

Daftar Pustaka

- Abdollahpour, I., Nedjat, S., Noroozian, M., Salimi, Y., & Majdzadeh, R. (2017). Positive Aspects of Caregiving Questionnaire: A Validation Study in Caregivers of Patients with Dementia. *Journal of Geriatric Psychiatry and Neurology*, 30(2), 77–83. <https://doi.org/10.1177/0891988716686831>
- Abdollahpour, I., Nedjat, S., & Salimi, Y. (2018). Positive Aspects of Caregiving and Caregiver Burden: A Study of Caregivers of Patients With Dementia. *Journal of Geriatric Psychiatry and Neurology*, 31(1), 34–38. <https://doi.org/10.1177/0891988717743590>
- Abdullah, N. N., Idris, I. B., Shamsuddin, K., & Abdullah, N. M. A. (2019). Health-Related Quality of Life (HRQOL) of Gastrointestinal Cancer Caregivers: The Impact of Caregiving. *Asian Pacific Journal of Cancer Prevention : APJCP*, 20(4), 1191–1197. <https://doi.org/10.31557/APJCP.2019.20.4.1191>
- Adashek, J. J., & Subbiah, I. M. (2020). Caring for the caregiver: A systematic review characterising the experience of caregivers of older adults with advanced cancers. *ESMO Open*, 5(5). <https://doi.org/10.1136/esmoopen-2020-000862>
- Adelman, R. D., Tmanova, L. L., Delgado, D., Dion, S., & Lachs, M. S. (2014). Caregiver burden: A clinical review. *JAMA - Journal of the American Medical Association*, 311(10), 1052–1059. <https://doi.org/10.1001/jama.2014.304>
- Ahlam, A., Hind, M., Haddou Rahou, B., Rachid, R., & Hassan, E. (2019). Quality of life of Moroccan patients on the palliative phase of advanced cancer. *BMC Research Notes*, 12(1), 1–6. <https://doi.org/10.1186/s13104-019-4390-1>
- Aizer, A. A., Chen, M. H., McCarthy, E. P., Mendu, M. L., Koo, S., Wilhite, T. J., Graham, P. L., Choueiri, T. K., Hoffman, K. E., Martin, N. E., Hu, J. C., & Nguyen, P. L. (2013). Marital status and survival in patients with cancer. *Journal of Clinical Oncology*, 31(31), 3869–3876. <https://doi.org/10.1200/JCO.2013.49.6489>
- Akhtari-Zavare, M., Mohd-Sidik, S., Periasamy, U., Rampal, L., Fadhilah, S. I., & Mahmud, R. (2018). Determinants of quality of life among Malaysian cancer patients: a cross-sectional study. *Health and Quality of Life Outcomes*, 16(1), 163. <https://doi.org/10.1186/s12955-018-0989-5>
- Akpan-Idiok, P. A., & Anarado, A. N. (2014). Caregivers' perceptions of benefits of caregiving to advanced cancer patients attending university of Calabar teaching hospital, Calabar, Nigeria. In *International Journal of Cancer Research* (Vol. 10, Issue 2, pp. 54–64). <https://doi.org/10.3923/rjcr.2014.74.80>

- Alam, S., Hannon, B., & Zimmermann, C. (2020). Palliative Care for Family Caregivers. *Journal of Clinical Oncology*, 38(9), 926–936. <https://doi.org/10.1200/JCO.19.00018>
- Albers, G., Echteld, M. A., De Vet, H. C., Onwuteaka-Philipsen, B. D., Van Der Linden, M. H., & Deliens, L. (2010). Evaluation of quality-of-life measures for use in palliative care: A systematic review. *Palliative Medicine*, 24(1), 17–37. <https://doi.org/10.1177/0269216309346593>
- American Cancer Society. (2020). Signs and Symptoms of Cancer. *American Cancer Society*, 1–3. Retrieved from <http://www.cancer.org/cancer/cancerbasics/signs-and-symptoms-of-cancer>
- Andini, N., & Gayatri, D. (2021). Relationship between individual and disease characteristics and quality of life of cancer patients in Indonesia. *Enfermeria Clinica*, 31, S408–S412. <https://doi.org/10.1016/j.enfcli.2020.09.035>
- Aoun, S. M., Gill, F. J., Phillips, M. B., Momber, S., Cuddeford, L., Deleuil, R., Stegmann, R., Howting, D., & Lyon, M. E. (2020). The profile and support needs of parents in paediatric palliative care: comparing cancer and non-cancer groups. *Palliative Care and Social Practice*, 14(September), 1–14. <https://doi.org/10.1177/2632352420958000>
- Aoun, S. M., Grande, G., Howting, D., Deas, K., Toye, C., Troeung, L., Stajduhar, K., & Ewing, G. (2015). The Impact of the Carer Support Needs Assessment Tool (CSNAT) in Community Palliative Care Using a Stepped Wedge Cluster Trial. *PLOS ONE*, 10(4), e0123012. <https://doi.org/10.1371/journal.pone.0123012>
- Arias-Rojas, M., Arredondo Holgín, E., Carreño Moreno, S., Posada López, C., & Tellez, B. (2022). Validation of the Latin American-Spanish Version of the scale ‘Quality of Life in Life-Threatening Illness–Family Caregiver Version’ (QOLLTI-F). *Health and Social Care in the Community*, 30(3), e832–e841. <https://doi.org/10.1111/hsc.13453>
- Axelsson, L., Alvariza, A., Carlsson, N., Cohen, S. R., Sawatzky, R., & Årestedt, K. (2020). Measuring quality of life in life-threatening illness-content validity and response processes of MQOL-E and QOLLTI-F in Swedish patients and family carers. *BMC Palliative Care*, 19(1), 1–9. <https://doi.org/10.1186/s12904-020-00549-6>
- Ayalew, T. L., Wale, B. G., Haile, K. E., Zewudie, B. T., & Feleke, M. G. (2022). Health-related quality of life and associated factors among cancer patients in Ethiopia: Systematic review and meta-analysis. *PLoS ONE*, 17(12), 1–20. <https://doi.org/10.1371/journal.pone.0277839>

- Bauer, R., Sterzinger, L., Koepke, F., & Spiessl, H. (2013). Rewards of caregiving and coping strategies of caregivers of patients with mental illness. *Psychiatric Services*, 64(2), 185–188. <https://doi.org/10.1176/appi.ps.001212012>
- Bauman, J. R., Schleicher, S. M., Nipp, R., El-Jawahri, A., Pirl, W. F., Greer, J. A., & Temel, J. S. (2018). Enhancing communication between oncology care providers and patient caregivers during hospice. *The Journal of Community and Supportive Oncology*, 16(2), e72–e80. <https://doi.org/10.12788/jcso.0391>
- Bentur, N., & Resnizky, S. (2005). Validation of the McGill Quality of Life Questionnaire in home hospice settings in Israel. *Palliative Medicine*, 19(7), 538–544. <https://doi.org/10.1191/0269216305pm1052oa>
- Bergin, R., Emery, J., Bollard, R., & White, V. (2017). How rural and urban patients in Australia with colorectal or breast cancer experience choice of treatment provider: A qualitative study. *European Journal of Cancer Care*, 26(6), 1–10. <https://doi.org/10.1111/ecc.12646>
- Bergin, R. J., Emery, J. D., Bollard, R., & White, V. (2020). Comparing Pathways to Diagnosis and Treatment for Rural and Urban Patients With Colorectal or Breast Cancer: A Qualitative Study. *Journal of Rural Health*, 36(4), 517–535. <https://doi.org/10.1111/jrh.12437>
- Broese van Groenou, M. I., de Boer, A., & Iedema, J. (2013). Positive and negative evaluation of caregiving among three different types of informal care relationships. *European Journal of Ageing*, 10(4), 301–311. <https://doi.org/10.1007/s10433-013-0276-6>
- Bužngová, R., Kozáková, R., & Sikorová, L. (2015). Assessment of Quality of Life of Family Members of Inpatients with End-Stage Disease. *Journal of Palliative Care*, 31(4), 250–257. <https://doi.org/10.1177/082585971503100407>
- Cancer Council. (2021). *Understanding Palliative Care*. Retrieved from <https://www.cancer.org.au/assets/pdf/understanding-palliative-care-booklet>
- Candy, B., Jones, L., Drake, R., Leurent, B., & King, M. (2011). Interventions for supporting informal caregivers of patients in the terminal phase of a disease. *Cochrane Database of Systematic Reviews*, 15(6), CD007617. <https://doi.org/10.1002/14651858.CD007617.pub2>
- Caputo, J., Pavalko, E. K., & Hardy, M. A. (2016). The Long-Term Effects of Caregiving on Women's Health and Mortality. *Journal of Marriage and Family*, 78(5), 1382–1398. <https://doi.org/10.1111/jomf.12332>

- Carbonneau, H., Caron, C. D., & Desrosiers, J. (2011). Effects of an adapted leisure education program as a means of support for caregivers of people with dementia. *Archives of Gerontology and Geriatrics*, 53(1), 31–39. <https://doi.org/10.1016/j.archger.2010.06.009>
- Chen, M. C., Chen, K. M., & Chu, T. P. (2015). Caregiver Burden, Health Status, and Learned Resourcefulness of Older Caregivers. *Western Journal of Nursing Research*, 37(6), 767–780. <https://doi.org/10.1177/0193945914525280>
- Cheng, S. T., Lam, L. C. W., Kwok, T., Ng, N. S. S., & Fung, A. W. T. (2013). The social networks of hong kong chinese family caregivers of alzheimer's disease: Correlates with positive gains and burden. *Gerontologist*, 53(6), 998–1008. <https://doi.org/10.1093/geront/gns195>
- Cherny, N. I., Paluch-Shimon, S., & Berner-Wygoda, Y. (2018). Palliative care: Needs of advanced breast cancer patients. *Breast Cancer: Targets and Therapy*, 10, 231–243. <https://doi.org/10.2147/BCTT.S160462>
- Choi, J. Y., Donahoe, M. P., & Hoffman, L. A. (2016). Psychological and physical health in family caregivers of intensive care unit survivors: Current knowledge and future research strategies. *Journal of Korean Academy of Nursing*, 46(2), 159–167. <https://doi.org/10.4040/jkan.2016.46.2.159>
- Choi, J. Y., Son, Y. J., & Tate, J. A. (2019). Exploring positive aspects of caregiving in family caregivers of adult ICU survivors from ICU to four months post-ICU discharge. *Heart and Lung*, 48(6), 553–559. <https://doi.org/10.1016/j.hrtlng.2019.09.001>
- Choi, Y. S., Hwang, S. W., Hwang, I. C., Lee, Y. J., Kim, Y. S., Kim, H. M., Youn, C. H., Ahn, H. Y., & Koh, S.-J. (2016). Factors Associated with Quality of Life Among Family Caregivers of Terminally Ill Cancer Patients. *Psycho-Oncology*, 25(2), 217–224. <https://doi.org/10.1002/pon.3904>
- Cohen, C. A., Colantonio, A., & Vernich, L. (2002). Positive aspects of caregiving: Rounding out the caregiver experience. *International Journal of Geriatric Psychiatry*, 17(2), 184–188. <https://doi.org/10.1002/gps.561>
- Cohen, R., Leis, A. M., Kuhl, D., Charbonneau, C., Ritvo, P., & Ashbury, F. D. (2006). QOLLTI-F: Measuring family carer quality of life. *Palliative Medicine*, 20(8), 755–767. <https://doi.org/10.1177/0269216306072764>
- Cohen, S. R., Sawatzky, R., Russell, L. B., Shahidi, J., Heyland, D. K., & Gadermann, A. M. (2016). Measuring the quality of life of people at the end of life: The McGill Quality of Life Questionnaire-Revised. *Palliative Medicine*, 31(2), 120–129. <https://doi.org/10.1177/0269216316659603>

- Connor, S., Morris, C., Jaramillo, E., Harding, R., Cleary, J., Haste, B.,... Knaul, F. (2020). Global Atlas of Palliative Care. In *The Worldwide Palliative Care Alliance*. The Worldwide Palliative Care Alliance. [https://cdn.who.int/media/docs/default-source/integrated-health-services-\(ihs\)/csy/palliative-care/whpca_global_atlas_p5_digital_final.pdf?sfvrsn=1b54423a_3](https://cdn.who.int/media/docs/default-source/integrated-health-services-(ihs)/csy/palliative-care/whpca_global_atlas_p5_digital_final.pdf?sfvrsn=1b54423a_3)
- Coolbrandt, A., Sterckx, W., Clement, P., Borgenon, S., Decruyenaere, M., De Vleeschouwer, S., Mees, A., & Dierckx De Casterlé, B. (2015). Family Caregivers of Patients with a High-Grade Glioma: A Qualitative Study of Their Lived Experience and Needs Related to Professional Care. *Cancer Nursing*, 38(5), 406–413. <https://doi.org/10.1097/NCC.0000000000000216>
- Cooper, B., Kinsella, G. J., & Picton, C. (2006). Development and initial validation of a family appraisal of caregiving questionnaire for palliative care. *Psycho-Oncology*, 15(7), 613–622. <https://doi.org/10.1002/pon.1001>
- Cui, J., Song, L. J., Zhou, L. J., Meng, H., & Zhao, J. J. (2014). Needs of family caregivers of advanced cancer patients: A survey in Shanghai of China. *European Journal of Cancer Care*, 23(4), 562–569. <https://doi.org/10.1111/ecc.12174>
- Dadsetan, F., Shahrababaki, P. M., Mirzai, M., & Nouhi, E. (2021). Palliative care needs of patients with multiple sclerosis in southeast Iran. *BMC Palliative Care*, 20(1), 1–7. <https://doi.org/10.1186/s12904-021-00867-3>
- Dahlan, M. S. (2015). *Statistik Untuk Kedokteran dan Kesehatan Deskriptif, Bivariat, dan Multivariat Dilengkapi Aplikasi Menggunakan SPSS* (6th ed.). Epidemiologi Indonesia.
- Dahlan, M. S. (2016). *Langkah-Langkah Membuat Proposal Penelitian Bidang Kedokteran dan Kesehatan*. Sagung Seto.
- Dahlan, M. S. (2019). *Besar Sampel Dalam Penelitian Kedokteran dan Kesehatan* (5th ed.). Sagung Seto.
- Damen, A., Raijmakers, N. J. H., van Roij, J., Visser, A., Beuken-Everdingen, M. van den, Kuip, E., van Laarhoven, H. W. M., van Leeuwen-Snoeks, L., van der Padt-Pruijsten, A., Smilde, T. J., Leget, C., & Fitchett, G. (2022). Spiritual Well-Being and Associated Factors in Dutch Patients With Advanced Cancer. *Journal of Pain and Symptom Management*, 63(3), 404–414. <https://doi.org/10.1016/j.jpainsymman.2021.10.004>
- de Labra, C., Millán-Calenti, J. C., Buján, A., Núñez-Naveira, L., Jensen, A. M., Peersen, M. C., Mojs, E., Samborski, W., & Maseda, A. (2015). Predictors of caregiving satisfaction in informal caregivers of people with dementia. *Archives of Gerontology and Geriatrics*, 60(3), 380–388.

<https://doi.org/10.1016/j.archger.2015.03.002>

Deshields, T. L., Rihanek, A., Potter, P., Zhang, Q., Kuhrik, M., Kuhrik, N., & O'Neill, J. (2012). Psychosocial aspects of caregiving: Perceptions of cancer patients and family caregivers. *Supportive Care in Cancer*, 20(2), 349–356. <https://doi.org/10.1007/s00520-011-1092-1>

Dionne-Odom, J. N., Azuero, A., Taylor, R. A., Wells, R. D., Hendricks, B. A., Bechthold, A. C., Reed, R. D., Harrell, E. R., Dosse, C. K., Engler, S., McKie, P., Ejem, D., Bakitas, M. A., & Rosenberg, A. R. (2021). Resilience, preparedness, and distress among family caregivers of patients with advanced cancer. *Supportive Care in Cancer*, 29(11), 6913–6920. <https://doi.org/10.1007/s00520-021-06265-y>

Dogbey, D. M., Burger, H., Edge, J., Mihalik, M., & Savieri, P. (2022). Identification of Palliative Care Needs in Cancer Patients in a Surgical Emergency Center. *Journal of Pain and Symptom Management*, 63(2), 260–270. <https://doi.org/10.1016/j.jpainsymman.2021.08.008>

Effendy, C., Vernooij-Dassen, M., Setiyarini, S., Kristanti, M. S., Tejawinata, S., Vissers, K., & Engels, Y. (2015). Family caregivers' involvement in caring for a hospitalized patient with cancer and their quality of life in a country with strong family bonds. *Psycho-Oncology*, 24(5), 585–591. <https://doi.org/10.1002/pon.3701>

Effendy, C., Vissers, K., Tejawinata, S., Ecu, P., Vernooij-dassen, M., & Engels, Y. (2014). Dealing with Symptoms and Issues of Hospitalized Patients with Cancer in Indonesia: The Role of Families, Nurses, and Physicians. *World Institute of Pain*, 15(5), 441–446. <https://doi.org/10.1111/papr.12203>

EJ, S., Varghese, Y., DSilva, F., Bhat, S., Sukseh, & Nalini, M. (2023). Factors Associated with Caregivers Burden of Breast Cancer Patients – Review. *HIV Nursing*, 23(3), 340–346. <https://www.hivnursing.net/index.php/hiv/article/view/1721>

Epps, F. (2015). A Closer Look at Religiosity Among Family Caregivers. *Journal of Religion, Spirituality and Aging*, 27(2–3), 166–182. <https://doi.org/10.1080/15528030.2014.971141>

Essadi, I., Kaakoua, M., Lalya, I., Bejjou, F. E., Zyani, M., & Belbaraka, R. (2021). Factors influencing Quality of life in Moroccan patients with cancer: A monocentric Prospective study. *The Gulf Journal of Oncology*, 1(36), 30–35.

Fayers, P. M., & Machin, D. (2016). *Quality of Life: The Assessment, Analysis and Reporting of Patient-reported Outcomes* (3rd ed.). John Wiley & Sons, Ltd. <https://doi.org/10.1002/9781118758991>

- Fereidouni, A., Ebadi, A., Rassouli, M., Hosseini, S. M., Mollahadi, M., Khorshidvand, A., Javid, M., Ansari, B., Rezaei, M. S., & Barasteh, S. (2022). Psychometric Properties of the “Quality of Life in Life-Threatening Illness—Family Carer Version” (QOLLTI-F) in Persian-Speaking Carers of COVID-19 Patients. *Frontiers in Psychology*, 13(May), 1–8. <https://doi.org/10.3389/fpsyg.2022.838074>
- Ferrell, B., Hanson, J., & Grant, M. (2013). An overview and evaluation of the oncology family caregiver project: improving quality of life and quality of care for oncology family caregivers. *Psycho-Oncology*, 22(7), 1645–1652. <https://doi.org/10.1002/pon.3198>
- Franchini, L., Ercolani, G., Ostan, R., Raccichini, M., Samolsky-Dekel, A., Malerba, M. B., Melis, A., Varani, S., & Pannuti, R. (2020). Caregivers in home palliative care: gender, psychological aspects, and patient’s functional status as main predictors for their quality of life. *Supportive Care in Cancer*, 28(7), 3227–3235. <https://doi.org/10.1007/s00520-019-05155-8>
- Fu, W., Li, J., Fang, F., Zhao, D., Hao, W., & Li, S. (2021). Subjective burdens among informal caregivers of critically ill patients: a cross-sectional study in rural Shandong, China. *BMC Palliative Care*, 20(1), 1–11. <https://doi.org/10.1186/s12904-021-00858-4>
- Gan, G. G., Tey, K. W. F., Mat, S., Saad, M., Bee, P. C., Malik, R. A., Ho, G. F., & Ng, C. G. (2022). Quality of Life of Family Caregivers of Cancer Patients in a Developing Nation. *Asian Pacific Journal of Cancer Prevention*, 23(11), 3905–3914. <https://doi.org/10.31557/APJCP.2022.23.11.3905>
- Ganesh Kumar, S., Majumdar, A., & Pavithra, G. (2014). Quality of Life and Its Associated Factors Using WHOQOL- BREF Among Elderly in Urban Puducherry, India. *Journal of Clinical and Diagnostic Research*, 8(1), 54–57. <https://doi.org/10.7860/JCDR/2014/6996.3917>
- García-Mochón, L., Peña-Longobardo, L. M., Río-Lozano, M. Del, Oliva-Moreno, J., Larrañaga-Padilla, I., & Del Mar García-Calvente, M. (2019). Determinants of burden and satisfaction in informal caregivers: Two sides of the same coin? the CUIDAR-SE study. *International Journal of Environmental Research and Public Health*, 16(22), 4378. <https://doi.org/10.3390/ijerph16224378>
- Girgis, A., Lambert, S., Johnson, C., Waller, A., & Currow, D. (2013). Physical, psychosocial, relationship, and economic burden of caring for people with cancer: A review. *Journal of Oncology Practice*, 9(4), 197–202. <https://doi.org/10.1200/JOP.2012.000690>
- Given, B. A., Given, C. W., & Sherwood, P. (2012). The challenge of quality cancer care for family caregivers. *Seminars in Oncology Nursing*, 28(4), 205–212. <https://doi.org/10.1016/j.soncn.2012.09.002>

- Glajchen, M. (2004). The emerging role and needs of family caregivers in cancer care. *The Journal of Supportive Oncology*, 2(2), 145–155.
- Goswami, S., & Gupta, S. S. (2020). How cancer of oral cavity affects the family caregivers? – A cross-sectional study in Wardha, India, using the Caregiver Quality of Life Index – Cancer questionnaire. *South Asian Journal of Cancer*, 09(01), 62–65. https://doi.org/10.4103/sajc.sajc_331_18
- Gupta, S., Isherwood, G., Jones, K., & Van Impe, K. (2015). Assessing health status in informal schizophrenia caregivers compared with health status in non-caregivers and caregivers of other conditions. *BMC Psychiatry*, 15(1), 1–11. <https://doi.org/10.1186/s12888-015-0547-1>
- Haan, M. M., Olthuis, G., & van Gorp, J. L. P. (2021). Feeling called to care: a qualitative interview study on normativity in family caregivers' experiences in Dutch home settings in a palliative care context. *BMC Palliative Care*, 20(1), 1–15. <https://doi.org/10.1186/s12904-021-00868-2>
- Hacialioglu, N., Özer, N., Yilmaz karabulutlu, E., Erdem, N., & Erci, B. (2010). The quality of life of family caregivers of cancer patients in the East of Turkey. *European Journal of Oncology Nursing*, 14(3), 211–217. <https://doi.org/10.1016/j.ejon.2010.01.017>
- Han, K.-T., Park, E.-C., Kim, J.-H., Kim, S. J., & Park, S. (2014). Is marital status associated with quality of life? *Health and Quality of Life Outcomes*, 12, 109. <https://doi.org/10.1186/s12955-014-0109-0>
- Hansen, T., & Slagsvold, B. (2015). Feeling the squeeze? The effects of combining work and informal caregiving on psychological well-being. *European Journal of Ageing*, 12(1), 51–60. <https://doi.org/10.1007/s10433-014-0315-y>
- Haraldstad, K., Wahl, A., Andenæs, R., Andersen, J. R., Andersen, M. H., Beisland, E., Borge, C. R., Engebretsen, E., Eisemann, M., Halvorsrud, L., Hanssen, T. A., Haugstvedt, A., Haugland, T., Johansen, V. A., Larsen, M. H., Løvereide, L., Løyland, B., Kvarme, L. G., Moons, P., ... Helseth, S. (2019). A systematic review of quality of life research in medicine and health sciences. *Quality of Life Research: An International Journal of Quality of Life Aspects of Treatment, Care and Rehabilitation*, 28(10), 2641–2650. <https://doi.org/10.1007/s11136-019-02214-9>
- Hasan, R. Z., Ng, B. K., Phon, S. E., Karim, A. K. A., Lim, P. S., & Azurah, A. G. N. (2022). Quality of Life of Gynaecology Patients and Family Caregivers. *International Journal of Environmental Research and Public Health*, 19(11). <https://doi.org/10.3390/ijerph19116450>

- Haun, M. W., Sklenarova, H., Brechtel, A., Herzog, W., & Hartmann, M. (2014). Distress in cancer patients and their caregivers and association with the caregivers' perception of dyadic communication. *Oncology Research and Treatment*, 37(7–8), 384–388. <https://doi.org/10.1159/000364885>
- Heale, R., & Twycross, A. (2015). Validity and reliability in quantitative studies. *Evidence-Based Nursing*, 18(3), 66–67. <https://doi.org/10.1136/eb-2015-102129>
- Hidayat, A. A. A. (2009). *Metode penelitian keperawatan dan teknik analisis data*. Salemba Medika.
- Higginson, I. J., Yi, D., Johnston, B. M., Ryan, K., McQuillan, R., Selman, L., Pantilat, S. Z., Daveson, B. A., Morrison, R. S., & Normand, C. (2020). Associations between informal care costs, care quality, carer rewards, burden and subsequent grief: the international, access, rights and empowerment mortality follow-back study of the last 3 months of life (IARE I study). *BMC Medicine*, 18(1), 1–13. <https://doi.org/10.1186/s12916-020-01768-7>
- Holgín, E. A., Arias-Rojas, M., & Moreno, S. C. (2021). Quality of life of family caregivers of people with cancer in palliative care. *Revista Da Escola de Enfermagem*, 55, 1–8. <https://doi.org/10.1590/S1980-220X2020015103740>
- Hsu, T., Loscalzo, M., Ramani, R., Forman, S., Popplewell, L., Clark, K., Katheria, V., Feng, T., Strowbridge, R., Rinehart, R., Smith, D., Matthews, K., Dillehunt, J., & Hurria, A. (2014). Factors associated with high burden in caregivers of older adults with cancer. *Cancer*, 120(18), 2927–2935. <https://doi.org/10.1002/cncr.28765>
- Imaiso, J., Tsukasaki, K., & Okoshi, F. (2012). Differences in Home-based Family Caregiving Appraisal for Caregivers of the Elderly in Rural and Urban Japanese Communities. *Journal of Community Health Nursing*, 29(1), 25–38. <https://doi.org/10.1080/07370016.2012.645733>
- Irfan, B., Irfan, O., Ansari, A., Qidwai, W., & Nanji, K. (2017). Impact of Caregiving on Various Aspects of the Lives of Caregivers. *Cureus*, 9(5), e1213. <https://doi.org/10.7759/cureus.1213>
- Janssen, D. J. A., Spruit, M. A., Wouters, E. F. M., & Schols, J. M. G. A. (2012). Family Caregiving in Advanced Chronic Organ Failure. *Journal of the American Medical Directors Association*, 13(4), 394–399. <https://doi.org/10.1016/j.jamda.2011.04.017>
- Jones, P. S., Lee, J. W., Phillips, L. R., Zhang, X. E., & Jaceldo, K. B. (2001). An Adaptation of Brislin's Translation Model for Cross-cultural Research. *Nursing Research*, 50(5), 300–304. <https://doi.org/10.1097/00006199-200109000-00008>

- Keir, S. T., Farland, M. M., Lipp, E. S., & Friedman, H. S. (2009). Family appraisal of caregiving in a brain cancer model. *Journal of Hospice and Palliative Nursing*, 11(1), 60–66. <https://doi.org/10.1097/NJH.0b013e3181917e35>
- Kemenkes. (2017). Petunjuk Teknis: Paliatif Kanker Pada Dewasa. In *Kemenkes RI*. Retrieved from http://p2ptm.kemkes.go.id/uploads/VHcrbkVobjRzUDN3UCs4eUJ0dVBndz09/2017/08/PETUNJUK_TEKNIS_PALIATIF_KANKER_PADA_DEWASA.pdf
- Kemenkes. (2019). Kementerian Kesehatan Republik Indonesia. In *Kementerian Kesehatan RI*. Kementerian Kesehatan Republik Indonesia.
- Kementerian Kesehatan Republik Indonesia. (2015). Pedoman Nasional Program Paliatif Kanker. In *Kementerian Kesehatan Republik Indonesia*.
- Kent, E. E., Lee, S., Asad, S., Dobbins, E. E., Aimone, E. V., & Park, E. M. (2023). “If I wasn’t in a rural area, I would definitely have more support”: social needs identified by rural cancer caregivers and hospital staff. *Journal of Psychosocial Oncology*, 41(4), 393–410. <https://doi.org/10.1080/07347332.2022.2129547>
- Kent, E. E., Rowland, J. H., Northouse, L., Litzelman, K., Chou, W.-Y. S., Shelburne, N., Timura, C., O’Mara, A., & Huss, K. (2016). Caring for caregivers and patients: Research and clinical priorities for informal cancer caregiving. *Cancer*, 122(13), 1987–1995. <https://doi.org/10.1002/cncr.29939>
- Kilic, S. T., & Oz, F. (2019). Family caregivers’ involvement in caring with cancer and their quality of life. *Asian Pacific Journal of Cancer Prevention*, 20(6), 1735–1741. <https://doi.org/10.31557/APJCP.2019.20.6.1735>
- Kim, S. H., Gu, S. K., Yun, Y. H., Lee, C. G., Choi, Y. S., Lee, W. S., Kim, S. Y., & Heo, D. S. (2007). Validation study of the Korean version of the McGill Quality of Life Questionnaire. *Palliative Medicine*, 21(5), 441–447. <https://doi.org/10.1177/0269216307079816>
- Kim, Y., Shaffer, K. M., Carver, C. S., & Cannady, R. S. (2016). Quality of life of family caregivers 8 years after a relative’s cancer diagnosis: Follow-up of the National Quality of Life Survey for Caregivers. *Psycho-Oncology*, 25(3), 266–274. <https://doi.org/10.1002/pon.3843>
- Kumar, K., Mittal, A., Kamboj, D., Goel, G. S., Mittal, A., & Garg, S. K. (2023). Factors Affecting Quality of Life of Cancer Patients in North India. *Asian Pacific Journal of Cancer Prevention: APJCP*, 24(2), 641–648. <https://doi.org/10.31557/APJCP.2023.24.2.641>

- Kung'u, M., Onsongo, L., & Ogutu, J. O. (2022). Factors influencing quality of life among cancer survivors in Kenya. *African Health Sciences*, 22(4), 87–95. <https://doi.org/10.4314/ahs.v22i4.12>
- Kunkle, R., Chaperon, C., & Berger, A. M. (2021). Formal Caregiver Burden in Nursing Homes: An Integrative Review. *Western Journal of Nursing Research*, 43(9), 877–893. <https://doi.org/10.1177/0193945920979691>
- Lai, Y.-H. Y.-H., Liao, C.-T. C.-T., Huang, B.-S. B.-S., Lin, C.-Y. C.-Y., Fan, K.-H. K.-H., Chang, J. T.-C. T.-C., Chen, S.-C., Lai, Y.-H. Y.-H., Liao, C.-T. C.-T., Huang, B.-S. B.-S., Lin, C.-Y. C.-Y., Fan, K.-H. K.-H., & Chang, J. T.-C. T.-C. (2014). Unmet supportive care needs and characteristics of family caregivers of patients with oral cancer after surgery. *Psycho-Oncology*, 577(January), 569–577.
- Laidsaar-Powell, R. C., Butow, P. N., Bu, S., Charles, C., Gafni, A., Lam, W. W. T., Jansen, J., McCaffery, K. J., Shepherd, H. L., Tattersall, M. H. N., & Juraskova, I. (2013). Physician-patient-companion communication and decision-making: A systematic review of triadic medical consultations. *Patient Education and Counseling*, 91(1), 3–13. <https://doi.org/10.1016/j.pec.2012.11.007>
- Lam, K., Chow, E., Zhang, L., Wong, E., Bedard, G., Fairchild, A., Vassiliou, V., El-Din, M. A., Jesus-Garcia, R., Kumar, A., Forges, F., Tseng, L. M., Hou, M. F., Chie, W. C., & Bottomley, A. (2013). Determinants of quality of life in advanced cancer patients with bone metastases undergoing palliative radiation treatment. *Supportive Care in Cancer*, 21(11), 3021–3030. <https://doi.org/10.1007/s00520-013-1876-6>
- Lambert, S. D., Harrison, J. D., Smith, E., Bonevski, B., Carey, M., Laws, C., Paul, C., & Girgis, A. (2012). The unmet needs of partners and caregivers of adults diagnosed with cancer: A systematic review. *BMJ Supportive and Palliative Care*, 2(3), 224–230. <https://doi.org/10.1136/bmjspcare-2012-000226>
- Lamore, K., Montalescot, L., & Untas, A. (2017). Treatment decision-making in chronic diseases: What are the family members' roles, needs and attitudes? A systematic review. *Patient Education and Counseling*, 100(12), 2172–2181. <https://doi.org/10.1016/j.pec.2017.08.003>
- Lapau, B. (2015). *Metode penelitian kesehatan: metode ilmiah penulisan skripsi, tesis, dan disertasi* (2nd ed.). Yayasan Pustaka Obor Indonesia.
- Lee, J., Yoo, M. S., & Jung, D. (2010). Caregiving appraisal of family caregivers for older stroke patients in Korea. *International Nursing Review*, 57(1), 107–112. <https://doi.org/10.1111/j.1466-7657.2009.00751.x>

- Lee, K. C., Yiin, J. J., & Chao, Y. F. (2016). Effect of integrated caregiver support on caregiver burden for people taking care of people with cancer at the end of life: A cohort and quasi-experimental clinical trial. *International Journal of Nursing Studies*, 56(April), 17–26. <https://doi.org/10.1016/j.ijnurstu.2016.01.002>
- Leggett, A. N., Meyer, O. L., Bugajski, B. C., & Polenick, C. A. (2020). Accentuate the Positive: The Association Between Informal and Formal Supports and Caregiving Gains. *Journal of Applied Gerontology*, 40(7), 763–771. <https://doi.org/10.1177/0733464820914481>
- Li, Q., & Loke, A. Y. (2013). The positive aspects of caregiving for cancer patients: A critical review of the literature and directions for future research. *Psycho-Oncology*, 22(11), 2399–2407. <https://doi.org/10.1002/pon.3311>
- Li, Q., Xu, Y., Zhou, H., & Loke, A. Y. (2016). Factors Influencing The Health-Related Quality of Life of Chinese Advanced Cancer Patients and Their Spousal Caregivers: A Cross-Sectional Study. *BMC Palliative Care*, 15(1), 1–14. <https://doi.org/10.1186/s12904-016-0142-3>
- Lim, H. A., Tan, J. Y., Chua, J., Yoong, R. K., Lim, S. E., Kua, E. H., & Mahendran, R. (2017). Quality of Life of Family Caregivers of Cancer Patients in Singapore and Globally. *Singapore Medical Journal*, 58(5), 258–261. <https://doi.org/10.11622/smedj.2016083>
- Liu, H. Y., & Huang, L. H. (2018). The relationship between family functioning and caregiving appraisal of dementia family caregivers: caregiving self-efficacy as a mediator. *Aging and Mental Health*, 22(4), 558–567. <https://doi.org/10.1080/13607863.2016.1269148>
- Liu, Z., Heffernan, C., & Tan, J. (2020). Caregiver burden: A concept analysis. *International Journal of Nursing Sciences*, 7(4), 438–445. <https://doi.org/10.1016/j.ijnss.2020.07.012>
- Lo, R. S. K., Woo, J., Zhoc, K. C. H., Li, C. Y. P., Yeo, W., Johnson, P., Mak, Y., & Lee, J. (2001). Cross-cultural validation of the McGill quality of life questionnaire in Hong Kong Chinese. *Palliative Medicine*, 15(5), 387–397. <https://doi.org/10.1191/026921601680419438>
- Lobb, E. A., Bindley, K., Sanderson, C., MacLeod, R., & Mowll, J. (2019). Navigating the path to care and death at home—it is not always smooth: a qualitative examination of the experiences of bereaved family caregivers in palliative care. *Journal of Psychosocial Oncology Research & Practice*, 1(1), e3. <https://doi.org/10.1097/OR9.0000000000000003>

- McCaffrey, N., Bradley, S., Ratcliffe, J., & Currow, D. C. (2016). What Aspects of Quality of Life Are Important From Palliative Care Patients' Perspectives? A Systematic Review of Qualitative Research. *Journal of Pain and Symptom Management*, 52(2), 318–328.e5. <https://doi.org/10.1016/j.jpainsymman.2016.02.012>
- McCauley, R., McQuillan, R., Ryan, K., & Foley, G. (2021). Mutual support between patients and family caregivers in palliative care: A systematic review and narrative synthesis. *Palliative Medicine*, 35(5), 875–885. <https://doi.org/10.1177/0269216321999962>
- McDonald, J., Swami, N., Pope, A., Hales, S., Nissim, R., Rodin, G., Hannon, B., & Zimmermann, C. (2018). Caregiver quality of life in advanced cancer: Qualitative results from a trial of early palliative care. *Palliative Medicine*, 32(1), 69–78. <https://doi.org/10.1177/0269216317739806>
- Meecharoen, W., Sirapo-ngam, Y., Monkong, S., Oratai, P., & Northouse L., L. (2013). Factors Influencing Quality of Life among Family Caregivers of Patients with Advanced Cancer: A Causal Model. *Pacific Rim International Journal of Nursing Research*, 17(4), 304–316. <http://search.ebscohost.com/login.aspx?direct=true&db=jlh&AN=2012359696&site=ehost-live>
- Mehta, A., Cohen, S. R., Ezer, H., Carnevale, F. A., & Ducharme, F. (2011). Striving to respond to palliative care patients' pain at home: A puzzle for family caregivers. *Oncology Nursing Forum*, 38(1), 37–45. <https://doi.org/10.1188/11.ONF.E37-E45>
- Miyamoto, Y., Tachimori, H., & Ito, H. (2010). Formal Caregiver Burden in Dementia: Impact of Behavioral and Psychological Symptoms of Dementia and Activities of Daily Living. *Geriatric Nursing*, 31(4), 246–253. <https://doi.org/10.1016/j.gerinurse.2010.01.002>
- Morishita-Kawahara, M., Koike, K., Kawahara, T., & Kamibeppu, K. (2022). The relationship between satisfaction with care and quality of life among family caregivers of patients living with terminal cancer in general wards: A cross-sectional study. *International Journal of Palliative Nursing*, 28(12), 575–582. <https://doi.org/10.12968/ijpn.2022.28.12.575>
- Mosher, C. E., Bakas, T., & Champion, V. L. (2013). Physical health, mental health, and life changes among family caregivers of patients with lung cancer. *Oncology Nursing Forum*, 40(1), 53–61. <https://doi.org/10.1188/13.ONF.53-61>
- Mosquera, I., Vergara, I., Larrañaga, I., Machón, M., del Río, M., & Calderón, C. (2016). Measuring the impact of informal elderly caregiving: a systematic review of tools. *Quality of Life Research*, 25(5), 1059–1092.

<https://doi.org/10.1007/s11136-015-1159-4>

Nakken, N., Spruit, M. A., Wouters, E. F. M., Schols, J. M. G. A., & Janssen, D. J. A. (2015). Family caregiving during 1-year follow-up in individuals with advanced chronic organ failure. *Scandinavian Journal of Caring Sciences*, 29(4), 734–744. <https://doi.org/10.1111/scs.12204>

National Cancer Institute. (2015). *Cancer Staging*. Retrieved from <https://www.cancer.gov/about-cancer/diagnosis-staging/staging>

National Cancer Institute. (2019). *Symptoms of cancer*. 1–2. Retrieved from <https://www.cancer.gov/about-cancer/diagnosis-staging/symptoms>

National Cancer Institute. (2021). *Palliative Care in Cancer*. 1–4. Retrieved from <https://www.cancer.gov/about-cancer/advanced-cancer/care-choices/palliative-care-fact-sheet>

Nayak, M., George, A., Vidyasagar, M., Mathew, S., Nayak, S., Nayak, B., Shashidhara, Y., & Kamath, A. (2017). Quality of life among cancer patients. *Indian Journal of Palliative Care*, 23(4), 445. https://doi.org/10.4103/IJPC.IJPC_82_17

Nguyen, H. T. H., Duong, K. L., Nguyen, S. T., Trinh, Q., Hoang, H. T. L., Phung, T. Q., Lin, H. W., & Nguyen, H. T. L. (2022). Quality of Life and Its Associated Factors Among Cancer Patients Receiving Chemotherapy at Oncology Hospitals in Vietnam After the Third Wave of the COVID-19 Pandemic. *Cancer Management and Research*, 14(July), 2429–2444. <https://doi.org/10.2147/CMAR.S370034>

Norinder, M., Årestedt, K., Lind, S., Axelsson, L., Grande, G., Ewing, G., Holm, M., Öhlén, J., Benkel, I., & Alvariza, A. (2021). Higher levels of unmet support needs in spouses are associated with poorer quality of life – a descriptive cross-sectional study in the context of palliative home care. *BMC Palliative Care*, 20(1), 1–11. <https://doi.org/10.1186/s12904-021-00829-9>

Oechsle, K. (2019). Current Advances in Palliative & Hospice Care: Problems and Needs of Relatives and Family Caregivers During Palliative and Hospice Care—An Overview of Current Literature. *Medical Sciences*, 7(3), 43. <https://doi.org/10.3390/medsci7030043>

Oncology Nursing Society. (2019). *Caregiver strain and burden*. Retrieved from https://www.ons.org/pep/caregiver-strain-and-burden?display=pepnavigator&sort_by=created&items_per_page=50

Park, J. H., Jung, Y. S., Kim, J. Y., & Bae, S. H. (2021). Determinants of quality of life in women immediately following the completion of primary treatment of breast cancer: A cross-sectional study. *PLoS ONE*, 16(10), 1–13.

<https://doi.org/10.1371/journal.pone.0258447>

- Pinquart, M., & Sörensen, S. (2003). Associations of stressors and uplifts of caregiving with caregiver burden and depressive mood: A meta-analysis. *Journals of Gerontology - Series B Psychological Sciences and Social Sciences*, 58(2), 112–128. <https://doi.org/10.1093/geronb/58.2.P112>
- Polit, D. F., & Beck, heryl T. (2006). The content validity index: Are you sure you know what's being reported? critique and recommendations. *Research in Nursing & Health*, 29(5), 489–497. <https://doi.org/10.1002/nur.20147>
- Prince, M., Brodaty, H., Uwakwe, R., Acosta, D., Ferri, C. P., Guerra, M., Huang, Y., Jacob, K., Llibre Rodriguez, J. J., Salas, A., Sosa, A. L., Williams, J. D., Jotheeswaran, A. T., & Liu, Z. (2012). Strain and its correlates among carers of people with dementia in low-income and middle-income countries. A 10/66 Dementia Research Group population-based survey. *International Journal of Geriatric Psychiatry*, 27(7), 670–682. <https://doi.org/10.1002/gps.2727>
- Ramasubbu, S. K., Pasricha, R. K., Nath, U. K., Rawat, V. S., & Das, B. (2021). Quality of life and factors affecting it in adult cancer patients undergoing cancer chemotherapy in a tertiary care hospital. *Cancer Reports*, 4(2), 1–11. <https://doi.org/10.1002/cnr2.1312>
- Ran, M., Peng, L., Liu, Q., Pender, M., He, F., & Wang, H. (2018). The association between quality of life(QOL) and health literacy among junior middle school students: a cross-sectional study. *BMC Public Health*, 18(1), 1183. <https://doi.org/10.1186/s12889-018-6082-5>
- Reeve, B. B., Cai, J., Zhang, H., Weissler, M. C., Wisniewski, K., Gross, H., & Olshan, A. F. (2016). Factors That Impact Health-Related Quality of Life Over Time For Individuals with Head and Neck Cancer. *Laryngoscope*, 126(12), 2718–2725. <https://doi.org/10.1002/lary.26073>
- Rha, S. Y., Park, Y., Song, S. K., Lee, C. E., & Lee, J. (2015). Caregiving Burden and The Quality of Life of Family Caregivers of Cancer Patients: The Relationship and Correlates. *European Journal of Oncology Nursing*, 19(4), 376–382. <https://doi.org/10.1016/j.ejon.2015.01.004>
- Rochmawati, E., Wiechula, R., & Cameron, K. (2016). Current status of palliative care services in Indonesia: a literature review. *International Nursing Review*, 63(2), 180–190. <https://doi.org/10.1111/inr.12236>
- Rodríguez-Sánchez, E., Pérez-Péaranda, A., Losada-Baltar, A., Pérez-Arechaederra, D., Gámez-Marcos, M. Á., Patino-Alonso, M. C., & García-Ortiz, L. (2011). Relationships between quality of life and family function in caregiver. *BMC Family Practice*, 12, 1–7. <https://doi.org/10.1186/1471-2296-12-19>

- Roick, J., Danker, H., Kersting, A., Dietrich, A., Dietz, A., Papsdorf, K., Meixensberger, J., Stolzenburg, J.-U., Wirtz, H., & Singer, S. (2019). The association of socioeconomic status with quality of life in cancer patients over a 6-month period using individual growth models. *Supportive Care in Cancer : Official Journal of the Multinational Association of Supportive Care in Cancer*, 27(9), 3347–3355. <https://doi.org/10.1007/s00520-018-4634-y>
- Sannes, T. S., Simoneau, T. L., Mikulich-Gilbertson, S. K., Natvig, C. L., Brewer, B. W., Kilbourn, K., & Laudenslager, M. L. (2019). Distress and quality of life in patient and caregiver dyads facing stem cell transplant: identifying overlap and unique contributions. *Supportive Care in Cancer*, 27(6), 2329–2337. <https://doi.org/10.1007/s00520-018-4496-3>
- Schulz, R., & Eden, J. (2016). Families caring for an aging America. In *Families Caring for an Aging America*. <https://doi.org/10.17226/23606>
- Schur, S., Ebert-Vogel, A., Amering, M., Masel, E. K., Neubauer, M., Schrott, A., Sibitz, I., Watzke, H., & Schrank, B. (2014). Validation of the “Quality of Life in Life-Threatening Illness—Family Carer Version” (QOLLI-F) in German-speaking carers of advanced cancer patients. *Supportive Care in Cancer*, 22(10), 2783–2791. <https://doi.org/10.1007/s00520-014-2272-6>
- Schwartz, K., Beebe-Dimmer, J., Hastert, T. A., Ruterbusch, J. J., Mantey, J., Harper, F., Thompson, H., Pandolfi, S., & Schwartz, A. G. (2021). Caregiving Burden Among Informal Caregivers of African American Cancer Survivors. *Journal of Cancer Survivorship*, 15(4), 630–640. <https://doi.org/10.1007/s11764-020-00956-x>
- Seidel, D., & Thyrian, J. R. (2019). Burden of caring for people with dementia – Comparing family caregivers and professional caregivers. A descriptive study. *Journal of Multidisciplinary Healthcare*, 12, 655–663. <https://doi.org/10.2147/JMDH.S209106>
- Semiatin, A. M., & O’Connor, M. K. (2012). The relationship between self-efficacy and positive aspects of caregiving in Alzheimer’s disease caregivers. *Aging and Mental Health*, 16(6), 683–688. <https://doi.org/10.1080/13607863.2011.651437>
- Senden, C., Vandecasteele, T., Vandenbergh, E., Versluys, K., Piers, R., Grypdonck, M., & Van Den Noortgate, N. (2015). The interaction between lived experiences of older patients and their family caregivers confronted with a cancer diagnosis and treatment: A qualitative study. *International Journal of Nursing Studies*, 52(1), 197–206. <https://doi.org/10.1016/j.ijnurstu.2014.07.012>

- Seow, H., & Bainbridge, D. (2018). A Review of the Essential Components of Quality Palliative Care in the Home. *Journal of Palliative Medicine*, 21(S1), S-37-S-44. <https://doi.org/10.1089/jpm.2017.0392>
- Sklenarova, H., Krümpelmann, A., Haun, M. W., Friederich, H. C., Huber, J., Thomas, M., Winkler, E. C., Herzog, W., & Hartmann, M. (2015). When do we need to care about the caregiver? Supportive care needs, anxiety, and depression among informal caregivers of patients with cancer and cancer survivors. *Cancer*, 121(9), 1513–1519. <https://doi.org/10.1002/cncr.29223>
- Somanna, S. N., Sastry, N. B., Cheluvaryaswamy, R., & Malila, N. (2022). Quality of life and Its Determinants among Cervical Cancer Patients in South India. *Asian Pacific Journal of Cancer Prevention*, 23(8), 2727–2733. <https://doi.org/10.31557/APJCP.2022.23.8.2727>
- Sottile, P. D., Lynch, Y., Mealer, M., & Moss, M. (2016). Association Between Resilience and Family Member Psychologic Symptoms in Critical Illness. *Critical Care Medicine*, 44(8), e721–e727. <https://doi.org/10.1097/CCM.0000000000001673>
- Sugiyono. (2016). *Metode Penelitian Kuantitatif, Kualitatif, dan R & D*. Alfabeta.
- Sukmawati, L. (2018). *Validitas dan reliabilitas kuisi MCGill Quality of Life revised versi Bahasa Indonesia pada pasien kanker*. (Skripsi Sarjana, Universitas Gadjah Mada)
- Sung, P., Goh, V. S., Azman, N. D., Visaria, A., & Malhotra, R. (2022). Types of Caregiving Experience and Their Association With Caregiver Depressive Symptoms and Quality of Life. *Journal of Aging and Health*, 34(4–5), 591–601. <https://doi.org/10.1177/08982643211051568>
- Takahashi, M., Tanaka, K., & Miyaoka, H. (2005). Depression and associated factors of informal caregivers versus professional caregivers of demented patients. *Psychiatry and Clinical Neurosciences*, 59(4), 473–480. <https://doi.org/10.1111/j.1440-1819.2005.01401.x>
- Talepasand, S., Pooragha, F., & Kazemi, M. (2013). Resiliency and quality of life in patients with cancer: Moderating role of duration of awareness of cancer. *Iranian Journal of Cancer Prevention*, 6(4), 222–226.
- Tan, J. Y., Molassiotis, A., Lloyd-Williams, M., & Yorke, J. (2018). Burden, emotional distress and quality of life among informal caregivers of lung cancer patients: An exploratory study. *European Journal of Cancer Care*, 27(1), 1–11. <https://doi.org/10.1111/ecc.12691>

- Tarlow, B. J., Wisniewski, S. R., Belle, S. H., Rubert, M., Ory, M. G., & Gallagher-Thompson, D. (2004). Positive aspects of caregiving: Contributions of the REACH project to the development of new measures for Alzheimer's caregiving. *Research on Aging*, 26(4), 429–453. <https://doi.org/10.1177/0164027504264493>
- Tate, J. A., & Choi, J. (2020). Positive Appraisal of Caregiving for Intensive Care Unit Survivors: A Qualitative Secondary Analysis. *American Journal of Critical Care*, 29(5), 340–349. <https://doi.org/10.4037/ajcc2020953>
- Toohy, M. J., Muralidharan, A., Medoff, D., Lucksted, A., & Dixon, L. (2016). Caregiver Positive and Negative Appraisals. *Journal of Nervous & Mental Disease*, 204(2), 156–159. <https://doi.org/10.1097/NMD.0000000000000447>
- Tsang, S., Royse, C. F., & Terkawi, A. S. (2017). Guidelines for developing, translating, and validating a questionnaire in perioperative and pain medicine. *Saudi Journal of Anaesthesia*, 11(5), S80–S89. https://doi.org/10.4103/sja.SJA_203_17
- Tsujikawa, M., Yokoyama, K., Urakawa, K., & Onishi, K. (2009). Reliability and validity of Japanese version of the McGill Quality of Life Questionnaire assessed by application in palliative care wards. *Palliative Medicine*, 23(7), 659–664. <https://doi.org/10.1177/0269216309106875>
- Turkoglu, N., & Kilic, D. (2012). Effects of care burdens of caregivers of cancer patients on their quality of life. *Asian Pacific Journal of Cancer Prevention*, 13(8), 4141–4145. <https://doi.org/10.7314/APJCP.2012.13.8.4141>
- Üstündag, S., & Zencirci, A. D. (2015). Factors affecting the quality of life of cancer patients undergoing chemotherapy: A questionnaire study. *Asia-Pacific Journal of Oncology Nursing*, 2(1), 17–25. <https://doi.org/10.4103/2347-5625.152402>
- Valero-Cantero, I., Casals, C., Carrión-Velasco, Y., Barón-López, F. J., Martínez-Valero, F. J., & Vázquez-Sánchez, M. Á. (2022). The influence of symptom severity of palliative care patients on their family caregivers. *BMC Palliative Care*, 21(1), 1–7. <https://doi.org/10.1186/s12904-022-00918-3>
- Van Lancker, A., Velghe, A., Van Hecke, A., Verbrugghe, M., Van Den Noortgate, N., Grypdonck, M., Verhaeghe, S., Bekkering, G., & Beeckman, D. (2014). Prevalence of Symptoms in Older Cancer Patients Receiving Palliative Care: A Systematic Review and Meta-Analysis. *Journal of Pain and Symptom Management*, 47(1), 90–104. <https://doi.org/10.1016/j.jpainsymman.2013.02.016>

- Vincent-Onabajo, G., Gayus, P., Masta, M. A., Ali, M. U., Gujba, F. K., Modu, A., & Hassan, S. U. (2018). Caregiving Appraisal by Family Caregivers of Stroke Survivors in Nigeria. *Journal of Caring Sciences*, 7(4), 183–188. <https://doi.org/10.15171/jcs.2018.028>
- Vitaliano, P. P., Strachan, E., Dansie, E., Goldberg, J., & Buchwald, D. (2014). Does caregiving cause psychological distress? the case for familial and genetic vulnerabilities in female twins. *Annals of Behavioral Medicine*, 47(2), 198–207. <https://doi.org/10.1007/s12160-013-9538-y>
- Wang, T., Molassiotis, A., Chung, B. P. M., & Tan, J. Y. (2018). Unmet care needs of advanced cancer patients and their informal caregivers: a systematic review. *BMC Palliative Care*, 17(1), 1–29. <https://doi.org/10.1186/s12904-018-0346-9>
- Wantonoro, W., Suryaningsih, E. K., Anita, D. C., & Nguyen, T. Van. (2022). Palliative Care: A Concept Analysis Review. *SAGE Open Nursing*, 8(63), 1–9. <https://doi.org/10.1177/23779608221117379>
- Wentlandt, K., Seccareccia, D., Kevork, N., Workentin, K., Blacker, S., Grossman, D., & Zimmermann, C. (2016). Quality of Care and Satisfaction with Care on Palliative Care Units. *Journal of Pain and Symptom Management*, 51(2), 184–192. <https://doi.org/10.1016/j.jpainsymman.2015.10.006>
- WHO. (2020). *Assesing National Capacity For The Prevention and Control of Noncommunicable Diseases : Report of the 2019 Global Survey*. Retrieved from <https://www.who.int/publications/i/item/ncd-ccs-2019>
- Williams, A. L., & McCorkle, R. (2011). Cancer family caregivers during the palliative, hospice, and bereavement phases: a review of the descriptive psychosocial literature. *Palliative & Supportive Care*, 9(3), 315–325. <https://doi.org/10.1017/S1478951511000265>
- World Health Organization. (2012). Programme on Mental Health WHOQoL User Manual. In *World Health Organization*. Retrieved from <https://apps.who.int/iris/rest/bitstreams/110129/retrieve>
- World Health Organization. (2018). Non-Communicable Diseases Country Profiles. In *World Health Organization*. Retrieved from <https://apps.who.int/iris/handle/10665/274512>
- World Health Organization. (2020). *Palliative care*. Retrieved from <https://www.who.int/news-room/fact-sheets/detail/palliative-care>
- World Health Organization. (2022). *Cancer*. Retrieved from <https://www.who.int/news-room/fact-sheets/detail/cancer>

- Yan, B., Yang, L. M., Hao, L. P., Yang, C., Quan, L., Wang, L. H., Wu, Z., Li, X. P., Gao, Y. T., Sun, Q., & Yuan, J. M. (2016). Determinants of quality of life for breast cancer patients in Shanghai, China. *PLoS ONE*, *11*(4), 1–14. <https://doi.org/10.1371/journal.pone.0153714>
- Yifru, T. A., Kisa, S., Dinegde, N. G., & Atnafu, N. T. (2021). Dysphagia and its impact on the quality of life of head and neck cancer patients: institution-based cross-sectional study. *BMC Research Notes*, *14*(1), 1–7. <https://doi.org/10.1186/s13104-020-05440-4>
- Yihedego, E., Aga, F., Gela, D., & Boka, A. (2020). Quality of life and associated factors among family caregivers of adult cancer patients in Addis Ababa, Ethiopia. *Cancer Management and Research*, *12*, 10047–10054. <https://doi.org/10.2147/CMAR.S266416>
- Yu, H., Li, L., Liu, C., Huang, W., Zhou, J., Fu, W., Ma, Y., Li, S., Chang, Y., Liu, G., & Wu, Q. (2017). Factors associated with the quality of life of family caregivers for leukemia patients in China. *Health and Quality of Life Outcomes*, *15*(1), 55. <https://doi.org/10.1186/s12955-017-0628-6>