INTEGRATED CANCER RESEARCH CENTER

Patient: Kardashian, Kimberly

Clinic : Grady Memorial Hospital, Atlanta, GA

Gender : Female DOB : 10/21/1980

HIGH RISK

Gene	Mutation	Known Associations
MSH2	c.8C>G	Lynch-syndrome
BRCA1	c.9A>T	Breast-cancer
BRCA2	c.10G>C	Ovarian-cancer

Gene Details: MSH2 c.8C>G : NM-000251.2

Functional Significance: Deleterious

The heterozycous germline MSH2 c.8C>G is predicted to result in the premature truncation of the MSH2 protein due to abnormal protein production and/or function.

Clinical Significance: High Cancer Risk

This mutation is associated with increased cancer risk and should be regarded as clinically significant.

Additional Information

Genes Analyzed: APC, ATM, BARD1, BMPR1A, BRCA1, BRCA2, BRIP1, CDH1, CDK4, CDKN2A, CHEK2, EPCAM, MLH1, MSH2, MSH6, MUTYH, NBN, PALB2, PMS2, PTEN, RAD51C, SMAD4, STK11, TP53

Disclaimer: The technical specifications summary of ICRC Gene Expose describes the analysis, method, performance, nomenclature, and interpretive criteria of this test. The classification and interpretation of variants identified reflect the current state of scientific understanding at the time of this report.

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