Joellen Schildkraut

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Lack of association between modifiable exposures and glioma risk: A Mendelian randomisation analysis. Neuro-Oncology, 2020, 22, 207-215.	1.2	19
2	Mendelian randomisation study of the relationship between vitamin D and risk of glioma. Scientific Reports, 2018, 8, 2339.	3.3	23
3	Impact of atopy on risk of glioma: a Mendelian randomisation study. BMC Medicine, 2018, 16, 42.	5.5	38
4	Influence of obesity-related risk factors in the aetiology of glioma. British Journal of Cancer, 2018, 118, 1020-1027.	6.4	32
5	Correlation between germline mutations in MMR genes and microsatellite instability in ovarian cancer specimens. Familial Cancer, 2017, 16, 351-355.	1.9	18
6	History of chickenpox in glioma risk: a report from the glioma international case–control study (<scp>GICC</scp>). Cancer Medicine, 2016, 5, 1352-1358.	2.8	36
7	Approaching a Scientific Consensus on the Association between Allergies and Glioma Risk: A Report from the Glioma International Case-Control Study. Cancer Epidemiology Biomarkers and Prevention, 2016, 25, 282-290.	2.5	89
8	Targeted Sequencing in Chromosome 17q Linkage Region Identifies Familial Glioma Candidates in the Gliogene Consortium. Scientific Reports, 2015, 5, 8278.	3.3	22
9	Maternal cadmium, iron and zinc levels, DNA methylation and birth weight. BMC Pharmacology & Toxicology, 2015, 16, 20.	2.4	95
10	Cross Cancer Genomic Investigation of Inflammation Pathway for Five Common Cancers: Lung, Ovary, Prostate, Breast, and Colorectal Cancer. Journal of the National Cancer Institute, 2015, 107, djv246.	6.3	63
11	Variation in NF-κB Signaling Pathways and Survival in Invasive Epithelial Ovarian Cancer. Cancer Epidemiology Biomarkers and Prevention, 2014, 23, 1421-1427.	2.5	13
12	Erythrocyte folate concentrations, CpG methylation at genomically imprinted domains, and birth weight in a multiethnic newborn cohort. Epigenetics, 2014, 9, 1120-1130.	2.7	73
13	Maternal Stress, Preterm Birth, and DNA Methylation at Imprint Regulatory Sequences in Humans. Genetics & Epigenetics, 2014, 6, GEG.S18067.	2.5	93
14	Cigarette smoking and risk of ovarian cancer: a pooled analysis of 21 case–control studies. Cancer Causes and Control, 2013, 24, 989-1004.	1.8	84
15	Gene Set Analysis of Survival Following Ovarian Cancer Implicates Macrolide Binding and Intracellular Signaling Genes. Cancer Epidemiology Biomarkers and Prevention, 2012, 21, 529-536.	2.5	7
16	Breast and Ovarian Cancer Risk and Risk Reduction in Jewish <i>BRCA1/2</i> Mutation Carriers. Journal of Clinical Oncology, 2012, 30, 1321-1328.	1.6	31
17	Common variants at 19p13 are associated with susceptibility to ovarian cancer. Nature Genetics, 2010, 42, 880-884.	21.4	235
18	A genome-wide association study identifies susceptibility loci for ovarian cancer at 2q31 and 8q24. Nature Genetics, 2010, 42, 874-879.	21.4	321

#	Article	IF	CITATIONS
19	Association of Risk-Reducing Surgery in <emph type="ital">BRCA1</emph> or <emph type="ital">BRCA2 Mutation Carriers With Cancer Risk and Mortality. JAMA - Journal of the American Medical Association, 2010, 304, 967.</emph 	7.4	1,241
20	Attitudes, Knowledge, and Risk Perceptions of Women With Breast and/or Ovarian Cancer Considering Testing for BRCA1 and BRCA2. Journal of Clinical Oncology, 1999, 17, 1040-1040.	1.6	127
21	Ethical ramifications of alternative means of recruiting research participants from cancer registries. , 1999, 86, 647-651.		22
22	Effect of BRCA1 and BRCA2 on the Association Between Breast Cancer Risk and Family History. Journal of the National Cancer Institute, 1998, 90, 1824-1829.	6.3	102
23	Probability of Carrying a Mutation of Breast-Ovarian Cancer Gene BRCA1 Based on Family History. Journal of the National Cancer Institute, 1997, 89, 227-237.	6.3	344