Yun-Ching Chen

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/4953322/publications.pdf

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16 papers	719 citations	933447 10 h-index	996975 15 g-index
16	16	16	2090
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	Ultra-Small Lung Cysts Impair Diffusion Without Obstructing Air Flow in Lymphangioleiomyomatosis. Chest, 2021, 160, 199-208.	0.8	7
2	Toward best practice in cancer mutation detection with whole-genome and whole-exome sequencing. Nature Biotechnology, 2021, 39, 1141-1150.	17.5	66
3	Aberrant DNA methylation defines isoform usage in cancer, with functional implications. PLoS Computational Biology, 2019, 15, e1007095.	3.2	16
4	Diverging Clonal Evolution during Sequential Therapy with Chemoimmunotherapy Followed By BTK Inhibitors. Blood, 2019, 134, 850-850.	1.4	1
5	Spatial Genomic Heterogeneity in Chronic Lymphocytic Leukemia. Blood, 2019, 134, 3017-3017.	1.4	0
6	Human retinoic acid–regulated CD161+ regulatory T cells support wound repair in intestinal mucosa. Nature Immunology, 2018, 19, 1403-1414.	14.5	86
7	High density lipoprotein proteome is associated with cardiovascular risk factors and atherosclerosis burden as evaluated by coronary CT angiography. Atherosclerosis, 2018, 278, 278-285.	0.8	39
8	Matching phenotypes to whole genomes: Lessons learned from four iterations of the personal genome project community challenges. Human Mutation, 2017, 38, 1266-1276.	2.5	14
9	Significant associations between driver gene mutations and DNA methylation alterations across many cancer types. PLoS Computational Biology, 2017, 13, e1005840.	3.2	39
10	CpG island methylator phenotype in adenocarcinomas from the digestive tract: Methods, conclusions, and controversies. World Journal of Gastrointestinal Oncology, 2017, 9, 105.	2.0	9
11	Exome Sequencing of Familial Bipolar Disorder. JAMA Psychiatry, 2016, 73, 590.	11.0	97
12	Whole Genome Sequencing Defines the Genetic Heterogeneity of Familial Pancreatic Cancer. Cancer Discovery, 2016, 6, 166-175.	9.4	282
13	A Probabilistic Model to Predict Clinical Phenotypic Traits from Genome Sequencing. PLoS Computational Biology, 2014, 10, e1003825.	3.2	10
14	A Hybrid Likelihood Model for Sequence-Based Disease Association Studies. PLoS Genetics, 2013, 9, e1003224.	3.5	19
15	Exonic DNA Sequencing of ERBB4 in Bipolar Disorder. PLoS ONE, 2011, 6, e20242.	2.5	13
16	Profile analysis of expressed sequence tags derived from the ovary of tilapia, Oreochromis mossambicus. Aquaculture, 2006, 251, 537-548.	3.5	21