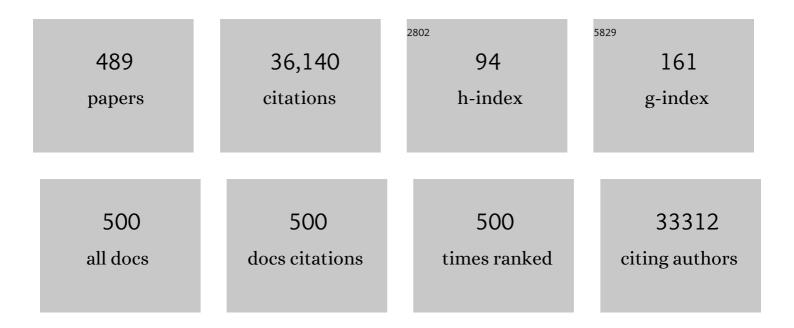
## Eleonora Aronica

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/4091646/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	Genetic pathogenesis of the epileptogenic lesions in Tuberous Sclerosis Complex: Therapeutic targeting of the mTOR pathway. Epilepsy and Behavior, 2022, 131, 107713.	1.7	10
2	Interactome screening of <i>C9orf72</i> dipeptide repeats reveals VCP sequestration and functional impairment by polyGA. Brain, 2022, 145, 684-699.	7.6	15
3	Evolution of electroencephalogram in infants with tuberous sclerosis complex and neurodevelopmental outcome: a prospective cohort study. Developmental Medicine and Child Neurology, 2022, 64, 495-501.	2.1	3
4	Distinct DNA Methylation Patterns of Subependymal Giant Cell Astrocytomas in Tuberous Sclerosis Complex. Cellular and Molecular Neurobiology, 2022, 42, 2863-2892.	3.3	1
5	A Novel SCA3 Knock-in Mouse Model Mimics the Human SCA3 Disease Phenotype Including Neuropathological, Behavioral, and Transcriptional Abnormalities Especially in Oligodendrocytes. Molecular Neurobiology, 2022, 59, 495-522.	4.0	22
6	Down-regulation of the brain-specific cell-adhesion molecule contactin-3 in tuberous sclerosis complex during the early postnatal period. Journal of Neurodevelopmental Disorders, 2022, 14, 8.	3.1	4
7	Increased expression of complement components in tuberous sclerosis complex and focal cortical dysplasia type 2B brain lesions. Epilepsia, 2022, 63, 364-374.	5.1	10
8	Evidence of SARS-CoV-2 in nasal brushings and olfactory mucosa biopsies of COVID-19 patients. PLoS ONE, 2022, 17, e0266740.	2.5	6
9	Pathophysiological Mechanisms and Treatment of Dermatomyositis and Immune Mediated Necrotizing Myopathies: A Focused Review. International Journal of Molecular Sciences, 2022, 23, 4301.	4.1	16
10	Clinical characteristics and outcome in muscular sarcoidosis: a retrospective cohort study and literature review. Neuromuscular Disorders, 2022, 32, 557-563.	0.6	5
11	Differentially Expressed miRNAs in Age-Related Neurodegenerative Diseases: A Meta-Analysis. Genes, 2022, 13, 1034.	2.4	4
12	Unexpected Effect of IL- $1\hat{1}^2$ on the Function of GABAA Receptors in Pediatric Focal Cortical Dysplasia. Brain Sciences, 2022, 12, 807.	2.3	5
13	The <scp>ILAE</scp> consensus classification of focal cortical dysplasia: An update proposed by an ad hoc task force of the <scp>ILAE</scp> diagnostic methods commission. Epilepsia, 2022, 63, 1899-1919.	5.1	88
14	Subependymal giant cell astrocytomas are characterized by mTORC1 hyperactivation, a very low somatic mutation rate, and a unique gene expression profile. Modern Pathology, 2021, 34, 264-279.	5.5	16
15	Primary mismatch repair deficient IDH-mutant astrocytoma (PMMRDIA) is a distinct type with a poor prognosis. Acta Neuropathologica, 2021, 141, 85-100.	7.7	52
16	Prevention of Epilepsy in Infants with Tuberous Sclerosis Complex in the <scp>EPISTOP</scp> Trial. Annals of Neurology, 2021, 89, 304-314.	5.3	137
17	Intravenous immunoglobulins as first-line treatment in idiopathic inflammatory myopathies: a pilot study. Rheumatology, 2021, 60, 1784-1792.	1.9	25
18	Seizure activity and brain damage in a model of focal nonâ€convulsive <i>status epilepticus</i> . Neuropathology and Applied Neurobiology, 2021, 47, 679-693.	3.2	9

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19	Frequent SLC35A2 brain mosaicism in mild malformation of cortical development with oligodendroglial hyperplasia in epilepsy (MOGHE). Acta Neuropathologica Communications, 2021, 9, 3.	5.2	62
20	GATOR1-related focal cortical dysplasia in epilepsy surgery patients and their families: A possible gradient in severity?. European Journal of Paediatric Neurology, 2021, 30, 88-96.	1.6	16
21	Immunohistochemical Detection of Neural Stem Cells and Glioblastoma Stem Cells in the Subventricular Zone of Glioblastoma Patients. Journal of Histochemistry and Cytochemistry, 2021, 69, 349-364.	2.5	12
22	Human brain pathology in myotonic dystrophy type 1: A systematic review. Neuropathology, 2021, 41, 3-20.	1.2	21
23	Early epileptiform EEG activity in infants with tuberous sclerosis complex predicts epilepsy and neurodevelopmental outcomes. Epilepsia, 2021, 62, 1208-1219.	5.1	19
24	Upregulation of the pathogenic transcription factor SPI1/PU.1 in tuberous sclerosis complex and focal cortical dysplasia by oxidative stress. Brain Pathology, 2021, 31, e12949.	4.1	11
25	Angiocentric glioma-associated seizures: The possible role of EATT2, pyruvate carboxylase and glutamine synthetase. Seizure: the Journal of the British Epilepsy Association, 2021, 86, 152-154.	2.0	8
26	Dissecting the Molecular Determinants of GABAA Receptors Current Rundown, a Hallmark of Refractory Human Epilepsy. Brain Sciences, 2021, 11, 441.	2.3	4
27	Hsp90â€mediated regulation of DYRK3 couples stress granule disassembly and growth via mTORC1 signaling. EMBO Reports, 2021, 22, e51740.	4.5	41
28	A serum microRNA sequence reveals fragile X protein pathology in amyotrophic lateral sclerosis. Brain, 2021, 144, 1214-1229.	7.6	8
29	Proteomics and Transcriptomics of the Hippocampus and Cortex in SUDEP and High-Risk SUDEP Patients. Neurology, 2021, 96, e2639-e2652.	1.1	24
30	Altered perivascular fibroblast activity precedes ALS disease onset. Nature Medicine, 2021, 27, 640-646.	30.7	69
31	Toward a better definition of focal cortical dysplasia: An iterative histopathological and genetic agreement trial. Epilepsia, 2021, 62, 1416-1428.	5.1	54
32	Pathomechanisms of ALS8: altered autophagy and defective RNA binding protein (RBP) homeostasis due to the VAPB P56S mutation. Cell Death and Disease, 2021, 12, 466.	6.3	13
33	Expression and Cell Type-specific Localization of Inflammasome Sensors in the Spinal Cord of SOD1(G93A) Mice and Sporadic Amyotrophic lateral sclerosis Patients. Neuroscience, 2021, 463, 288-302.	2.3	8
34	Balloon cells promote immune system activation in focal cortical dysplasia type 2b. Neuropathology and Applied Neurobiology, 2021, 47, 826-839.	3.2	14
35	MicroRNAâ€34a activation in tuberous sclerosis complex during early brain development may lead to impaired corticogenesis. Neuropathology and Applied Neurobiology, 2021, 47, 796-811.	3.2	5
36	Impaired myelin production due to an intrinsic failure of oligodendrocytes in mTORpathies. Neuropathology and Applied Neurobiology, 2021, 47, 812-825.	3.2	13

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37	Toward a refined genotype–phenotype classification scheme for the international consensus classification of Focal Cortical Dysplasia. Brain Pathology, 2021, 31, e12956.	4.1	22
38	Seizure-mediated iron accumulation and dysregulated iron metabolism after status epilepticus and in temporal lobe epilepsy. Acta Neuropathologica, 2021, 142, 729-759.	7.7	31
39	Interplay between immunity and amyotrophic lateral sclerosis: Clinical impact. Neuroscience and Biobehavioral Reviews, 2021, 127, 958-978.	6.1	22
40	ApoE4 disrupts interaction of sortilin with fatty acid-binding protein 7 essential to promote lipid signaling. Journal of Cell Science, 2021, 134, .	2.0	11
41	Complement factor C1q mediates sleep spindle loss and epileptic spikes after mild brain injury. Science, 2021, 373, eabj2685.	12.6	55
42	CXCL1-CXCR1/2 signaling is induced in human temporal lobe epilepsy and contributes to seizures in a murine model of acquired epilepsy. Neurobiology of Disease, 2021, 158, 105468.	4.4	15
43	A Selective Competitive Inhibitor of Aldehyde Dehydrogenase 1A3 Hinders Cancer Cell Growth, Invasiveness and Stemness In Vitro. Cancers, 2021, 13, 356.	3.7	21
44	Neurite Outgrowth Inhibitor (NogoA) Is Upregulated in White Matter Lesions of Complex Cortical Malformations. Journal of Neuropathology and Experimental Neurology, 2021, 80, 274-282.	1.7	0
45	The matrix metalloproteinase inhibitor IPR-179 has antiseizure and antiepileptogenic effects. Journal of Clinical Investigation, 2021, 131, .	8.2	35
46	High frequency oscillations associate with neuroinflammation in low-grade epilepsy associated tumors. Clinical Neurophysiology, 2021, , .	1.5	8
47	CXCR2 increases in ALS cortical neurons and its inhibition prevents motor neuron degeneration in vitro and improves neuromuscular function in SOD1G93A mice. Neurobiology of Disease, 2021, 160, 105538.	4.4	9
48	OptimisAtion of Diagnostic Accuracy in idioPathic inflammaTory myopathies (ADAPT study): a protocol for a prospective diagnostic accuracy study of multimodality testing in patients suspected of a treatable idiopathic inflammatory myopathy. BMJ Open, 2021, 11, e053594.	1.9	1
49	No evidence of aberrant amyloid β and phosphorylated tau expression in herpes simplex virusâ€infected neurons of the trigeminal ganglia and brain. Brain Pathology, 2021, , e13044.	4.1	6
50	Increased matrix metalloproteinases expression in tuberous sclerosis complex: modulation by microRNA 146a and 147b <i>in vitro</i> . Neuropathology and Applied Neurobiology, 2020, 46, 142-159.	3.2	17
51	Rare Diseases of Neurodevelopment: Maintain the Mystery or Use a Dazzling Tool for Investigation? The Case of Rett Syndrome. Neuroscience, 2020, 439, 146-152.	2.3	10
52	Erythropoietin Increases GABAA Currents in Human Cortex from TLE Patients. Neuroscience, 2020, 439, 153-162.	2.3	7
53	Coding and non-coding transcriptome of mesial temporal lobe epilepsy: Critical role of small non-coding RNAs. Neurobiology of Disease, 2020, 134, 104612.	4.4	33
54	Isomorphic diffuse glioma is a morphologically and molecularly distinct tumour entity with recurrent gene fusions of MYBL1 or MYB and a benign disease course. Acta Neuropathologica, 2020, 139, 193-209.	7.7	83

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55	microRNAâ€132 is overexpressed in glia in temporal lobe epilepsy and reduces the expression of proâ€epileptogenic factors in human cultured astrocytes. Glia, 2020, 68, 60-75.	4.9	49
56	MEK/MELK inhibition and blood–brain barrier deficiencies in atypical teratoid/rhabdoid tumors. Neuro-Oncology, 2020, 22, 58-69.	1.2	21
57	Neurosurgical treatment of subependymal giant cell astrocytomas in tuberous sclerosis complex: a series of 44 surgical procedures in 31 patients. Child's Nervous System, 2020, 36, 951-960.	1.1	14
58	Complement C5 Contributes to Brain Injury After Subarachnoid Hemorrhage. Translational Stroke Research, 2020, 11, 678-688.	4.2	24
59	The coding and non-coding transcriptional landscape of subependymal giant cell astrocytomas. Brain, 2020, 143, 131-149.	7.6	24
60	Loss of UGP2 in brain leads to a severe epileptic encephalopathy, emphasizing that bi-allelic isoform-specific start-loss mutations of essential genes can cause genetic diseases. Acta Neuropathologica, 2020, 139, 415-442.	7.7	38
61	Diffuse glioneuronal tumour with oligodendrogliomaâ€like features and nuclear clusters (DGONC) – a molecularly defined glioneuronal CNS tumour class displaying recurrent monosomy 14. Neuropathology and Applied Neurobiology, 2020, 46, 422-430.	3.2	51
62	Chronic activation of antiâ€oxidant pathways and iron accumulation in epileptogenic malformations. Neuropathology and Applied Neurobiology, 2020, 46, 546-563.	3.2	21
63	SorCS2 facilitates release of endostatin from astrocytes and controls postâ€stroke angiogenesis. Glia, 2020, 68, 1304-1316.	4.9	27
64	Editorial: Epilepsy and Neurodevelopmental Diseases. Frontiers in Cellular Neuroscience, 2020, 14, 255.	3.7	0
65	Myelin Pathology Beyond White Matter in Tuberous Sclerosis Complex (TSC) Cortical Tubers. Journal of Neuropathology and Experimental Neurology, 2020, 79, 1054-1064.	1.7	21
66	Tuberous Sclerosis Complex as Disease Model for Investigating mTOR-Related Gliopathy During Epileptogenesis. Frontiers in Neurology, 2020, 11, 1028.	2.4	25
67	Reduced expression of the glucocorticoid receptor in the hippocampus of patients with drugâ€resistant temporal lobe epilepsy and comorbid depression. Epilepsia, 2020, 61, 1595-1605.	5.1	22
68	Astrocytes as Guardians of Neuronal Excitability: Mechanisms Underlying Epileptogenesis. Frontiers in Neurology, 2020, 11, 591690.	2.4	83
69	CXCL13/CXCR5 signalling is pivotal to preserve motor neurons in amyotrophic lateral sclerosis. EBioMedicine, 2020, 62, 103097.	6.1	16
70	Modulation of GABAergic dysfunction due to SCN1A mutation linked to Hippocampal Sclerosis. Annals of Clinical and Translational Neurology, 2020, 7, 1726-1731.	3.7	4
71	Identification of Specific Circular RNA Expression Patterns and MicroRNA Interaction Networks in Mesial Temporal Lobe Epilepsy. Frontiers in Genetics, 2020, 11, 564301.	2.3	11
72	Long-lasting blood-brain barrier dysfunction and neuroinflammation after traumatic brain injury. Neurobiology of Disease, 2020, 145, 105080.	4.4	92

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73	International consensus recommendations on the diagnostic work-up for malformations of cortical development. Nature Reviews Neurology, 2020, 16, 618-635.	10.1	53
74	ls autism driven by epilepsy in infants with Tuberous Sclerosis Complex?. Annals of Clinical and Translational Neurology, 2020, 7, 1371-1381.	3.7	23
75	Seizure outcome and use of antiepileptic drugs after epilepsy surgery according to histopathological diagnosis: a retrospective multicentre cohort study. Lancet Neurology, The, 2020, 19, 748-757.	10.2	177
76	Prediction of Neurodevelopment in Infants With Tuberous Sclerosis Complex Using Early EEG Characteristics. Frontiers in Neurology, 2020, 11, 582891.	2.4	19
77	Aggregates of RNA Binding Proteins and ER Chaperones Linked to Exosomes in Granulovacuolar Degeneration of the Alzheimer's Disease Brain. Journal of Alzheimer's Disease, 2020, 75, 139-156.	2.6	22
78	Dysregulation of the MMP/TIMP Proteolytic System in Subependymal Giant Cell Astrocytomas in Patients With Tuberous Sclerosis Complex: Modulation of MMP by MicroRNA-320d In Vitro. Journal of Neuropathology and Experimental Neurology, 2020, 79, 777-790.	1.7	12
79	TSC2 pathogenic variants are predictive of severe clinical manifestations in TSC infants: results of the EPISTOP study. Genetics in Medicine, 2020, 22, 1489-1497.	2.4	51
80	Increased expression of miR142 and miR155 in glial and immune cells after traumatic brain injury may contribute to neuroinflammation via astrocyte activation. Brain Pathology, 2020, 30, 897-912.	4.1	23
81	pCREB expression in human tissues from epilepsy surgery. Epilepsia, 2020, 61, 1240-1252.	5.1	1
82	Adenosine kinase inhibition promotes proliferation of neural stem cells after traumatic brain injury. Brain Communications, 2020, 2, fcaa017.	3.3	15
83	Calcineurin Controls Expression of EAAT1/GLAST in Mouse and Human Cultured Astrocytes through Dynamic Regulation of Protein Synthesis and Degradation. International Journal of Molecular Sciences, 2020, 21, 2213.	4.1	9
84	Neonatal bacterial meningitis versus ventriculitis: a cohort-based overview of clinical characteristics, microbiology and imaging. European Journal of Pediatrics, 2020, 179, 1969-1977.	2.7	7
85	Expression and Cellular Distribution of P-Glycoprotein and Breast Cancer Resistance Protein in Amyotrophic Lateral Sclerosis Patients. Journal of Neuropathology and Experimental Neurology, 2020, 79, 266-276.	1.7	17
86	Same same but different: A Webâ€based deep learning application revealed classifying features for the histopathologic distinction of cortical malformations. Epilepsia, 2020, 61, 421-432.	5.1	17
87	Brain tumour diagnostics using a DNA methylationâ€based classifier as a diagnostic support tool. Neuropathology and Applied Neurobiology, 2020, 46, 478-492.	3.2	59
88	Splicing Players Are Differently Expressed in Sporadic Amyotrophic Lateral Sclerosis Molecular Clusters and Brain Regions. Cells, 2020, 9, 159.	4.1	21
89	Phytocannabinoids in Neurological Diseases: Could They Restore a Physiological GABAergic Transmission?. International Journal of Molecular Sciences, 2020, 21, 723.	4.1	38
90	A nicotinamide phosphoribosyltransferase–GAPDH interaction sustains the stress-induced NMN/NAD+ salvage pathway in the nucleus. Journal of Biological Chemistry, 2020, 295, 3635-3651.	3.4	21

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91	Chronic Regulation of miR-124-3p in the Perilesional Cortex after Experimental and Human TBI. International Journal of Molecular Sciences, 2020, 21, 2418.	4.1	20
92	Natural killer cells modulate motor neuron-immune cell cross talk in models of Amyotrophic Lateral Sclerosis. Nature Communications, 2020, 11, 1773.	12.8	93
93	CpG and non-CpG Presenilin1 methylation pattern in course of neurodevelopment and neurodegeneration is associated with gene expression in human and murine brain. Epigenetics, 2020, 15, 781-799.	2.7	39
94	The adult human subventricular zone: partial ependymal coverage and proliferative capacity of cerebrospinal fluid. Brain Communications, 2020, 2, fcaa150.	3.3	10
95	Increased expression of myelin-associated genes in frontal cortex of overexpressing rats and Parkinson's disease patients. Aging, 2020, 12, 18889-18906.	3.1	1
96	Review: Challenges in the histopathological classification of ganglioglioma and DNT: microscopic agreement studies and a preliminary genotypeâ€phenotype analysis. Neuropathology and Applied Neurobiology, 2019, 45, 95-107.	3.2	46
97	Precise detection of low-level somatic mutation in resected epilepsy brain tissue. Acta Neuropathologica, 2019, 138, 901-912.	7.7	92
98	Integrative multi-omic analysis identifies new drivers and pathways in molecularly distinct subtypes of ALS. Scientific Reports, 2019, 9, 9968.	3.3	28
99	Group I mGluR-Mediated Activation of Martinotti Cells Inhibits Local Cortical Circuitry in Human Cortex. Frontiers in Cellular Neuroscience, 2019, 13, 315.	3.7	15
100	TLR3 preconditioning induces anti-inflammatory and anti-ictogenic effects in mice mediated by the IRF3/IFN-β axis. Brain, Behavior, and Immunity, 2019, 81, 598-607.	4.1	14
101	Building Bridges Between the Clinic and the Laboratory: A Meeting Review – Brain Malformations: A Roadmap for Future Research. Frontiers in Cellular Neuroscience, 2019, 13, 434.	3.7	3
102	Postmortem Cortex Samples Identify Distinct Molecular Subtypes of ALS: Retrotransposon Activation, Oxidative Stress, and Activated Glia. Cell Reports, 2019, 29, 1164-1177.e5.	6.4	184
103	Phenotypes and malignancy risk of different <i>FUS</i> mutations in genetic amyotrophic lateral sclerosis. Annals of Clinical and Translational Neurology, 2019, 6, 2384-2394.	3.7	49
104	Response to: Diagnostic value of additional histopathological fascia examination in idiopathic inflammatory myopathies. European Journal of Neurology, 2019, 26, e95.	3.3	0
105	Idiopathic inflammatory myopathy. Neurology, 2019, 93, e889-e894.	1.1	17
106	Cognitive functioning after epilepsy surgery in children with mild malformation of cortical development and focal cortical dysplasia. Epilepsy and Behavior, 2019, 94, 209-215.	1.7	21
107	Diagnostic value of additional histopathological fascia examination in idiopathic inflammatory myopathies. European Journal of Neurology, 2019, 26, 1494-1496.	3.3	3
108	Rosette-forming glioneuronal tumors share a distinct DNA methylation profile and mutations in FGFR1, with recurrent co-mutation of PIK3CA and NF1. Acta Neuropathologica, 2019, 138, 497-504.	7.7	57

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109	Mutant FUS and ELAVL4 (HuD) Aberrant Crosstalk in Amyotrophic Lateral Sclerosis. Cell Reports, 2019, 27, 3818-3831.e5.	6.4	51
110	Early Clinical Predictors of Autism Spectrum Disorder in Infants with Tuberous Sclerosis Complex: Results from the EPISTOP Study. Journal of Clinical Medicine, 2019, 8, 788.	2.4	42
111	Targeting oxidative stress improves disease outcomes in a rat model of acquired epilepsy. Brain, 2019, 142, e39-e39.	7.6	137
112	Therapeutic effect of Anakinra in the relapsing chronic phase of febrile infection–related epilepsy syndrome. Epilepsia Open, 2019, 4, 344-350.	2.4	85
113	The Roof is Leaking and a Storm is Raging: Repairing the Blood–Brain Barrier in the Fight Against Epilepsy. Epilepsy Currents, 2019, 19, 177-181.	0.8	40
114	Genomic <scp>DNA</scp> methylation distinguishes subtypes of human focal cortical dysplasia. Epilepsia, 2019, 60, 1091-1103.	5.1	61
115	Quantitative Third Harmonic Generation Microscopy for Assessment of Glioma in Human Brain Tissue. Advanced Science, 2019, 6, 1900163.	11.2	24
116	New insights into a spectrum of developmental malformations related to <scp>mTOR</scp> dysregulations: challenges and perspectives. Journal of Anatomy, 2019, 235, 521-542.	1.5	63
117	SorCS2 Controls Functional Expression of Amino Acid Transporter EAAT3 and Protects Neurons from Oxidative Stress and Epilepsy-Induced Pathology. Cell Reports, 2019, 26, 2792-2804.e6.	6.4	39
118	HR23B pathology preferentially co-localizes with p62, pTDP-43 and poly-GA in C9ORF72-linked frontotemporal dementia and amyotrophic lateral sclerosis. Acta Neuropathologica Communications, 2019, 7, 39.	5.2	9
119	FUS pathology in ALS is linked to alterations in multiple ALS-associated proteins and rescued by drugs stimulating autophagy. Acta Neuropathologica, 2019, 138, 67-84.	7.7	94
120	Longâ€ŧerm seizure outcome after epilepsy surgery in patients with mild malformation of cortical development and focal cortical dysplasia. Epilepsia Open, 2019, 4, 170-175.	2.4	17
121	Curcumin reduces development of seizurelike events in the hippocampalâ€entorhinal cortex slice culture model for epileptogenesis. Epilepsia, 2019, 60, 605-614.	5.1	13
122	Changes in vascular density in resected tissue of 97 patients with mild malformation of cortical development, focal cortical dysplasia or TSCâ€related cortical tubers. International Journal of Developmental Neuroscience, 2019, 79, 96-104.	1.6	5
123	Rapidly progressive amyotrophic lateral sclerosis is associated with microglial reactivity and small heat shock protein expression in reactive astrocytes. Neuropathology and Applied Neurobiology, 2019, 45, 459-475.	3.2	23
124	GSK3Î <sup>2</sup> activity alleviates epileptogenesis and limits GluA1 phosphorylation. EBioMedicine, 2019, 39, 377-387.	6.1	17
125	Oxidative stress and inflammation in a spectrum of epileptogenic cortical malformations: molecular insights into their interdependence. Brain Pathology, 2019, 29, 351-365.	4.1	54

126 First Results of the EPISTOP Study. , 2019, 50, .

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127	Differential Fatty Acid-Binding Protein Expression in Persistent Radial Glia in the Human and Sheep Subventricular Zone. Developmental Neuroscience, 2018, 40, 145-161.	2.0	10
128	Sushi repeatâ€containing protein Xâ€linked 2: A novel phylogenetically conserved hypothalamoâ€pituitary protein. Journal of Comparative Neurology, 2018, 526, 1806-1819.	1.6	4
129	A novel action of lacosamide on GABA A currents sets the ground for a synergic interaction with levetiracetam in treatment of epilepsy. Neurobiology of Disease, 2018, 115, 59-68.	4.4	26
130	Neuropathology of epilepsy. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2018, 145, 193-216.	1.8	57
131	mi <scp>R</scp> 147b: <scp>A</scp> novel key regulator of interleukin 1 betaâ€mediated inflammation in human astrocytes. Glia, 2018, 66, 1082-1097.	4.9	28
132	mTOR dysregulation and tuberous sclerosis-related epilepsy. Expert Review of Neurotherapeutics, 2018, 18, 185-201.	2.8	68
133	Commonalities in epileptogenic processes from different acute brain insults: Do they translate?. Epilepsia, 2018, 59, 37-66.	5.1	206
134	Hippocampal Radial Glial Subtypes and Their Neurogenic Potential in Human Fetuses and Healthy and Alzheimer's Disease Adults. Cerebral Cortex, 2018, 28, 2458-2478.	2.9	128
135	DNA methylation-based classification of central nervous system tumours. Nature, 2018, 555, 469-474.	27.8	1,872
136	Architectural B-cell organization in skeletal muscle identifies subtypes of dermatomyositis. Neurology: Neuroimmunology and NeuroInflammation, 2018, 5, e451.	6.0	19
137	The physiological phosphorylation of tau is critically changed in fetal brains of individuals with Down syndrome. Neuropathology and Applied Neurobiology, 2018, 44, 314-327.	3.2	22
138	Review: Neuroinflammatory pathways as treatment targets and biomarker candidates in epilepsy: emerging evidence from preclinical and clinical studies. Neuropathology and Applied Neurobiology, 2018, 44, 91-111.	3.2	186
139	Multinodular and vacuolating neuronal tumors in epilepsy: dysplasia or neoplasia?. Brain Pathology, 2018, 28, 155-171.	4.1	54
140	PACAP and PAC1R are differentially expressed in motor cortex of amyotrophic lateral sclerosis patients and support survival of iPSCâ€derived motor neurons. Journal of Cellular Physiology, 2018, 233, 3343-3351.	4.1	25
141	Golgin A4 in CSF and granulovacuolar degenerations of patients with Alzheimer disease. Neurology, 2018, 91, e1799-e1808.	1.1	11
142	n-3 Docosapentaenoic acid-derived protectin D1 promotes resolution of neuroinflammation and arrests epileptogenesis. Brain, 2018, 141, 3130-3143.	7.6	55
143	A novel GABAergic dysfunction in human Dravet syndrome. Epilepsia, 2018, 59, 2106-2117.	5.1	46
144	TBIO-17. IMPLEMENTATION OF METHYLATION PROFILING FOR CNS TUMOR DIAGNOSIS IN THE PRINCESS MÃXIMA CENTER FOR PEDIATRIC ONCOLOGY, THE NETHERLANDS. Neuro-Oncology, 2018, 20, i183-i184.	1.2	0

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145	Histopathology of Listeria Meningitis. Journal of Neuropathology and Experimental Neurology, 2018, 77, 950-957.	1.7	19
146	Activation of the innate immune system is evident throughout epileptogenesis and is associated with bloodâ€brain barrier dysfunction and seizure progression. Epilepsia, 2018, 59, 1931-1944.	5.1	67
147	miR-124-3p is a chronic regulator of gene expression after brain injury. Cellular and Molecular Life Sciences, 2018, 75, 4557-4581.	5.4	40
148	Management of epilepsy associated with tuberous sclerosis complex: Updated clinical recommendations. European Journal of Paediatric Neurology, 2018, 22, 738-748.	1.6	151
149	Increased expression of matrix metalloproteinase 3 can be attenuated by inhibition of microRNA-155 in cultured human astrocytes. Journal of Neuroinflammation, 2018, 15, 211.	7.2	36
150	Effects of rapamycin and curcumin on inflammation and oxidative stress in vitro and in vivo — in search of potential anti-epileptogenic strategies for temporal lobe epilepsy. Journal of Neuroinflammation, 2018, 15, 212.	7.2	48
151	Role of miR-146a in neural stem cell differentiation and neural lineage determination: relevance for neurodevelopmental disorders. Molecular Autism, 2018, 9, 38.	4.9	70
152	Delayed cerebral thrombosis complicating pneumococcal meningitis: an autopsy study. Annals of Intensive Care, 2018, 8, 20.	4.6	21
153	Contribution of Altered Endocannabinoid System to Overactive mTORC1 Signaling in Focal Cortical Dysplasia. Frontiers in Pharmacology, 2018, 9, 1508.	3.5	8
154	MicroRNA519d and microRNA4758 can identify gangliogliomas from dysembryoplastic neuroepithelial tumours and astrocytomas. Oncotarget, 2018, 9, 28103-28115.	1.8	5
155	Modulation of GABAA Receptors in the Treatment of Epilepsy. Current Pharmaceutical Design, 2018, 23, 5563-5568.	1.9	27
156	Common data elements and data management: Remedy to cure underpowered preclinical studies. Epilepsy Research, 2017, 129, 87-90.	1.6	35
157	Seven tesla <scp>MRI</scp> improves detection of focal cortical dysplasia in patients with refractory focal epilepsy. Epilepsia Open, 2017, 2, 162-171.	2.4	47
158	Proteomic profiling of the spinal cord in ALS: decreased ATP5D levels suggest synaptic dysfunction in ALS pathogenesis. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2017, 18, 210-220.	1.7	25
159	The ALS-linked E102Q mutation in Sigma receptor-1 leads to ER stress-mediated defects in protein homeostasis and dysregulation of RNA-binding proteins. Cell Death and Differentiation, 2017, 24, 1655-1671.	11.2	77
160	A combinatorial approach to identify calpain cleavage sites in the Machado-Joseph disease protein ataxin-3. Brain, 2017, 140, 1280-1299.	7.6	33
161	FCD Type II and mTOR pathway: Evidence for different mechanisms involved in the pathogenesis of dysmorphic neurons. Epilepsy Research, 2017, 129, 146-156.	1.6	21
162	Blockade of the IL-1R1/TLR4 pathway mediates disease-modification therapeutic effects in a model of acquired epilepsy. Neurobiology of Disease, 2017, 99, 12-23.	4.4	149

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