

Alan Donaldson

List of Publications by Year in descending order

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Version: 2024-02-01

11
papers

977
citations

1163117

8
h-index

1281871

11
g-index

11
all docs

11
docs citations

11
times ranked

2904
citing authors

#	ARTICLE	IF	CITATIONS
1	Association of Type and Location of <i>BRCA1</i> and <i>BRCA2</i> Mutations With Risk of Breast and Ovarian Cancer. <i>JAMA - Journal of the American Medical Association</i> , 2015, 313, 1347.	7.4	390
2	High Rate of Recurrent De Novo Mutations in Developmental and Epileptic Encephalopathies. <i>American Journal of Human Genetics</i> , 2017, 101, 664-685.	6.2	337
3	Prostate Cancer Risks for Male <i>BRCA1</i> and <i>BRCA2</i> Mutation Carriers: A Prospective Cohort Study. <i>European Urology</i> , 2020, 77, 24-35.	1.9	124
4	Association of Genomic Domains in <i>BRCA1</i> and <i>BRCA2</i> with Prostate Cancer Risk and Aggressiveness. <i>Cancer Research</i> , 2020, 80, 624-638.	0.9	39
5	A case-only study to identify genetic modifiers of breast cancer risk for <i>BRCA1/BRCA2</i> mutation carriers. <i>Nature Communications</i> , 2021, 12, 1078.	12.8	19
6	Breast and Prostate Cancer Risks for Male <i>BRCA1</i> and <i>BRCA2</i> Pathogenic Variant Carriers Using Polygenic Risk Scores. <i>Journal of the National Cancer Institute</i> , 2022, 114, 109-122.	6.3	19
7	Psychosocial effects of whole-body MRI screening in adult high-risk pathogenic <i>TP53</i> mutation carriers: a case-controlled study (SIGNIFY). <i>Journal of Medical Genetics</i> , 2020, 57, 226-236.	3.2	15
8	<i>SOX11</i> variants cause a neurodevelopmental disorder with infrequent ocular malformations and hypogonadotropic hypogonadism and with distinct DNA methylation profile. <i>Genetics in Medicine</i> , 2022, 24, 1261-1273.	2.4	14
9	Phenotypic expansion of the <i>BPTF</i> -related neurodevelopmental disorder with dysmorphic facies and distal limb anomalies. <i>American Journal of Medical Genetics, Part A</i> , 2021, 185, 1366-1378.	1.2	8
10	Prostate Cancer Risk by <i>BRCA2</i> Genomic Regions. <i>European Urology</i> , 2020, 78, 494-497.	1.9	6
11	<i>RRM1</i> variants cause a mitochondrial DNA maintenance disorder via impaired de novo nucleotide synthesis. <i>Journal of Clinical Investigation</i> , 2022, 132, .	8.2	6