Stephane Decramer

List of Publications by Year
in descending order

96	5,215	81900 39	91884 69
papers	citations	h-index	g-index

114	114	114	5710
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Preemptive Kidney Transplantation Is Associated With Transplantation Outcomes in Children: Results From the French Kidney Replacement Therapy Registry. Transplantation, 2022, 106, 401-411.	1.0	9
2	Gitelman-Like Syndrome Caused by Pathogenic Variants in mtDNA. Journal of the American Society of Nephrology: JASN, 2022, 33, 305-325.	6.1	26
3	Definition, diagnosis and management of fetal lower urinary tract obstruction: consensus of the ERKNet CAKUT-Obstructive Uropathy Work Group. Nature Reviews Urology, 2022, 19, 295-303.	3.8	27
4	Amniotic fluid peptides predict postnatal kidney survival in developmental kidney disease. Kidney International, 2021, 99, 737-749.	5.2	15
5	Mycophenolic acid area under the concentration-time curve is associated with therapeutic response in childhood-onset lupus nephritis. Pediatric Nephrology, 2021, 36, 341-347.	1.7	21
6	Association of kidney biopsy findings with short- and medium-term outcomes in children with moderate-to-severe IgA vasculitis nephritis. European Journal of Pediatrics, 2021, 180, 3209-3218.	2.7	7
7	Mapping of the amniotic fluid proteome of fetuses with congenital anomalies of the kidney and urinary tract identifies plastin 3 as a protein involved in glomerular integrity. Journal of Pathology, 2021, 254, 575-588.	4.5	4
8	The European Rare Kidney Disease Registry (ERKReg): objectives, design and initial results. Orphanet Journal of Rare Diseases, 2021, 16, 251.	2.7	26
9	To biopsy or not to biopsy: Henoch-SchÃ¶nlein nephritis in children, a 5-year follow-up study. Pediatric Nephrology, 2021, , 1.	1.7	4
10	Hepatocyte nuclear factorâ€1Î² shapes the energetic homeostasis of kidney tubule cells. FASEB Journal, 2021, 35, e21931.	0.5	8
11	School level of children carrying a HNF1B variant or a deletion. European Journal of Human Genetics, 2020, 28, 56-63.	2.8	9
12	A single-center study to evaluate the efficacy of a fetal urine peptide signature predicting postnatal renal outcome in fetuses with posterior urethral valves. Pediatric Nephrology, 2020, 35, 469-475.	1.7	17
13	The ANTENATAL multicentre study to predict postnatal renal outcome in fetuses with posterior urethral valves: objectives and design. CKJ: Clinical Kidney Journal, 2020, 13, 371-379.	2.9	18
14	The low affinity p75 neurotrophin receptor is down-regulated in congenital anomalies of the kidney and the urinary tract: Possible involvement in early nephrogenesis. Biochemical and Biophysical Research Communications, 2020, 533, 786-791.	2.1	3
15	Comparison of the amniotic fluid and fetal urine peptidome for biomarker discovery in renal developmental disease. Scientific Reports, 2020, 10, 21706.	3.3	8
16	Quality of life in children with severe forms of idiopathic nephrotic syndrome in stable remissionâ€”A crossâ€sectional study. Acta Paediatrica, International Journal of Paediatrics, 2019, 108, 2267-2273.	1.5	18
17	Ageâ€specific characteristics of neutrophilic dermatoses and neutrophilic diseases in children. Journal of the European Academy of Dermatology and Venereology, 2019, 33, 2179-2187.	2.4	9
18	Social deprivation is associated with poor kidney transplantation outcome in children. Kidney International, 2019, 96, 769-776.	5.2	25

#	Article	IF	Citations
19	Consensus recommendations for diagnosis, management and treatment of Fabry disease in paediatric patients. Clinical Genetics, 2019, 96, 107-117.	2.0	87
20	Adverse events associated with currently used medical treatments for cystinuria and treatment goals: results from a series of 442 patients in France. BJU International, 2019, 124, 849-861.	2.5	30
21	Treatment and long-term outcome in primary distal renal tubular acidosis. Nephrology Dialysis Transplantation, 2019, 34, 981-991.	0.7	75
22	Urinary proteome signature of Renal Cysts and Diabetes syndrome in children. Scientific Reports, 2019, 9, 2225.	3.3	15
23	Patterns of Clinical Response to Eculizumab in Patients With C3 Glomerulopathy. American Journal of Kidney Diseases, 2018, 72, 84-92.	1.9	94
24	Prevalence of Novel MAGED2 Mutations in Antenatal Bartter Syndrome. Clinical Journal of the American Society of Nephrology: CJASN, 2018, 13, 242-250.	4.5	45
25	Combination of the fetal urinary metabolome and peptidome for the prediction of postnatal renal outcome in fetuses with PUV. Journal of Proteomics, 2018, 184, 1-9.	2.4	16
26	Clinical outcomes in children with Henochâ€“SchÃ¶nlein purpura nephritis without crescents. Pediatric Nephrology, 2017, 32, 1193-1199.	1.7	40
27	Systems biology combining human- and animal-data miRNA and mRNA data identifies new targets in ureteropelvic junction obstruction. BMC Systems Biology, 2017, 11, 31.	3.0	12
28	Clinical and Genetic Spectrum of Bartter Syndrome Type 3. Journal of the American Society of Nephrology: JASN, 2017, 28, 2540-2552.	6.1	92
29	Cryptosporidium spp. Infection in Solid Organ Transplantation. Transplantation, 2017, 101, 826-830.	1.0	44
30	Familial and syndromic lupus share the same phenotype as other early-onset forms of lupus. Joint Bone Spine, 2017, 84, 589-593.	1.6	7
31	Age-Dependent Risk of Graft Failure in Young Kidney Transplant Recipients. Transplantation, 2017, 101, 1327-1335.	1.0	43
32	Hepatocyte Nuclear Factor-1Î² Controls Mitochondrial Respiration in Renal Tubular Cells. Journal of the American Society of Nephrology: JASN, 2017, 28, 3205-3217.	6.1	43
33	Atypical hemolytic uremic syndrome triggered by varicella infection. IDCases, 2017, 9, 89-90.	0.9	6
34	Alternative complement pathway hemolytic assays reveal incomplete complement blockade in patients treated with eculizumab. Clinical Immunology, 2017, 183, 1-7.	3.2	16
35	Calcineurin Inhibitors Downregulate HNF-1Î² and May Affect the Outcome of HNF1B Patients After Renal Transplantation. Transplantation, 2016, 100, 1970-1978.	1.0	6
36	A capillary electrophoresis coupled to mass spectrometry pipeline for long term comparable assessment of the urinary metabolome. Scientific Reports, 2016, 6, 34453.	3.3	28

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37	Mycophenolic Acid Pharmacokinetics and Relapse in Children with Steroidâ€“Dependent Idiopathic Nephrotic Syndrome. Clinical Journal of the American Society of Nephrology: CJASN, 2016, 11, 1777-1782.	4.5	35
38	Hyperechogenic kidneys and polyhydramnios associated with HNF1B gene mutation. Pediatric Nephrology, 2016, 31, 1705-1708.	1.7	16
39	Initial presentation and outcome of pediatric-onset mixed connective tissue disease: A French multicenter retrospective study. Joint Bone Spine, 2016, 83, 369-371.	1.6	17
40	Comprehensive PKD1 and PKD2 Mutation Analysis in Prenatal Autosomal Dominant Polycystic Kidney Disease. Journal of the American Society of Nephrology: JASN, 2016, 27, 722-729.	6.1	68
41	Even mild cases of paediatric <scp>H</scp>enochâ€<scp>S</scp>chÃ¶nlein purpura nephritis show significant longâ€term proteinuria. Acta Paediatrica, International Journal of Paediatrics, 2015, 104, 843-848.	1.5	21
42	CKD and Its Risk Factors among Patients with Cystinuria. Clinical Journal of the American Society of Nephrology: CJASN, 2015, 10, 842-851.	4.5	71
43	Clinical characteristics and outcomes of childhood-onset ANCA-associated vasculitis: a French nationwide study. Nephrology Dialysis Transplantation, 2015, 30 Suppl 1, i104-12.	0.7	45
44	The Authors Reply. Kidney International, 2015, 87, 1259.	5.2	2
45	Clinical proteomics in obstetrics and neonatology. Expert Review of Proteomics, 2014, 11, 75-89.	3.0	31
46	Pharmacokinetics of mycophenolate mofetil in children with lupus and clinical findings in favour of therapeutic drug monitoring. British Journal of Clinical Pharmacology, 2014, 78, 867-876.	2.4	42
47	Label-free Quantitative Urinary Proteomics Identifies the Arginase Pathway as a New Player in Congenital Obstructive Nephropathy. Molecular and Cellular Proteomics, 2014, 13, 3421-3434.	3.8	24
48	SFP P-054 â€“ Penser HÃ©mochromatose nÃ©onatale devant un tableau de SHU du nouveau-nÃ©. Archives De Pediatrie, 2014, 21, 764.	1.0	0
49	The HNF1B score is a simple tool to select patients for HNF1B gene analysis. Kidney International, 2014, 86, 1007-1015.	5.2	104
50	Fetal Urinary Peptides to Predict Postnatal Outcome of Renal Disease in Fetuses with Posterior Urethral Valves (PUV). Science Translational Medicine, 2013, 5, 198ra106.	12.4	86
51	Protocol biopsies in pediatric renal transplant recipients on cyclosporine versus tacrolimus-based immunosuppression. Pediatric Nephrology, 2013, 28, 493-498.	1.7	8
52	Long-term outcome of children treated with rituximab for idiopathic nephrotic syndrome. Pediatric Nephrology, 2013, 28, 911-918.	1.7	42
53	Diagnosis of Streptococcus pneumoniaeâ€“associated Hemolytic Uremic Syndrome. Pediatric Infectious Disease Journal, 2013, 32, 1045-1049.	2.0	27
54	Should SIX2 be routinely tested in patients with isolated congenital abnormalities of kidneys and/or urinary tract (CAKUT)?. European Journal of Medical Genetics, 2012, 55, 688-689.	1.3	7

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55	Urinary Proteome Analysis at 5-Year Followup of Patients With Nonoperated Ureteropelvic Junction Obstruction Suggests Ongoing Kidney Remodeling. Journal of Urology, 2012, 187, 1006-1011.	0.4	31
56	Expression of Renal Cystic Genes in Patients with <i>HNF1B </i>Mutations. Nephron Clinical Practice, 2012, 120, c71-c78.	2.3	8
57	A rare cause of anaemia associated with hypertension in a 14-year-old girl. Pediatric Radiology, 2012, 42, 624-626.	2.0	3
58	Update of PAX2 mutations in renal coloboma syndrome and establishment of a locus-specific database. Human Mutation, 2012, 33, 457-466.	2.5	109
59	Severe neonatal hypercalcemia related to maternal exposure to nutritional supplement containing Spirulina. European Journal of Clinical Pharmacology, 2012, 68, 221-222.	1.9	7
60	Mycophenolate mofetil for steroid-dependent nephrotic syndrome: a phase II Bayesian trial. Pediatric Nephrology, 2012, 27, 389-396.	1.7	48
61	Diagnosis, management, and prognosis of HNF1B nephropathy in adulthood. Kidney International, 2011, 80, 768-776.	5.2	154
62	Development of a Bayesian estimator for the therapeutic drug monitoring of mycophenolate mofetil in children with idiopathic nephrotic syndrome. Pharmacological Research, 2011, 63, 423-431.	7.1	44
63	A 17q12 chromosomal duplication associated with renal disease and esophageal atresia. European Journal of Medical Genetics, 2011, 54, e437-e440.	1.3	28
64	Congenital ureteropelvic junction obstruction: human disease and animal models. International Journal of Experimental Pathology, 2011, 92, 168-192.	1.3	81
65	Long-term effects of cyclophosphamide therapy in steroid-dependent or frequently relapsing idiopathic nephrotic syndrome. Nephrology Dialysis Transplantation, 2011, 26, 178-184.	0.7	37
66	Varicella-zoster virus meningoencephalitis without skin lesions in a paediatric kidney recipient. Nephrology Dialysis Transplantation, 2011, 26, 378-379.	0.7	2
67	Advances in urinary proteome analysis and biomarker discovery in pediatric renal disease. Pediatric Nephrology, 2010, 25, 27-35.	1.7	66
68	Cryptosporidiosis in pediatric renal transplantation. Pediatric Nephrology, 2010, 25, 571-572.	1.7	1
69	Urinary proteome analysis identifies infants but not older children requiring pyeloplasty. Pediatric Nephrology, 2010, 25, 1673-1678.	1.7	58
70	Blockade of the Kinin B1 Receptor Ameloriates Glomerulonephritis. Journal of the American Society of Nephrology: JASN, 2010, 21, 1157-1164.	6.1	47
71	Clinical features and management of arterial hypertension in children with Williams-Beuren syndrome. Nephrology Dialysis Transplantation, 2010, 25, 434-438.	0.7	41
72	Spectrum of HNF1B Mutations in a Large Cohort of Patients Who Harbor Renal Diseases. Clinical Journal of the American Society of Nephrology: CJASN, 2010, 5, 1079-1090.	4.5	236

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73	Naturally Occurring Human Urinary Peptides for Use in Diagnosis of Chronic Kidney Disease. Molecular and Cellular Proteomics, 2010, 9, 2424-2437.	3.8	434
74	Phenotypeâ€“genotype correlation in antenatal and neonatal variants ofÂBartter syndrome. Nephrology Dialysis Transplantation, 2009, 24, 1455-1464.	0.7	137
75	Mutations in the RARE and MARE regulatory sequences of HNF1Â are not a frequent cause of kidney/urinary tract malformation. CKJ: Clinical Kidney Journal, 2009, 2, 333-334.	2.9	0
76	Cryptosporidiosis in paediatric renal transplantation. Pediatric Nephrology, 2009, 24, 2245-2255.	1.7	43
77	The human urinary proteome reveals high similarity between kidney aging and chronic kidney disease. Proteomics, 2009, 9, 2108-2117.	2.2	82
78	Steroid-responsive nephrotic syndrome in a child with juvenile idiopathic arthritis. Pediatric Nephrology, 2008, 23, 651-654.	1.7	7
79	CEâ€MS analysis of the human urinary proteome for biomarker discovery and disease diagnostics. Proteomics - Clinical Applications, 2008, 2, 964-973.	1.6	178
80	Urine in Clinical Proteomics. Molecular and Cellular Proteomics, 2008, 7, 1850-1862.	3.8	368
81	Nephrin Mutations Can Cause Childhood-Onset Steroid-Resistant Nephrotic Syndrome. Journal of the American Society of Nephrology: JASN, 2008, 19, 1871-1878.	6.1	119
82	Prospective, Randomized Trial Comparing Short and Long Intravenous Antibiotic Treatment of Acute Pyelonephritis in Children: Dimercaptosuccinic Acid Scintigraphic Evaluation at 9 Months. Pediatrics, 2008, 121, e553-e560.	2.1	46
83	Identification of Urinary Biomarkers by Proteomics in Newborns: Use in Obstructive Nephropathy. , 2008, 160, 127-141.		36
84	Anomalies of the TCF2 Gene Are the Main Cause of Fetal Bilateral Hyperechogenic Kidneys. Journal of the American Society of Nephrology: JASN, 2007, 18, 923-933.	6.1	216
85	Thrombotic Microangiopathy in a Child With Acute Pancreatitis. Journal of Pediatric Gastroenterology and Nutrition, 2007, 44, 149-151.	1.8	0
86	A database of naturally occurring human urinary peptides and proteins for use in clinical applications. Nature Precedings, 2007, , .	0.1	0
87	Non-invasive markers of ureteropelvic junction obstruction. World Journal of Urology, 2007, 25, 457-465.	2.2	49
88	Renal Phenotypes Related to Hepatocyte Nuclear Factor-1Î² (TCF2) Mutations in a Pediatric Cohort. Journal of the American Society of Nephrology: JASN, 2006, 17, 497-503.	6.1	238
89	Predicting the clinical outcome of congenital unilateral ureteropelvic junction obstruction in newborn by urinary proteome analysis. Nature Medicine, 2006, 12, 398-400.	30.7	248
90	Urinary aquaporin-2 excretion during early human development. Pediatric Nephrology, 2006, 21, 947-952.	1.7	19

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91	GFR is better estimated by considering both serum cystatin C and creatinine levels. Pediatric Nephrology, 2006, 21, 1299-1306.	1.7	114
92	Long-Term Effects of In Utero Exposure to Cyclosporin A on Renal Function in the Rabbit. Journal of the American Society of Nephrology: JASN, 2004, 15, 2687-2693.	6.1	52
93	Renal outcome of children exposed to cyclosporine in utero. Transplantation Proceedings, 2004, 36, S208-S210.	0.6	47
94	49 Urinary Excretion of Aquaporin 2 In term and Preterm Infants. Pediatric Research, 2004, 56, 472-472.	2.3	0
95	Cyclosporin A Administration during Pregnancy Induces a Permanent Nephron Deficit in Young Rabbits. Journal of the American Society of Nephrology: JASN, 2003, 14, 3188-3196.	6.1	44
96	In utero exposure to immunosuppressive drugs: experimental and clinical studies. Pediatric Nephrology, 2002, 17, 121-130.	1.7	94