

Justine Bacchetta

List of Publications by Year in descending order

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Version: 2024-02-01

161
papers

5,049
citations

126907

33
h-index

118850

62
g-index

192
all docs

192
docs citations

192
times ranked

5760
citing authors

#	ARTICLE	IF	CITATIONS
1	Genetic assessment in primary hyperoxaluria: why it matters. <i>Pediatric Nephrology</i> , 2023, 38, 625-634.	1.7	14
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	Tubular phosphate handling: references from child to adulthood in the era of standardized serum creatinine. <i>Nephrology Dialysis Transplantation</i> , 2022, 37, 2150-2156.	0.7	18
4	Primary hyperoxaluria type 1: novel therapies at a glance. <i>CKJ: Clinical Kidney Journal</i> , 2022, 15, i17-i22.	2.9	10
5	Muscle and Bone Impairment in Infantile Nephropathic Cystinosis: New Concepts. <i>Cells</i> , 2022, 11, 170.	4.1	2
6	Primary hyperoxaluria type 1: time for prime time?. <i>CKJ: Clinical Kidney Journal</i> , 2022, 15, i1-i3.	2.9	4
7	Parathyroid hormone and phosphate homeostasis in patients with Bartter and Gitelman syndrome: an international cross-sectional study. <i>Nephrology Dialysis Transplantation</i> , 2022, 37, 2474-2486.	0.7	5
8	Nephrocalcinosis in very low birth weight infants: incidence, associated factors, and natural course. <i>Pediatric Nephrology</i> , 2022, 37, 3093-3104.	1.7	3
9	Copper Isotope Evidence of Oxidative Stress-Induced Hepatic Breakdown and the Transition to Hepatocellular Carcinoma. , 2022, 1, 480-486.		5
10	Idiopathic nephrotic syndrome relapse following COVID-19 vaccination: a series of 25 cases. <i>CKJ: Clinical Kidney Journal</i> , 2022, 15, 1574-1582.	2.9	7
11	Naturally occurring stable calcium isotope ratios are a novel biomarker of bone calcium balance in chronic kidney disease. <i>Kidney International</i> , 2022, 102, 613-623.	5.2	12
12	Inactivation of Osteoblast PKC Signaling Reduces Cortical Bone Mass and Density and Aggravates Renal Osteodystrophy in Mice with Chronic Kidney Disease on High Phosphate Diet. <i>International Journal of Molecular Sciences</i> , 2022, 23, 6404.	4.1	4
13	Fluconazole in hypercalciuric patients with increased 1,25(OH)2D levels: the prospective, randomized, placebo-controlled, double-blind FLUCOLITH trial. <i>Trials</i> , 2022, 23, .	1.6	1
14	Reloxaliase in Enteric Hyperoxaluria – The Recent Brake. , 2022, 1, .		1
15	Review: Neonatal dialysis is technically feasible but ethical and global issues need to be addressed. <i>Acta Paediatrica, International Journal of Paediatrics</i> , 2021, 110, 781-788.	1.5	10
16	Bone evaluation in paediatric chronic kidney disease: clinical practice points from the European Society for Paediatric Nephrology CKD-MBD and Dialysis working groups and CKD-MBD working group of the ERA-EDTA. <i>Nephrology Dialysis Transplantation</i> , 2021, 36, 413-425.	0.7	30
17	Active vitamin D is cardioprotective in experimental uraemia but not in children with CKD Stages 3-5. <i>Nephrology Dialysis Transplantation</i> , 2021, 36, 442-451.	0.7	5
18	Big data and outcomes in paediatric haemodialysis: how can nephrologists use these new tools in daily practice?. <i>Nephrology Dialysis Transplantation</i> , 2021, 36, 387-391.	0.7	1

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19	Rare diseases of phosphate and calcium metabolism: Crossing glances between nephrology and endocrinology. <i>Annales D'Endocrinologie</i> , 2021, 82, 30-35.	1.4	3
20	European Consensus Statement on the diagnosis and management of osteoporosis in chronic kidney disease stages G4â€“G5D. <i>Nephrology Dialysis Transplantation</i> , 2021, 36, 42-59.	0.7	107
21	The Management of CKD-MBD in Pediatric Dialysis Patients. , 2021, , 541-558.		0
22	Chronic Kidney Disease â€“ Mineral and Bone Disorder (CKD-MBD). , 2021, , 1-29.		0
23	Combined use of creatinine and cystatin C improves the detection of renal dysfunction in children undergoing home parenteral nutrition. <i>Journal of Parenteral and Enteral Nutrition</i> , 2021, , .	2.6	1
24	A prospective caseâ€“control pilot study to evaluate bone microarchitecture in children and teenagers on long-term parenteral nutrition using HR-pQCT. <i>Scientific Reports</i> , 2021, 11, 9151.	3.3	3
25	Tyrosinemia type 1 in pediatric nephrology: Not always straightforward. <i>Archives De Pediatrie</i> , 2021, 28, 338-341.	1.0	2
26	The European Rare Kidney Disease Registry (ERKReg): objectives, design and initial results. <i>Orphanet Journal of Rare Diseases</i> , 2021, 16, 251.	2.7	26
27	Calcium isotope fractionation by osteoblasts and osteoclasts, across endothelial and epithelial cell barriers, and with binding to proteins. <i>American Journal of Physiology - Regulatory Integrative and Comparative Physiology</i> , 2021, 321, R29-R40.	1.8	5
28	Hemodiafiltration Is Associated With Reduced Inflammation and Increased Bone Formation Compared With Conventional Hemodialysis in Children: The HDF, Hearts and Heights (3H) Study. <i>Kidney International Reports</i> , 2021, 6, 2358-2370.	0.8	11
29	A third of premature neonates displayed inadequate 25â€“hydroxyvitamin D levels before being discharged from a French neonatal intensive care unit. <i>Acta Paediatrica, International Journal of Paediatrics</i> , 2021, , .	1.5	3
30	Local protocol helped to deliver vitamin D levels more accurately in preterm infants. <i>Acta Paediatrica, International Journal of Paediatrics</i> , 2021, , .	1.5	4
31	Response to Cysteamine in Osteoclasts Obtained from Patients with Nephropathic Cystinosis: A Genotype/Phenotype Correlation. <i>Cells</i> , 2021, 10, 2498.	4.1	4
32	A report from the European Hyperoxaluria Consortium (OxalEurope) Registry on a large cohort of patients with primary hyperoxaluria type 3. <i>Kidney International</i> , 2021, 100, 621-635.	5.2	26
33	X-linked hypophosphatemia and burosumab: Practical clinical points from the French experience. <i>Joint Bone Spine</i> , 2021, 88, 105208.	1.6	14
34	Bone marrow adiposity inversely correlates with bone turnover in pediatric renal osteodystrophy. <i>Bone Reports</i> , 2021, 15, 101104.	0.4	0
35	FGF23 and infectious diseases. , 2021, , 175-182.		1
36	Hyperphosphatemia and Chronic Kidney Disease: A Major Daily Concern Both in Adults and in Children. <i>Calcified Tissue International</i> , 2021, 108, 116-127.	3.1	17

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37	Intermittent Bi-Daily Sub-cutaneous Teriparatide Administration in Children With Hypoparathyroidism: A Single-Center Experience. <i>Frontiers in Pediatrics</i> , 2021, 9, 764040.	1.9	2
38	Teenagers and young adults with a past of allogenic hematopoietic stem cell transplantation are at significant risk of chronic kidney disease. <i>Pediatric Nephrology</i> , 2021, , 1.	1.7	0
39	Peripheral Blood Mononuclear Cells (PBMCs) to Dissect the Underlying Mechanisms of Bone Disease in Chronic Kidney Disease and Rare Renal Diseases. <i>Current Osteoporosis Reports</i> , 2021, 19, 553.	3.6	2
40	Physiology of FGF23 and overview of genetic diseases associated with renal phosphate wasting. <i>Metabolism: Clinical and Experimental</i> , 2020, 103, 153865.	3.4	55
41	Treatment of hyperphosphatemia: the dangers of high PTH levels. <i>Pediatric Nephrology</i> , 2020, 35, 493-500.	1.7	15
42	Intermittent cholecalciferol supplementation in children and teenagers followed in pediatric nephrology: data from a prospective single-center single-arm open trial. <i>European Journal of Pediatrics</i> , 2020, 179, 661-669.	2.7	6
43	Association between 25(OH) vitamin D and graft survival in renal transplanted children. <i>Pediatric Transplantation</i> , 2020, 24, e13809.	1.0	3
44	Long-term outcomes of peritoneal dialysis started in infants below 6 months of age: An experience from two tertiary centres. <i>Nephrologie Et Therapeutique</i> , 2020, 16, 424-430.	0.5	3
45	Bone Disease in Nephropathic Cystinosis: Beyond Renal Osteodystrophy. <i>International Journal of Molecular Sciences</i> , 2020, 21, 3109.	4.1	15
46	Cinacalcet studies in pediatric subjects with secondary hyperparathyroidism receiving dialysis. <i>Pediatric Nephrology</i> , 2020, 35, 1679-1697.	1.7	12
47	The use of cinacalcet after pediatric renal transplantation: an international CERTAIN Registry analysis. <i>Pediatric Nephrology</i> , 2020, 35, 1707-1718.	1.7	9
48	The interest of oral calcium loads test in the diagnosis and management of pediatric nephrolithiasis with hypercalciuria: Experience from a tertiary pediatric centre. <i>Journal of Pediatric Urology</i> , 2020, 16, 489.e1-489.e9.	1.1	4
49	Inhibition of Osteoclast Differentiation by 1,25(OH) ₂ D ₃ and the Calcimimetic KP2326 Reveals 1,25(OH) ₂ D ₃ Resistance in Advanced CKD. <i>Journal of Bone and Mineral Research</i> , 2020, 35, 2265-2274.	2.8	8
50	Developing Consensus-Based Outcome Domains for Trials in Children and Adolescents With CKD: An International Delphi Survey. <i>American Journal of Kidney Diseases</i> , 2020, 76, 533-545.	1.9	19
51	An Expert Perspective on Phosphate Dysregulation With a Focus on Chronic Hypophosphatemia. <i>Journal of Bone and Mineral Research</i> , 2020, 37, 12-20.	2.8	11
52	Management of children with congenital nephrotic syndrome: challenging treatment paradigms. <i>Nephrology Dialysis Transplantation</i> , 2019, 34, 1369-1377.	0.7	32
53	Determinants of Statural Growth in European Children With Chronic Kidney Disease: Findings From the Cardiovascular Comorbidity in Children With Chronic Kidney Disease (4C) Study. <i>Frontiers in Pediatrics</i> , 2019, 7, 278.	1.9	19
54	Cinacalcet use in paediatric dialysis: a position statement from the European Society for Paediatric Nephrology and the Chronic Kidney Disease-Mineral and Bone Disorders Working Group of the ERA-EDTA. <i>Nephrology Dialysis Transplantation</i> , 2019, 35, 47-64.	0.7	18

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55	Defects in t6A tRNA modification due to GON7 and YRDC mutations lead to Galloway-Mowat syndrome. <i>Nature Communications</i> , 2019, 10, 3967.	12.8	66
56	Assessment of mineral and bone biomarkers highlights a high frequency of hypercalciuria in asymptomatic healthy teenagers. <i>Acta Paediatrica, International Journal of Paediatrics</i> , 2019, 108, 2253-2260.	1.5	15
57	Management of bone disease in cystinosis: Statement from an international conference. <i>Journal of Inherited Metabolic Disease</i> , 2019, 42, 1019-1029.	3.6	39
58	Clinical practice recommendations for growth hormone treatment in children with chronic kidney disease. <i>Nature Reviews Nephrology</i> , 2019, 15, 577-589.	9.6	103
59	Clinical practice recommendations for the diagnosis and management of X-linked hypophosphataemia. <i>Nature Reviews Nephrology</i> , 2019, 15, 435-455.	9.6	318
60	Escherichia coli-associated hemolytic uremic syndrome and severe chronic hepatocellular cholestasis: complication or side effect of eculizumab?. <i>Pediatric Nephrology</i> , 2019, 34, 1289-1293.	1.7	6
61	Fluconazole as a New Therapeutic Tool to Manage Patients With NPT1c (SLC34A3) Mutation: A Case Report. <i>American Journal of Kidney Diseases</i> , 2019, 73, 886-889.	1.9	14
62	High Incidence of Cranial Synostosis and Chiari I Malformation in Children With X-Linked Hypophosphatemic Rickets (XLHR). <i>Journal of Bone and Mineral Research</i> , 2019, 34, 490-496.	2.8	53
63	Infants with congenital nephrotic syndrome have comparable outcomes to infants with other renal diseases. <i>Pediatric Nephrology</i> , 2019, 34, 649-655.	1.7	16
64	Skin microvascular dysfunction as an early cardiovascular marker in primary hyperoxaluria type I. <i>Pediatric Nephrology</i> , 2019, 34, 319-327.	1.7	4
65	Towards adulthood with a solitary kidney. <i>Pediatric Nephrology</i> , 2019, 34, 2311-2323.	1.7	28
66	Treatment and outcome of congenital nephrotic syndrome. <i>Nephrology Dialysis Transplantation</i> , 2019, 34, 458-467.	0.7	42
67	Longitudinal Bone Mineralization Assessment in Children Treated With Long-Term Parenteral Nutrition for Severe Intestinal Failure. <i>Journal of Parenteral and Enteral Nutrition</i> , 2018, 42, 613-622.	2.6	14
68	Teenagers and young adults with nephropathic cystinosis display significant bone disease and cortical impairment. <i>Pediatric Nephrology</i> , 2018, 33, 1165-1172.	1.7	16
69	Renal transplantation in children under 3 years of age: Experience from a single-center study. <i>Pediatric Transplantation</i> , 2018, 22, e13116.	1.0	7
70	Bone disease in nephropathic cystinosis is related to cystinosis-induced osteoclastic dysfunction. <i>Nephrology Dialysis Transplantation</i> , 2018, 33, 1525-1532.	0.7	16
71	Patient and transplant outcome in infants starting renal replacement therapy before 2 years of age. <i>Nephrology Dialysis Transplantation</i> , 2018, 33, 1459-1465.	0.7	15
72	Hyperphosphatemic tumoral calcinosis caused by FGF23 compound heterozygous mutations: what are the therapeutic options for a better control of phosphatemia?. <i>Pediatric Nephrology</i> , 2018, 33, 1263-1267.	1.7	17

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73	Treatment by immunoadsorption for recurrent focal segmental glomerulosclerosis after paediatric kidney transplantation: a multicentre French cohort. <i>Nephrology Dialysis Transplantation</i> , 2018, 33, 954-963.	0.7	36
74	Skeletal impairment in Pierson syndrome: Is there a role for laminin α 2 in bone physiology?. <i>Bone</i> , 2018, 106, 187-193.	2.9	8
75	Standardization of pediatric urological terms: a multidisciplinary European glossary. <i>Pediatric Radiology</i> , 2018, 48, 291-303.	2.0	11
76	Vitamin D deficiency is associated with mortality in maintenance dialysis: moving forward from epidemiology to clinical trials. <i>Nephrology Dialysis Transplantation</i> , 2018, 33, 1679-1682.	0.7	8
77	The interplay between bone and vessels in pediatric CKD: lessons from a single-center study. <i>Pediatric Nephrology</i> , 2018, 33, 1565-1575.	1.7	14
78	C3 glomerulopathy and eculizumab: a report on four paediatric cases. <i>Pediatric Nephrology</i> , 2017, 32, 1023-1028.	1.7	29
79	Bone biopsy practice patterns across Europe: the European renal osteodystrophy initiative—a position paper. <i>Nephrology Dialysis Transplantation</i> , 2017, 32, 1608-1613.	0.7	41
80	Clinical practice recommendations for treatment with active vitamin D analogues in children with chronic kidney disease Stages 2–5 and on dialysis. <i>Nephrology Dialysis Transplantation</i> , 2017, 32, 1114-1127.	0.7	51
81	Clinical practice recommendations for native vitamin D therapy in children with chronic kidney disease Stages 2–5 and on dialysis. <i>Nephrology Dialysis Transplantation</i> , 2017, 32, 1098-1113.	0.7	84
82	Evidence for Bone and Mineral Metabolism Alterations in Children With Autosomal Dominant Polycystic Kidney Disease. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2017, 102, 4210-4217.	3.6	15
83	Standardization of pediatric urological terms: A multidisciplinary European glossary. <i>Journal of Pediatric Urology</i> , 2017, 13, 641-650.	1.1	5
84	The Relationship between Body Composition and Bone Quality Measured with HR-pQCT in Peritoneal Dialysis Patients. <i>Peritoneal Dialysis International</i> , 2017, 37, 548-555.	2.3	3
85	Worldwide view of nephropathic cystinosis: results from a survey from 30 countries. <i>BMC Nephrology</i> , 2017, 18, 210.	1.8	16
86	L'hyperoxalurie primitive, aujourd'hui et demain. <i>Bulletin De L'Academie Nationale De Medecine</i> , 2017, 201, 1361-1375.	0.0	1
87	Efficacy of extracorporeal albumin dialysis for acute kidney injury due to cholestatic jaundice nephrotoxicity. <i>BMJ Case Reports</i> , 2016, 2016, bcr2015213257.	0.5	11
88	French law: what about a reasoned reimbursement of serum vitamin D assays?. <i>Psychologie & Neuropsychiatrie Du Vieillessement</i> , 2016, 14, 377-382.	0.2	7
89	Impact of a change in protected environment on the occurrence of severe bacterial and fungal infections in children undergoing hematopoietic stem cell transplantation. <i>European Journal of Haematology</i> , 2016, 97, 70-77.	2.2	12
90	Pediatric combined liver–kidney transplantation: a single-center experience of 18 cases. <i>Pediatric Nephrology</i> , 2016, 31, 1517-1529.	1.7	36

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91	Combining exercise and growth hormone therapy: how can we translate from animal models to chronic kidney disease children?. <i>Nephrology Dialysis Transplantation</i> , 2016, 31, 1191-1194.	0.7	1
92	Genetic, Environmental, and Disease-Associated Correlates of Vitamin D Status in Children with CKD. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2016, 11, 1145-1153.	4.5	10
93	Osteocalcin Signaling in Myofibers Is Necessary and Sufficient for Optimum Adaptation to Exercise. <i>Cell Metabolism</i> , 2016, 23, 1078-1092.	16.2	302
94	Bone impairment in primary hyperoxaluria: a review. <i>Pediatric Nephrology</i> , 2016, 31, 1-6.	1.7	34
95	FGF23 in chronic kidney disease: are we lost in translation?. <i>BoneKEY Reports</i> , 2016, 5, 770.	2.7	0
96	Skeletal implications and management of cystinosis: three case reports and literature review. <i>BoneKEY Reports</i> , 2016, 5, 828.	2.7	15
97	Vitamin D in Children with Chronic Kidney Disease: A Focus on Longitudinal Bone Growth. , 2016, , 229-245.		0
98	Early-onset hypoparathyroidism and chronic keratitis revealing <scp>APECED</scp>. <i>Clinical Case Reports (discontinued)</i> , 2015, 3, 809-813.	0.5	4
99	Pediatric combined liver-kidney transplantation. <i>Current Opinion in Organ Transplantation</i> , 2015, 20, 543-549.	1.6	19
100	Markers of Bone Metabolism Are Affected by Renal Function and Growth Hormone Therapy in Children with Chronic Kidney Disease. <i>PLoS ONE</i> , 2015, 10, e0113482.	2.5	33
101	SP665THE RELATION BETWEEN ADIPOKINES, BODY COMPOSITION, AND BONE HEALTH MEASURED WITH HR-~PQCT IN HEMODIALYSIS PATIENTS. <i>Nephrology Dialysis Transplantation</i> , 2015, 30, iii598-iii598.	0.7	1
102	Renal function can be impaired in children with primary hyperoxaluria type 3. <i>Pediatric Nephrology</i> , 2015, 30, 1807-1813.	1.7	29
103	Biphasic Effects of Vitamin D and FGF23 on Human Osteoclast Biology. <i>Calcified Tissue International</i> , 2015, 97, 69-79.	3.1	33
104	Paediatric liver transplanted patients and prevalence of hepatitis E virus.. <i>Journal of Clinical Virology</i> , 2015, 69, 22-26.	3.1	14
105	Bone impairment in oxalosis: An ultrastructural bone analysis. <i>Bone</i> , 2015, 81, 161-167.	2.9	23
106	Mutation Update of the <i>CLCN5</i> Gene Responsible for Dent Disease 1. <i>Human Mutation</i> , 2015, 36, 743-752.	2.5	66
107	Primary disease recurrence effects on paediatric renal transplantation outcomes. <i>Nature Reviews Nephrology</i> , 2015, 11, 371-384.	9.6	34
108	Serum sclerostin: the missing link in the bone-vessel cross-talk in hemodialysis patients?. <i>Osteoporosis International</i> , 2015, 26, 2165-2174.	3.1	46

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109	<i>CYP24A1</i> Mutations in a Cohort of Hypercalcemic Patients: Evidence for a Recessive Trait. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2015, 100, E1343-E1352.	3.6	116
110	Calcium balance in pediatric online hemodiafiltration: Beware of sodium and bicarbonate in the dialysate. <i>Nephrologie Et Therapeutique</i> , 2015, 11, 483-486.	0.5	4
111	Nephropathic Cystinosis – A Gap between Developing and Developed Nations. <i>New England Journal of Medicine</i> , 2014, 370, 1366-1367.	27.0	27
112	Immune, metabolic and epidemiological aspects of vitamin D in chronic kidney disease and transplant patients. <i>Clinical Biochemistry</i> , 2014, 47, 509-515.	1.9	8
113	2014 update of recommendations on the prevention and treatment of glucocorticoid-induced osteoporosis. <i>Joint Bone Spine</i> , 2014, 81, 493-501.	1.6	92
114	Rapid access to renal transplant waiting list in children: impact of patient and centre characteristics in France. <i>Nephrology Dialysis Transplantation</i> , 2014, 29, 1973-1979.	0.7	18
115	Suppression of Iron-Regulatory Heparin by Vitamin D. <i>Journal of the American Society of Nephrology: JASN</i> , 2014, 25, 564-572.	6.1	252
116	Fludrocortisone as a new tool for managing tubulopathy after pediatric renal transplantation: a series of cases. <i>Pediatric Nephrology</i> , 2014, 29, 2061-2064.	1.7	6
117	Eculizumab in neonatal hemolytic uremic syndrome with homozygous factor H deficiency. <i>Pediatric Nephrology</i> , 2014, 29, 2415-2419.	1.7	18
118	Antibacterial Responses by Peritoneal Macrophages Are Enhanced Following Vitamin D Supplementation. <i>PLoS ONE</i> , 2014, 9, e116530.	2.5	26
119	Idiopathic juvenile osteoporosis: a cross-sectional single-centre experience with bone histomorphometry and quantitative computed tomography. <i>Pediatric Rheumatology</i> , 2013, 11, 6.	2.1	20
120	From bone abnormalities to mineral metabolism dysregulation in autosomal dominant polycystic kidney disease. <i>Pediatric Nephrology</i> , 2013, 28, 2089-2096.	1.7	14
121	Beyond mineral metabolism, is there an interplay between FGF23 and vitamin D in innate immunity?. <i>Pediatric Nephrology</i> , 2013, 28, 577-582.	1.7	25
122	Fibroblast growth factor 23 inhibits extrarenal synthesis of 1,25-dihydroxyvitamin D in human monocytes. <i>Journal of Bone and Mineral Research</i> , 2013, 28, 46-55.	2.8	163
123	Long-term critical issues in pediatric renal transplant recipients: a single-center experience. <i>Transplant International</i> , 2013, 26, 154-161.	1.6	28
124	The Skeletal Consequences of Growth Hormone Therapy in Dialyzed Children. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2013, 8, 824-832.	4.5	28
125	The consequences of pediatric renal transplantation on bone metabolism and growth. <i>Current Opinion in Organ Transplantation</i> , 2013, 18, 555-562.	1.6	19
126	What about the renal function during childhood of children born from dialysed mothers?. <i>Nephrology Dialysis Transplantation</i> , 2012, 27, 2365-2369.	0.7	24

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127	Bone microarchitecture is more severely affected in patients on hemodialysis than in those receiving peritoneal dialysis. <i>Kidney International</i> , 2012, 82, 581-588.	5.2	34
128	GFR Estimation in Adolescents and Young Adults. <i>Journal of the American Society of Nephrology: JASN</i> , 2012, 23, 989-996.	6.1	74
129	The consequences of chronic kidney disease on bone metabolism and growth in children. <i>Nephrology Dialysis Transplantation</i> , 2012, 27, 3063-3071.	0.7	88
130	Uric acid and IGF1 as possible determinants of FGF23 metabolism in children with normal renal function. <i>Pediatric Nephrology</i> , 2012, 27, 1131-1138.	1.7	31
131	Evaluation of Hypophosphatemia: Lessons From Patients With Genetic Disorders. <i>American Journal of Kidney Diseases</i> , 2012, 59, 152-159.	1.9	39
132	Re: Imaging Strategy for Infants With Urinary Tract Infection: A New Algorithm. <i>Journal of Urology</i> , 2011, 186, 2506-2507.	0.4	0
133	Evolution of renal oxygen content measured by BOLD MRI downstream a chronic renal artery stenosis. <i>Nephrology Dialysis Transplantation</i> , 2011, 26, 1205-1210.	0.7	40
134	Bone assessment in children with chronic kidney disease: data from two new bone imaging techniques in a single-center pilot study. <i>Pediatric Nephrology</i> , 2011, 26, 587-595.	1.7	36
135	CKD-MBD after kidney transplantation. <i>Pediatric Nephrology</i> , 2011, 26, 2143-2151.	1.7	26
136	FGF23 and paediatric transplantation: a single-centre French experience. <i>Nephrology Dialysis Transplantation</i> , 2011, 26, 3421-3422.	0.7	5
137	Congenital versus acquired solitary kidney: is the difference relevant?. <i>Nephrology Dialysis Transplantation</i> , 2011, 26, 2188-2194.	0.7	66
138	Which Creatinine and Cystatin C Equations Can Be Reliably Used in Children?. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2011, 6, 552-560.	4.5	114
139	Primary Hyperoxaluria. <i>International Journal of Nephrology</i> , 2011, 2011, 1-11.	1.3	76
140	Early impairment of trabecular microarchitecture assessed with HR-pQCT in patients with stage II-IV chronic kidney disease. <i>Journal of Bone and Mineral Research</i> , 2010, 25, 849-857.	2.8	87
141	Assessment of Bone Microarchitecture in Chronic Kidney Disease: A Comparison of 2D Bone Texture Analysis and High-Resolution Peripheral Quantitative Computed Tomography at the Radius and Tibia. <i>Calcified Tissue International</i> , 2010, 87, 385-391.	3.1	11
142	Nephrolithiasis related to inborn metabolic diseases. <i>Pediatric Nephrology</i> , 2010, 25, 415-424.	1.7	77
143	Bone metabolism in oxalosis: a single-center study using new imaging techniques and biomarkers. <i>Pediatric Nephrology</i> , 2010, 25, 1081-1089.	1.7	31
144	The Influence of Glomerular Filtration Rate and Age on Fibroblast Growth Factor 23 Serum Levels in Pediatric Chronic Kidney Disease. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2010, 95, 1741-1748.	3.6	112

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145	Assessment of hand bone loss in rheumatoid arthritis by high-resolution peripheral quantitative CT. <i>Annals of the Rheumatic Diseases</i> , 2010, 69, 1671-1676.	0.9	95
146	What is the best alternative to inulin clearance to estimate GFR in patients with decompensated alcoholic cirrhosis?. <i>Nephrology Dialysis Transplantation</i> , 2010, 25, 3569-3575.	0.7	24
147	The Case of Severe voiding dysfunction: ask the child to smile. <i>Kidney International</i> , 2010, 78, 225-226.	5.2	3
148	Both extrauterine and intrauterine growth restriction impair renal function in children born very preterm. <i>Kidney International</i> , 2009, 76, 445-452.	5.2	119
149	Precocious puberty and unlicensed paediatric drugs for severe hyperparathyroidism. <i>Nephrology Dialysis Transplantation</i> , 2009, 24, 2595-2598.	0.7	13
150	The relationship between adipokines, osteocalcin and bone quality in chronic kidney disease. <i>Nephrology Dialysis Transplantation</i> , 2009, 24, 3120-3125.	0.7	53
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