

# SÃ©bastien Lebon

## List of Publications by Year in descending order

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Version: 2024-02-01

27  
papers

849  
citations

840776

11  
h-index

580821

25  
g-index

28  
all docs

28  
docs citations

28  
times ranked

1973  
citing authors

#	ARTICLE	IF	CITATIONS
1	Prenatal diagnosis of Aicardi syndrome based on a suggestive imaging pattern: A multicenter case series. <i>Prenatal Diagnosis</i> , 2022, 42, 484-494.	2.3	8
2	Biallelic variants in ZNF526 cause a severe neurodevelopmental disorder with microcephaly, bilateral cataract, epilepsy and simplified gyration. <i>Journal of Medical Genetics</i> , 2021, , jmedgenet-2020-107430.	3.2	5
3	Exposure to alirocumab during the first trimester of pregnancy: A case report. <i>Birth Defects Research</i> , 2021, 113, 1156-1160.	1.5	5
4	Effects of eight neuropsychiatric copy number variants on human brain structure. <i>Translational Psychiatry</i> , 2021, 11, 399.	4.8	18
5	Agenesis of the Corpus Callosum with Facial Dysmorphism and Intellectual Disability in Sibs Associated with Compound Heterozygous KDM5B Variants. <i>Genes</i> , 2021, 12, 1397.	2.4	1
6	Structural brain abnormalities in epilepsy with myoclonic atonic seizures. <i>Epilepsy Research</i> , 2021, 177, 106771.	1.6	2
7	Pediatric Encephalopathy: Clinical, Biochemical and Cellular Insights into the Role of Gln52 of GNAO1 and GNAI1 for the Dominant Disease. <i>Cells</i> , 2021, 10, 2749.	4.1	16
8	De novo variants in CACNA1E found in patients with intellectual disability, developmental regression and social cognition deficit but no seizures. <i>Molecular Autism</i> , 2021, 12, 69.	4.9	12
9	Clinical Reasoning: Rapidly progressive gait disorder and cranial nerves involvement in a 9-year-old boy. <i>Neurology</i> , 2020, 94, e330-e334.	1.1	0
10	Childhood neurodegeneration associated with a specific UBTF variant: a new case report and review of the literature. <i>BMC Neurology</i> , 2020, 20, 17.	1.8	15
11	Sultiame pharmacokinetic profile in plasma and erythrocytes after single oral doses: A pilot study in healthy volunteers. <i>Pharmacology Research and Perspectives</i> , 2020, 8, e00558.	2.4	8
12	Developmental trajectories of neuroanatomical alterations associated with the 16p11.2 Copy Number Variations. <i>NeuroImage</i> , 2019, 203, 116155.	4.2	9
13	Quantifying the Effects of 16p11.2 Copy Number Variants on Brain Structure: A Multisite Genetic-First Study. <i>Biological Psychiatry</i> , 2018, 84, 253-264.	1.3	56
14	Periinsular anterior quadrantotomy: technical note. <i>Journal of Neurosurgery: Pediatrics</i> , 2018, 21, 124-132.	1.3	11
15	Autosomal recessive primary microcephaly due to ASPM mutations: An update. <i>Human Mutation</i> , 2018, 39, 319-332.	2.5	53
16	Midazolam as a first-line treatment for neonatal seizures: Retrospective study. <i>Pediatrics International</i> , 2018, 60, 498-500.	0.5	12
17	Prenatal Brainstem Disruptions: Small Lesions – Big Problems. <i>Neuropediatrics</i> , 2017, 48, 350-355.	0.6	1
18	SCN1A Does Not Determine Seizure Duration in Children Unaffected by Dravet's Syndrome. <i>Journal of Pediatric Epilepsy</i> , 2017, 06, 111-114.	0.2	0

#	ARTICLE	IF	CITATIONS
19	Perinatal arterial ischemic stroke related to carotid artery occlusion. <i>European Journal of Paediatric Neurology</i> , 2016, 20, 639-648.	1.6	26
20	Defining the Effect of the 16p11.2 Duplication on Cognition, Behavior, and Medical Comorbidities. <i>JAMA Psychiatry</i> , 2016, 73, 20.	11.0	195
21	Early-onset encephalopathy with epilepsy associated with a novel splice site mutation in <i>SMC1A</i> . <i>American Journal of Medical Genetics, Part A</i> , 2015, 167, 3076-3081.	1.2	22
22	When should clinicians search for GLUT1 deficiency syndrome in childhood generalized epilepsies?. <i>European Journal of Paediatric Neurology</i> , 2015, 19, 170-175.	1.6	9
23	16p11.2 600 kb Duplications confer risk for typical and atypical Rolandic epilepsy. <i>Human Molecular Genetics</i> , 2014, 23, 6069-6080.	2.9	61
24	When is a child with status epilepticus likely to have Dravet syndrome?. <i>Epilepsy Research</i> , 2014, 108, 740-747.	1.6	16
25	A 600â€¦kb deletion syndrome at 16p11.2 leads to energy imbalance and neuropsychiatric disorders. <i>Journal of Medical Genetics</i> , 2012, 49, 660-668.	3.2	251
26	Anti-N-Methyl-Aspartate (NMDA) Receptor Encephalitis Mimicking a Primary Psychiatric Disorder in an Adolescent. <i>Journal of Child Neurology</i> , 2012, 27, 1607-1610.	1.4	29
27	An initial MRI picture of limbic encephalitis in subacute sclerosing panencephalitis. <i>European Journal of Paediatric Neurology</i> , 2011, 15, 544-546.	1.6	8