

Francesco Gavazzi

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/4745574/publications.pdf>

Version: 2024-02-01

14
papers

371
citations

1163065

8
h-index

1058452

14
g-index

15
all docs

15
docs citations

15
times ranked

438
citing authors

#	ARTICLE	IF	CITATIONS
1	Hodgkin lymphoma in an individual with <i>TREX1</i> -mediated Aicardi Goutières syndrome. <i>Pediatric Blood and Cancer</i> , 2022, 69, e29322.	1.5	1
2	The 2021 European Alliance of Associations for Rheumatology/American College of Rheumatology points to consider for diagnosis and management of autoinflammatory type I interferonopathies: CANDLE/PRAAS, SAVI and AGS. <i>Annals of the Rheumatic Diseases</i> , 2022, 81, 601-613.	0.9	31
3	The 2021 European Alliance of Associations for Rheumatology/American College of Rheumatology Points to Consider for Diagnosis and Management of Autoinflammatory Type I Interferonopathies: CANDLE, PRAAS, SAVI, and AGS. <i>Arthritis and Rheumatology</i> , 2022, 74, 735-751.	5.6	23
4	Late-Onset Aicardi-Goutières Syndrome: A Characterization of Presenting Clinical Features. <i>Pediatric Neurology</i> , 2021, 115, 1-6.	2.1	18
5	Hepatic Involvement in Aicardi-Goutières Syndrome. <i>Neuropediatrics</i> , 2021, 52, 441-447.	0.6	6
6	The epileptology of Aicardi-Goutières syndrome: electro-clinical-radiological findings. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2021, 86, 197-209.	2.0	2
7	Acquisition of Developmental Milestones in Hypomyelination With Atrophy of the Basal Ganglia and Cerebellum and Other TUBB4A-Related Leukoencephalopathy. <i>Journal of Child Neurology</i> , 2021, , 088307382100097.	1.4	1
8	Acquisition of Developmental Milestones in Hypomyelination With Atrophy of the Basal Ganglia and Cerebellum and Other TUBB4A-Related Leukoencephalopathy. <i>Journal of Child Neurology</i> , 2021, 36, 805-811.	1.4	3
9	Reliability of the Telemedicine Application of the Gross Motor Function Measure-88 in Patients With Leukodystrophy. <i>Pediatric Neurology</i> , 2021, 125, 34-39.	2.1	6
10	Developmental Outcomes of Aicardi Goutières Syndrome. <i>Journal of Child Neurology</i> , 2020, 35, 7-16.	1.4	40
11	Genetic and phenotypic spectrum associated with <i>IFIH1</i> gain-of-function. <i>Human Mutation</i> , 2020, 41, 837-849.	2.5	63
12	Janus Kinase Inhibition in the Aicardi-Goutières Syndrome. <i>New England Journal of Medicine</i> , 2020, 383, 986-989.	27.0	109
13	Development of a neurologic severity scale for Aicardi Goutières Syndrome. <i>Molecular Genetics and Metabolism</i> , 2020, 130, 153-160.	1.1	25
14	Neonatal detection of Aicardi Goutières Syndrome by increased C26:0 lysophosphatidylcholine and interferon signature on newborn screening blood spots. <i>Molecular Genetics and Metabolism</i> , 2017, 122, 134-139.	1.1	43