Daniel Duran

List of Publications by Year in descending order

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Πληίει Πιίβλη

#	Article	IF	CITATIONS
1	Associations of meningioma molecular subgroup and tumor recurrence. Neuro-Oncology, 2021, 23, 783-794.	1.2	83
2	Protein kinase <scp>D1</scp> variant associated with human epilepsy and peripheral nerve hypermyelination. Clinical Genetics, 2021, 100, 176-186.	2.0	1
3	Coronavirus Disease 2019 and Pituitary Apoplexy: A Single-Center Case Series and Review of the Literature. World Neurosurgery, 2021, 152, e678-e687.	1.3	20
4	<i>DIAPH1</i> Variants in Non–East Asian Patients With Sporadic Moyamoya Disease. JAMA Neurology, 2021, 78, 993.	9.0	33
5	Modulation of brain cation-Clâ^' cotransport via the SPAK kinase inhibitor ZT-1a. Nature Communications, 2020, 11, 78.	12.8	69
6	Genomic alterations in Turcot syndrome: Insights from whole exome sequencing. Journal of the Neurological Sciences, 2020, 417, 117056.	0.6	1
7	Correlations between genomic subgroup and clinical features in a cohort of more than 3000 meningiomas. Journal of Neurosurgery, 2020, 133, 1345-1354.	1.6	83
8	PATH-39. ASSOCIATIONS OF GENOMIC SUBGROUP WITH RECURRENCE IN LOW-GRADE MENINGIOMAS. Neuro-Oncology, 2020, 22, ii172-ii173.	1.2	1
9	EphrinB2-EphB4-RASA1 Signaling in Human Cerebrovascular Development and Disease. Trends in Molecular Medicine, 2019, 25, 265-286.	6.7	39
10	GENE-56. MENINGIOMA GENOMIC SUBGROUP AS A PREDICTOR OF POST-OPERATIVE PATIENT OUTCOMES: IMPLICATIONS FOR TREATMENT AND FOLLOW-UP. Neuro-Oncology, 2019, 21, vi109-vi110.	1.2	0
11	Mutations in Chromatin Modifier and Ephrin Signaling Genes in Vein of Galen Malformation. Neuron, 2019, 101, 429-443.e4.	8.1	56
12	MNGI-09. MENINGIOMA WITH MULTIPLE DRIVERS: GENOMIC LANDSCAPE AND CLINICAL CORRELATIONS. Neuro-Oncology, 2019, 21, vi141-vi141.	1.2	0
13	9p24 triplication in syndromic hydrocephalus with diffuse villous hyperplasia of the choroid plexus. Journal of Physical Education and Sports Management, 2018, 4, a003145.	1.2	8
14	A novel association of campomelic dysplasia and hydrocephalus with an unbalanced chromosomal translocation upstream of <i>SOX9</i> . Journal of Physical Education and Sports Management, 2018, 4, a002766.	1.2	8
15	De Novo Mutation in Genes Regulating Neural Stem Cell Fate in Human Congenital Hydrocephalus. Neuron, 2018, 99, 302-314.e4.	8.1	112
16	De novo <i>MYH9</i> mutation in congenital scalp hemangioma. Journal of Physical Education and Sports Management, 2018, 4, a002998.	1.2	9
17	Integrated genomic analyses of de novo pathways underlying atypical meningiomas. Nature Communications, 2017, 8, 14433.	12.8	156
18	Xp22.2 Chromosomal Duplication in Familial Intracranial Arachnoid Cyst. JAMA Neurology, 2017, 74, 1503.	9.0	6

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19	Inflammation-dependent cerebrospinal fluid hypersecretion by the choroid plexus epithelium in posthemorrhagic hydrocephalus. Nature Medicine, 2017, 23, 997-1003.	30.7	256
20	Malignant Cerebellar Edema Subsequent to Accidental Prescription Opioid Intoxication in Children. Frontiers in Neurology, 2017, 8, 362.	2.4	16
21	Digenic mutations of human OCRL paralogs in Dent's disease type 2 associated with Chiari I malformation. Human Genome Variation, 2016, 3, 16042.	0.7	8
22	Recurrent somatic mutations in POLR2A define a distinct subset of meningiomas. Nature Genetics, 2016, 48, 1253-1259.	21.4	265
23	Cerebrospinal fluid hypersecretion in pediatric hydrocephalus. Neurosurgical Focus, 2016, 41, E10.	2.3	59
24	Functional kinomics establishes a critical node of volume-sensitive cation-Clâ^ cotransporter regulation in the mammalian brain. Scientific Reports, 2016, 6, 35986.	3.3	38
25	Peripheral motor neuropathy is associated with defective kinase regulation of the KCC3 cotransporter. Science Signaling, 2016, 9, ra77.	3.6	46
26	Prognostic Factors in Patients with Primary Hemangiopericytomas of the Central Nervous System: A Series of 103 Cases at a Single Institution. World Neurosurgery, 2016, 90, 414-419.	1.3	14