

Alison Meynert

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/401574/publications.pdf>

Version: 2024-02-01

18
papers

4,626
citations

687363

13
h-index

888059

17
g-index

20
all docs

20
docs citations

20
times ranked

12429
citing authors

#	ARTICLE	IF	CITATIONS
1	A promoter-level mammalian expression atlas. <i>Nature</i> , 2014, 507, 462-470.	27.8	1,838
2	Genetic mechanisms of critical illness in COVID-19. <i>Nature</i> , 2021, 591, 92-98.	27.8	1,014
3	A small-cell lung cancer genome with complex signatures of tobacco exposure. <i>Nature</i> , 2010, 463, 184-190.	27.8	972
4	A prenylated dsRNA sensor protects against severe COVID-19. <i>Science</i> , 2021, 374, eabj3624.	12.6	124
5	Small latin squares, quasigroups, and loops. <i>Journal of Combinatorial Designs</i> , 2007, 15, 98-119.	0.6	107
6	Heterozygous Loss-of-Function Mutations in YAP1 Cause Both Isolated and Syndromic Optic Fissure Closure Defects. <i>American Journal of Human Genetics</i> , 2014, 94, 295-302.	6.2	93
7	Monoallelic and Biallelic Mutations in MAB21L2 Cause a Spectrum of Major Eye Malformations. <i>American Journal of Human Genetics</i> , 2014, 94, 915-923.	6.2	79
8	Heterochromatin delays CRISPR-Cas9 mutagenesis but does not influence the outcome of mutagenic DNA repair. <i>PLoS Biology</i> , 2018, 16, e2005595.	5.6	75
9	Pan-ancestry exome-wide association analyses of COVID-19 outcomes in 586,157 individuals. <i>American Journal of Human Genetics</i> , 2021, 108, 1350-1355.	6.2	72
10	DNA Polymerase Epsilon Deficiency Causes IMAGE Syndrome with Variable Immunodeficiency. <i>American Journal of Human Genetics</i> , 2018, 103, 1038-1044.	6.2	71
11	Structural Variants at the <i>BRCA1/2</i> Loci are a Common Source of Homologous Repair Deficiency in High-grade Serous Ovarian Carcinoma. <i>Clinical Cancer Research</i> , 2021, 27, 3201-3214.	7.0	27
12	ITPase deficiency causes a Martsolf-like syndrome with a lethal infantile dilated cardiomyopathy. <i>PLoS Genetics</i> , 2019, 15, e1007605.	3.5	25
13	Increased ultra-rare variant load in an isolated Scottish population impacts exonic and regulatory regions. <i>PLoS Genetics</i> , 2019, 15, e1008480.	3.5	17
14	Dynamic epigenetic changes to <i>VHL</i> occur with sunitinib in metastatic clear cell renal cancer. <i>Oncotarget</i> , 2016, 7, 25241-25250.	1.8	14
15	Ancient DNA at the edge of the world: Continental immigration and the persistence of Neolithic male lineages in Bronze Age Orkney. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2022, 119, .	7.1	12
16	Picking Pyknons out of the Human Genome. <i>Cell</i> , 2006, 125, 836-838.	28.9	9
17	<i>In Vivo</i> Modeling of Patient Genetic Heterogeneity Identifies New Ways to Target Cholangiocarcinoma. <i>Cancer Research</i> , 2022, 82, 1548-1559.	0.9	8
18	Molecular stratification of endometrioid ovarian carcinomas.. <i>Journal of Clinical Oncology</i> , 2019, 37, 5553-5553.	1.6	0