Svjetlana Lovric

List of Publications by Year
in descending order

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30 papers	3,416 citations	257101 24 h-index	454577 30 g-index
31 all docs	31 docs citations	31 times ranked	4588 citing authors

#	Article	IF	Citations
1	A Single-Gene Cause in 29.5% of Cases of Steroid-Resistant Nephrotic Syndrome. Journal of the American Society of Nephrology: JASN, 2015, 26, 1279-1289.	3.0	499
2	ADCK4 mutations promote steroid-resistant nephrotic syndrome through CoQ10 biosynthesis disruption. Journal of Clinical Investigation, 2013, 123, 5179-5189.	3.9	275
3	ARHGDIA mutations cause nephrotic syndrome via defective RHO GTPase signaling. Journal of Clinical Investigation, 2013, 123, 3243-3253.	3.9	196
4	ZMYND10 Is Mutated in Primary Ciliary Dyskinesia and Interacts with LRRC6. American Journal of Human Genetics, 2013, 93, 336-345.	2.6	183
5	Safety and clinical outcomes of rituximab therapy in patients with different autoimmune diseases: experience from a national registry (GRAID). Arthritis Research and Therapy, 2011, 13, R75.	1.6	170
6	Whole Exome Sequencing of Patients with Steroid-Resistant Nephrotic Syndrome. Clinical Journal of the American Society of Nephrology: CJASN, 2018, 13, 53-62.	2.2	170
7	Mutations in KEOPS-complex genes cause nephrotic syndrome with primary microcephaly. Nature Genetics, 2017, 49, 1529-1538.	9.4	164
8	Mutations in sphingosine-1-phosphate lyase cause nephrosis with ichthyosis and adrenal insufficiency. Journal of Clinical Investigation, 2017, 127, 912-928.	3.9	160
9	Genetic testing in steroid-resistant nephrotic syndrome: when and how?. Nephrology Dialysis Transplantation, 2016, 31, 1802-1813.	0.4	159
10	KANK deficiency leads to podocyte dysfunction and nephrotic syndrome. Journal of Clinical Investigation, 2015, 125, 2375-2384.	3.9	159
11	Mutations in nuclear pore genes NUP93, NUP205 and XPO5 cause steroid-resistant nephrotic syndrome. Nature Genetics, 2016, 48, 457-465.	9.4	149
12	Detection of circulating microparticles by flow cytometry: influence of centrifugation, filtration of buffer, and freezing. Vascular Health and Risk Management, 2010, 6, 1125.	1.0	123
13	Whole-Exome Sequencing Enables a Precision Medicine Approach for Kidney Transplant Recipients. Journal of the American Society of Nephrology: JASN, 2019, 30, 201-215.	3.0	110
14	Mutations in EMP2 Cause Childhood-Onset Nephrotic Syndrome. American Journal of Human Genetics, 2014, 94, 884-890.	2.6	101
15	FAT1 mutations cause a glomerulotubular nephropathy. Nature Communications, 2016, 7, 10822.	5.8	99
16	Mutations in six nephrosis genes delineate a pathogenic pathway amenable to treatment. Nature Communications, 2018, 9, 1960.	5.8	90
17	Mutations in multiple components of the nuclear pore complex cause nephrotic syndrome. Journal of Clinical Investigation, 2018, 128, 4313-4328.	3.9	89
18	Defects of CRB2 Cause Steroid-Resistant Nephrotic Syndrome. American Journal of Human Genetics, 2015, 96, 153-161.	2.6	88

#	Article	IF	Citations
19	Spectrum of mutations in Chinese children with steroid-resistant nephrotic syndrome. Pediatric Nephrology, 2017, 32, 1181-1192.	0.9	81
20	Rituximab as rescue therapy in anti-neutrophil cytoplasmic antibody-associated vasculitis: a single-centre experience with 15 patients. Nephrology Dialysis Transplantation, 2008, 24, 179-185.	0.4	76
21	Rapid Detection of Monogenic Causes of Childhood-Onset Steroid-Resistant Nephrotic Syndrome. Clinical Journal of the American Society of Nephrology: CJASN, 2014, 9, 1109-1116.	2.2	74
22	Advillin acts upstream of phospholipase C Ïµ1 in steroid-resistant nephrotic syndrome. Journal of Clinical Investigation, 2017, 127, 4257-4269.	3.9	39
23	Circulating ADAM17 Level Reflects Disease Activity in Proteinase-3 ANCA-Associated Vasculitis. Journal of the American Society of Nephrology: JASN, 2015, 26, 2860-2870.	3.0	38
24	Efficacy and Safety of Rituximab Treatment in Patients with Antineutrophil Cytoplasmic Antibody-associated Vasculitides: Results from a German Registry (GRAID). Journal of Rheumatology, 2012, 39, 2153-2156.	1.0	32
25	Analysis of 24 genes reveals a monogenic cause in 11.1% of cases with steroid-resistant nephrotic syndrome at a single center. Pediatric Nephrology, 2018, 33, 305-314.	0.9	30
26	Combination of everolimus with calcineurin inhibitor medication resulted in post-transplant haemolytic uraemic syndrome in lung transplant recipients--a case series. Nephrology Dialysis Transplantation, 2011, 26, 3032-3038.	0.4	18
27	Mutations in PRDM15 Are a Novel Cause of Galloway-Mowat Syndrome. Journal of the American Society of Nephrology: JASN, 2021, 32, 580-596.	3.0	15
28	Panel sequencing distinguishes monogenic forms of nephritis from nephrosis in children. Nephrology Dialysis Transplantation, 2019, 34, 474-485.	0.4	13
29	Removal of elevated circulating angiopoietin-2 by plasma exchange â€“ A pilot study in critically ill patients with thrombotic microangiopathy and anti-glomerular basement membrane disease. Thrombosis and Haemostasis, 2010, 104, 1038-1043.	1.8	11
30	Longâ€term B cell depletion associates with regeneration of kidney function. Immunity, Inflammation and Disease, 2021, 9, 1479-1488.	1.3	5