Tai-Heng Chen

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/1299264/publications.pdf

Version: 2024-02-01

516561 189801 2,691 61 16 50 citations h-index g-index papers 61 61 61 3117 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Nusinersen versus Sham Control in Infantile-Onset Spinal Muscular Atrophy. New England Journal of Medicine, 2017, 377, 1723-1732.	13.9	1,533
2	Nusinersen initiated in infants during the presymptomatic stage of spinal muscular atrophy: Interim efficacy and safety results from the Phase 2 NURTURE study. Neuromuscular Disorders, 2019, 29, 842-856.	0.3	401
3	Randomized, double-blind, placebo-controlled trial of hydroxyurea in spinal muscular atrophy. Neurology, 2010, 75, 2190-2197.	1.5	69
4	New and Developing Therapies in Spinal Muscular Atrophy: From Genotype to Phenotype to Treatment and Where Do We Stand?. International Journal of Molecular Sciences, 2020, 21, 3297.	1.8	67
5	Neurological involvement associated with COVID-19 infection in children. Journal of the Neurological Sciences, 2020, 418, 117096.	0.3	64
6	Posterior Reversible Encephalopathy Syndrome in Children. Journal of Child Neurology, 2013, 28, 1378-1386.	0.7	59
7	Prevalence and Risk Factors for Feeding and Swallowing Difficulties in Spinal Muscular Atrophy Types Il and III. Journal of Pediatrics, 2012, 160, 447-451.e1.	0.9	54
8	Infantile facioscapulohumeral muscular dystrophy revisited: Expansion of clinical phenotypes in patients with a very short EcoRI fragment. Neuromuscular Disorders, 2013, 23, 298-305.	0.3	42
9	Childhood Posterior Reversible Encephalopathy Syndrome: Clinicoradiological Characteristics, Managements, and Outcome. Frontiers in Pediatrics, 2020, 8, 585.	0.9	38
10	Aicardi syndrome in a 47, XXY male neonate with lissencephaly and holoprosencephaly. Journal of the Neurological Sciences, 2009, 278, 138-140.	0.3	28
11	Combined noninvasive ventilation and mechanical inâ€exsufflator in the treatment of pediatric acute neuromuscular respiratory failure. Pediatric Pulmonology, 2014, 49, 589-596.	1.0	28
12	Identification of bidirectional gene conversion between SMN1 and SMN2 by simultaneous analysis of SMN dosage and hybrid genes in a Chinese population. Journal of the Neurological Sciences, 2011, 308, 83-87.	0.3	24
13	Selective Neuromuscular Denervation in Taiwanese Severe SMA Mouse Can Be Reversed by Morpholino Antisense Oligonucleotides. PLoS ONE, 2016, 11, e0154723.	1.1	23
14	Multifaceted roles of microRNAs: From motor neuron generation in embryos to degeneration in spinal muscular atrophy. ELife, $2019, 8, .$	2.8	23
15	Metabolic and Nutritional Issues Associated with Spinal Muscular Atrophy. Nutrients, 2020, 12, 3842.	1.7	20
16	Correlation between muscle involvement, phenotype and D4Z4 fragment size in facioscapulohumeral muscular dystrophy. Neuromuscular Disorders, 2012, 22, 331-338.	0.3	19
17	The use of hospital medical records for child injury surveillance in northern Malawi. Tropical Doctor, 2009, 39, 170-172.	0.2	16
18	A Taiwanese Boy With Congenital Generalized Lipodystrophy Caused by Homozygous Ile262fs Mutation in the <i>BSCL2</i> Gene. Kaohsiung Journal of Medical Sciences, 2010, 26, 615-620.	0.8	16

#	Article	IF	CITATIONS
19	Care for Patients With Neuromuscular Disorders in the COVID-19 Pandemic Era. Frontiers in Neurology, 2021, 12, 607790.	1.1	16
20	Identification of <i> KLHL40 < /i > mutations by targeted next-generation sequencing facilitated a prenatal diagnosis in a family with three consecutive affected fetuses with fetal akinesia deformation sequence. Prenatal Diagnosis, 2016, 36, 1135-1138.</i>	1.1	15
21	Combined noninvasive ventilation and mechanical insufflator–exsufflator for acute respiratory failure in patients with neuromuscular disease: effectiveness and outcome predictors. Therapeutic Advances in Respiratory Disease, 2019, 13, 175346661987592.	1.0	13
22	Hydroxychloroquine was associated with reduced risk of new-onset diabetes mellitus in patients with SjA¶gren syndrome. QJM - Monthly Journal of the Association of Physicians, 2019, 112, 757-762.	0.2	13
23	Posterior reversible encephalopathy syndrome in critically ill children: a case series. Intensive Care Medicine, 2013, 39, 155-156.	3.9	12
24	Comparison of Clinical Characteristics Between Febrile and Afebrile Seizures Associated With Acute Gastroenteritis in Childhood. Frontiers in Pediatrics, 2020, 8, 167.	0.9	10
25	Gastrointestinal Involvements in Children With COVID-related Multisystem Inflammatory Syndrome. Gastroenterology, 2021, 160, 1887-1888.	0.6	10
26	Symmetric Atrophy of Bilateral Distal Upper Extremities and HyperlgEaemia in a Male Adolescent With Hirayama Disease. Journal of Child Neurology, 2010, 25, 371-374.	0.7	9
27	Posterior Reversible Encephalopathy Syndrome With Spinal Cord Involvement in Children. Journal of Child Neurology, 2017, 32, 112-119.	0.7	7
28	Early-Onset Infantile Facioscapulohumeral Muscular Dystrophy: A Timely Review. International Journal of Molecular Sciences, 2020, 21, 7783.	1.8	7
29	Circulating microRNAs as potential biomarkers and therapeutic targets in spinal muscular atrophy. Therapeutic Advances in Neurological Disorders, 2020, 13, 175628642097995.	1.5	7
30	Reliability and Validity of Outcome Measures of In-Hospital and At-Home Visits in a Randomized, Double-Blind, Placebo-Controlled Trial for Spinal Muscular Atrophy. Journal of Child Neurology, 2014, 29, 1680-1684.	0.7	6
31	Emergency room visits and admission rates of children with neuromuscular disorders: A 10-year experience in a medical center in Taiwan. Pediatrics and Neonatology, 2019, 60, 405-410.	0.3	5
32	Noninvasive Ventilation and Mechanical Insufflator-Exsufflator for Acute Respiratory Failure in Children With Neuromuscular Disorders. Frontiers in Pediatrics, 2020, 8, 593282.	0.9	5
33	Letters to the Editor. Journal of Paediatrics and Child Health, 2008, 44, 154-155.	0.4	4
34	Prostaglandin I2 Analogues Enhance Growth-Related Oncogene-α Expression in Human Monocyte-Derived Dendritic Cells. Inflammation, 2010, 33, 334-343.	1.7	3
35	Hemifacial Atrophy With Intracranial Calcification. Pediatric Neurology, 2013, 49, 72-73.	1.0	3
36	Benign reversible course in infants manifesting clinicopathological features of fatal mitochondrial myopathy due to m.14674 T>C mt-tRNAGlu mutation. QJM - Monthly Journal of the Association of Physicians, 2013, 106, 953-954.	0.2	3

3

#	Article	IF	CITATIONS
37	Noninvasive airway approaches for acute neuromuscular respiratory failure in emergency departments. Pediatric Pulmonology, 2017, 52, E55-E57.	1.0	3
38	Spontaneous hemopneumothorax simulating acute abdominal affections. Pediatric Pulmonology, 2014, 49, E1-4.	1.0	2
39	Transient Cerebral Arteriopathy in a Child Associated With Cytomegalovirus Infection. Child Neurology Open, 2015, 2, 2329048X1560202.	0.5	2
40	Neonatal lupus. QJM - Monthly Journal of the Association of Physicians, 2020, 113, 905-906.	0.2	2
41	Massive cerebral air embolism in a preterm with fetal alcohol syndrome. Neurology India, 2009, 57, 227.	0.2	1
42	Increasing Recognition of Cases With Male Aicardi Syndrome. Journal of Child Neurology, 2010, 25, 129-129.	0.7	1
43	Bluish discoloration of hemiscrotum. Journal of Paediatrics and Child Health, 2011, 47, 309-309.	0.4	1
44	Hair-on-End Skull in an Infant without Anemia. Journal of Pediatrics, 2012, 161, 367.	0.9	1
45	A child with severe stridor. Emergency Medicine Journal, 2013, 30, 603-603.	0.4	1
46	The football sign in a neonate. QJM - Monthly Journal of the Association of Physicians, 2014, 107, 237-238.	0.2	1
47	An intra-abdominal oval-shaped mass. Journal of Paediatrics and Child Health, 2014, 50, 162-162.	0.4	1
48	Respiratory management for patients with neuromuscular disorders during the COVID-19 pandemic. Therapeutic Advances in Respiratory Disease, 2020, 14, 175346662095378.	1.0	1
49	Comment on: "Myostatin inhibition in combination with antisense oligonucleotide therapy improves outcomes in spinal muscular atrophy―by Zhou et al Journal of Cachexia, Sarcopenia and Muscle, 2020, 11, 1377-1378.	2.9	1
50	Emerging Innovative Therapies of Spinal Muscular Atrophy: Current Knowledge and Perspectives. Frontiers in Clinical Drug Research CNS and Neurological Disorders, 2020, , 1-34.	0.1	1
51	Is the male Aicardi's characterized by 46 XXY karyotpe?. Journal of the Neurological Sciences, 2009, 284, 222.	0.3	0
52	Letters to the Editor. Journal of Paediatrics and Child Health, 2010, 46, 280-281.	0.4	0
53	P1.52 Infantile facioscapulohumeral muscular dystrophy with a very short 4q35 fragment: additional features. Neuromuscular Disorders, 2010, 20, 616.	0.3	0
54	P3.36 Gene conversion between SMN1 and SMN2 in normals, carriers and spinal muscular atrophy patients. Neuromuscular Disorders, 2010, 20, 652.	0.3	O

Tai-Heng Chen

#	Article	IF	CITATIONS
55	P2.36 Correlation between muscle involvement, phenotype and D4Z4 fragment size in facioscapulohumeral muscular dystrophy. Neuromuscular Disorders, 2011, 21, 671.	0.3	О
56	Elevated serum αâ€fetoprotein in a neonate with cutaneous infantile hemangioendothelioma. Pediatrics International, 2011, 53, 258-261.	0.2	0
57	An Unusual Orbital Mass with Dural Tail Signs. Internal Medicine, 2012, 51, 2063-2064.	0.3	0
58	The Peeling Dermatitis With a Peculiar Demarcation. American Journal of the Medical Sciences, 2013, 346, 421.	0.4	0
59	Methodological considerations in combined noninvasive ventilation and mechanical inâ€exsufflator. Pediatric Pulmonology, 2014, 49, 1045-1046.	1.0	O
60	Community-acquired pneumonia in pediatric patients with acute neuromuscular respiratory failure: A microbiologic perspective. Pediatric Pulmonology, 2014, 49, 827-828.	1.0	0
61	The "wrench-head―appearance of thigh muscle CT in infantile facioscapulohumeral muscular dystrophy. Acta Neurologica Belgica, 2021, , 1.	0.5	0