

Two-Stage Breast Cancer Screening in the Developing World

Charalabos Batsis

Published online: 21 September 2010
© Société Internationale de Chirurgie 2010

Approximately two-thirds of women with a new diagnosis of breast cancer in the United States and northwestern Europe have an early-stage breast cancer, and most have a good prognosis with long-term disease-free survival. National mammographic screening programs have resulted in this improved early detection, saving the lives of millions of women in the industrialized world. However, the lack of such technology and screening programs in non-industrialized countries may be the cause of high breast cancer mortality in these countries.

Denewer and colleagues' [1] article in the *World Journal of Surgery* evaluated whether the surgeon's clinical breast examination could improve the rate of detecting early-stage breast cancer in Egypt. The authors reported that the mean tumor size in Egyptian women at the time of diagnosis was 4.5 cm, and the median age was approximately 46 years. From a total of 57,500 women aged 25–65 years in the targeted population, the voluntary participation rate for the surgeon's clinical examination was 10.2%. Abnormal clinical findings were found in 3.2% (191/5900). These 191 women then underwent a second stage of examination that included repeat clinical examination plus ultrasonography and/or mammography. A total of 18 breast cancers were detected using this two-stage approach, and the median tumor size was 1.5 cm. The cost of screening per cancer case detected was approximately \$415 (US), and the overall cost of treating a screen-detected cancer was \$1015–\$1215 (US). The authors concluded that this two-stage screening approach is effective and reduces the cost of managing breast cancer in Egypt.

The study has some limitations. Only 18 breast cancers were detected, and about 90% of the targeted population did not participate. Moreover, given the limitation of the clinical examination (as shown in Western large-scale studies compared to mammographic screening), a large number of Egyptian women with early-stage breast cancer may not be detected by this two-stage approach. However, given the limitations of performing mass screening programs in developing countries, the study by Denewer and colleagues provides an appropriate effort to reduce late diagnosis and death from breast cancer in the real world of developing countries.

Despite advances in screening and treatment in the developed world, many women are still diagnosed at more advanced stages, and a substantial proportion of women with early breast cancer develop a recurrence and die of the disease. The latest research in cancer genetics, genomics, molecular biology, DNA sequencing technology, and translational oncology is focused on personalized risk prediction and individualized primary prevention and treatment. Novel therapeutic strategies aimed at the development of valid biomarkers and highly effective targeted drugs may improve outcomes of patients with breast cancer or other major cancer types [2–15].

References

1. Denewer A, Hussein O, Farouk O et al (2010) Cost-effectiveness of clinical breast assessment-based screening in rural Egypt. *World J Surg* 34:2204–2210
2. Wacholder S, Hartge P, Prentice R et al (2010) Performance of common genetic variants in breast-cancer risk models. *N Engl J Med* 362:986–993
3. Ziogas D, Roukos DH (2009) Genetics and personal genomics for personalized breast cancer surgery: progress and challenges in research and clinical practice. *Ann Surg Oncol* 16(7):1771–1782

C. Batsis (✉)
Department of Surgery, School of Medicine,
University of Ioannina, 451 10 Ioannina, Greece
e-mail: chbatsis@hotmail.com

4. Katsios C, Roukos DH (2010) Individual genomes and personalized medicine: life diversity and complexity. *Per Med* 7(4): 347–350
5. Roukos DH, Ziogas D (2010) From tumor size and HER2 status to systems oncology for very early breast cancer treatment. *Expert Rev Anticancer Ther* 10(2):123–128
6. Roukos DH (2010) Complete genome sequencing and network modeling to overcome trastuzumab resistance. *Pharmacogenomics* 11(8):1039–1043
7. Roukos DH (2010) Next-generation, genome sequencing-based biomarkers: concerns and challenges for medical practice. *Biomark Med* 4(4):583–586
8. Roukos DH, Katsios C, Liakakos T (2010) Genotype-phenotype map and molecular networks: a promising solution in overcoming colorectal cancer resistance to targeted treatment. *Expert Rev Mol Diagn* 10(5):541–545
9. Roukos DH (2010) Bionetworks-based personalized medicine versus comparative-effectiveness research or harmonization of both in cancer management? *Expert Rev Mol Diagn* 10(3): 247–250
10. Roukos DH (2010) Systems medicine: a real approach for future personalized oncology? *Pharmacogenomics* 11(3):283–287
11. Roukos DH (2010) Novel clinico-genome network modeling for revolutionizing genotype-phenotype-based personalized cancer care. *Expert Rev Mol Diagn* 10(1):33–48
12. Roukos DH (2009) Isolated tumor cells in breast cancer. *N Engl J Med* 361:1994–1995; author reply 1995–6
13. Roukos DH, Tzakos A, Zografos G (2009) Current concerns and challenges towards tailored anti-angiogenic therapy in cancer. *Expert Rev Anticancer Ther* 9(10):1413–1416
14. Roukos DH, Ziogas D (2009) Human genetic and structural genomic variation: would genome-wide association studies be the solution for cancer complexity like Alexander the Great for the “Gordian Knot”? *Ann Surg Oncol* 16(3):774–775
15. Roukos DH (2010) Targeting gastric cancer with trastuzumab: new clinical practice and innovative developments to overcome resistance. *Ann Surg Oncol* 17:14–17