

# Miguel Angel Cortez

## List of Publications by Year in descending order

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77  
papers

2,717  
citations

186265

28  
h-index

206112

48  
g-index

78  
all docs

78  
docs citations

78  
times ranked

2955  
citing authors

#	ARTICLE	IF	CITATIONS
1	Seizure frequency discrepancy between subjective and objective ictal electroencephalography data in dogs. <i>Journal of Veterinary Internal Medicine</i> , 2021, 35, 1819-1825.	1.6	16
2	Effect of prior general anesthesia or sedation and antiseizure drugs on the diagnostic utility of wireless video electroencephalography in dogs. <i>Journal of Veterinary Internal Medicine</i> , 2020, 34, 1967-1974.	1.6	12
3	Neurexin 2 regulates absence seizures and behavioral arrests through GABAergic transmission within the thalamocortical circuitry. <i>Nature Communications</i> , 2020, 11, 3744.	12.8	18
4	EEG before and after total corpus callosotomy for pharmacoresistant infantile spasms: Fast oscillations and slow-wave connectivity in hypsarrhythmia. <i>Epilepsia</i> , 2019, 60, 1849-1860.	5.1	16
5	Pharmacologically induced absence seizures versus kindling in Wistar rats. <i>İstanbul Kültür Sanat Akademisi Dergisi</i> , 2019, 7, 25-34.	0.3	1
6	Hypsarrhythmia in epileptic spasms: Synchrony in chaos. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2018, 58, 55-61.	2.0	6
7	Prolonged rhythmic mid-temporal discharges (RMTD) in a 5-year old child. <i>Journal of Clinical Neuroscience</i> , 2018, 48, 81-82.	1.5	1
8	Absence Seizures as a Feature of Juvenile Myoclonic Epilepsy in Rhodesian Ridgeback Dogs. <i>Journal of Veterinary Internal Medicine</i> , 2018, 32, 428-432.	1.6	16
9	Latitudinal differences on the global epidemiology of infantile spasms: systematic review and meta-analysis. <i>Orphanet Journal of Rare Diseases</i> , 2018, 13, 216.	2.7	29
10	Perineal stimulation triggering seizures in a child with Dravet syndrome. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2018, 62, 106-107.	2.0	2
11	Methodologic recommendations and possible interpretations of video-EEG recordings in immature rodents used as experimental controls: TASK1/WG2 report of the ILAE/AES Joint Translational Task Force. <i>Epilepsia Open</i> , 2018, 3, 437-459.	2.4	12
12	Activation of Entorhinal Cortical Projections to the Dentate Gyrus Underlies Social Memory Retrieval. <i>Cell Reports</i> , 2018, 23, 2379-2391.	6.4	56
13	Kcnj6 (GIRK2) trisomy is not sufficient for conferring the susceptibility to infantile spasms seen in the Ts65Dn mouse model of down syndrome. <i>Epilepsy Research</i> , 2018, 145, 82-88.	1.6	11
14	5. Prospective pre-emptive EEG study prior to west syndrome. <i>Clinical Neurophysiology</i> , 2018, 129, e46.	1.5	0
15	Generalized myoclonic epilepsy with photosensitivity in juvenile dogs caused by a defective DIRAS family GTPase 1. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2017, 114, 2669-2674.	7.1	39
16	Diagnostic Utility of Wireless Video-EEG in Unsedated Dogs. <i>Journal of Veterinary Internal Medicine</i> , 2017, 31, 1469-1476.	1.6	34
17	Neurexin 3 R451C mutation alters electroencephalography spectral activity in an animal model of autism spectrum disorders. <i>Molecular Brain</i> , 2017, 10, 10.	2.6	24
18	Infantile spasms in down syndrome: Rescue by knockdown of the GIRK2 channel. <i>Annals of Neurology</i> , 2016, 80, 511-521.	5.3	22

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19	Systemic availability of guanidinoacetate affects GABAA receptor function and seizure threshold in GAMT deficient mice. <i>Amino Acids</i> , 2016, 48, 2041-2047.	2.7	13
20	Acute and chronic pharmacological models of generalized absence seizures. <i>Journal of Neuroscience Methods</i> , 2016, 260, 175-184.	2.5	14
21	Targeting the GABAB Receptor for the Treatment of Epilepsy. <i>Receptors</i> , 2016, , 175-195.	0.2	3
22	The $\text{GIRK}_2$ subunit is involved in IS $\delta$ -like seizures induced by $\text{GABA}_B$ receptor agonists. <i>Epilepsia</i> , 2015, 56, 1081-1087.	5.1	19
23	LIMK1 Regulates Long-Term Memory and Synaptic Plasticity via the Transcriptional Factor CREB. <i>Molecular and Cellular Biology</i> , 2015, 35, 1316-1328.	2.3	62
24	Paroxysmal Alpha Activity in Rett Syndrome: A Case Report. <i>Pediatric Neurology</i> , 2014, 51, 421-425.	2.1	6
25	EEG and neuroimaging correlations in children with lissencephaly. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2013, 22, 189-193.	2.0	12
26	Electrophysiological Recording Techniques. 2011. Edited by Robert P. Vertes, Robert W. Stackman Jr. Published by Humana Press. 284 pages. C\$120 approx.. <i>Canadian Journal of Neurological Sciences</i> , 2013, 40, 271-272.	0.5	0
27	Environmental Enrichment Improves Behavioral Outcome in the AY-9944 Model of Childhood Atypical Absence Epilepsy. <i>International Journal of Neuroscience</i> , 2012, 122, 449-457.	1.6	10
28	Circadian profiles of focal epileptic seizures: A need for reappraisal. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2012, 21, 412-416.	2.0	36
29	Mania-like behavior induced by genetic dysfunction of the neuron-specific $\text{Na}^+, \text{K}^+$ -ATPase $\text{I}\pm 3$ sodium pump. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011, 108, 18144-18149.	7.1	127
30	Evidence that clozapine directly interacts on the GABAB receptor. <i>NeuroReport</i> , 2011, 22, 637-641.	1.2	43
31	Arrhythmia and sudden death associated with elevated cardiac chloride channel activity. <i>Journal of Cellular and Molecular Medicine</i> , 2011, 15, 2307-2316.	3.6	14
32	Treatment of Infantile Spasms. <i>Journal of Child Neurology</i> , 2011, 26, 1411-1421.	1.4	63
33	PTG Depletion Removes Lafora Bodies and Rescues the Fatal Epilepsy of Lafora Disease. <i>PLoS Genetics</i> , 2011, 7, e1002037.	3.5	185
34	The Significance of Frontal Intermittent Rhythmic Delta Activity in Children. <i>Canadian Journal of Neurological Sciences</i> , 2010, 37, 656-661.	0.5	5
35	Absence seizures with myoclonic features in a juvenile Chihuahua dog. <i>Epileptic Disorders</i> , 2010, 12, 138-141.	1.3	23
36	GABAB receptors in absence epilepsy. <i>Epilepsia</i> , 2010, 51, 24-24.	5.1	5

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37	Disruption of ClC-2 expression is associated with progressive neurodegeneration in aging mice. <i>Neuroscience</i> , 2010, 167, 154-162.	2.3	17
38	Mutation I810N in the $\beta 3$ isoform of Na <sup>+</sup> ,K <sup>+</sup> -ATPase causes impairments in the sodium pump and hyperexcitability in the CNS. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2009, 106, 14085-14090.	7.1	128
39	Predictive Value of Clinical and EEG Features in the Diagnosis of Stroke and Hypoxic Ischemic Encephalopathy in Neonates With Seizures. <i>Stroke</i> , 2009, 40, 2402-2407.	2.0	52
40	Infantile Spasms and Down Syndrome: A New Animal Model. <i>Pediatric Research</i> , 2009, 65, 499-503.	2.3	76
41	The circuitry of atypical absence seizures in GABABR1a transgenic mice. <i>Pharmacology Biochemistry and Behavior</i> , 2009, 94, 124-130.	2.9	25
42	Succinic semialdehyde dehydrogenase deficiency: Lessons from mice and men. <i>Journal of Inherited Metabolic Disease</i> , 2009, 32, 343-352.	3.6	97
43	Monoamine variability in the chronic model of atypical absence seizures. <i>Epilepsia</i> , 2009, 50, 768-775.	5.1	3
44	GABA receptor proteins within lipid rafts in the AY9944 model of atypical absence seizures. <i>Epilepsia</i> , 2009, 50, 776-788.	5.1	11
45	Severity of atypical absence phenotype in GABAB transgenic mice is subunit specific. <i>Epilepsy and Behavior</i> , 2009, 14, 577-581.	1.7	20
46	Circadian distribution of generalized tonic-clonic seizures associated with murine succinic semialdehyde dehydrogenase deficiency, a disorder of GABA metabolism. <i>Epilepsy and Behavior</i> , 2008, 13, 290-294.	1.7	18
47	A ketogenic diet rescues the murine succinic semialdehyde dehydrogenase deficient phenotype. <i>Experimental Neurology</i> , 2008, 210, 449-457.	4.1	54
48	5-HT <sub>2</sub> modulation of AY-9944 induced atypical absence seizures. <i>Neuroscience Letters</i> , 2007, 418, 13-17.	2.1	15
49	Transgenic mice over-expressing GABABR1a receptors acquire an atypical absence epilepsy-like phenotype. <i>Neurobiology of Disease</i> , 2007, 26, 439-451.	4.4	33
50	Chronobiometry of Behavioral Activity in the Ts65Dn Model of Down Syndrome. <i>Behavior Genetics</i> , 2007, 37, 388-398.	2.1	31
51	Daily rhythms of seizure activity and behavior in a model of atypical absence epilepsy. <i>Epilepsy and Behavior</i> , 2006, 9, 564-572.	1.7	26
52	Serotonin Depletion Attenuates AY-9944-Mediated Atypical Absence Seizures. <i>Epilepsia</i> , 2006, 47, 240-246.	5.1	23
53	Nonconvulsive Seizures in the Pediatric Intensive Care Unit: Etiology, EEG, and Brain Imaging Findings. <i>Epilepsia</i> , 2006, 47, 1510-1518.	5.1	97
54	GABAB receptor antagonism abolishes the learning impairments in rats with chronic atypical absence seizures. <i>European Journal of Pharmacology</i> , 2006, 541, 64-72.	3.5	34

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55	Succinic Semialdehyde Dehydrogenase Deficiency: GABAB receptor-mediated function. <i>Brain Research</i> , 2006, 1090, 15-22.	2.2	62
56	Status epilepticus in mice deficient for succinate semialdehyde dehydrogenase: GABAA receptor-mediated mechanisms. <i>Annals of Neurology</i> , 2006, 59, 42-52.	5.3	61
57	Pharmacologic Models of Generalized Absence Seizures in Rodents. , 2006, , 111-126.		10
58	Animal models of epilepsy and progressive effects of seizures. <i>Advances in Neurology</i> , 2006, 97, 293-304.	0.8	16
59	A Reappraisal of Rhythmic Coma Patterns in Children. <i>Canadian Journal of Neurological Sciences</i> , 2005, 32, 518-523.	0.5	11
60	Clinical and Neurophysiologic Spectrum Associated with Atypical Absence Seizures in Children with Intractable Epilepsy. <i>Journal of Child Neurology</i> , 2005, 20, 404-410.	1.4	29
61	Reactive EEG Patterns in Pediatric Coma. <i>Pediatric Neurology</i> , 2005, 33, 345-349.	2.1	44
62	Lovastatin exacerbates atypical absence seizures with only minimal effects on brain sterols. <i>Journal of Lipid Research</i> , 2004, 45, 2038-2043.	4.2	18
63	Laforin preferentially binds the neurotoxic starch-like polyglucosans, which form in its absence in progressive myoclonus epilepsy. <i>Human Molecular Genetics</i> , 2004, 13, 1117-1129.	2.9	101
64	Absence seizures in succinic semialdehyde dehydrogenase deficient mice: a model of juvenile absence epilepsy. <i>Pharmacology Biochemistry and Behavior</i> , 2004, 79, 547-553.	2.9	65
65	Hormonal regulation of atypical absence seizures. <i>Annals of Neurology</i> , 2004, 55, 353-361.	5.3	22
66	Refractory atypical absence seizures in rat: a two hit model. <i>Epilepsy Research</i> , 2004, 62, 53-63.	1.6	26
67	Learning and memory impairment in rats with chronic atypical absence seizures. <i>Experimental Neurology</i> , 2004, 190, 328-336.	4.1	48
68	Dynamical regimes underlying epileptiform events: role of instabilities and bifurcations in brain activity. <i>Physica D: Nonlinear Phenomena</i> , 2003, 186, 205-220.	2.8	49
69	Anticonvulsant properties of acetone, a brain ketone elevated by the ketogenic diet. <i>Annals of Neurology</i> , 2003, 54, 219-226.	5.3	160
70	Brain Sterols in the AY-9944 Rat Model of Atypical Absence Seizures. <i>Epilepsia</i> , 2002, 43, 3-8.	5.1	24
71	A chronic model of atypical absence seizures: studies of developmental and gender sensitivity. <i>Epilepsy Research</i> , 2002, 48, 111-119.	1.6	38
72	Alteration of GLUR2 expression in the rat brain following absence seizures induced by $\delta^3$ -hydroxybutyric acid. <i>Epilepsy Research</i> , 2001, 44, 41-51.	1.6	14

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73	$\hat{1}^3$ -Hydroxybutyric acid-induced absence seizures in GluR2 null mutant mice. Brain Research, 2001, 897, 27-35.	2.2	19
74	Type I Diabetes and Multiple Sclerosis Patients Target Islet Plus Central Nervous System Autoantigens; Nonimmunized Nonobese Diabetic Mice Can Develop Autoimmune Encephalitis. Journal of Immunology, 2001, 166, 2831-2841.	0.8	84
75	Interactions of Clobazam With Conventional Antiepileptics in Children. Journal of Child Neurology, 1997, 12, 208-213.	1.4	47
76	Infantile spasms: Seasonal onset differences and zeitgebers. Pediatric Neurology, 1997, 16, 220-224.	2.1	17
77	Recurrent seizures in metachromatic leukodystrophy. Pediatric Neurology, 1997, 17, 150-154.	2.1	36