Luis GonzÃ¡lez GutiÃ©rrez-Solana

List of Publications by Year
in descending order

13 papers	619 citations	1307594 7 h-index	677142 22 g-index
27 all docs	27 docs citations	27 times ranked	980 citing authors

#	Article	IF	Citations
1	A Fatal Mitochondrial Disease Is Associated with Defective NFU1 Function in the Maturation of a Subset of Mitochondrial Fe-S Proteins. American Journal of Human Genetics, 2011, 89, 656-667.	6.2	262
2	Glial fibrillary acidic protein mutations in infantile, juvenile, and adult forms of Alexander disease. Annals of Neurology, 2005, 57, 310-326.	5.3	220
3	Clinical, biochemical, and molecular studies in pyridoxineâ€dependent epilepsy. Antisense therapy as possible new therapeutic option. Epilepsia, 2013, 54, 239-248.	5.1	43
4	Phenytoin-induced visual disturbances mimicking Delirium Tremens in a child. European Journal of Paediatric Neurology, 2010, 14, 460-463.	1.6	12
5	Urinary sulphatoxymelatonin as a biomarker of serotonin status in biogenic amine-deficient patients. Scientific Reports, 2017, 7, 14675.	3.3	8
6	Classic Ketogenic Diet and Modified Atkins Diet in SLC2A1 Positive and Negative Patients with Suspected GLUT1 Deficiency Syndrome: A Single Center Analysis of 18 Cases. Nutrients, 2021, 13, 840.	4.1	8
7	Efficacy and safety of intrathecal idursulfase in pediatric patients with mucopolysaccharidosis type II and early cognitive impairment: Design and methods of a controlled, randomized, phase II/III multicenter study. Molecular Genetics and Metabolism, 2018, 123, S99-S100.	1.1	7
8	Procalcitonin, a high acute phase reactant in antiepileptic hypersentivity syndrome in pediatric age. European Journal of Paediatric Neurology, 2012, 16, 200-202.	1.6	5
9	Efficacy of Brivaracetam in children with epilepsy. Epilepsy Research, 2021, 177, 106757.	1.6	5
10	Tratamiento de las enfermedades lisosomales en la poblaciÃ³n pediÃ¡trica. Anales De Pediatria Continuada, 2013, 11, 159-161.	0.1	3
11	Long-term normalization of cognitive and psychopathological alterations in a juvenile Niemannâ€“Pick type C case. Neurodegenerative Disease Management, 2020, 10, 69-76.	2.2	3
12	Transition process from paediatric to adult care in patients with inborn errors of metabolism. Consensus statement. Medicina ClÃnica (English Edition), 2016, 147, 506.e1-506.e7.	0.2	2
13	Neurodevelopmental status and adaptive behavior of pediatric patients with Hunter syndrome: A longitudinal observational study. Molecular Genetics and Metabolism, 2019, 126, S103.	1.1	0