6), 988–996. <https://doi.org/10.1188/02.ONF.988-996>
- 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. <https://doi.org/10.1542/peds.2005-0144>
- Mitchell, M. L., Coyer, F., Kean, S., Stone, R., Murfield, J., & Dwan, T. (2016). Patient, family-centred care interventions within the adult ICU setting: An integrative review. *Australian Critical Care : Official Journal of the Confederation of Australian Critical Care Nurses*, 29(4), 179–193. <https://doi.org/10.1016/j.aucc.2016.08.002>
- Morita, T., Akechi, T., Ikenaga, M., Kizawa, Y., Kohara, H., Mukaiyama, T., Nakaho, T., Nakashima, N., Shima, Y., Matsubara, T., Fujimori, M., & Uchitomi, Y. (2004). Communication about the ending of anticancer treatment and transition to palliative care. *Annals of Oncology*, 15(10), 1551–1557. <https://doi.org/10.1093/annonc/mdh386>
- Murphy, J. F. (2010). Trends in Cancer Immunotherapy. *Clinical Medicine Insights: Oncology*, 4, CMO.S4795. <https://doi.org/10.4137/CMO.S4795>
- National Advisory Committee on Palliative Care. (2017). Report from National Policy Advisory Committee Meeting : 17 November 2017. *Palliative Care*, March, 2018.
- National Center for Cultural Competence. (2006). *A Guide for Advancing Family-Centered and Culturally and Linguistically Competent Care*. <https://nccc.georgetown.edu/documents/fccleguide.pdf>
- National Institutes of Health. (2019). *National Cancer Institute (NCI)*. National Institutes of Health. <https://www.nih.gov/about-nih/what-we-do/nih-almanac/national-cancer-institute-nci>
- Neilson, S. J., Gibson, F., & Greenfield, S. M. (2015). Pediatric Oncology Palliative Care: Experiences of General Practitioners and Bereaved Parents. *Journal of Palliative Care & Medicine*, 5(2), 214. <https://doi.org/10.4172/2165-7386.1000214>
- Ngwenya, N., Ambler, J., & Archary, M. (2019). Qualitative situational analysis of palliative care for adolescents with cancer and HIV in South Africa: healthcare worker perceptions. *BMJ Open*, 9(1), e023225. <https://doi.org/10.1136/bmjopen-2018-023225>
- Ogle, S. (2006). Clinical Application of Family Management Styles to Families of Children With Cancer. *Journal of Pediatric Oncology Nursing : Official Journal of the Association of Pediatric Oncology Nurses*, 23, 28–31. <https://doi.org/10.1177/1043454205283586>
- Olding, M., McMillan, S. E., Reeves, S., Schmitt, M. H., Puntillo, K., & Kitto, S. (2016). Patient and family involvement in adult critical and intensive care settings: a scoping review. *Health Expectations : An International Journal of Public Participation in Health Care and Health Policy*, 19(6), 1183–1202. <https://doi.org/10.1111/hex.12402>
- Pan, H. T., Wu, L. M., & Wen, S. H. (2017). Quality of Life and Its Predictors Among Children and Adolescents With Cancer. *Cancer nursing*, 40(5), 343–351. <https://doi.org/10.1097/NCC.0000000000000433>
- Perry, M., Draxkovic, I., Achterberg, T. van, Eijken, M. van, Lucassen, P., Vernooij-Dassen, M., & Rikkert, M. O. (2010). Development and Validation of Quality Indicators for Dementia Diagnosis and Management in a Primary Care Setting Diagnosis and Management in a Primary Care Setting. *J Am Geriatr Soc*, 58(3), 557–563. <https://doi.org/10.1111/j.1532-5415.2010.02726.x>

- Potter, & Perry. (2005). *Buku Ajar Fundamental Keperawatan Konsep, Proses dan Praktik* (Ed.4). EGC.
- Potts, & Mandleco. (2007). *Pediatric Nursing. Caring for Children and Their Families* (2nd ed.). Thomson.
- Purmailani. (2014). Pengaruh pendekatan family centered care terhadap penurunan kecemasan pasien anak toddler di Rumah Sakit Emanuel Klampok Banjarnegara. *Jurnal Universitas Muhammadiyah Purwokerto*.
- Raffi, F., Oskouie, F., & Shoghi, M. (2014). Caring for a child with cancer: impact on mother's health. *Asian Pacific Journal of Cancer Prevention : APJCP*, 15(4), 1731–1738. <https://doi.org/10.7314/apjcp.2014.15.4.1731>
- Riskesdas. (2013). *RISET KESEHATAN DASAR*.
- Saeui, W., Chintanadilok, N., Sriussadaporn, P., & Sanasuttipun, W. (2009). The Effects of an Empowerment Program on the Competence of Caregivers in Caring for Preschool Children with Acute Leukemia Undergoing Chemotherapy. *J Nurs Sci SI*, 27(2), 8–17.
- Saleeba, A. (2008). *Importance of Family-Centered Care in Pediatric Nursing*.
- Santo, E., Gaiva, M., Espinosa, M., Barbosa, D., & Belasco, A. (2011). Taking Care of Children With Cancer: Evaluation of the Caregivers' Burden and Quality of Life. *Revista Latino-Americana de Enfermagem*, 19, 515–522. <https://doi.org/10.1590/S0104-11692011000300010>
- Sav, A., King, M. A., Whitty, J. A., Kendall, E., McMillan, S. S., Kelly, F., Hunter, B., & Wheeler, A. J. (2015). Burden of treatment for chronic illness: a concept analysis and review of the literature. *Health Expectations : An International Journal of Public Participation in Health Care and Health Policy*, 18(3), 312–324. <https://doi.org/10.1111/hex.12046>
- Segaric, C. A., & Hall, W. A. (2015). Progressively engaging: constructing nurse, patient, and family relationships in acute care settings. *Journal of Family Nursing*, 21(1), 35–56. <https://doi.org/10.1177/1074840714564787>
- Sen-Ngam, K., Pratepchaikul, L., & Phuwathananon, K. (2011). *Maternal knowledge and practice development in caring for children with leukemia*. 23. https://www.researchgate.net/publication/279465239_Maternal_knowledge_and_practice_development_in_caring_for_children_with_leukemia
- Siegel, R. L., Miller, K. D., & Jemal, A. (2017). Cancer statistics, 2017. *CA: A Cancer Journal for Clinicians*, 67(1), 7–30. <https://doi.org/10.3322/caac.21387>
- Sloper, T., Larcombe, I. J., & Charlton, A. (1994). Psychosocial adjustment of five-year survivors of childhood cancer. *Journal of Cancer Education : The Official Journal of the American Association for Cancer Education*, 9(3), 163–169. <https://doi.org/10.1080/08858199409528300>
- Smith, J., Swallow, V., & Coyne, I. (2015). Involving parents in managing their child's long-term condition-a concept synthesis of family-centered care and partnership-in-care. *Journal of Pediatric Nursing*, 30(1), 143–159. <https://doi.org/10.1016/j.pedn.2014.10.014>
- Stienen, J. J. C., Tabbers, M. M., Benninga, M. A., Harmsen, M., & Ouwens, M. M. T. J. (2011). Development of quality indicators based on a multidisciplinary, evidence-based guideline on pediatric constipation. *Eur J Pediatr*, 170, 1513–1519. <https://doi.org/10.1007/s00431-011-1458-x>
- Sudiharto. (2007). *Asuhan Keperawatan Keluarga dengan Pendekatan Keperawatan Transkultural*. EGC.
- Supartini, Y. (2004). *Buku Ajar Konsep Dasar Keperawatan Anak*. EGC.
- Thompson, K., Dyson, G., Holland, L., & Joubert, L. (2013). An exploratory study of oncology specialists' understanding of the preferences of young people living with cancer. *Social*



- Work in Health Care*, 52(2–3), 166–190. <https://doi.org/10.1080/00981389.2012.737898>
- Van Dongen-Melman, J. E. (2000). Developing psychosocial aftercare for children surviving cancer and their families. *Acta Oncologica (Stockholm, Sweden)*, 39(1), 23–31. <https://doi.org/10.1080/028418600430932>
- Vedel, I., Ghadi, V., Lapointe, L., Routelous, C., Aegerter, P., & Guirimand, F. (2014). Patients', family caregivers', and professionals' perspectives on quality of palliative care: a qualitative study. *Palliative Medicine*, 28(9), 1128–1138. <https://doi.org/10.1177/0269216314532154>
- Wiener, L., & Pao, M. (2012). Comprehensive and Family-Centered Psychosocial Care in Pediatric Oncology: Integration of Clinical Practice and Research. In *Pediatric Psycho-Oncology* (pp. 7–17). <https://doi.org/10.1002/9781119941033.ch2>
- Wong, D. L. (2009). *Buku Ajar Keperawatan Pediatrik*. EGC.
- Woodgate, R. (2000). Part II: A critical review of qualitative research related to children's experiences with cancer. *Journal of Pediatric Oncology Nursing*, 17(4), 207–228. <https://doi.org/10.1053/jpon.2000.16397>
- Young, B., Dixon-Woods, M., Findlay, M., & Heney, D. (2002). Parenting in a crisis: conceptualising mothers of children with cancer. *Social Science & Medicine (1982)*, 55(10), 1835–1847. [https://doi.org/10.1016/s0277-9536\(01\)00318-5](https://doi.org/10.1016/s0277-9536(01)00318-5)