

REFERENCES

1. Alsop, K., Fereday, S., Meldrum, C., deFazio, A., Emmanuel, C., George, J., et al. (2012). BRCA mutation frequency and patterns of treatment response in BRCA mutation-positive women with ovarian cancer: A report from the Australian ovarian cancer study group. *Journal of Clinical Oncology*, 30(21), 2654–2663.
2. Candido-dos-Reis, F. J., Song, H., Goode, E. L., Cunningham, J. M., Fridley, B. L., Larson, M. C., et al., Australian Ovarian Cancer Study, G. (2015). Germline mutation in BRCA1 or BRCA2 and ten-year survival for women diagnosed with epithelial ovarian cancer. *Clinical Cancer Research*, 21(3), 652–657.
3. Domchek, S. M., Jhaveri, K., Patil, S., Stopfer, J. E., Hudis, C., Powers, J., et al. (2013). Risk of metachronous breast cancer after BRCA mutation-associated ovarian cancer. *Cancer*, 119(7), 1344–1348.
4. Easton, D. F., Pharoah, P. D., Antoniou, A. C., Tischkowitz, M., Tavtigian, S. V., Nathanson, K. L., & Foulkes, W. D. (2015). Gene-panel sequencing and the prediction of breast-cancer risk. *The New England Journal of Medicine*, 372(23), 2243–2257.
5. Gangi, A., Cass, I., Paik, D., Barmparas, G., Karlan, B., Dang, C., & Amersi, F. F. (2014). Breast cancer following ovarian cancer in BRCA mutation carriers. *JAMA Surgery*, 149(12), 1306–1313.
6. Hartmann, L. C., & Lindor, N. M. (2015). The role of risk-reducing surgery in hereditary breast and ovarian cancer. *New England Journal of Medicine*, 374(5), 454–468.
7. Konecny, G. E., & Kristeleit, R. S. (2016). PARP inhibitors for BRCA1/2-mutated and sporadic ovarian cancer: Current practice and future directions. *British Journal of Cancer*, 115(10), 1157–1173.
8. Kriege, M., Brekelmans, C. T., Boetes, C., Besnard, P. E., Zonderland, H. M., Obdeijn, I. M., & Magnetic Resonance Imaging Screening Study, G. (2004). Efficacy of MRI and mammography for breast cancer screening in women with a familial or genetic predisposition. *The New England Journal of Medicine*, 351(5), 427–437.
9. Mavaddat, N., Peock, S., Frost, D., Ellis, S., Platte, R., Fineberg, E., & Embrace. (2013). Cancer risks for BRCA1 and BRCA2 mutation carriers: Results from prospective analysis of EMBRACE. *Journal of the National Cancer Institute*, 105(11), 812–822.