Taina T Nieminen

List of Publications by Year
in descending order

18	635	567281 15	888059 17
papers	citations	h-index	g-index

19	19	19	1214
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	A novel essential splice site variant in SPTB in a large hereditary spherocytosis family. Molecular Genetics & Genomic Medicine, 2021, 9, e1641.	1.2	2
2	Updates in the field of hereditary nonpolyposis colorectal cancer. Expert Review of Gastroenterology and Hepatology, 2020, 14, 707-720.	3.0	18
3	Thyroid Carcinomas That Occur in Familial Adenomatous Polyposis Patients Recurrently Harbor Somatic Variants inAPC,BRAF, andKTM2D. Thyroid, 2020, 30, 380-388.	4.5	18
4	A Truncating Germline Mutation of TINF2 in Individuals with Thyroid Cancer or Melanoma Results in Longer Telomeres. Thyroid, 2020, 30, 204-213.	4.5	27
5	Identification of Rare Variants Predisposing to Thyroid Cancer. Thyroid, 2019, 29, 946-955.	4.5	41
6	Biallelic germline nonsense variant of MLH3 underlies polyposis predisposition. Genetics in Medicine, 2019, 21, 1868-1873.	2.4	39
7	Sequencing of Lynch syndrome tumors reveals the importance of epigenetic alterations. Oncotarget, 2017, 8, 108020-108030.	1.8	18
8	Desmoid tumor patients carry an elevated risk of familial adenomatous polyposis. Journal of Surgical Oncology, 2016, 113, 209-212.	1.7	29
9	Pseudoexons provide a mechanism for allele-specific expression of APC in familial adenomatous polyposis. Oncotarget, 2016, 7, 70685-70698.	1.8	17
10	Methyltransferase expression and tumor suppressor gene methylation in sporadic and familial colorectal cancer. Genes Chromosomes and Cancer, 2015, 54, 776-787.	2.8	14
11	Reply. Gastroenterology, 2015, 148, 259.	1.3	0
12	Germline Mutation of RPS20, Encoding a Ribosomal Protein, Causes Predisposition to Hereditary Nonpolyposis Colorectal Carcinoma Without DNA Mismatch Repair Deficiency. Gastroenterology, 2014, 147, 595-598.e5.	1.3	143
13	Promoterâ€specific alterations of APC are a rare cause for mutationâ€negative familial adenomatous polyposis. Genes Chromosomes and Cancer, 2014, 53, 857-864.	2.8	38
14	Breast carcinoma and Lynch syndrome: molecular analysis of tumors arising in mutation carriers, non-carriers, and sporadic cases. Breast Cancer Research, 2012, 14, R90.	5.0	39
15	Distinct Genetic and Epigenetic Signatures of Colorectal Cancers According to Ethnic Origin. Cancer Epidemiology Biomarkers and Prevention, 2012, 21, 202-211.	2.5	24
16	LINE-1 hypomethylation in familial and sporadic cancer. Journal of Molecular Medicine, 2012, 90, 827-835.	3.9	40
17	BMPR1A Mutations in Hereditary Nonpolyposis Colorectal Cancer Without Mismatch Repair Deficiency. Gastroenterology, 2011, 141, e23-e26.	1.3	47
18	Molecular Analysis of Endometrial Tumorigenesis: Importance of Complex Hyperplasia Regardless of Atypia. Clinical Cancer Research, 2009, 15, 5772-5783.	7.0	80