

Nir Pillar

List of Publications by Year in descending order

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Version: 2024-02-01

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	Assessing the involvement of the placental microbiome and virome in preeclampsia using non coding RNA sequencing. <i>Journal of Perinatal Medicine</i> , 2021, 49, 1071-1083.	1.4	0
2	Blood transcriptional response to treatment-resistant depression during electroconvulsive therapy. <i>Journal of Psychiatric Research</i> , 2021, 141, 92-103.	3.1	5
3	The Contribution of MicroRNAs to the Inflammatory and Neoplastic Characteristics of Erdheim-Chester Disease. <i>Cancers</i> , 2020, 12, 3240.	3.7	5
4	The black sheep of the family- whole-exome sequencing in family of lithium response discordant bipolar monozygotic twins. <i>European Neuropsychopharmacology</i> , 2020, 34, 19-27.	0.7	8
5	MicroRNAs in ascending thoracic aortic aneurysms. <i>Bioscience Reports</i> , 2020, 40, .	2.4	10
6	Characterization of MicroRNA and Gene Expression Profiles Following Ricin Intoxication. <i>Toxins</i> , 2019, 11, 250.	3.4	8
7	Dual inhibition of ABCE1 and LCP1 by microRNA-96 results in an additive effect in breast cancer mouse model. <i>Oncotarget</i> , 2019, 10, 2086-2094.	1.8	11
8	Comparison of breast cancer metastasis models reveals a possible mechanism of tumor aggressiveness. <i>Cell Death and Disease</i> , 2018, 9, 1040.	6.3	36
9	Loss of <i>Protocadherin12</i> leads to <i>Diencephalic Mesencephalic Junction Dysplasia Syndrome</i> . <i>Annals of Neurology</i> , 2018, 84, 638-647.	5.3	19
10	Refining the phenotype associated with <i>GNB1</i> mutations: Clinical data on 18 newly identified patients and review of the literature. <i>American Journal of Medical Genetics, Part A</i> , 2018, 176, 2259-2275.	1.2	47
11	Analysis of microRNAs in familial Mediterranean fever. <i>PLoS ONE</i> , 2018, 13, e0197829.	2.5	22
12	<i>SMYD1</i> is the underlying gene for the AnWj-negative blood group phenotype. <i>European Journal of Haematology</i> , 2018, 101, 496-501.	2.2	7
13	Circulating MicroRNAs: a Potential Biomarker for Cardiac Damage, Inflammatory Response, and Left Ventricular Function Recovery in Pediatric Viral Myocarditis. <i>Journal of Cardiovascular Translational Research</i> , 2018, 11, 319-328.	2.4	34
14	X-linked elliptocytosis with impaired growth is related to mutated AMMECR1. <i>Gene</i> , 2017, 606, 47-52.	2.2	6
15	A rare variant in the <i>FHL1</i> gene associated with X-linked recessive hypoparathyroidism. <i>Human Genetics</i> , 2017, 136, 835-845.	3.8	17
16	Mutations in the <i>NEB</i> gene cause fetal akinesia/arthrogryposis multiplex congenita. <i>Prenatal Diagnosis</i> , 2017, 37, 144-150.	2.3	18
17	Exosomal microRNAs derived from colorectal cancer-associated fibroblasts: role in driving cancer progression. <i>Aging</i> , 2017, 9, 2666-2694.	3.1	112
18	MicroRNAs as predictors for CNS relapse of systemic diffuse large B-cell lymphoma. <i>Oncotarget</i> , 2017, 8, 86020-86030.	1.8	13

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19	Distinctive pattern of let-7 family microRNAs in aggressive carcinoma of the oral tongue in young patients. <i>Oncology Letters</i> , 2016, 12, 1729-1736.	1.8	34
20	Actionable clinical decisions based on comprehensive genomic evaluation in asymptomatic adults. <i>Molecular Genetics & Genomic Medicine</i> , 2015, 3, 433-439.	1.2	7
21	The possible involvement of microRNAs in preeclampsia and gestational diabetes mellitus. <i>Best Practice and Research in Clinical Obstetrics and Gynaecology</i> , 2015, 29, 176-182.	2.8	30
22	Sequencing your genome: your future is here, but are you sure you want to know it?. <i>Genetical Research</i> , 2014, 96, e006.	0.9	2
23	Hypereosinophilic Syndrome With Cardiac Involvement: Early Diagnosis by Cardiac Magnetic Resonance Imaging. <i>Canadian Journal of Cardiology</i> , 2012, 28, 515.e11-515.e13.	1.7	11