Tracy A Briggs

List of Publications by Year
in descending order

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28 papers	1,596 citations	471061 17 h-index	476904 29 g-index
31 all docs	31 docs citations	31 times ranked	3131 citing authors

#	Article	IF	Citations
1	Detection of interferon alpha protein reveals differential levels and cellular sources in disease. Journal of Experimental Medicine, 2017, 214, 1547-1555.	4.2	288
2	Tartrate-resistant acid phosphatase deficiency causes a bone dysplasia with autoimmunity and a type I interferon expression signature. Nature Genetics, 2011, 43, 127-131.	9.4	214
3	Assessment of Type I Interferon Signaling in Pediatric Inflammatory Disease. Journal of Clinical Immunology, 2017, 37, 123-132.	2.0	163
4	Blood RNA analysis can increase clinical diagnostic rate and resolve variants of uncertain significance. Genetics in Medicine, 2020, 22, 1005-1014.	1.1	99
5	Oculo-auriculo-vertebral spectrum: Clinical and molecular analysis of 51 patients. European Journal of Medical Genetics, 2015, 58, 455-465.	0.7	83
6	Severe type I interferonopathy and unrestrained interferon signaling due to a homozygous germline mutation in STAT2. Science Immunology, 2019, 4, .	5.6	80
7	Spondyloenchondrodysplasia Due to Mutations in ACP5: A Comprehensive Survey. Journal of Clinical Immunology, 2016, 36, 220-234.	2.0	71
8	Hereditary Alpha-Tryptasemia: UK Prevalence and Variability in Disease Expression. Journal of Allergy and Clinical Immunology: in Practice, 2020, 8, 3549-3556.	2.0	70
9	FOXP2variants in 14 individuals with developmental speech and language disorders broaden the mutational and clinical spectrum. Journal of Medical Genetics, 2017, 54, 64-72.	1.5	67
10	Genetic and phenotypic spectrum associated with IFIH1 gainâ€ofâ€function. Human Mutation, 2020, 41, 837-849.	1.1	63
11	Genetic, Phenotypic, and Interferon Biomarker Status in ADAR1-Related Neurological Disease. Neuropediatrics, 2017, 48, 166-184.	0.3	62
12	'Reluctant pioneer': A qualitative study of doctors' experiences as patients with long COVID. Health Expectations, 2021, 24, 833-842.	1.1	54
13	Long COVID risk - a signal to address sex hormones and women's health. Lancet Regional Health - Europe, The, 2021, 11, 100242.	3.0	48
14	Tartrateâ€Resistant Acid Phosphatase Deficiency in the Predisposition to Systemic Lupus Erythematosus. Arthritis and Rheumatology, 2017, 69, 131-142.	2.9	47
15	Brief Report: Vitamin D Deficiency Is Associated With Endothelial Dysfunction and Increases Type I Interferon Gene Expression in a Murine Model of Systemic Lupus Erythematosus. Arthritis and Rheumatology, 2016, 68, 2929-2935.	2.9	30
16	Temple syndrome as a result of isolated hypomethylation of the 14q32 imprinted DLK1/MEG3 region. American Journal of Medical Genetics, Part A, 2016, 170, 170-175.	0.7	25
17	Disease modeling of core pre-mRNA splicing factor haploinsufficiency. Human Molecular Genetics, 2019, 28, 3704-3723.	1.4	24
18	Type I interferon in patients with systemic autoimmune rheumatic disease is associated with haematological abnormalities and specific autoantibody profiles. Arthritis Research and Therapy, 2019, 21, 147.	1.6	20

#	Article	IF	Citations
19	Curation and expansion of Human Phenotype Ontology for defined groups of inborn errors of immunity. Journal of Allergy and Clinical Immunology, 2022, 149, 369-378.	1.5	16
20	Personalised virtual gene panels reduce interpretation workload and maintain diagnostic rates of proband-only clinical exome sequencing for rare disorders. Journal of Medical Genetics, 2022, 59, 393-398.	1.5	14
21	Differential levels of IFNÎ± subtypes in autoimmunity and viral infection. Cytokine, 2021, 144, 155533.	1.4	12
22	MRSD: A quantitative approach for assessing suitability of RNA-seq in the investigation of mis-splicing in Mendelian disease. American Journal of Human Genetics, 2022, 109, 210-222.	2.6	12
23	Childhood-onset autoimmune cytopenia as the presenting feature of biallelicACP5mutations. Pediatric Blood and Cancer, 2017, 64, 306-310.	0.8	6
24	The (Orf)ull truth about IRF5 and type I interferons in SLE. Nature Reviews Rheumatology, 2020, 16, 543-544.	3.5	6
25	Ligase IV syndrome can present with microcephaly and radial ray anomalies similar to Fanconi anaemia plus fatal kidney malformations. European Journal of Medical Genetics, 2020, 63, 103974.	0.7	5
26	Biallelic Mutations in MTPAP Associated with a Lethal Encephalopathy. Neuropediatrics, 2020, 51, 178-184.	0.3	3
27	The diagnostic utility of clinical exome sequencing in 60 patients with hearing loss disorders: A singleâ€institution experience. Clinical Otolaryngology, 2021, 46, 1257-1262.	0.6	3
28	Further delineation of phenotypic spectrum of <scp>SCN2A</scp>â€related disorder. American Journal of Medical Genetics, Part A, 2022, 188, 867-877.	0.7	3