

# Ronald L Thibert

## List of Publications by Year in descending order

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Version: 2024-02-01

43  
papers

1,569  
citations

394421

19  
h-index

315739

38  
g-index

45  
all docs

45  
docs citations

45  
times ranked

1959  
citing authors

#	ARTICLE	IF	CITATIONS
1	Parent Description of Anxiety in Angelman Syndrome. <i>Journal of Autism and Developmental Disorders</i> , 2022, 52, 3612-3625.	2.7	8
2	A multidisciplinary approach and consensus statement to establish standards of care for Angelman syndrome. <i>Molecular Genetics &amp; Genomic Medicine</i> , 2022, 10, e1843.	1.2	14
3	Longitudinal EEG model detects antisense oligonucleotide treatment effect and increased UBE3A in Angelman syndrome. <i>Brain Communications</i> , 2022, 4, .	3.3	5
4	Angelman syndrome genotypes manifest varying degrees of clinical severity and developmental impairment. <i>Molecular Psychiatry</i> , 2021, 26, 3625-3633.	7.9	41
5	Delta power robustly predicts cognitive function in Angelman syndrome. <i>Annals of Clinical and Translational Neurology</i> , 2021, 8, 1433-1445.	3.7	23
6	Biochemical Analysis of a Patient with Fumarate Hydratase Deficiency. <i>FASEB Journal</i> , 2021, 35, .	0.5	0
7	Teaching NeuroImage: Notched Delta and Angelman Syndrome. <i>Neurology</i> , 2021, 97, e1543-e1544.	1.1	3
8	The STARS Phase 2 Study. <i>Neurology</i> , 2021, 96, e1024-e1035.	1.1	12
9	Behavioral characterization of dup15q syndrome: Toward meaningful endpoints for clinical trials. <i>American Journal of Medical Genetics, Part A</i> , 2020, 182, 71-84.	1.2	21
10	Clinical report and biochemical analysis of a patient with fumarate hydratase deficiency. <i>American Journal of Medical Genetics, Part A</i> , 2020, 182, 504-507.	1.2	2
11	Biochemical Analysis of a Patient with Fumarate Hydratase Deficiency. <i>FASEB Journal</i> , 2020, 34, 1-1.	0.5	0
12	Buspirone for the treatment of anxiety-related symptoms in Angelman syndrome. <i>Psychiatric Genetics</i> , 2019, 29, 51-56.	1.1	11
13	How do we diagnose and treat epilepsy with myoclonic-atonic seizures (Doose syndrome)? Results of the Pediatric Epilepsy Research Consortium survey. <i>Epilepsy Research</i> , 2018, 144, 14-19.	1.6	14
14	Angelman syndrome in adolescence and adulthood: A retrospective chart review of 53 cases. <i>American Journal of Medical Genetics, Part A</i> , 2018, 176, 1327-1334.	1.2	27
15	Diazepam for outpatient treatment of nonconvulsive status epilepticus in pediatric patients with Angelman syndrome. <i>Epilepsy and Behavior</i> , 2018, 82, 74-80.	1.7	15
16	Myoclonus in Angelman syndrome. <i>Epilepsy and Behavior</i> , 2018, 82, 170-174.	1.7	20
17	Case 27-2018: A 3-Year-Old Boy with Seizures. <i>New England Journal of Medicine</i> , 2018, 379, 870-878.	27.0	4
18	Low glycemic index treatment for seizure control in Angelman syndrome: A case series from the Center for Dietary Therapy of Epilepsy at the Massachusetts General Hospital. <i>Epilepsy and Behavior</i> , 2017, 68, 45-50.	1.7	43

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19	Validation of a smartphone-based EEG among people with epilepsy: A prospective study. <i>Scientific Reports</i> , 2017, 7, 45567.	3.3	39
20	Delta rhythmicity is a reliable EEG biomarker in Angelman syndrome: a parallel mouse and human analysis. <i>Journal of Neurodevelopmental Disorders</i> , 2017, 9, 17.	3.1	74
21	Prevalence of gastrointestinal symptoms in Angelman syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2017, 173, 2703-2709.	1.2	21
22	Multisite Semiautomated Clinical Data Repository for Duplication 15q Syndrome: Study Protocol and Early Uses. <i>JMIR Research Protocols</i> , 2017, 6, e194.	1.0	4
23	A Rare Inherited 15q11.2-q13.1 Interstitial Duplication with Maternal Somatic Mosaicism, Renal Carcinoma, and Autism. <i>Frontiers in Genetics</i> , 2016, 7, 205.	2.3	4
24	Seizure treatment in Angelman syndrome: A case series from the Angelman Syndrome Clinic at Massachusetts General Hospital. <i>Epilepsy and Behavior</i> , 2016, 60, 138-141.	1.7	32
25	Novel Compound Heterozygous Mutations Expand the Recognized Phenotypes of <i>FARS2</i> -Linked Disease. <i>Journal of Child Neurology</i> , 2016, 31, 1127-1137.	1.4	36
26	Medication prescribing and patient-reported outcome measures in people with epilepsy in Bhutan. <i>Epilepsy and Behavior</i> , 2016, 59, 122-127.	1.7	11
27	Identification of a distinct developmental and behavioral profile in children with Dup15q syndrome. <i>Journal of Neurodevelopmental Disorders</i> , 2016, 8, 19.	3.1	47
28	The Use of Magnetic Resonance Spectroscopy in the Evaluation of Pediatric Patients With Seizures. <i>Pediatric Neurology</i> , 2016, 58, 57-66.	2.1	9
29	Electroencephalographic patterns during sleep in children with chromosome 15q11.2-13.1 duplications (Dup15q). <i>Epilepsy and Behavior</i> , 2016, 57, 133-136.	1.7	11
30	Rufinamide treatment for refractory epilepsy in a largely pediatric population. <i>Journal of Pediatric Epilepsy</i> , 2015, 01, 097-101.	0.2	2
31	Hippocampal Abnormalities in Magnetic Resonance Imaging (MRI) of 15q Duplication Syndromes. <i>Journal of Child Neurology</i> , 2015, 30, 333-338.	1.4	20
32	Gastrointestinal problems in 15q duplication syndrome. <i>European Journal of Medical Genetics</i> , 2015, 58, 191-193.	1.3	13
33	Early and effective treatment of <i>KCNQ2</i> encephalopathy. <i>Epilepsia</i> , 2015, 56, 685-691.	5.1	229
34	Angelman syndrome in adulthood. <i>American Journal of Medical Genetics, Part A</i> , 2015, 167, 331-344.	1.2	68
35	A survey of seizures and current treatments in 15q duplication syndrome. <i>Epilepsia</i> , 2014, 55, 396-402.	5.1	80
36	If not Angelman, what is it? a review of Angelman-like syndromes. <i>American Journal of Medical Genetics, Part A</i> , 2014, 164, 975-992.	1.2	80

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37	Neurologic Manifestations of Angelman Syndrome. <i>Pediatric Neurology</i> , 2013, 48, 271-279.	2.1	150
38	The Interstitial Duplication 15q11.2â€“q13 Syndrome Includes Autism, Mild Facial Anomalies and a Characteristic EEG Signature. <i>Autism Research</i> , 2013, 6, 268-279.	3.8	130
39	Low glycemic index treatment for seizures in Angelman syndrome. <i>Epilepsia</i> , 2012, 53, 1498-1502.	5.1	52
40	Levodopa Response Reveals Sepiapterin Reductase Deficiency in a Female Heterozygote with Adrenoleukodystrophy. <i>JIMD Reports</i> , 2011, 3, 79-82.	1.5	6
41	Reversible uncal herniation in a neonate with a large MCA infarct. <i>Brain and Development</i> , 2009, 31, 763-765.	1.1	5
42	Epilepsy in Angelman syndrome: A questionnaireâ€“based assessment of the natural history and current treatment options. <i>Epilepsia</i> , 2009, 50, 2369-2376.	5.1	115
43	Epilepsy and the sleepâ€“wake patterns found in Angelman syndrome. <i>Epilepsia</i> , 2009, 50, 2497-2500.	5.1	65