Nisha Limaye

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/7094934/publications.pdf

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22	623	1478505 6	1058476 14
papers	citations	h-index	g-index

22	22	22	981
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Somatic Activating PIK3CA Mutations Cause Venous Malformation. American Journal of Human Genetics, 2015, 97, 914-921.	6.2	244
2	Blue Rubber Bleb Nevus (BRBN) Syndrome Is Caused by Somatic TEK (TIE2) Mutations. Journal of Investigative Dermatology, 2017, 137, 207-216.	0.7	148
3	PDGFRB gain-of-function mutations in sporadic infantile myofibromatosis. Human Molecular Genetics, 2017, 26, 1801-1810.	2.9	77
4	Association of PDGFRB Mutations With Pediatric Myofibroma and Myofibromatosis. JAMA Dermatology, 2019, 155, 946.	4.1	43
5	Liquid biopsy for mutational profiling of locoregional recurrent and/or metastatic head and neck squamous cell carcinoma. Oral Oncology, 2020, 104, 104631.	1.5	42
6	Liquid Biopsy to Detect Minimal Residual Disease: Methodology and Impact. Cancers, 2021, 13, 5364.	3.7	31
7	Common Transcriptomic Effects of Abatacept and Other DMARDs on Rheumatoid Arthritis Synovial Tissue. Frontiers in Immunology, 2021, 12, 724895.	4.8	8
8	Development of SARS-CoV2 humoral response including neutralizing antibodies is not sufficient to protect patients against fatal infection. Scientific Reports, 2022, 12, 2077.	3.3	8
9	High p16^{INK4a}, a marker of cellular senescence, is associated with renal injury, impairment and outcome in lupus nephritis. RMD Open, 2021, 7, e001844.	3.8	7
10	Tumor sequencing is useful to refine the analysis of germline variants in unexplained high-risk breast cancer families. Breast Cancer Research, 2020, 22, 36.	5.0	6
11	Novel human pathological mutations. Gene symbol: KRIT1. Disease: cerebral cavernous malformation. Human Genetics, 2007, 122, 549-50.	3.8	4
12	Novel human pathological mutations. Gene symbol: KRIT1. Disease: cerebral cavernous malformation. Human Genetics, 2007, 122, 552.	3.8	3
13	Atypical phenotype? The answerâ€™s in the genotype: AGS caused by a novel RNASEH2C variant combined with XLA caused by a BTK deficiency. Rheumatology, 2021, 60, e240-e242.	1.9	1
14	Novel human pathological mutations. Gene symbol: KRIT1. Disease: cerebral cavernous malformation. Human Genetics, 2007, 122, 552.	3.8	1
15	Novel human pathological mutations. Gene symbol: KRIT1. Disease: cerebral cavernous malformation. Human Genetics, 2007, 122, 549.	3.8	0
16	Novel human pathological mutations. Gene symbol: KRIT1. Disease: cerebral cavernous malformation. Human Genetics, 2007, 122, 550.	3.8	0
17	Novel human pathological mutations. Gene symbol: KRIT1. Disease: cerebral cavernous malformation. Human Genetics, 2007, 122, 550.	3.8	0
18	Novel human pathological mutations. Gene symbol: KRIT1. Disease: cerebral cavernous malformation. Human Genetics, 2007, 122, 550.	3.8	0

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19	Novel human pathological mutations. Gene symbol: KRIT1. Disease: cerebral cavernous malformation. Human Genetics, 2007, 122, 551.	3.8	0
20	Novel human pathological mutations. Gene symbol: KRIT1. Disease: cerebral cavernous malformation. Human Genetics, 2007, 122, 551.	3.8	0
21	Novel human pathological mutations. Gene symbol: KRIT1. Disease: cerebral cavernous malformation. Human Genetics, 2007, 122, 551.	3.8	0
22	Novel human pathological mutations. Gene symbol: KRIT1. Disease: cerebral cavernous malformation. Human Genetics, 2007, 122, 552.	3.8	0