Junya Shimizu

List of Publications by Year
in descending order

10 papers	98 citations	1684188 5 h-index	1720034 7 g-index
10 all docs	10 docs citations	10 times ranked	360 citing authors

#	Article	IF	Citations
1	X-Linked Alport Syndrome Caused by Splicing Mutations in COL4A5. Clinical Journal of the American Society of Nephrology: CJASN, 2014, 9, 1958-1964.	4.5	31
2	Progressive brain atrophy in Schinzelâ€“Giedion syndrome with a SETBP1 mutation. European Journal of Medical Genetics, 2015, 58, 369-371.	1.3	22
3	The identification of a novel splicing mutation in C1qB in a Japanese family with C1q deficiency: a case report. Pediatric Rheumatology, 2013, 11, 41.	2.1	15
4	Clinical spectrum of male patients with OFD1 mutations. Journal of Human Genetics, 2019, 64, 3-9.	2.3	12
5	Lupus anticoagulant-hypoprothrombinemia syndrome and immunoglobulin-A vasculitis: a report of Japanese sibling cases and review of the literature. Rheumatology International, 2019, 39, 1811-1819.	3.0	9
6	A 9-year-old girl with Kawasaki disease and pulmonary nodules. Clinical Rheumatology, 2020, 39, 3139-3140.	2.2	5
7	Bone marrow transplantation from a human leukocyte antigen-mismatched unrelated donor in a case with C1q deficiency associated with refractory systemic lupus erythematosus. International Journal of Hematology, 2021, 113, 302-307.	1.6	4
8	Infantile leukocytoclastic vasculitis caused by enterotoxinâ€producing methicillinâ€sensitive Staphylococcus aureus. Pediatric Dermatology, 2021, 38, 1288-1291.	0.9	0
9	A case of acute poststreptococcal glomerulonephritis following surgery for biliary atresia in a patient with biliary cirrhosis. Japanese Journal of Pediatric Nephrology, 2018, 31, 63-68.	0.0	0
10	Acute pancreatitis during longâ€term peritoneal dialysis management associated with the OFDâ€1 mutation. Pediatrics International, 2022, 64, e14925.	0.5	0