Anne-Bine S Skytte

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/2426932/publications.pdf

Version: 2024-02-01

37 papers	2,623 citations	430874 18 h-index	345221 36 g-index
37 all docs	37 docs citations	37 times ranked	6867 citing authors

#	Article	IF	Citations
1	Multiple independent variants at the TERT locus are associated with telomere length and risks of breast and ovarian cancer. Nature Genetics, 2013, 45, 371-384.	21.4	493
2	Association of Type and Location ofBRCA1andBRCA2Mutations With Risk of Breast and Ovarian Cancer. JAMA - Journal of the American Medical Association, 2015, 313, 1347.	7.4	390
3	Cancer Risks Associated With Germline PALB2 Pathogenic Variants: An International Study of 524 Families. Journal of Clinical Oncology, 2020, 38, 674-685.	1.6	270
4	Genome-Wide Association Study in BRCA1 Mutation Carriers Identifies Novel Loci Associated with Breast and Ovarian Cancer Risk. PLoS Genetics, 2013, 9, e1003212.	3.5	244
5	Mutational spectrum in a worldwide study of 29,700 families withBRCA1orBRCA2mutations. Human Mutation, 2018, 39, 593-620.	2.5	224
6	Targeted Prostate Cancer Screening in BRCA1 and BRCA2 Mutation Carriers: Results from the Initial Screening Round of the IMPACT Study. European Urology, 2014, 66, 489-499.	1.9	195
7	Prediction of Breast and Prostate Cancer Risks in Male BRCA1 and BRCA2 Mutation Carriers Using Polygenic Risk Scores. Journal of Clinical Oncology, 2017, 35, 2240-2250.	1.6	152
8	The Extended Clinical Phenotype of 26 Patients with Chronic Mucocutaneous Candidiasis due to Gain-of-Function Mutations in STAT1. Journal of Clinical Immunology, 2016, 36, 73-84.	3.8	124
9	Comprehensive Cancer-Predisposition Gene Testing in an Adult Multiple Primary Tumor Series Shows a Broad Range of Deleterious Variants and Atypical Tumor Phenotypes. American Journal of Human Genetics, 2018, 103, 3-18.	6.2	46
10	Multilocus Inherited Neoplasia Alleles Syndrome. JAMA Oncology, 2016, 2, 373.	7.1	43
11	Estimating the effect size of the 15Q11.2 BP1â€“BP2 deletion and its contribution to neurodevelopmental symptoms: recommendations for practice. Journal of Medical Genetics, 2019, 56, 701-710.	3.2	43
12	Dosage Changes of a Segment at 17p13.1 Lead to Intellectual Disability and Microcephaly as a Result of Complex Genetic Interaction of Multiple Genes. American Journal of Human Genetics, 2014, 95, 565-578.	6.2	40
13	Association of Genomic Domains in BRCA1 and BRCA2 with Prostate Cancer Risk and Aggressiveness. Cancer Research, 2020, 80, 624-638.	0.9	39
14	Ovarian cancer susceptibility alleles and risk of ovarian cancer inBRCA1andBRCA2mutation carriers. Human Mutation, 2012, 33, 690-702.	2.5	34
15	Transcriptomeâ€wide association study of breast cancer risk by estrogenâ€receptor status. Genetic Epidemiology, 2020, 44, 442-468.	1.3	32
16	Populationâ€based survey of cancer risks in chromosome 3 translocation carriers. Genes Chromosomes and Cancer, 2010, 49, 52-58.	2.8	27
17	Birtâ€“Haggâ€“DubÃ© syndrome: a case report and a review of the literature. European Clinical Respiratory Journal, 2017, 4, 1292378.	1.5	24
18	Genetic screening of the FLCN gene identify six novel variants and a Danish founder mutation. Journal of Human Genetics, 2017, 62, 151-157.	2.3	19

#	Article	IF	Citations
19	Revised Danish guidelines for the cancer surveillance of patients with Cowden Syndrome. European Journal of Medical Genetics, 2020, 63, 103873.	1.3	19
20	<scp>UBE</scp> 2 <scp>QL</scp> 1 is Disrupted by a Constitutional Translocation Associated with Renal Tumor Predisposition and is a Novel Candidate Renal Tumor Suppressor Gene. Human Mutation, 2013, 34, 1650-1661.	2.5	18
21	New Pathogenic Germline Variants in Very Early Onset and Familial Colorectal Cancer Patients. Frontiers in Genetics, 2020, 11, 566266.	2.3	16
22	Identification of BRCA1-deficient ovarian cancers. Acta Obstetricia Et Gynecologica Scandinavica, 2011, 90, 593-599.	2.8	15
23	Childhood pneumothorax in Birtâ€Hoggâ€DubÃ© syndrome: A cohort study and review of the literature. Molecular Genetics & Genomic Medicine, 2018, 6, 332-338.	1.2	15
24	The use of expanded carrier screening of gamete donors. Human Reproduction, 2021, 36, 1702-1710.	0.9	14
25	Identification of eight novel SDHB, SDHC, SDHD germline variants in Danish pheochromocytoma/paraganglioma patients. Hereditary Cancer in Clinical Practice, 2016, 14, 13.	1.5	13
26	Increased risk of male cancer and identification of a potential prostate cancer cluster region in BRCA2. Acta OncolÃ³gica, 2016, 55, 38-44.	1.8	13
27	Diagnosis and Treatment of Blau Syndrome/Early-onset Sarcoidosis, an Autoinflammatory Granulomatous Disease, in an Infant. Acta Dermato-Venereologica, 2017, 97, 126-127.	1.3	13
28	BRCA1/BRCA2 founder mutations and cancer risks: impact in the western Danish population. Familial Cancer, 2016, 15, 507-512.	1.9	9
29	Impact of genetic counseling on the uptake of contralateral prophylactic mastectomy among younger women with breast cancer. Acta OncolÃ³gica, 2020, 59, 60-65.	1.8	9
30	Population frequencies of pathogenic alleles of BRCA1 and BRCA2: analysis of 173 Danish breast cancer pedigrees using the BOADICEA model. Familial Cancer, 2019, 18, 381-388.	1.9	8
31	Attitudes of sperm donors towards offspring, identity release and extended genetic screening. Reproductive BioMedicine Online, 2021, 43, 700-707.	2.4	6
32	Clinical characteristics and registry-validated extended pedigrees of germline TP53 mutation carriers in Denmark. PLoS ONE, 2018, 13, e0190050.	2.5	6
33	Familial idiopathic pulmonary fibrosis in a young female. Respiratory Medicine Case Reports, 2018, 24, 1-4.	0.4	4
34	No evidence of increased breast cancer risk for proven noncarriers from BRCA1 and BRCA2 families. Familial Cancer, 2016, 15, 523-528.	1.9	3
35	Genetically diagnosed Birtâ€“Hoggâ€“DubÃ© syndrome and familial cerebral cavernous malformations in the same individual: a case report. Familial Cancer, 2017, 16, 139-142.	1.9	2
36	A case of penta X syndrome caused by nondisjunction in maternal meiosis 1 and 2. Clinical Case Reports (discontinued), 2017, 5, 1136-1140.	0.5	1

#	Article	IF	Citations
37	AN IMPROVED METHOD FOR ZIKA VIRUS SCREENING IN A GAMETE BANK SETTING. Fertility and Sterility, 2020, 114, e277.	1.0	0