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Date of request: September 21, 2015

NCCN Guidelines Panel: Prostate Cancer Early Detection

<u>Specific Changes</u>: Incorporate recommendation and relevant citations for genetic risk assessment based on existing NCCN guidelines (Genetic/Familial High-Risk Assessment: Breast and Ovarian) into the Baseline Evaluation flowchart on page PROSD-2 and the discussion of Prostate Cancer Risk in Genetic Syndromes on page MS-11.

FDA Clearance: Not applicable.

Rationale: The NCCN Guideline for Genetic/Familial High-Risk Assessment: Breast and Ovarian provides specific criteria for genetic evaluation/referral and testing of individuals with a personal and/or family history of prostate cancer (pages BR/OV-1 and HBOC-1), as well as prostate screening recommendations for men identified with pathogenic mutations in the genes *BRCA1* and *BRCA2* as a result of genetic testing. Attention to these guidelines is a key component of the baseline evaluation and risk assessment process by which informed decisions are made regarding prostate cancer screening.

<u>Citations:</u> The following articles are submitted in support of this proposed change. We would like to acknowledge the contributions of NCCN panel members who are also co-authors or co-contributors of some of these publications.

Daly M et al. NCCN Clinical Practice Guidelines in Oncology®: Genetic/Familial High-Risk Assessment: Breast and Ovarian. V 2.2015. June 25. Available at http://www.nccn.org.

Sincerely,

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