Adriana A De Jesus

List of Publications by Year
in descending order

18 papers	2,082 citations	687363 13 h-index	839539 18 g-index
19 all docs	19 docs citations	19 times ranked	3133 citing authors

#	Article	IF	Citations
1	NEMO-NDAS: A Panniculitis in the Young Representing an Autoinflammatory Disorder in Disguise. American Journal of Dermatopathology, 2022, 44, e64-e66.	0.6	3
2	Protein kinase R is an innate immune sensor of proteotoxic stress via accumulation of cytoplasmic IL-24. Science Immunology, 2022, 7, eabi6763.	11.9	22
3	Immunopathological signatures in multisystem inflammatory syndrome in children and pediatric COVID-19. Nature Medicine, 2022, 28, 1050-1062.	30.7	144
4	Post-SARS-CoV-2 Vaccine Monitoring of Disease Flares in Autoinflammatory Diseases. Journal of Clinical Immunology, 2022, 42, 732-735.	3.8	3
5	Genetically programmed alternative splicing of NEMO mediates an autoinflammatory disease phenotype. Journal of Clinical Investigation, 2022, 132, .	8.2	15
6	The 2021 EULAR/American College of Rheumatology points to consider for diagnosis, management and monitoring of the interleukin-1 mediated autoinflammatory diseases: cryopyrin-associated periodic syndromes, tumour necrosis factor receptor-associated periodic syndrome, mevalonate kinase deficiency, and deficiency of the interleukin-1 receptor antagonist. Annals of the Rheumatic Diseases, 2022, 81, 907-921.	0.9	38
7	Case Report: Novel SAVI-Causing Variants in STING1 Expand the Clinical Disease Spectrum and Suggest a Refined Model of STING Activation. Frontiers in Immunology, 2021, 12, 636225.	4.8	18
8	Immunodeficiency and bone marrow failure with mosaic and germline TLR8 gain of function. Blood, 2021, 137, 2450-2462.	1.4	47
9	A clinical score to guide in decision making for monogenic type I IFNopathies. Pediatric Research, 2020, 87, 745-752.	2.3	16
10	A novel STING1 variant causes a recessive form of STING-associated vasculopathy with onset in infancy (SAVI). Journal of Allergy and Clinical Immunology, 2020, 146, 1204-1208.e6.	2.9	45
11	Expression of interferon-regulated genes in juvenile dermatomyositis versus Mendelian autoinflammatory interferonopathies. Arthritis Research and Therapy, 2020, 22, 69.	3.5	39
12	Distinct interferon signatures and cytokine patterns define additional systemic autoinflammatory diseases. Journal of Clinical Investigation, 2020, 130, 1669-1682.	8.2	142
13	Severe autoinflammation in 4 patients with C-terminal variants in cell division control protein 42 homolog (CDC42) successfully treated with IL-1Î² inhibition. Journal of Allergy and Clinical Immunology, 2019, 144, 1122-1125.e6.	2.9	85
14	Novel proteasome assembly chaperone mutations in PSMG2/PAC2 cause the autoinflammatory interferonopathy CANDLE/PRAAS4. Journal of Allergy and Clinical Immunology, 2019, 143, 1939-1943.e8.	2.9	82
15	Development of a Validated Interferon Score Using NanoString Technology. Journal of Interferon and Cytokine Research, 2018, 38, 171-185.	1.2	120
16	Interleukin-18 diagnostically distinguishes and pathogenically promotes human and murine macrophage activation syndrome. Blood, 2018, 131, 1442-1455.	1.4	288
17	JAK1/2 inhibition with baricitinib in the treatment of autoinflammatory interferonopathies. Journal of Clinical Investigation, 2018, 128, 3041-3052.	8.2	387
18	An activating NLRC4 inflammasome mutation causes autoinflammation with recurrent macrophage activation syndrome. Nature Genetics, 2014, 46, 1140-1146.	21.4	585