

Informed Consent for Cancer Treatment and Research

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Ethical Principles of Informed Consent Process

- Angelos, P. (2000). *Ethical issues in cancer patient care*. Norwell, MA: Kluwer Academic Publishers.
- Berry, D.L., Dodd, M.J., Hinds, P.S., & Ferrell, B.R. (1996). Informed consent: Process and clinical issues. *Oncology Nursing Forum*, 23, 507–512.
- Clark, S. (2003). Informed consent without bureaucracy. *Journal of Clinical Neuroscience*, 10, 35–36.
- Dalla-Vorgia, P., Lascaratos, J., Skiadas, P., & Garanis-Papadatos, T. (2001). Is consent in medicine a concept only of modern times? *Journal of Medical Ethics*, 27, 59–61.
- English, D.C. (2002). Valid informed consent: A process, not a signature. *American Surgeon*, 68, 45–48.
- Jerrold, L. (2002). The limits of informed consent. *American Journal of Orthodontics and Dentofacial Orthopedics*, 121, 542–544.
- Leino-Kilpi, H., Valimaki, M., Arndt, M., Dassen, T., Gasull, M., Lemonidou, C., et al. (2000). *Patient's autonomy, privacy and informed consent*. Amsterdam: IOS Press.
- Loyal, L., & Tobias, J.S. (2001). *Informed consent in medical research*. London: BMJ Books.
- Manning, S.S., & Gaul, C.E. (1997). The ethics of informed consent: A critical variable in the self-determination of health and mental health clients. *Social Work in Health Care*, 25, 103–117.
- Neumann, J.L. (2001). Ethical issues confronting oncology nurses. *Nursing Clinics of North America*, 36, 827–841.
- O'Neill, O. (2003). Some limits of informed consent. *Journal of Medical Ethics*, 29, 4–7.
- Pape, T. (1997). Legal and ethical considerations of informed consent. *AORN Journal*, 65, 1122–1127.
- Rousseau, P. (2001). Ethical and legal issues in palliative care. *Primary care: Clinics in office practice*, 28, 391–400.
- Sims, J., & Miracle, V. (2002). Elements of an informed consent. *Dimensions of Critical Care Nursing*, 21, 242–245.

- Thall, P.F. (2002). Ethical issues in oncology biostatistics. *Statistical Methods in Medical Research*, 11, 428–448.

Disclosing Information About Options, Risks, Benefits, and Alternatives

- Albrecht, T.L., Blanchard, C., Ruckdeschel, J.C., Coovert, M., & Strongbow, R. (1999). Strategic physician communication and oncology clinical trials. *Journal of Clinical Oncology*, 17, 3324–3332.
- Baylis, F., & Downie, J. (2001). Professional recommendations: Disclosing facts and values. *Journal of Medical Ethics*, 27, 20–24.
- Bjorn, E., Rossel, P., & Holm, S. (1999). Can the written information to research subjects be improved?: An empirical study. *Journal of Medical Ethics*, 25, 263–267.
- Butow, P.N., Dunn, S.M., & Tattersall, M.H. (2000). Denial, misinformation and the “assault of truth.” In E. Bruera & R.K. Portenoy (Eds.), *Topics in palliative care* (4th ed., pp. 263–278). Oxford, UK: Oxford University Press.
- Calman, K.C. (2002). Communication of risk: Choice, consent, and trust. *Lancet*, 360, 166–168.
- Carrie, C., & Freeman, C. (2001). Are all consents really informed? *International Journal of Radiation Oncology, Biology, Physics*, 51, 287.
- Chan, Y., Irish, J.C., Wood, S.J., Rotstein, L.E., Brown, D.H., Gullane, P.J., et al. (2002). Patient education and informed consent in head and neck surgery. *Archives of Otolaryngology and Head and Neck Surgery*, 128, 1269–1274.
- Davis, T.C., Williams, M.V., Marin, E., Parker, R.M., & Glass, J. (2002). Health literacy and cancer communication. *CA: A Cancer Journal for Clinicians*, 52, 134–159.
- Fleissig, A., Jenkins, V., & Fallowfield, L. (2001). Results of an intervention study to improve communication about randomized clinical trials of cancer therapy. *European Journal of Cancer*, 37, 322–331.
- Fried, E. (2001). The therapeutic misconception, beneficence, and respect. *Accountability in Research*, 8, 331–348.
- Garrud, P., Wood, M., & Stainsby, L. (2001). Impact of risk information in a patient education leaflet. *Patient Education and Counseling*, 43, 301–304.
- Gattellari, M., Voigt, K.J., Butow, P.N., & Tattersall, M.H. (2002). When the treatment goal is not cure: Are cancer patients equipped to make informed decisions? *Journal of Clinical Oncology*, 20, 503–513.
- Holmes-Rovner, M., & Wills, C.E. (2002). Improving consent: Insights from behavioural decision research. *Medical Care*, 40(Suppl. 9), 30–38.
- Jefford, M., & Tattersall, M.H. (2002). Informing and involving cancer patients in their own care. *Lancet Oncology*, 3, 629–637.
- Lewis, D. (1999). Computer-based approaches to patient education: A review of the literature. *Journal of the American Medical Informatics Association*, 6, 272–282.
- Tattersall, M.H., & Butow, P.N. (2002). Consultation audio tapes: An underused cancer patient information aid and clinical research tool. *Lancet Oncology*, 3, 431–437.
- Tattersall, M.H., Gattellari, M., Voigt, K., & Butow, P.N. (2002). When the treatment goal is not cure: Are patients informed adequately? *Supportive Care in Cancer*, 10, 314–321.
- Wilt, T. (2002). Clarifying uncertainty regarding detection and treatment of early-stage prostate cancer. *Seminars in Urologic Oncology*, 20, 10–17.
- Weston, J., Hannah, M., & Downes, J. (1997). Evaluating the benefits of a patient information video during the informed consent process. *Patient Education and Counseling*, 30, 239–245.
- Ziegler, D.K., Mosier, M.C., Buenaver, M., & Okuyemi, K. (2001). How much information about adverse effects of medication do patients want from physicians? *Archives of Internal Medicine*, 161, 706–713.
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