



Kenneth Offit, MD

Research in the laboratory focuses on the discovery and characterization of novel cancer predisposing genes in humans, including both common and rare variants. We conduct studies to describe phenotype, penetrance, modifying effect, and clinical outcomes associated with germline genetic alterations in cancer patients and their families. In addition, our research focuses on the interpretation and clinical translation of results of massively parallel sequencing of germline genomes in cancer-prone kindreds.

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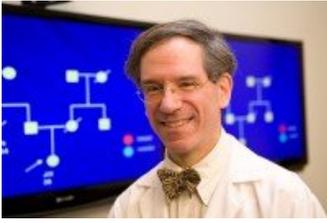
FINDING



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New findings could lead to a different kind of cancer drug that targets a DNA damage repair pathway called nucleotide excision repair.

IN THE CLINIC



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MSK's new Robert and Kate Niehaus Center for Inherited Cancer Genomics is using the latest in gene sequencing technologies to discover the inherited causes of cancer.

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Publications Highlights

[Wei X, Calvo-Vidal MN, Chen S, Wu G, Revuelta MV, Sun J, Zhang J, Walsh MF, Nichols KE, Joseph V, Snyder C, Vachon CM, McKay JD, Wang SP, Jayabalan DS, Jacobs LM, Becirovic D, Waller RG, Artomov M, Viale A, Patel J, Phillip J, Chen-Kiang S, Curtin K, Salama M, Atanackovic D, Niesvizky R, Landgren O, Slager SL, Godley LA, Churpek J, Garber JE, Anderson KC, Daly MJ, Roeder RG, Dumontet C, Lynch HT, Mullighan CG, Camp NJ, Offit K, Klein RJ, Yu H, Cerchiatti L, Lipkin SM. Germline Lysine-Specific Demethylase 1 \(*LSD1/KDM1A*\) Mutations Confer Susceptibility to Multiple Myeloma. *Cancer Res.* 2018 May 15;78\(10\):2747-2759.](#)

[Katona BW, Yurgelun MB, Garber JE, Offit K, Domchek SM, Robson ME, Stadler ZK. A counseling framework for moderate-penetrance colorectal cancer susceptibility genes. *Genet Med.* 2018 Mar 1.](#)

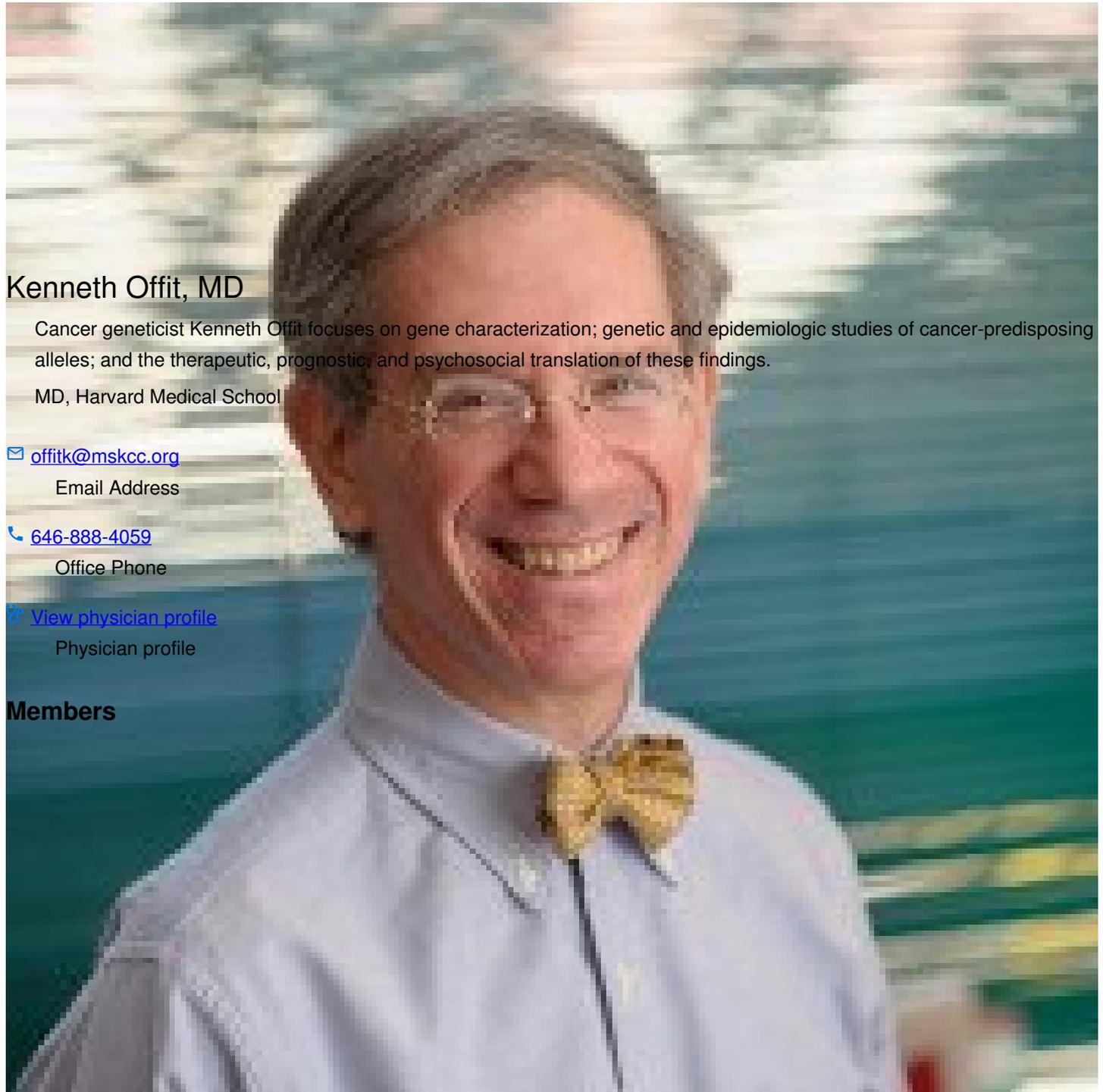
[Walsh MF, Kennedy J, Harlan M, Kentsis A, Shukla N, Musinsky J, Roberts S, Kung AL, Robson M, Kushner BH, Meyers P, Offit K. Germline *BRCA2* mutations detected in pediatric sequencing studies impact parents' evaluation and care. *Cold Spring Harb Mol Case Stud.* 2017 Nov 21;3\(6\).](#)

[Mandelker D, Zhang L, Kemel Y, Stadler ZK, Joseph V, Zehir A, Pradhan N, Arnold A, Walsh MF, Li Y, Balakrishnan AR, Syed A, Prasad M, Nafa K, Carlo MI, Cadoo KA, Sheehan M, Fleischut MH, Salo-Mullen E, Trottier M, Lipkin SM, Lincoln A, Mukherjee S, Ravichandran V, Cambria R, Galle J, Abida W, Arcila ME, Benayed R, Shah R, Yu K, Bajorin DF, Coleman JA, Leach SD, Lowery MA, Garcia-Aguilar J, Kantoff PW, Sawyers CL, Dickler MN, Saltz L, Motzer RJ, O'Reilly EM, Scher HI, Baselga J, Klimstra DS, Solit DB, Hyman DM, Berger MF, Ladanyi M, Robson ME, Offit K. Mutation Detection in Patients With Advanced Cancer by Universal Sequencing of Cancer-Related Genes in Tumor and Normal DNA vs Guideline-Based Germline Testing. *JAMA.* 2017 Sep 5;318\(9\):825-835. Doi: 10.1001/jama.2017.11137](#)

[Hamilton JG, Shuk E, Genoff MC, Rodríguez VM, Hay JL, Offit K, Robson ME. Interest and Attitudes of Patients With Advanced Cancer With Regard to Secondary Germline Findings From Tumor Genomic Profiling. *J Oncol Pract.* 2017 Jul;13\(7\):e590-e601.](#)

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People



Kenneth Offit, MD

Cancer geneticist Kenneth Offit focuses on gene characterization; genetic and epidemiologic studies of cancer-predisposing alleles; and the therapeutic, prognostic, and psychosocial translation of these findings.
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Kenneth Offit discloses the following relationships and financial interests:

AnaNeo Therapeutics

Equity

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