SÃ©bastien Lebon

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/3192999/publications.pdf

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27	849	840776	580821
papers	citations	11 h-index	25 g-index

28	28	28	1973
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	A 600â€…kb deletion syndrome at 16p11.2 leads to energy imbalance and neuropsychiatric disorders. Journal of Medical Genetics, 2012, 49, 660-668.	3.2	251
2	Defining the Effect of the 16p11.2 Duplication on Cognition, Behavior, and Medical Comorbidities. JAMA Psychiatry, 2016, 73, 20.	11.0	195
3	16p11.2 600 kb Duplications confer risk for typical and atypical Rolandic epilepsy. Human Molecular Genetics, 2014, 23, 6069-6080.	2.9	61
4	Quantifying the Effects of 16p11.2 Copy Number Variants on Brain Structure: A Multisite Genetic-First Study. Biological Psychiatry, 2018, 84, 253-264.	1.3	56
5	Autosomal recessive primary microcephaly due to ASPM mutations: An update. Human Mutation, 2018, 39, 319-332.	2.5	53
6	Anti-N-Methyl-<scp>d</scp>-Aspartate (NMDA) Receptor Encephalitis Mimicking a Primary Psychiatric Disorder in an Adolescent. Journal of Child Neurology, 2012, 27, 1607-1610.	1.4	29
7	Perinatal arterial ischemic stroke related to carotid artery occlusion. European Journal of Paediatric Neurology, 2016, 20, 639-648.	1.6	26
8	Earlyâ€onset encephalopathy with epilepsy associated with a novel splice site mutation in SMC1A. American Journal of Medical Genetics, Part A, 2015, 167, 3076-3081.	1.2	22
9	Effects of eight neuropsychiatric copy number variants on human brain structure. Translational Psychiatry, 2021, 11, 399.	4.8	18
10	When is a child with status epilepticus likely to have Dravet syndrome?. Epilepsy Research, 2014, 108, 740-747.	1.6	16
11	Pediatric Encephalopathy: Clinical, Biochemical and Cellular Insights into the Role of Gln52 of GNAO1 and GNAI1 for the Dominant Disease. Cells, 2021, 10, 2749.	4.1	16
12	Childhood neurodegeneration associated with a specific UBTF variant: a new case report and review of the literature. BMC Neurology, 2020, 20, 17.	1.8	15
13	Midazolam as a firstâ€line treatment for neonatal seizures: Retrospective study. Pediatrics International, 2018, 60, 498-500.	0.5	12
14	De novo variants in CACNA1E found in patients with intellectual disability, developmental regression and social cognition deficit but no seizures. Molecular Autism, 2021, 12, 69.	4.9	12
15	Periinsular anterior quadrantotomy: technical note. Journal of Neurosurgery: Pediatrics, 2018, 21, 124-132.	1.3	11
16	When should clinicians search for GLUT1 deficiency syndrome in childhood generalized epilepsies?. European Journal of Paediatric Neurology, 2015, 19, 170-175.	1.6	9
17	Developmental trajectories of neuroanatomical alterations associated with the 16p11.2 Copy Number Variations. NeuroImage, 2019, 203, 116155.	4.2	9
18	An initial MRI picture of limbic encephalitis in subacute sclerosing panencephalitis. European Journal of Paediatric Neurology, 2011, 15, 544-546.	1.6	8

#	Article	IF	Citations
19	Sultiame pharmacokinetic profile in plasma and erythrocytes after single oral doses: A pilot study in healthy volunteers. Pharmacology Research and Perspectives, 2020, 8, e00558.	2.4	8
20	Prenatal diagnosis of Aicardi syndrome based on a suggestive imaging pattern: A multicenter caseâ€series. Prenatal Diagnosis, 2022, 42, 484-494.	2.3	8
21	Biallelic variants in ZNF526 cause a severe neurodevelopmental disorder with microcephaly, bilateral cataract, epilepsy and simplified gyration. Journal of Medical Genetics, 2021, , jmedgenet-2020-107430.	3.2	5
22	Exposure to alirocumab during the first trimester ofÂpregnancy: A case report. Birth Defects Research, 2021, 113, 1156-1160.	1.5	5
23	Structural brain abnormalities in epilepsy with myoclonic atonic seizures. Epilepsy Research, 2021, 177, 106771.	1.6	2
24	Prenatal Brainstem Disruptions: Small Lesionsâ€“Big Problems. Neuropediatrics, 2017, 48, 350-355.	0.6	1
25	Agenesis of the Corpus Callosum with Facial Dysmorphism and Intellectual Disability in Sibs Associated with Compound Heterozygous KDM5B Variants. Genes, 2021, 12, 1397.	2.4	1
26	SCN1A Does Not Determine Seizure Duration in Children Unaffected by Dravet's Syndrome. Journal of Pediatric Epilepsy, 2017, 06, 111-114.	0.2	0
27	Clinical Reasoning: Rapidly progressive gait disorder and cranial nerves involvement in a 9-year-old boy. Neurology, 2020, 94, e330-e334.	1.1	0