Katherine Dixon

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/5905747/publications.pdf

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14	405	1163117 8	1125743 13
papers	citations	h-index	g-index

15	15	15	1078
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Specifications of the ACMG/AMP variant curation guidelines for the analysis of germline CDH1 sequence variants. Human Mutation, 2018, 39, 1553-1568.	2.5	138
2	Pan-cancer analysis of advanced patient tumors reveals interactions between therapy and genomic landscapes. Nature Cancer, 2020, 1, 452-468.	13.2	103
3	Improved structural variant interpretation for hereditary cancer susceptibility using long-read sequencing. Genetics in Medicine, 2020, 22, 1892-1897.	2.4	42
4	Endometrial Cancer Molecular Risk Stratification is Equally Prognostic for Endometrioid Ovarian Carcinoma. Clinical Cancer Research, 2020, 26, 5400-5410.	7.0	41
5	Burden of hereditary cancer susceptibility in unselected patients with pancreatic ductal adenocarcinoma referred for germline screening. Cancer Medicine, 2020, 9, 4004-4013.	2.8	25
6	Insights into interplay between rexinoid signaling and myogenic regulatory factor-associated chromatin state in myogenic differentiation. Nucleic Acids Research, 2017, 45, 11236-11248.	14.5	18
7	Modelling hereditary diffuse gastric cancer initiation using transgenic mouseâ€derived gastric organoids and singleâ€cell sequencing. Journal of Pathology, 2021, 254, 254-264.	4.5	11
8	Retinoid X Receptor-selective Signaling in the Regulation of Akt/Protein Kinase B Isoform-specific Expression. Journal of Biological Chemistry, 2016, 291, 3090-3099.	3.4	10
9	Establishing a Framework for the Clinical Translation of Germline Findings in Precision Oncology. JNCI Cancer Spectrum, 2020, 4, pkaa045.	2.9	6
10	Longâ€read genome sequencing resolves a complex 13q structural variant associated with syndromic anophthalmia. American Journal of Medical Genetics, Part A, 2022, 188, 1589-1594.	1.2	4
11	Gene expression profiling discerns molecular pathways elicited by ligand signaling to enhance the specification of embryonic stem cells into skeletal muscle lineage. Cell and Bioscience, 2017, 7, 23.	4.8	3
12	An infant with congenital respiratory insufficiency and diaphragmatic paralysis: A novel <scp>BICD2</scp> phenotype?. American Journal of Medical Genetics, Part A, 2022, 188, 926-930.	1.2	3
13	Integrating Tumor Sequencing Into Clinical Practice for Patients With Mismatch Repair-Deficient Lynch Syndrome Spectrum Cancers. Clinical and Translational Gastroenterology, 2021, 12, e00397.	2.5	1
14	Rare APC promoter 1B variants in gastric cancer kindreds unselected for fundic gland polyposis. Gut, 2021, 70, 1415-1416.	12.1	0