Rodney D. Gilbert

List of Publications by Year
in descending order

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42 papers	1,565 citations	430442 18 h-index	377514 34 g-index
43 all docs	43 docs citations	43 times ranked	3219 citing authors

#	Article	IF	Citations
1	Clonal myelopoiesis promotes adverse outcomes in chronic kidney disease. Leukemia, 2022, 36, 507-515.	3.3	49
2	Daily low-dose prednisolone to prevent relapse of steroid-sensitive nephrotic syndrome in children with an upper respiratory tract infection: PREDNOS2 RCT. Health Technology Assessment, 2022, 26, 1-94.	1.3	4
3	Evaluation of Daily Low-Dose Prednisolone During Upper Respiratory Tract Infection to Prevent Relapse in Children With Relapsing Steroid-Sensitive Nephrotic Syndrome. JAMA Pediatrics, 2022, 176, 236.	3.3	20
4	MO048PATHOGENIC VARIANTS IN CHLORIDE VOLTAGE-GATED CHANNEL 5 (CLCN5), ASSOCIATED WITH DENT DISEASE TYPE 1, SHOULD BE CONSIDERED IN END-STAGE KIDNEY DISEASE OF UNKNOWN AETIOLOGY. Nephrology Dialysis Transplantation, 2021, 36, .	0.4	0
5	Ectopic vortex veins and varices in Donnai Barrow syndrome. Ophthalmic Genetics, 2021, , 1-5.	0.5	1
6	Vitamin B6 in Pediatric Renal Transplant Recipients. , 2019, 29, 205-208.		2
7	Thrombotic microangiopathy following haematopoietic stem cell transplant. Pediatric Nephrology, 2018, 33, 1489-1500.	0.9	29
8	Management of Denys-Drash syndrome: A case series based on an international survey. Clinical Nephrology Case Studies, 2018, 6, 36-44.	0.3	11
9	AMMECR1: a single point mutation causes developmental delay, midface hypoplasia and elliptocytosis. Journal of Medical Genetics, 2017, 54, 269-277.	1.5	12
10	Progressive myoclonic epilepsy with Fanconi syndrome. JRSM Open, 2016, 7, 205427041562314.	0.2	3
11	Basal metabolic rate in children with chronic kidney disease and healthy control children. Pediatric Nephrology, 2015, 30, 1995-2001.	0.9	12
12	Factors influencing success of clinical genome sequencing across a broad spectrum of disorders. Nature Genetics, 2015, 47, 717-726.	9.4	310
13	Eculizumab in atypical haemolytic uraemic syndrome with severe cardiac and neurological involvement. Pediatric Nephrology, 2014, 29, 1103-1106.	0.9	23
14	Short course daily prednisolone therapy during an upper respiratory tract infection in children with relapsing steroid-sensitive nephrotic syndrome (PREDNOS 2): protocol for a randomised controlled trial. Trials, 2014, 15, 147.	0.7	24
15	Children with nephrotic syndrome have greater bone area but similar volumetric bone mineral density to healthy controls. Bone, 2014, 58, 108-113.	1.4	12
16	Bilineal inheritance of PKD1 abnormalities mimicking autosomal recessive polycystic disease. Pediatric Nephrology, 2013, 28, 2217-2220.	0.9	18
17	Eculizumab therapy for atypical haemolytic uraemic syndrome due to a gain-of-function mutation of complement factor B. Pediatric Nephrology, 2013, 28, 1315-1318.	0.9	21
18	Mutations in PIK3R1 Cause SHORT Syndrome. American Journal of Human Genetics, 2013, 93, 158-166.	2.6	156

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19	Does dysregulated complement activation contribute to haemolytic uraemic syndrome secondary to Streptococcus pneumoniae?. Medical Hypotheses, 2013, 81, 400-403.	0.8	38
20	Mosaicism of the UDP-Galactose Transporter SLC35A2 Causes a Congenital Disorder of Glycosylation. American Journal of Human Genetics, 2013, 92, 632-636.	2.6	114
21	Simultaneous Sequencing of 24 Genes Associated with Steroid-Resistant Nephrotic Syndrome. Clinical Journal of the American Society of Nephrology: CJASN, 2013, 8, 637-648.	2.2	152
22	Cisplatin-induced haemolytic uraemic syndrome associated with a novel intronic mutation ofCD46treated with eculizumab. CKJ: Clinical Kidney Journal, 2013, 6, 421-425.	1.4	29
23	Exome analysis resolves differential diagnosis of familial kidney disease and uncovers a potential confounding variant. Genetical Research, 2013, 95, 165-173.	0.3	14
24	Understanding and managing hyponatraemia. Paediatrics and Child Health (United Kingdom), 2010, 20, 261-265.	0.2	3
25	Mutations in phospholipase C epsilon 1 are not sufficient to cause diffuse mesangial sclerosis. Kidney International, 2009, 75, 415-419.	2.6	35
26	Professionals Against Child Abuse express support for David Southall. Lancet, The, 2009, 373, 2021.	6.3	0
27	The glomerulonephritides. Paediatrics and Child Health (United Kingdom), 2008, 18, 354-357.	0.2	0
28	Hemolytic Uremic Syndrome Associated with Invasive Pneumococcal Disease: The United Kingdom Experience. Journal of Pediatrics, 2007, 151, 140-144.	0.9	153
29	Acute renal failure in a patient with paroxysmal cold hemoglobinuria. Pediatric Nephrology, 2007, 22, 593-596.	0.9	14
30	Patient with an EYA1 mutation with features of branchio-oto-renal and oto-facio-cervical syndrome. Clinical Dysmorphology, 2006, 15, 211-212.	0.1	9
31	Rituximab therapy for steroid-dependent minimal change nephrotic syndrome. Pediatric Nephrology, 2006, 21, 1698-1700.	0.9	102
32	Laparoscopic renal biopsy in obese children. Pediatric Nephrology, 2005, 20, 495-498.	0.9	15
33	Renal function, renal failure and renal transplantation. , 2005, , 441-451.		0
34	Urinary L-lactate excretion is increased in renal Fanconi syndrome. Nephrology Dialysis Transplantation, 2004, 19, 1767-1773.	0.4	15
35	Bilateral multicystic kidneys ? an unusual case. Pediatric Nephrology, 2002, 17, 964-965.	0.9	3
36	STREPTOCOCCUS PNEUMONIAE-ASSOCIATED HEMOLYTIC UREMIC SYNDROME. Pediatric Infectious Disease Journal, 1998, 17, 530-532.	1.1	20

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37	Pearson's Syndrome Presenting with Fanconi Syndrome. Ultrastructural Pathology, 1996, 20, 473-475.	0.4	22
38	Interaction between Clonidine and Cyclosporine A. Nephron, 1995, 71, 105-105.	0.9	7
39	The glomerulonephritides in children: new thoughts on aetiology and treatment. Current Paediatrics, 1995, 5, 75-79.	0.2	0
40	The clinical course of hepatitis B virus-associated nephropathy. Pediatric Nephrology, 1994, 8, 11-14.	0.9	88
41	Acute promyelocytic leukemia. A childhood cluster. Cancer, 1987, 59, 933-935.	2.0	25
42	Hemizygous loss of function mutations in CLCN5 causing end-stage kidney disease without Dent disease phenotype. CKJ: Clinical Kidney Journal, 0, , .	1.4	0