## Fine-mapping of the HNF1B multicancer locus identifies endometrial cancer risk

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**Citation Report** 

#	Article	IF	CITATIONS
1	Meta-analysis of genome-wide association studies identifies common susceptibility polymorphisms for colorectal and endometrial cancer near SH2B3 and TSHZ1. Scientific Reports, 2015, 5, 17369.	1.6	35
2	Candidate locus analysis of the TERT–CLPTM1L cancer risk region on chromosome 5p15 identifies multiple independent variants associated with endometrial cancer risk. Human Genetics, 2015, 134, 231-245.	1.8	34
3	Limited evidence that cancer susceptibility regions are preferential targets for somatic mutation. Genome Biology, 2015, 16, 193.	3.8	19
4	Comprehensive genetic assessment of the ESR1 locus identifies a risk region for endometrial cancer. Endocrine-Related Cancer, 2015, 22, 851-861.	1.6	25
5	Five endometrial cancer risk loci identified through genome-wide association analysis. Nature Genetics, 2016, 48, 667-674.	9.4	77
6	Genetic Risk Score Mendelian Randomization Shows that Obesity Measured as Body Mass Index, but not Waist:Hip Ratio, Is Causal for Endometrial Cancer. Cancer Epidemiology Biomarkers and Prevention, 2016, 25, 1503-1510.	1.1	64
8	A Common Variant at the 14q32 Endometrial Cancer Risk Locus Activates AKT1 through YY1 Binding. American Journal of Human Genetics, 2016, 98, 1159-1169.	2.6	32
9	CYP19A1 fine-mapping and Mendelian randomization: estradiol is causal for endometrial cancer. Endocrine-Related Cancer, 2016, 23, 77-91.	1.6	62
10	Expression, Epigenetic and Genetic Changes of HNF1B in Endometrial Lesions. Pathology and Oncology Research, 2016, 22, 523-530.	0.9	19
11	GWAS meta-analysis of 16 852 women identifies new susceptibility locus for endometrial cancer. Human Molecular Genetics, 2016, 25, ddw092.	1.4	19
12	Diabetes mellitus and gynecologic cancer: molecular mechanisms, epidemiological, clinical and prognostic perspectives. Archives of Gynecology and Obstetrics, 2016, 293, 239-246.	0.8	50
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14	Endometrial cancer gene panels: clinical diagnostic vs research germline DNA testing. Modern Pathology, 2017, 30, 1048-1068.	2.9	37
15	Current strategies in the diagnosis of endometrial cancer. Archives of Gynecology and Obstetrics, 2017, 296, 5-14.	0.8	12
17	Family history of cancer predicts endometrial cancer risk independently of Lynch Syndrome: Implications for genetic counselling. Gynecologic Oncology, 2017, 147, 381-387.	0.6	30
18	Pathology and Molecular Pathology of Uterine and Ovarian Cancers. , 2017, , 247-278.		0
19	The OncoArray Consortium: A Network for Understanding the Genetic Architecture of Common Cancers. Cancer Epidemiology Biomarkers and Prevention, 2017, 26, 126-135.	1.1	278
20	Genetic overlap between endometriosis and endometrial cancer: evidence from crossâ€disease genetic correlation and GWAS metaâ€analyses. Cancer Medicine, 2018, 7, 1978-1987.	1.3	62

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21	Genome-wide analysis of PDX1 target genes in human pancreatic progenitors. Molecular Metabolism, 2018, 9, 57-68.	3.0	67
22	Genome-wide meta-analysis identifies five new susceptibility loci for pancreatic cancer. Nature Communications, 2018, 9, 556.	5.8	188
23	Common Genetic Variation and Susceptibility to Ovarian Cancer: Current Insights and Future Directions. Cancer Epidemiology Biomarkers and Prevention, 2018, 27, 395-404.	1.1	33
24	Risk and prognostic factors for endometrial carcinoma after diagnosis of breast or Lynchâ€associated cancers—A populationâ€based analysis. Cancer Medicine, 2018, 7, 6411-6422.	1.3	9
25	Cumulative evidence for relationships between multiple variants of HNF1B and the risk of prostate and endometrial cancers. BMC Medical Genetics, 2018, 19, 128.	2.1	5
26	Identification of nine new susceptibility loci for endometrial cancer. Nature Communications, 2018, 9, 3166.	5.8	178
27	Genome-Wide Association Studies of Endometrial Cancer: Latest Developments and Future Directions. Cancer Epidemiology Biomarkers and Prevention, 2019, 28, 1095-1102.	1.1	32
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30	Germline genetic variation in prostate susceptibility does not predict outcomes in the chemoprevention trials PCPT and SELECT. Prostate Cancer and Prostatic Diseases, 2020, 23, 333-342.	2.0	10
31	Analysis of expression, epigenetic, and genetic changes of HNF1B in 130 kidney tumours. Scientific Reports, 2020, 10, 17151.	1.6	5
32	HNF1B, EZH2 and ECI2 in prostate carcinoma. Molecular, immunohistochemical and clinico-pathological study. Scientific Reports, 2020, 10, 14365.	1.6	6
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34	Expression, Epigenetic, and Genetic Changes of HNF1B in Colorectal Lesions: an Analysis of 145 Cases. Pathology and Oncology Research, 2020, 26, 2337-2350.	0.9	7
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41	The prospect of discovering new biomarkers for ovarian cancer based on current knowledge ofi¿½susceptibility loci and genetic variation (Review). International Journal of Molecular Medicine, 2019, 44, 1599-1608.	1.8	2
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