

Georges Deschenes

List of Publications by Year in descending order

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Version: 2024-02-01

57
papers

1,582
citations

331670

21
h-index

330143

37
g-index

57
all docs

57
docs citations

57
times ranked

1731
citing authors

#	ARTICLE	IF	CITATIONS
1	Anti-rituximab antibodies in pediatric steroid-dependent nephrotic syndrome. <i>Pediatric Nephrology</i> , 2022, 37, 357-365.	1.7	11
2	Long-Term Transplantation Outcomes in Patients With Primary Hyperoxaluria Type 1 Included in the European Hyperoxaluria Consortium (OxalEurope) Registry. <i>Kidney International Reports</i> , 2022, 7, 210-220.	0.8	19
3	FC038: Efficacy of Levamisole for Maintaining Remission after the First Flare of Steroid Sensitive Nephrotic Syndrome in Children: The Nephrovir-3 Randomized Controlled Trial. <i>Nephrology Dialysis Transplantation</i> , 2022, 37, .	0.7	0
4	Telomere aberrations, including telomere loss, doublets, and extreme shortening, are increased in patients with infertility. <i>Fertility and Sterility</i> , 2021, 115, 164-173.	1.0	14
5	A global antiB cell strategy combining obinutuzumab and daratumumab in severe pediatric nephrotic syndrome. <i>Pediatric Nephrology</i> , 2021, 36, 1175-1182.	1.7	21
6	Adherence to cysteamine in nephropathic cystinosis: A unique electronic monitoring experience for a better understanding. A prospective cohort study: CrYSTobs. <i>Pediatric Nephrology</i> , 2021, 36, 581-589.	1.7	7
7	An international cohort study spanning five decades assessed outcomes of nephropathic cystinosis. <i>Kidney International</i> , 2021, 100, 1112-1123.	5.2	31
8	SOLUBLE CD89 IS A CRITICAL FACTOR FOR MESANGIAL PROLIFERATION IN CHILDHOOD IgA NEPHROPATHY. <i>Kidney International</i> , 2021, , .	5.2	8
9	Steroid therapy in children with IgA nephropathy. <i>Pediatric Nephrology</i> , 2020, 35, 359-366.	1.7	19
10	Histological prognostic factors in children with Henoch-Schönlein purpura nephritis. <i>Pediatric Nephrology</i> , 2020, 35, 313-320.	1.7	21
11	Efficacy and safety of intravenous immunoglobulin with rituximab versus rituximab alone in childhood-onset steroid-dependent and frequently relapsing nephrotic syndrome: protocol for a multicentre randomised controlled trial. <i>BMJ Open</i> , 2020, 10, e037306.	1.9	0
12	Telomere and Centromere Staining Followed by M-FISH Improves Diagnosis of Chromosomal Instability and Its Clinical Utility. <i>Genes</i> , 2020, 11, 475.	2.4	17
13	Sodiumâ€”not harmful?. <i>Pediatric Nephrology</i> , 2020, 35, 1771-1776.	1.7	2
14	Treating the idiopathic nephrotic syndrome: are steroids the answer?. <i>Pediatric Nephrology</i> , 2019, 34, 777-785.	1.7	6
15	Effect of different rituximab regimens on B cell depletion and time to relapse in children with steroid-dependent nephrotic syndrome. <i>Pediatric Nephrology</i> , 2019, 34, 253-259.	1.7	39
16	Hyponatremia in children under 100Âdays old: incidence and etiologies. <i>European Journal of Pediatrics</i> , 2019, 178, 1353-1361.	2.7	16
17	Quality of life in children with severe forms of idiopathic nephrotic syndrome in stable remissionâ€”A crossâ€”sectional study. <i>Acta Paediatrica, International Journal of Paediatrics</i> , 2019, 108, 2267-2273.	1.5	18
18	Social deprivation is associated with poor kidney transplantation outcome in children. <i>Kidney International</i> , 2019, 96, 769-776.	5.2	25

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19	The Urinary Excretion of Uromodulin is Regulated by the Potassium Channel ROMK. <i>Scientific Reports</i> , 2019, 9, 19517.	3.3	21
20	Remission of proteinuria in multidrug-resistant idiopathic nephrotic syndrome following immunoglobulin immunoadsorption. <i>Acta Paediatrica, International Journal of Paediatrics</i> , 2019, 108, 757-762.	1.5	3
21	Five-year outcome of children with idiopathic nephrotic syndrome: the NEPHROVIR population-based cohort study. <i>Pediatric Nephrology</i> , 2019, 34, 671-678.	1.7	25
22	How to improve response to rituximab treatment in children with steroid-dependent nephrotic syndrome: answer to Drs. Fujinaga and Nishino. <i>Pediatric Nephrology</i> , 2019, 34, 361-362.	1.7	3
23	Effect of nonsteroidal anti-inflammatory drugs in children with Bartter syndrome. <i>Pediatric Nephrology</i> , 2019, 34, 679-684.	1.7	17
24	Stiripentol protects against calcium oxalate nephrolithiasis and ethylene glycol poisoning. <i>Journal of Clinical Investigation</i> , 2019, 129, 2571-2577.	8.2	47
25	Immunosuppressive Treatment in Children With IgA Nephropathy and the Clinical Value of Podocytopathic Features. <i>Kidney International Reports</i> , 2018, 3, 916-925.	0.8	36
26	Severe neonatal hypertension revealing arterial tortuosity syndrome. <i>Kidney International</i> , 2018, 93, 526.	5.2	4
27	Autoantibodies against podocytic UCHL1 are associated with idiopathic nephrotic syndrome relapses and induce proteinuria in mice. <i>Journal of Autoimmunity</i> , 2018, 89, 149-161.	6.5	48
28	Clinical and genetic heterogeneity in familial steroid-sensitive nephrotic syndrome. <i>Pediatric Nephrology</i> , 2018, 33, 473-483.	1.7	34
29	Nephrotic-range proteinuria and brown urine in an 8-year-old girl: Questions. <i>Pediatric Nephrology</i> , 2018, 33, 1001-1002.	1.7	0
30	Nephrotic-range proteinuria and brown urine in an 8-year-old girl: Answers. <i>Pediatric Nephrology</i> , 2018, 33, 1003-1005.	1.7	0
31	Combination therapy of rituximab and mycophenolate mofetil in childhood lupus nephritis. <i>Pediatric Nephrology</i> , 2018, 33, 111-116.	1.7	17
32	Urine biochemistry to predict long-term outcomes in fetuses with posterior urethral valves. <i>Prenatal Diagnosis</i> , 2018, 38, 964-970.	2.3	22
33	Interdialytic weight gain and vasculopathy in children on hemodialysis: a single center study. <i>Pediatric Nephrology</i> , 2018, 33, 2329-2336.	1.7	10
34	Early cardiovascular manifestations in children and adolescents with autosomal dominant polycystic kidney disease: a single center study. <i>Pediatric Nephrology</i> , 2018, 33, 1513-1521.	1.7	15
35	Anti-Factor B and Anti-C3b Autoantibodies in C3 Glomerulopathy and Ig-Associated Membranoproliferative GN. <i>Journal of the American Society of Nephrology: JASN</i> , 2017, 28, 1603-1613.	6.1	83
36	Clinical outcomes in children with Henoch-Schönlein purpura nephritis without crescents. <i>Pediatric Nephrology</i> , 2017, 32, 1193-1199.	1.7	40

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37	Clinical and Genetic Spectrum of Bartter Syndrome Type 3. <i>Journal of the American Society of Nephrology: JASN</i> , 2017, 28, 2540-2552.	6.1	92
38	Variability of diagnostic criteria and treatment of idiopathic nephrotic syndrome across European countries. <i>European Journal of Pediatrics</i> , 2017, 176, 647-654.	2.7	18
39	C5 nephritic factors drive the biological phenotype of C3 glomerulopathies. <i>Kidney International</i> , 2017, 92, 1232-1241.	5.2	93
40	Transmission of Induced Chromosomal Aberrations through Successive Mitotic Divisions in Human Lymphocytes after In Vitro and <i>in Vivo</i> Radiation. <i>Scientific Reports</i> , 2017, 7, 3291.	3.3	27
41	Idiopathic nephrotic syndrome: the EBV hypothesis. <i>Pediatric Research</i> , 2017, 81, 233-239.	2.3	31
42	Fluid status evaluation by inferior vena cava diameter and bioimpedance spectroscopy in pediatric chronic hemodialysis. <i>BMC Nephrology</i> , 2017, 18, 373.	1.8	17
43	Long-term successful liver-kidney transplantation in a child with atypical hemolytic uremic syndrome caused by homozygous factor H deficiency. <i>Pediatric Nephrology</i> , 2016, 31, 2375-2378.	1.7	3
44	Mycophenolate mofetil in steroid-dependent idiopathic nephrotic syndrome. <i>Pediatric Nephrology</i> , 2016, 31, 2095-2101.	1.7	18
45	Observations of a large Dent disease cohort. <i>Kidney International</i> , 2016, 90, 430-439.	5.2	71
46	Hemolytic anemia and irreversible kidney and brain injuries after accidental intravenous injection of albendazole suspension in an infant. <i>Clinical Toxicology</i> , 2016, 54, 72-73.	1.9	4
47	Even mild cases of paediatric <i>Henoch-Schönlein purpura nephritis</i> show significant long-term proteinuria. <i>Acta Paediatrica, International Journal of Paediatrics</i> , 2015, 104, 843-848.	1.5	21
48	Cytomegalovirus infection can mimic genetic nephrotic syndrome: a case report. <i>BMC Nephrology</i> , 2015, 16, 156.	1.8	10
49	A new gel formulation of topical cysteamine for the treatment of corneal cystine crystals in cystinosis: The Cystadrops OCT-1 study. <i>Molecular Genetics and Metabolism</i> , 2014, 111, 314-320.	1.1	53
50	Prevalence of herpesviruses at onset of idiopathic nephrotic syndrome. <i>Pediatric Nephrology</i> , 2014, 29, 2325-2331.	1.7	37
51	Fulminant viral myocarditis after rituximab therapy in pediatric nephrotic syndrome. <i>Pediatric Nephrology</i> , 2013, 28, 1875-1879.	1.7	70
52	Rituximab in steroid-dependent idiopathic nephrotic syndrome in childhood--follow-up after CD19 recovery. <i>Nephrology Dialysis Transplantation</i> , 2012, 27, 1083-1089.	0.7	115
53	Mycophenolate mofetil for steroid-dependent nephrotic syndrome: a phase II Bayesian trial. <i>Pediatric Nephrology</i> , 2012, 27, 389-396.	1.7	48
54	Cyclophosphamide in steroid-dependent nephrotic syndrome. <i>Pediatric Nephrology</i> , 2011, 26, 927-932.	1.7	36

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55	Population pharmacokinetics and pharmacodynamics of cysteamine in nephropathic cystinosis patients. <i>Orphanet Journal of Rare Diseases</i> , 2011, 6, 86.	2.7	18
56	Rituximab efficiency in children with steroid-dependent nephrotic syndrome. <i>Pediatric Nephrology</i> , 2010, 25, 1109-1115.	1.7	84
57	Growth in boys with idiopathic nephrotic syndrome on long-term cyclosporin and steroid treatment. <i>Pediatric Nephrology</i> , 2009, 24, 2393-2400.	1.7	17