Remi Salomon

List of Publications by Year
in descending order

12 papers	426 citations	1163117 8 h-index	1281871 11 g-index
12 all docs	12 docs citations	12 times ranked	808 citing authors

#	Article	IF	Citations
1	Role of oculocerebrorenal syndrome of Lowe (OCRL) protein in megakaryocyte maturation, platelet production and functions: a study in patients with Lowe syndrome. British Journal of Haematology, 2021, 192, 909-921.	2.5	6
2	Fetal Cystoscopy and Vesicoamniotic Shunting in Lower Urinary Tract Obstruction: Long-Term Outcome and Current Technical Limitations. Fetal Diagnosis and Therapy, 2020, 47, 74-83.	1.4	33
3	Donor-targeted serotherapy as a rescue therapy for steroid-resistant acute GVHD after HLA-mismatched kidney transplantation. American Journal of Transplantation, 2020, 20, 2243-2253.	4.7	11
4	Electronic health records for the diagnosis of rare diseases. Kidney International, 2020, 97, 676-686.	5.2	37
5	Continuous ambulatory peritoneal dialysis (CAPD) in children: a successful case for a bright future in a developing country. Pan African Medical Journal, 2019, 33, 71.	0.8	2
6	Immunosuppressive Treatment in Children With IgA Nephropathy and the Clinical Value of Podocytopathic Features. Kidney International Reports, 2018, 3, 916-925.	0.8	36
7	A randomised Phase I/II trial to evaluate the efficacy and safety of orally administered Oxalobacter formigenes to treat primary hyperoxaluria. Pediatric Nephrology, 2017, 32, 781-790.	1.7	66
8	Blockade of C5 in Severe Acute Postinfectious Glomerulonephritis Associated With Antiâ€“Factor H Autoantibody. American Journal of Kidney Diseases, 2016, 68, 944-948.	1.9	6
9	FP820AGE DEPENDANT RISK OF GRAFT FAILURE IN YOUNG KIDNEY TRANSPLANT RECIPIENTS. Nephrology Dialysis Transplantation, 2015, 30, iii351-iii351.	0.7	0
10	Mutations of CEP83 Cause Infantile Nephronophthisis and Intellectual Disability. American Journal of Human Genetics, 2014, 94, 905-914.	6.2	90
11	From lowe syndrome to Dent disease: correlations between mutations of the OCRL1 gene and clinical and biochemical phenotypes. Human Mutation, 2011, 32, 379-388.	2.5	120
12	short report: Bleeding disorders in Lowe syndrome patients: evidence for a link between OCRL mutations and primary haemostasis disorders. British Journal of Haematology, 2010, 150, 685-688.	2.5	19