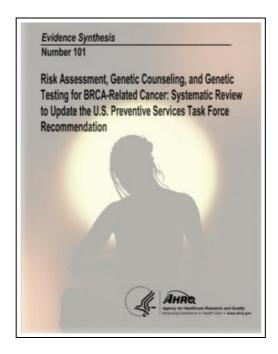
Risk Assessment, Genetic Counseling, and Genetic Testing for Brca-Related Cancer: Systematic Review to Update the U.S. Preventive Services Task Force Recommendation: Evidence Synthesis Number 101



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Reviews

Very useful for all group of people. It is amongst the most incredible pdf i actually have read through. Its been written in an extremely straightforward way and it is just right after i finished reading through this pdf by which basically modified me, change the way i think. (Felicia Nikolaus)

RISK ASSESSMENT, GENETIC COUNSELING, AND GENETIC TESTING FOR BRCA-RELATED CANCER: SYSTEMATIC REVIEW TO UPDATE THE U.S. PREVENTIVE SERVICES TASK FORCE RECOMMENDATION: EVIDENCE SYNTHESIS NUMBER 101



CreateSpace Independent Publishing Platform. Paperback. Book Condition: New. This item is printed on demand. Paperback. 368 pages. Dimensions: 11.0in. x 8.5in. x 0.8in.This systematic review is an update of the evidence for the U. S. Preventive Services Task Force (USPSTF) on the effectiveness and adverse effects of risk assessment, genetic counseling, and genetic testing for breast cancer susceptibility gene (BRCA)related cancer in women who do not have cancer but are potentially at increased risk. Its purpose is to evaluate and summarize evidence addressing specific key questions important to the USPSTF as it considers new recommendations for primary care practice. In 2005, based on results of a previous review, the USPSTF recommended against routine referral for genetic counseling or routine BRCA testing for women whose family histories are not associated with increased risks for deleterious mutations in breast cancer susceptibility gene 1 (BRCA1) or breast cancer susceptibility gene 2 (BRCA2) (D recommendation). The USPSTF also recommended that women whose family histories are associated with increased risks for mutations in the BRCA1 or BRCA2 genes be referred for genetic counseling and evaluation for BRCA testing (B recommendation). The USPSTF concluded that the potential harms of routine referral for genetic counseling or BRCA mutation testing in women without family history risk outweigh the benefits, and that the benefits of referring women with family history risk to suitably trained health care providers outweigh the harms. Benefits included improved accuracy of risk assessment and pretest probability for testing and improved patient knowledge, risk perception, and psychological and health outcomes. Potential harms included inaccurate risk assessment; inappropriate testing; misinterpretation of test results; and ethical, legal, and social implications; among others. The 2005 USPSTF recommendation was intended for the primary prevention of cancer and applied to women without previous diagnoses of brea

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