Cyril Gitiaux

List of Publications by Year
in descending order

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40	915	567144 15	501076 28
papers	citations	h-index	g-index

42	42	42	1233
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Carpal tunnel syndrome and finger deformities in children with mucopolysaccharidoses and mucolipidoses: a retrospective review of 52 patients. Journal of Hand Surgery: European Volume, 2022, 47, 469-474.	0.5	1
2	FDX2 and ISCU Gene Variations Lead to Rhabdomyolysis With Distinct Severity and Iron Regulation. Neurology: Genetics, 2022, 8, e648.	0.9	4
3	Efficacy and tolerance of corticosteroids and methotrexate in patients with juvenile dermatomyositis: a retrospective cohort study. Rheumatology, 2022, , .	0.9	0
4	Involvement of Type I Interferon Signaling in Muscle Stem Cell Proliferation During Dermatomyositis. Neurology, 2022, 98, .	1.5	13
5	Rituximab Therapy in the Treatment of Juvenile Myasthenia Gravis: The French Experience. Neurology, 2022, , 10.1212/WNL.0000000000200288.	1.5	10
6	Neuroinflammatory Disease following Severe Acute Respiratory Syndrome Coronavirus 2 Infection in Children. Journal of Pediatrics, 2022, 247, 22-28.e2.	0.9	15
7	From Diagnosis to Prognosis: Revisiting the Meaning of Muscle ISG15 Overexpression in Juvenile Inflammatory Myopathies. Arthritis and Rheumatology, 2021, 73, 1044-1052.	2.9	13
8	JAK inhibitors are effective in a subset of patients with juvenile dermatomyositis: a monocentric retrospective study. Rheumatology, 2021, 60, 5801-5808.	0.9	52
9	Dysregulation of the NRG1/ERBB pathway causes a developmental disorder with gastrointestinal dysmotility in humans. Journal of Clinical Investigation, 2021, 131, .	3.9	24
10	Electro-clinical features in epileptic children with chromosome 15q duplication syndrome. Clinical Neurophysiology, 2021, 132, 1126-1137.	0.7	0
11	Loss of function mutations in GEMIN5 cause a neurodevelopmental disorder. Nature Communications, 2021, 12, 2558.	5.8	28
12	Reply. Arthritis and Rheumatology, 2021, 73, 1566-1567.	2.9	0
13	Postnatal Diagnostic Workup in Children With Arthrogryposis: A Series of 82 Patients. Journal of Child Neurology, 2021, 36, 088307382110229.	0.7	0
14	Inflammatory myopathies in childhood. Neuromuscular Disorders, 2021, 31, 1051-1061.	0.3	6
15	Sebelipase alfa enzyme replacement therapy in Wolman disease: a nationwide cohort with up to ten years of follow-up. Orphanet Journal of Rare Diseases, 2021, 16, 507.	1.2	8
16	Circulating Interferonâ€Î± Measured With a Highly Sensitive Assay as a Biomarker for Juvenile Inflammatory Myositis Activity: Comment on the Article by Mathian et al. Arthritis and Rheumatology, 2020, 72, 195-197.	2.9	15
17	239th ENMC International Workshop: Classification of dermatomyositis, Amsterdam, the Netherlands, 14â€“16 December 2018. Neuromuscular Disorders, 2020, 30, 70-92.	0.3	148
18	Inhibition of IFNÎ± secretion in cells from patients with juvenile dermatomyositis under TBK1 inhibitor treatment revealed by single-molecular assay technology. Rheumatology, 2020, 59, 1171-1174.	0.9	5

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19	Severe Abdominal Manifestations in Juvenile Dermatomyositis. Journal of Pediatric Gastroenterology and Nutrition, 2020, 70, 247-251.	0.9	12
20	Anti-MDA5 juvenile idiopathic inflammatory myopathy: a specific subgroup defined by differentially enhanced interferon-Î± signalling. Rheumatology, 2020, 59, 1927-1937.	0.9	26
21	Diagnosis and Management of Carpal Tunnel Syndrome in Children with Mucopolysaccharidosis: A 10 Year Experience. Diagnostics, 2020, 10, 5.	1.3	13
22	Cardiac valve involvement in ADAR-related type I interferonopathy. Journal of Medical Genetics, 2020, 57, 475-478.	1.5	19
23	Deciphering the natural history of SCA7 in children. European Journal of Neurology, 2020, 27, 2267-2276.	1.7	12
24	Movement disorders in patients with alternating hemiplegia. Neurology, 2020, 94, e1378-e1385.	1.5	14
25	Severe Acute Flaccid Myelitis Associated With Enterovirus in Children: Two Phenotypes for Two Evolution Profiles?. Frontiers in Neurology, 2020, 11, 343.	1.1	6
26	Muscle ischaemia associated with NXP2 autoantibodies: a severe subtype of juvenile dermatomyositis. Rheumatology, 2018, 57, 873-879.	0.9	44
27	Myogenic Progenitor Cells Exhibit Type I Interferonâ€“Driven Proangiogenic Properties and Molecular Signature During Juvenile Dermatomyositis. Arthritis and Rheumatology, 2018, 70, 134-145.	2.9	38
28	Clinical features and evolution of juvenile myasthenia gravis in a French cohort. Muscle and Nerve, 2018, 57, 603-609.	1.0	22
29	A child with severe juvenile dermatomyositis treated with ruxolitinib. Brain, 2018, 141, e80-e80.	3.7	58
30	PLA2G6-associated neurodegeneration: Lessons from neurophysiological findings. European Journal of Paediatric Neurology, 2018, 22, 854-861.	0.7	9
31	Prenatal and postnatal presentations of corpus callosum agenesis with polymicrogyria caused by EGP5 mutation. American Journal of Medical Genetics, Part A, 2017, 173, 706-711.	0.7	12
32	Clinical, laboratory and molecular findings and long-term follow-up data in 96 French patients with PMM2-CDG (phosphomannomutase 2-congenital disorder of glycosylation) and review of the literature. Journal of Medical Genetics, 2017, 54, 843-851.	1.5	88
33	Acute axonal neuropathy subtype of Guillain BarrÃ© syndrome in a French pediatric series: Adequate follow-up may require repetitive electrophysiological studies. European Journal of Paediatric Neurology, 2017, 21, 891-897.	0.7	5
34	Presentations and outcomes of juvenile dermatomyositis patients admitted to intensive care units. Rheumatology, 2017, 56, 1814-1816.	0.9	7
35	Vasculopathy-related clinical and pathological features are associated with severe juvenile dermatomyositis. Rheumatology, 2016, 55, kev359.	0.9	21
36	Motor neuropathy contributes to crouching in patients with Dravet syndrome. Neurology, 2016, 87, 277-281.	1.5	37

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37	From splitting GLUT1 deficiency syndromes to overlapping phenotypes. European Journal of Medical Genetics, 2015, 58, 443-454.	0.7	52
38	Brain magnetic resonance imaging pattern and outcome in children with haemolyticâ€uraemic syndrome and neurological impairment treated with eculizumab. Developmental Medicine and Child Neurology, 2013, 55, 758-765.	1.1	41
39	Cerebral vasculitis in severe Kawasaki disease: early detection by magnetic resonance imaging and good outcome after intensive treatment. Developmental Medicine and Child Neurology, 2012, 54, 1160-1163.	1.1	35
40	A child with severe juvenile dermatomyositis treated with ruxolitinib. Journal of Financial Econometrics, 0, , .	0.8	0