Nir Pillar

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/9564922/publications.pdf

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23 papers	462 citations	840776 11 h-index	713466 21 g-index
24 all docs	24 docs citations	24 times ranked	1186 citing authors

#	Article	IF	Citations
1	Exosomal microRNAs derived from colorectal cancer-associated fibroblasts: role in driving cancer progression. Aging, 2017, 9, 2666-2694.	3.1	112
2	Refining the phenotype associated with GNB1 mutations: Clinical data on 18 newly identified patients and review of the literature. American Journal of Medical Genetics, Part A, 2018, 176, 2259-2275.	1.2	47
3	Comparison of breast cancer metastasis models reveals a possible mechanism of tumor aggressiveness. Cell Death and Disease, 2018, 9, 1040.	6.3	36
4	Distinctive pattern of let-7 family microRNAs in aggressive carcinoma of the oral tongue in young patients. Oncology Letters, 2016, 12, 1729-1736.	1.8	34
5	Circulating MicroRNAs: a Potential Biomarker for Cardiac Damage, Inflammatory Response, and Left Ventricular Function Recovery in Pediatric Viral Myocarditis. Journal of Cardiovascular Translational Research, 2018, 11, 319-328.	2.4	34
6	The possible involvement of microRNAs in preeclampsia and gestational diabetes mellitus. Best Practice and Research in Clinical Obstetrics and Gynaecology, 2015, 29, 176-182.	2.8	30
7	Analysis of microRNAs in familial Mediterranean fever. PLoS ONE, 2018, 13, e0197829.	2.5	22
8	Loss of Protocadherinâ€12<scp>L</scp>eads to <scp>D</scp>iencephalicâ€<scp>M</scp>esencephalic <scp>J</scp>unction <scp>D</scp>ysplasia <scp>S</scp>yndrome. Annals of Neurology, 2018, 84, 638-647.	5.3	19
9	Mutations in the NEB gene cause fetal akinesia/arthrogryposis multiplex congenita. Prenatal Diagnosis, 2017, 37, 144-150.	2.3	18
10	A rare variant in the FHL1 gene associated with X-linked recessive hypoparathyroidism. Human Genetics, 2017, 136, 835-845.	3.8	17
11	MicroRNAs as predictors for CNS relapse of systemic diffuse large B-cell lymphoma. Oncotarget, 2017, 8, 86020-86030.	1.8	13
12	Hypereosinophilic Syndrome With Cardiac Involvement: Early Diagnosis by Cardiac Magnetic Resonance Imaging. Canadian Journal of Cardiology, 2012, 28, 515.e11-515.e13.	1.7	11
13	Dual inhibition of ABCE1 and LCP1 by microRNA-96 results in an additive effect in breast cancer mouse model. Oncotarget, 2019, 10, 2086-2094.	1.8	11
14	MicroRNAs in ascending thoracic aortic aneurysms. Bioscience Reports, 2020, 40, .	2.4	10
15	Characterization of MicroRNA and Gene Expression Profiles Following Ricin Intoxication. Toxins, 2019, 11, 250.	3.4	8
16	The black sheep of the family- whole-exome sequencing in family of lithium response discordant bipolar monozygotic twins. European Neuropsychopharmacology, 2020, 34, 19-27.	0.7	8
17	Actionable clinical decisions based on comprehensive genomic evaluation in asymptomatic adults. Molecular Genetics & Genomic Medicine, 2015, 3, 433-439.	1.2	7
18	<scp>SMYD</scp>1 is the underlying gene for the AnWjâ€negative blood group phenotype. European Journal of Haematology, 2018, 101, 496-501.	2.2	7

#	Article	IF	Citations
19	X-linked elliptocytosis with impaired growth is related to mutated AMMECR1. Gene, 2017, 606, 47-52.	2.2	6
20	The Contribution of MicroRNAs to the Inflammatory and Neoplastic Characteristics of Erdheimâ€“Chester Disease. Cancers, 2020, 12, 3240.	3.7	5
21	Blood transcriptional response to treatment-resistant depression during electroconvulsive therapy. Journal of Psychiatric Research, 2021, 141, 92-103.	3.1	5
22	Sequencing your genome: your future is here, but are you sure you want to know it?. Genetical Research, 2014, 96, e006.	0.9	2
23	Assessing the involvement of the placental microbiome and virome in preeclampsia using non coding RNA sequencing. Journal of Perinatal Medicine, 2021, 49, 1071-1083.	1.4	0