## Zsofia Stadler

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/5738734/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	PD-1 Blockade in Mismatch Repair–Deficient, Locally Advanced Rectal Cancer. New England Journal of Medicine, 2022, 386, 2363-2376.	27.0	588
2	Cancer therapy shapes the fitness landscape of clonal hematopoiesis. Nature Genetics, 2020, 52, 1219-1226.	21.4	367
3	Identification of 12 new susceptibility loci for different histotypes of epithelial ovarian cancer. Nature Genetics, 2017, 49, 680-691.	21.4	356
4	Germline Variants in Targeted Tumor Sequencing Using Matched Normal DNA. JAMA Oncology, 2016, 2, 104.	7.1	270
5	Germline <i>BRCA</i> Mutations Denote a Clinicopathologic Subset of Prostate Cancer. Clinical Cancer Research, 2010, 16, 2115-2121.	7.0	263
6	Counselling framework for moderate-penetrance cancer-susceptibility mutations. Nature Reviews Clinical Oncology, 2016, 13, 581-588.	27.6	258
7	Reliable Pan-Cancer Microsatellite Instability Assessment by Using Targeted Next-Generation Sequencing Data. JCO Precision Oncology, 2017, 2017, 1-17.	3.0	209
8	Therapeutic Implications of Germline Testing in Patients With Advanced Cancers. Journal of Clinical Oncology, 2021, 39, 2698-2709.	1.6	83
9	Immunohistochemical Staining for DNA Mismatch Repair Proteins in Intestinal Tract Carcinoma. American Journal of Surgical Pathology, 2011, 35, 447-454.	3.7	82
10	Secondary mutation in a coding mononucleotide tract in MSH6 causes loss of immunoexpression of MSH6 in colorectal carcinomas with MLH1/PMS2 deficiency. Modern Pathology, 2013, 26, 131-138.	5.5	82
11	Germline EGFR T790M Mutation Found in Multiple Members of a Familial Cohort. Journal of Thoracic Oncology, 2014, 9, 554-558.	1.1	63
12	Risk of metachronous breast cancer after <i>BRCA</i> mutation–associated ovarian cancer. Cancer, 2013, 119, 1344-1348.	4.1	58
13	Susceptibility Loci Associated with Specific and Shared Subtypes of Lymphoid Malignancies. PLoS Genetics, 2013, 9, e1003220.	3.5	44
14	Assessment of SLX4 Mutations in Hereditary Breast Cancers. PLoS ONE, 2013, 8, e66961.	2.5	37
15	Clinical Calculator Based on Molecular and Clinicopathologic Characteristics Predicts Recurrence Following Resection of Stage I-III Colon Cancer. Journal of Clinical Oncology, 2021, 39, 911-919.	1.6	34
16	Risk-Reducing Bilateral Salpingo-Oophorectomy for Ovarian Cancer: A Review and Clinical Guide for Hereditary Predisposition Genes. JCO Oncology Practice, 2022, 18, 201-209.	2.9	34
17	Germline <i>SDHA</i> mutations in children and adults with cancer. Journal of Physical Education and Sports Management, 2018, 4, a002584.	1.2	33
18	Toward automation of germline variant curation in clinical cancer genetics. Genetics in Medicine, 2019, 21, 2116-2125.	2.4	27

ZSOFIA STADLER

#	Article	IF	CITATIONS
19	Breast and Prostate Cancer Risks for Male <i>BRCA1</i> and <i>BRCA2</i> Pathogenic Variant Carriers Using Polygenic Risk Scores. Journal of the National Cancer Institute, 2022, 114, 109-122.	6.3	19
20	Association of breast cancer risk in BRCA1 and BRCA2 mutation carriers with genetic variants showing differential allelic expression: identification of a modifier of breast cancer risk at locus 11q22.3. Breast Cancer Research and Treatment, 2017, 161, 117-134.	2.5	18
21	Tumor-Infiltrating Lymphocytes, Tumor Mutational Burden, and Genetic Alterations in Microsatellite Unstable, Microsatellite Stable, or Mutant <i>POLE/POLD1</i> Colon Cancer. JCO Precision Oncology, 2021, 5, 817-826.	3.0	18
22	The predictive ability of the 313 variant–based polygenic risk score for contralateral breast cancer risk prediction in women of European ancestry with a heterozygous BRCA1 or BRCA2 pathogenic variant. Genetics in Medicine, 2021, 23, 1726-1737.	2.4	16
23	Development and Assessment of a Clinical Calculator for Estimating the Likelihood of Recurrence and Survival Among Patients With Locally Advanced Rectal Cancer Treated With Chemotherapy, Radiotherapy, and Surgery. JAMA Network Open, 2021, 4, e2133457.	5.9	16
24	Prevalence and Characterization of Biallelic and Monoallelic <i>NTHL1</i> and <i>MSH3</i> Variant Carriers From a Pan-Cancer Patient Population. JCO Precision Oncology, 2021, 5, 455-465.	3.0	10
25	Variation in the Attitudes of Medical Oncologists Toward Research Biopsies in Patients With Metastatic Breast Cancer. Oncologist, 2015, 20, 992-1000.	3.7	8
26	A Coordinated Clinical Center for Young Onset Colorectal Cancer. Oncologist, 2021, 26, 625-629.	3.7	8
27	Factors Influencing Patient Preferences for Telehealth Cancer Genetic Counseling During the COVID-19 Pandemic. JCO Oncology Practice, 2022, 18, e462-e471.	2.9	8
28	Understanding Inherited Risk in Unselected Newly Diagnosed Patients With Endometrial Cancer. JCO Precision Oncology, 2019, 3, 1-15.	3.0	7
29	Paired Tumor-Normal Sequencing Provides Insights into TP53-Related Cancer Spectrum in Li-Fraumeni Patients. Journal of the National Cancer Institute, 2021, , .	6.3	6
30	Induction Chemotherapy Reduces Patient-reported Toxicities During Neoadjuvant Chemoradiation with Intensity Modulated Radiotherapy for Rectal Cancer. Clinical Colorectal Cancer, 2019, 18, 167-174.	2.3	3
31	Quantitative assessment of tumor-infiltrating lymphocytes in mismatch repair proficient colon cancer. Oncolmmunology, 2020, 9, 1841948.	4.6	3
32	<i>ATM</i> Germline-Mutated Gastroesophageal Junction Adenocarcinomas: Clinical Descriptors, Molecular Characteristics, and Potential Therapeutic Implications. Journal of the National Cancer Institute, 2022, 114, 761-770.	6.3	3
33	Intratumoral T-cell repertoires in DNA mismatch repair-proficient and -deficient colon tumors containing high or low numbers of tumor-infiltrating lymphocytes. Oncolmmunology, 2022, 11, 2054757.	4.6	3
34	Simplified Graded Infusion Strategy for Mitigation of Oxaliplatin Hypersensitivity. Clinical Colorectal Cancer, 2022, , .	2.3	2
35	Primary Clonal Loss of Mismatch Repair Protein on Immunohistochemistry: A Pattern of Abnormality That Warrants Genetic Workup. JCO Precision Oncology, 2022, ,	3.0	1