

# Svetlana Gataullina

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/479117/publications.pdf>

Version: 2024-02-01

15  
papers

365  
citations

933447

10  
h-index

1125743

13  
g-index

15  
all docs

15  
docs citations

15  
times ranked

704  
citing authors

#	ARTICLE	IF	CITATIONS
1	Epileptic spasms with terror during sleep in CDKL5 encephalopathy. <i>SLEEP Advances</i> , 2022, 3, .	0.2	2
2	<i>STXBP1</i> germline mutation and focal cortical dysplasia. <i>Epileptic Disorders</i> , 2021, 23, 143-147.	1.3	5
3	Lessons learned from 40 novel <i>PIGA</i> patients and a review of the literature. <i>Epilepsia</i> , 2020, 61, 1142-1155.	5.1	32
4	Neonatal SCN2A encephalopathy: A peculiar recognizable electroclinical sequence. <i>Epilepsy and Behavior</i> , 2020, 111, 107187.	1.7	12
5	Gene mutations in paediatric epilepsies cause NMDA $\hat{e}$ pathy, and phasic and tonic GABA $\hat{e}$ pathy. <i>Developmental Medicine and Child Neurology</i> , 2019, 61, 891-898.	2.1	20
6	Clinical study of 19 patients with <i>SCN8A</i> -related epilepsy: Two modes of onset regarding EEG and seizures. <i>Epilepsia</i> , 2019, 60, 845-856.	5.1	28
7	New insights in phenomenology and treatment of epilepsy in CDKL5 encephalopathy. <i>Epilepsy and Behavior</i> , 2019, 94, 308-311.	1.7	14
8	Is epilepsy the cause of comorbidities in Dravet syndrome?. <i>Developmental Medicine and Child Neurology</i> , 2018, 60, 8-8.	2.1	11
9	DMD and West syndrome. <i>Neuromuscular Disorders</i> , 2017, 27, 911-913.	0.6	5
10	From genotype to phenotype in Dravet disease. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2017, 44, 58-64.	2.0	142
11	Epilepsy in young <i>Tsc1</i> <sup>+/<math>\hat{a}</math>'</sup> mice exhibits age $\hat{e}$ dependent expression that mimics that of human tuberous sclerosis complex. <i>Epilepsia</i> , 2016, 57, 648-659.	5.1	25
12	Current and future treatment of infantile spasms. <i>Expert Opinion on Orphan Drugs</i> , 2015, 3, 1291-1297.	0.8	0
13	Seizures and epilepsy in hypoglycaemia caused by inborn errors of metabolism. <i>Developmental Medicine and Child Neurology</i> , 2015, 57, 194-199.	2.1	21
14	Topography of brain damage in metabolic hypoglycaemia is determined by age at which hypoglycaemia occurred. <i>Developmental Medicine and Child Neurology</i> , 2013, 55, 162-166.	2.1	28
15	Comorbidity and metabolic context are crucial factors determining neurological sequelae of hypoglycaemia. <i>Developmental Medicine and Child Neurology</i> , 2012, 54, 1012-1017.	2.1	20