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	239th ENMC International Workshop: Classification of dermatomyositis, Amsterdam, the Netherlands, 14â€“16 December 2018. Neuromuscular Disorders, 2020, 30, 70-92.	0.3	148
2	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
3	A child with severe juvenile dermatomyositis treated with ruxolitinib. Brain, 2018, 141, e80-e80.	3.7	58
4	From splitting GLUT1 deficiency syndromes to overlapping phenotypes. European Journal of Medical Genetics, 2015, 58, 443-454.	0.7	52
5	JAK inhibitors are effective in a subset of patients with juvenile dermatomyositis: a monocentric retrospective study. Rheumatology, 2021, 60, 5801-5808.	0.9	52
6	Muscle ischaemia associated with NXP2 autoantibodies: a severe subtype of juvenile dermatomyositis. Rheumatology, 2018, 57, 873-879.	0.9	44
7	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
8	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
9	Motor neuropathy contributes to crouching in patients with Dravet syndrome. Neurology, 2016, 87, 277-281.	1.5	37
10	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
11	Loss of function mutations in GEMIN5 cause a neurodevelopmental disorder. Nature Communications, 2021, 12, 2558.	5.8	28
12	Anti-MDA5 juvenile idiopathic inflammatory myopathy: a specific subgroup defined by differentially enhanced interferon-Î± signalling. Rheumatology, 2020, 59, 1927-1937.	0.9	26
13	Dysregulation of the NRG1/ERBB pathway causes a developmental disorder with gastrointestinal dysmotility in humans. Journal of Clinical Investigation, 2021, 131, .	3.9	24
14	Clinical features and evolution of juvenile myasthenia gravis in a French cohort. Muscle and Nerve, 2018, 57, 603-609.	1.0	22
15	Vasculopathy-related clinical and pathological features are associated with severe juvenile dermatomyositis. Rheumatology, 2016, 55, kev359.	0.9	21
16	Cardiac valve involvement in ADAR-related type I interferonopathy. Journal of Medical Genetics, 2020, 57, 475-478.	1.5	19
17	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
18	Neuroinflammatory Disease following Severe Acute Respiratory Syndrome Coronavirus 2 Infection in Children. Journal of Pediatrics, 2022, 247, 22-28.e2.	0.9	15

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19	Movement disorders in patients with alternating hemiplegia. Neurology, 2020, 94, e1378-e1385.	1.5	14
20	Diagnosis and Management of Carpal Tunnel Syndrome in Children with Mucopolysaccharidosis: A 10 Year Experience. Diagnostics, 2020, 10, 5.	1.3	13
21	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
22	Involvement of Type I Interferon Signaling in Muscle Stem Cell Proliferation During Dermatomyositis. Neurology, 2022, 98, .	1.5	13
23	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
24	Severe Abdominal Manifestations in Juvenile Dermatomyositis. Journal of Pediatric Gastroenterology and Nutrition, 2020, 70, 247-251.	0.9	12
25	Deciphering the natural history of SCA7 in children. European Journal of Neurology, 2020, 27, 2267-2276.	1.7	12
26	Rituximab Therapy in the Treatment of Juvenile Myasthenia Gravis: The French Experience. Neurology, 2022, , 10.1212/WNL.0000000000200288.	1.5	10
27	PLA2G6-associated neurodegeneration: Lessons from neurophysiological findings. European Journal of Paediatric Neurology, 2018, 22, 854-861.	0.7	9
28	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
29	Presentations and outcomes of juvenile dermatomyositis patients admitted to intensive care units. Rheumatology, 2017, 56, 1814-1816.	0.9	7
30	Severe Acute Flaccid Myelitis Associated With Enterovirus in Children: Two Phenotypes for Two Evolution Profiles?. Frontiers in Neurology, 2020, 11, 343.	1.1	6
31	Inflammatory myopathies in childhood. Neuromuscular Disorders, 2021, 31, 1051-1061.	0.3	6
32	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
33	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
34	FDX2 and ISCU Gene Variations Lead to Rhabdomyolysis With Distinct Severity and Iron Regulation. Neurology: Genetics, 2022, 8, e648.	0.9	4
35	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
36	A child with severe juvenile dermatomyositis treated with ruxolitinib. Journal of Financial Econometrics, 0, , .	0.8	0

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37	Electro-clinical features in epileptic children with chromosome 15q duplication syndrome. Clinical Neurophysiology, 2021, 132, 1126-1137.	0.7	0
38	Reply. Arthritis and Rheumatology, 2021, 73, 1566-1567.	2.9	0
39	Postnatal Diagnostic Workup in Children With Arthrogryposis: A Series of 82 Patients. Journal of Child Neurology, 2021, 36, 088307382110229.	0.7	0
40	Efficacy and tolerance of corticosteroids and methotrexate in patients with juvenile dermatomyositis: a retrospective cohort study. Rheumatology, 2022, , .	0.9	0