Michio Nagata

List of Publications by Year
in descending order

185 papers	8,769 citations	76326 40 h-index	46799 89 g-index
195 all docs	195 docs citations	195 times ranked	8170 citing authors

#	Article	IF	Citations
1	CCR2- and CCR5-mediated macrophage infiltration contributes to glomerular endocapillary hypercellularity in antibody-induced lupus nephritis. Rheumatology, 2022, 61, 3033-3048.	1.9	7
2	Clinical impact of urinary CD11b and CD163 on the renal outcomes of anti-neutrophil cytoplasmic antibody-associated glomerulonephritis. Nephrology Dialysis Transplantation, 2021, 36, 1452-1463.	0.7	5
3	Poststreptococcal acute glomerulonephritis in a girl with renal cell carcinoma: possible pathophysiological association. CEN Case Reports, 2021, 10, 139-144.	0.9	0
4	Literature review of allograft adenovirus nephritis and a case presenting as mass lesions in a transplanted kidney without symptoms of urinary tract infection or acute kidney injury. Transplant Infectious Disease, 2021, 23, e13468.	1.7	4
5	Nephrotic syndrome with focal segmental glomerular lesions unclassified by Columbia classification; Pathology and clinical implication. PLoS ONE, 2021, 16, e0244677.	2.5	3
6	Validation of the 2019 ACR/EULAR criteria for IgG4-related disease in a Japanese kidney disease cohort: a multicentre retrospective study by the IgG4-related kidney disease working group of the Japanese Society of Nephrology. Annals of the Rheumatic Diseases, 2021, 80, 956-957.	0.9	6
7	Glomerulonephritis with severe nephrotic syndrome induced by immune complexes composed of galactose-deficient IgA1 in primary SjÃ¶grenâ€™s syndrome: a case report. BMC Nephrology, 2021, 22, 108.	1.8	2
8	Glomerular Classification Using Convolutional Neural Networks Based on Defined Annotation Criteria and Concordance Evaluation Among Clinicians. Kidney International Reports, 2021, 6, 716-726.	0.8	16
9	Nationwide Survey of Post-Transplant Glomerular Diseases, Based on the Japan Renal Biopsy Registry (J-RBR). Annals of Transplantation, 2021, 26, e931873.	0.9	1
10	Unusual ischemic kidney injury presenting as slowly declining graft function and successful use of oral desmopressin in a kidney transplant recipient with subclinical central diabetes insipidus. Clinical Nephrology, 2021, 95, 208-214.	0.7	2
11	Coding practice in national and regional kidney biopsy registries. BMC Nephrology, 2021, 22, 193.	1.8	9
12	Genetic Background and Clinicopathologic Features of Adult-onset Nephronophthisis. Kidney International Reports, 2021, 6, 1346-1354.	0.8	14
13	Glomerular filtrate affects the dynamics of podocyte detachment in a model of diffuse toxic podocytopathy. Kidney International, 2021, 99, 1149-1161.	5.2	6
14	Caspase-3 regulates ureteric branching in mice via cell migration. Biochemical and Biophysical Research Communications, 2021, 559, 28-34.	2.1	2
15	Validation of the diagnostic criteria for IgG4-related kidney disease (IgG4-RKD) 2011, and proposal of a new 2020 version. Clinical and Experimental Nephrology, 2021, 25, 99-109.	1.6	20
16	The Clinical and Histopathological Feature of Renal Manifestation of TAFRO Syndrome. Kidney International Reports, 2020, 5, 1172-1179.	0.8	20
17	The revised version 2018 of the nationwide web-based registry system for kidney diseases in Japan: Japan Renal Biopsy Registry and Japan Kidney Disease Registry. Clinical and Experimental Nephrology, 2020, 24, 1058-1068.	1.6	5
18	Bidirectional, non-necrotizing glomerular crescents are the critical pathology in X-linked Alport syndrome mouse model harboring nonsense mutation of human COL4A5. Scientific Reports, 2020, 10, 18891.	3.3	5

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19	Biphasic MIF and SDF1 expression during podocyte injury promote CD44-mediated glomerular parietal cell migration in focal segmental glomerulosclerosis. American Journal of Physiology - Renal Physiology, 2020, 318, F741-F753.	2.7	9
20	Molecular mechanisms determining severity in patients with Pierson syndrome. Journal of Human Genetics, 2020, 65, 355-362.	2.3	8
21	A review of clinical characteristics and genetic backgrounds in Alport syndrome. Clinical and Experimental Nephrology, 2019, 23, 158-168.	1.6	135
22	Possible role of complement factor H in podocytes in clearing glomerular subendothelial immune complex deposits. Scientific Reports, 2019, 9, 7857.	3.3	21
23	A case report of crystalline light chain inclusion-associated kidney disease affecting podocytes but without Fanconi syndrome. Medicine (United States), 2019, 98, e13915.	1.0	8
24	Establishment of X-linked Alport syndrome model mice with a Col4a5 R471X mutation. Biochemistry and Biophysics Reports, 2019, 17, 81-86.	1.3	19
25	Histopathological classification of anti-neutrophil cytoplasmic antibody-associated glomerulonephritis in a nationwide Japanese prospective 2-year follow-up cohort study. Clinical and Experimental Nephrology, 2019, 23, 387-394.	1.6	9
26	Postâ€infectious acute glomerulonephritis with podocytopathy induced by parvovirus B19 infection. Pathology International, 2018, 68, 190-195.	1.3	6
27	Glomerulogenesis and the role of endothelium. Current Opinion in Nephrology and Hypertension, 2018, 27, 159-164.	2.0	6
28	Clinical features and pathogenesis of membranoproliferative glomerulonephritis: a nationwide analysis of the Japan renal biopsy registry from 2007 to 2015. Clinical and Experimental Nephrology, 2018, 22, 797-807.	1.6	19
29	AL amyloidosis with non-amyloid forming monoclonal immunoglobulin deposition; a case mimicking AHL amyloidosis. BMC Nephrology, 2018, 19, 337.	1.8	6
30	Immune-mediated acquired lecithin-cholesterol acyltransferase deficiency: A case report and literature review. Journal of Clinical Lipidology, 2018, 12, 888-897.e2.	1.5	10
31	Estimation of the number of histological diagnosis for IgG4-related kidney disease referred to the data obtained from the Japan Renal Biopsy Registry (J-RBR) questionnaire and cases reported in the Japanese Society of Nephrology Meetings. Clinical and Experimental Nephrology, 2017, 21, 97-103.	1.6	15
32	Focal segmental glomerulosclerosis and medullary nephrocalcinosis in children with ADCK4 mutations. Pediatric Nephrology, 2017, 32, 1547-1554.	1.7	27
33	Myeloperoxidaseâ€antineutrophil cytoplasmic antibody causes different renal diseases by immuneâ€complex formation and pauciâ€immune mechanism: A case report. Pathology International, 2017, 67, 419-424.	1.3	1
34	BMP7 dose-dependently stimulates proliferation and cadherin-11 expression via ERK and p38 in a murine metanephric mesenchymal cell line. Physiological Reports, 2017, 5, e13378.	1.7	7
35	Diversity of renal phenotypes in patients with WDR19 mutations: Two case reports. Nephrology, 2017, 22, 566-571.	1.6	12
36	Focal segmental glomerulosclerosis; why does it occur segmentally?. Pflugers Archiv European Journal of Physiology, 2017, 469, 983-988.	2.8	13

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37	Complex glomerular pathology of thrombotic microangiopathy and focal segmental glomerulosclerosis forms tumor-like mass in a renal transplant donor with severe renovascular hypertension. CEN Case Reports, 2017, 6, 12-17.	0.9	3
38	C3 glomerulopathy and current dilemmas. Clinical and Experimental Nephrology, 2017, 21, 541-551.	1.6	27
39	Clinical and histological features of lupus nephritis in Japan: A crossâ€sectional analysis of the Japan Renal Biopsy Registry (Jâ€RBR). Nephrology, 2017, 22, 885-891.	1.6	12
40	Effectiveness of Plasmapheresis in a Patient with Anti-glomerular Basement Membrane Antibody Glomerulonephritis with Advanced Kidney Dysfunction. Internal Medicine, 2017, 56, 2475-2479.	0.7	1
41	Anaemia is an essential complication of ANCA-associated renal vasculitis: a single center cohort study. BMC Nephrology, 2017, 18, 337.	1.8	21
42	Age-Related Renal Microvascular Changes: Evaluation by Three-Dimensional Digital Imaging of the Human Renal Microcirculation Using Virtual Microscopy. International Journal of Molecular Sciences, 2016, 17, 1831.	4.1	10
43	Temporal Changes in Post-Infectious Glomerulonephritis in Japan (1976-2009). PLoS ONE, 2016, 11, e0157356.	2.5	5
44	Heparan sulfate 6-O-endosulfatases, Sulf1 and Sulf2, regulate glomerular integrity by modulating growth factor signaling. American Journal of Physiology - Renal Physiology, 2016, 310, F395-F408.	2.7	19
45	Podocyte injury and its consequences. Kidney International, 2016, 89, 1221-1230.	5.2	342
46	A condition closely mimicking IgG4-related disease despite the absence of serum IgG4 elevation and IgG4-positive plasma cell infiltration. Modern Rheumatology, 2016, 26, 784-789.	1.8	16
47	Distribution and components of interstitial inflammation and fibrosis in IgG4-related kidney disease: analysis of autopsy specimens. Human Pathology, 2016, 55, 164-173.	2.0	14
48	Drug-induced kidney disease: a study of the Japan Renal Biopsy Registry from 2007 to 2015. Clinical and Experimental Nephrology, 2016, 20, 720-730.	1.6	18
49	Characteristic Tubulointerstitial Nephritis in IgG4-Related Kidney Disease. , 2016, , 105-113.		0
50	Characteristic Distribution of Inflammatory Lesions in IgG4-Related Kidney Disease: Findings from Autopsy Case Series. , 2016, , 187-191.		0
51	Effector and upstream mechanism of immune-mediated glomerulonephritis. Japanese Journal of Nephrology, 2016, 58, 614-21.	0.0	0
52	Rapidly progressive glomerulonephritis: recent advances. Japanese Journal of Nephrology, 2016, 58, 656-9.	0.0	0
53	Highâ€resolution threeâ€dimensional digital imaging of the human renal microcirculation: An aid to evaluating microvascular alterations in chronic kidney disease in humans. Pathology International, 2015, 65, 575-584.	1.3	3
54	Membranous Nephropathy with Solitary Immunoglobulin A Deposition. Internal Medicine, 2015, 54, 1081-1084.	0.7	4

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55	Rapid Deterioration of the Renal Function Caused by the Coexistence of Intratubular Amyloidosis and Myeloma Cast Nephropathy. Internal Medicine, 2015, 54, 3023-3028.	0.7	7
56	Podocyte injury-driven intracapillary plasminogen activator inhibitor type 1 accelerates podocyte loss via uPAR-mediated Î²1-integrin endocytosis. American Journal of Physiology - Renal Physiology, 2015, 308, F614-F626.	2.7	45
57	Podocyte Injuryâ€“Driven Lipid Peroxidation Accelerates the Infiltration of Glomerular Foam Cells in Focal Segmental Glomerulosclerosis. American Journal of Pathology, 2015, 185, 2118-2131.	3.8	39
58	Renal AH Amyloidosis Associated With a Truncated Immunoglobulin Heavy Chain Undetectable by Immunostaining. American Journal of Kidney Diseases, 2015, 66, 1095-1100.	1.9	20
59	Sustained Appearance of Urinary Podocytes Suggests Poor Renal Prognosis in Kidney Transplant Patients with Focal Segmental Glomerulosclerosis:. Clinical Laboratory, 2015, 61, 1961-6.	0.5	0
60	Diffuse Glomerular Nodular Lesions in Diabetic Pigs Carrying a Dominant-Negative Mutant Hepatocyte Nuclear Factor 1-Alpha, an Inheritant Diabetic Gene in Humans. PLoS ONE, 2014, 9, e92219.	2.5	13
61	The direction and role of phenotypic transition between podocytes and parietal epithelial cells in focal segmental glomerulosclerosis. American Journal of Physiology - Renal Physiology, 2014, 306, F98-F104.	2.7	41
62	Cyclosporine C2 Monitoring for the Treatment of Frequently Relapsing Nephrotic Syndrome in Children. Clinical Journal of the American Society of Nephrology: CJASN, 2014, 9, 271-278.	4.5	27
63	Graft versus host disease-dependent renal dysfunction after hematopoietic stem cell transplantation. CEN Case Reports, 2014, 3, 202-205.	0.9	3
64	Glomerular epithelial cell phenotype in diffuse mesangial sclerosis: a report of 2 cases with markedly increased urinary podocyte excretion. Human Pathology, 2014, 45, 1778-1783.	2.0	3
65	IgG4-Related Kidney Disease. , 2014, , 169-179.		1
66	Japan Renal Biopsy Registry and Japan Kidney Disease Registry: Committee Report for 2009 and 2010. Clinical and Experimental Nephrology, 2013, 17, 155-173.	1.6	111
67	Diabetic nephrotic syndrome 25 years after tumor-forming pancreatitis surgeryâ€”reply. Human Pathology, 2013, 44, 300.	2.0	0
68	Primary membranoproliferative glomerulonephritis on the decline: decreased rate from the 1970s to the 2000s in Japan. Clinical and Experimental Nephrology, 2013, 17, 248-254.	1.6	11
69	Aberrant Notch1-dependent effects on glomerular parietal epithelial cells promotes collapsing focal segmental glomerulosclerosis with progressive podocyte loss. Kidney International, 2013, 83, 1065-1075.	5.2	57
70	The podocyte's response to stress: the enigma of foot process effacement. American Journal of Physiology - Renal Physiology, 2013, 304, F333-F347.	2.7	231
71	Nephrotic Syndrome Caused by Immune-Mediated Acquired LCAT Deficiency. Journal of the American Society of Nephrology: JASN, 2013, 24, 1305-1312.	6.1	33
72	Henoch-SchÃ¶nlein purpura nephritis in a patient with IgG4-related disease: A possible association. Clinical Nephrology, 2013, 79, 246-252.	0.7	30

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73	Focal Segmental Glomerulosclerosis in Patients With Complete Deletion of One WT1 Allele. Pediatrics, 2012, 129, e1621-e1625.	2.1	11
74	Remission of proteinuria and preservation of renal function in patients with renal AA amyloidosis secondary to rheumatoid arthritis. Nephrology Dialysis Transplantation, 2012, 27, 633-639.	0.7	18
75	Renal disease in the elderly and the very elderly Japanese: analysis of the Japan Renal Biopsy Registry (J-RBR). Clinical and Experimental Nephrology, 2012, 16, 903-920.	1.6	91
76	Clinical and histological changes associated with corticosteroid therapy in IgG4-related tubulointerstitial nephritis. Modern Rheumatology, 2012, 22, 859-870.	1.8	44
77	Characteristic tubulointerstitial nephritis in IgG4-related disease. Human Pathology, 2012, 43, 536-549.	2.0	110
78	Histological heterogeneity of glomerular segmental lesions in focal segmental glomerulosclerosis. International Urology and Nephrology, 2012, 44, 183-196.	1.4	24
79	Clinical and histological changes associated with corticosteroid therapy in IgG4-related tubulointerstitial nephritis. Modern Rheumatology, 2012, 22, 859-870.	1.8	29
80	A case of multicentric Castleman's disease with membranoproliferative glomerulonephritis type 3â€like lesion. Pathology International, 2011, 61, 686-690.	1.3	5
81	Morphological and functional analyses of two infants with obstructive renal dysplasia. Clinical and Experimental Nephrology, 2011, 15, 602-606.	1.6	1
82	Japan Renal Biopsy Registry: the first nationwide, web-based, and prospective registry system of renal biopsies in Japan. Clinical and Experimental Nephrology, 2011, 15, 493-503.	1.6	127
83	Focal segmental glomerulosclerosis as a complication of hepatitis B virus infection. Nephrology Dialysis Transplantation, 2011, 26, 371-373.	0.7	21
84	Microscopic papillary tumor in a renal needle biopsy specimen for IgA nephropathy. CKJ: Clinical Kidney Journal, 2011, 4, 357-358.	2.9	0
85	Granular Swollen Epithelial Cells: A Histologic and Diagnostic Marker for Mitochondrial Nephropathy. American Journal of Surgical Pathology, 2010, 34, 262-270.	3.7	20
86	Renal involvement of monoclonal immunoglobulin deposition disease associated with an unusual monoclonal immunoglobulin A glycan profile. Clinical and Experimental Nephrology, 2010, 14, 389-395.	1.6	13
87	Successful treatment of collapsing focal segmental glomerulosclerosis with a combination of rituximab, steroids and ciclosporin. Pediatric Nephrology, 2010, 25, 957-959.	1.7	23
88	Resolution of Henoch-SchÃ¶nlein purpura nephritis after acquired IgA deficiency. Pediatric Nephrology, 2010, 25, 2355-2358.	1.7	3
89	A Case of Small-Cell Parotid Gland Carcinoma. Practica Otologica, 2010, 103, 729-735.	0.0	0
90	Amino Acid Transporter LAT3 Is Required for Podocyte Development and Function. Journal of the American Society of Nephrology: JASN, 2009, 20, 1586-1596.	6.1	34

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91	Acute and Transient Podocyte Loss and Proteinuria in Preeclampsia. Nephron Clinical Practice, 2009, 112, c65-c70.	2.3	54
92	Secondary membranous glomerulonephritis associated with recipient residual lymphoma cells after allogeneic bone marrow transplantation. Clinical and Experimental Nephrology, 2009, 13, 174-178.	1.6	6
93	Atypical phenotype of type I Bartter syndrome accompanied by focal segmental glomerulosclerosis. Pediatric Nephrology, 2009, 24, 415-418.	1.7	24
94	Reevaluation of glomerular charge selective protein-sieving function. Pediatric Nephrology, 2009, 24, 609-612.	1.7	5
95	Genetic Podocyte Lineage Reveals Progressive Podocytopenia with Parietal Cell Hyperplasia in a Murine Model of Cellular/Collapsing Focal Segmental Glomerulosclerosis. American Journal of Pathology, 2009, 174, 1675-1682.	3.8	46
96	Successful pregnancy in a female patient with congenital chloride diarrhea (CLD) and renal impairment. Journal of Nephrology, 2009, 22, 809-13.	2.0	1
97	Renal pathology of ANCA-related vasculitis: proposal for standardization of pathological diagnosis in Japan. Clinical and Experimental Nephrology, 2008, 12, 277-291.	1.6	25
98	An autopsy case of late-onset Epstein-Barr virus associated post-transplant lymphoproliferative disorders 11â€ƒyr after kidney transplantation. Clinical Transplantation, 2008, 22, 87-91.	1.6	0
99	Familial Focal Segmental Glomerulosclerosis Associated With an ACTN4 Mutation and Paternal Germline Mosaicism. American Journal of Kidney Diseases, 2008, 51, 834-838.	1.9	47
100	Hyporeninemic hypoaldosteronism from secondary amyloidosis. Kidney International, 2008, 74, 542.	5.2	7
101	A novel apolipoprotein E mutation, ApoE Tsukuba (Arg 114 Cys), in lipoprotein glomerulopathy. Nephrology Dialysis Transplantation, 2007, 23, 381-384.	0.7	22
102	Stable Transgene Expression in Mice Generated from Retrovirally Transduced Embryonic Stem Cells. Molecular Therapy, 2007, 15, 560-565.	8.2	9
103	Increased Expression of Vascular Endothelial Growth Factor in Kidney Leads to Progressive Impairment of Glomerular Functions. Journal of the American Society of Nephrology: JASN, 2007, 18, 2094-2104.	6.1	99
104	Deficiency of Endothelial Nitric-Oxide Synthase Confers Susceptibility to Diabetic Nephropathy in Nephropathy-Resistant Inbred Mice. American Journal of Pathology, 2007, 170, 1473-1484.	3.8	161
105	C4d Immunohistochemistry in glomerulonephritis with different antibodies. Clinical and Experimental Nephrology, 2007, 11, 287-291.	1.6	18
106	Triggers of relapse in steroid-dependent and frequently relapsing nephrotic syndrome. Pediatric Nephrology, 2007, 22, 232-236.	1.7	33
107	Murine Metanephric Mesenchyme Possesses Characteristics of Vascular Endothelial Cells in vitro. Nephron Experimental Nephrology, 2006, 102, e93-e98.	2.2	4
108	Unilateral renal angiodysplasia in a girl with hypertension. Pediatric Nephrology, 2006, 21, 292-294.	1.7	2

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109	Renal failure due to tubulointerstitial nephropathy in an infant with cranioectodermal dysplasia. Pediatric Nephrology, 2006, 21, 574-576.	1.7	6
110	Focal segmental glomerulosclerosis associated with essential thrombocythemia. Clinical and Experimental Nephrology, 2006, 10, 74-77.	1.6	20
111	Early recurrence of dense deposit disease with marked endocapillary proliferation after renal transplantation. Pathology International, 2006, 56, 101-109.	1.3	9
112	Charge Selective Function in Childhood Glomerular Diseases. Pediatric Research, 2006, 59, 336-340.	2.3	8
113	Tissue-specific distribution of an alternatively spliced COL4A5 isoform and non-random X chromosome inactivation reflect phenotypic variation in heterozygous X-linked Alport syndrome. Nephrology Dialysis Transplantation, 2006, 21, 1582-1587.	0.7	17
114	Role of integrin-linked kinase in epithelialâ€“mesenchymal transition in crescent formation of experimental glomerulonephritis. Nephrology Dialysis Transplantation, 2006, 21, 2380-2390.	0.7	51
115	MafB Is Essential for Renal Development and F4/80 Expression in Macrophages. Molecular and Cellular Biology, 2006, 26, 5715-5727.	2.3	189
116	Post-MRSA infection glomerulonephritis with marked Staphylococcus aureus cell envelope antigen deposition in glomeruli. Journal of Nephrology, 2006, 19, 215-9.	2.0	26
117	Primary Cerebral Angiitis Containing Marked Xanthoma Cells With Massive Intraparenchymal Involvement-Case Report-. Neurologia Medico-Chirurgica, 2005, 45, 156-160.	2.2	9
118	Retroperitoneal extragonadal germ cell tumor in a patient with Klinefelter's syndrome. International Journal of Urology, 2005, 12, 765-767.	1.0	1
119	Acute interstitial nephritis predisposed a six-year-old girl to minimal change nephrotic syndrome. Pediatric Nephrology, 2005, 20, 1168-1170.	1.7	8
120	A familial case of multicystic dysplastic kidney. Pediatric Nephrology, 2005, 20, 1245-1248.	1.7	11
121	Nephrotic Syndrome and Aberrant Expression of Laminin Isoforms in Glomerular Basement Membranes for an Infant With Herlitz Junctional Epidermolysis Bullosa. Pediatrics, 2005, 116, e601-e607.	2.1	32
122	MMF for Lupus Nephritis as a Possible First Aid Block. Nephron Clinical Practice, 2005, 100, c101-c102.	2.3	0
123	Confirmation of the occurrence of Parapenaeus Investigatoris Alcock & Anderson, 1899 (Decapoda,) Tj ETQq1 1 0.784314 rgBT /Over	0.3	1
124	Primary osteogenic sarcoma of the ethmoid sinus: A case report. Auris Nasus Larynx, 2005, 32, 411-413.	1.2	11
125	Pkd1 regulates immortalized proliferation of renal tubular epithelial cells through p53 induction and JNK activation. Journal of Clinical Investigation, 2005, 115, 910-918.	8.2	112
126	Spectrum of Germ Cell Tumors: From Head to Toe. Radiographics, 2004, 24, 387-404.	3.3	174

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127	First record of Metapenaeopsis Sibogae (De Man, 1907) (Decapoda, Penaeidae) from Japanese Waters. Crustaceana, 2004, 77, 333-339.	0.3	8
128	The classification of glomerulonephritis in systemic lupus erythematosus revisited. Kidney International, 2004, 65, 521-530.	5.2	1,272
129	Staphylococcus aureus cell envelope antigen is a new candidate for the induction of IgA nephropathy. Kidney International, 2004, 66, 121-132.	5.2	102
130	Cationic charge-preferential IgG reabsorption in the renal proximal tubules. Kidney International, 2004, 66, 1556-1560.	5.2	11
131	Treatment with the purine synthesis inhibitor mizoribine for ANCA-associated renal vasculitis. American Journal of Kidney Diseases, 2004, 44, 57-63.	1.9	37
132	The Classification of Glomerulonephritis in Systemic Lupus Erythematosus Revisited. Journal of the American Society of Nephrology: JASN, 2004, 15, 241-250.	6.1	1,626
133	In situ expression of connective tissue growth factor in human crescentic glomerulonephritis. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2004, 444, 257-263.	2.8	21
134	Podocytes, parietal cells, and glomerular pathology: the role of cell cycle proteins. Pediatric Nephrology, 2003, 18, 3-8.	1.7	24
135	Localization of intercellular adherens junction protein p120 catenin during podocyte differentiation. Anatomy and Embryology, 2003, 206, 175-184.	1.5	16
136	Simultaneous development of lymphoplasmacytic lymphoma and diffuse large B-cell lymphomaÂ-Âanalyses of the clonal relatedness by sequencing CDR3 in immunoglobulin heavy chain genes. European Journal of Haematology, 2003, 70, 119-124.	2.2	11
137	Variable expression of podocyte-related markers in the glomeruloid bodies in Wilms tumor. Pathology International, 2003, 53, 596-601.	1.3	8
138	Pathogenesis of human renal dysplasia: An alternative scenario to the major theories. Pediatrics International, 2003, 45, 605-609.	0.5	25
139	Connective Tissue Growth Factor Participates in Scar Formation of Crescentic Glomerulonephritis. Laboratory Investigation, 2003, 83, 1615-1625.	3.7	50
140	Cervical Intramedullary Gliofibroma in a Child. Pediatric Neurosurgery, 2002, 36, 105-110.	0.7	15
141	Pathogenesis of dysplastic kidney associated with urinary tract obstruction in utero. Nephrology Dialysis Transplantation, 2002, 17, 37-38.	0.7	26
142	Identity of Alpheus digitalis De Haan, 1844 and description of a new closely related species from the northwestern Pacific (Decapoda: Caridea: Alpheidae). Crustacean Research, 2002, 31, 73-90.	0.8	4
143	High Glucose Induced VEGF Expression via PKC and ERK in Glomerular Podocytes. Biochemical and Biophysical Research Communications, 2002, 290, 177-184.	2.1	115
144	Adenomyomatosis with Marked Subserosal Fibrosis and Lipomatosis of the Gallbladder: Mural Stratification Demonstrated with MR. Magnetic Resonance in Medical Sciences, 2002, 1, 125-128.	2.0	6

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145	Phenotypic changes and cell cycle activation in early tubulointerstitial injury of rat adriamycin nephrosis. Pathology International, 2002, 52, 214-223.	1.3	10
146	Role of cell cycle molecules in the pathophysiology of glomerular epithelial cells. Microscopy Research and Technique, 2002, 57, 203-207.	2.2	7
147	Glomerular differentiation in p27 and p57 double-mutant metanephroi. Anatomy and Embryology, 2002, 206, 31-36.	1.5	7
148	A decidualized endometrial cyst in a pregnant woman: a case observed with a steady-state free precession imaging sequence. Magnetic Resonance Imaging, 2002, 20, 301-304.	1.8	28
149	Pulmonary hypoplasia with an unusual prenatal history. Pediatric Pulmonology, 2002, 34, 218-221.	2.0	3
150	Podocyte Injury Promotes Progressive Nephropathy in Zucker Diabetic Fatty Rats. Laboratory Investigation, 2002, 82, 25-35.	3.7	189
151	Localization of Shiga toxins of enterohaemorrhagic Escherichia coli in kidneys of paediatric and geriatric patients with fatal haemolytic uraemic syndrome. Microbial Pathogenesis, 2001, 31, 59-67.	2.9	52
152	EFFECT OF PRE- AND POSTOPERATIVE PLASMAPHERESIS ON POSTTRANSPLANT RECURRENCE OF FOCAL SEGMENTAL GLOMERULOSCLEROSIS IN CHILDREN. Transplantation, 2001, 71, 628-633.	1.0	108
153	Heterogeneity of Prognosis in Adult IgA Nephropathy, Especially with Mild Proteinuria or Mild Histological Features.. Internal Medicine, 2001, 40, 697-702.	0.7	24
154	Initial pathological events in renal dysplasia with urinary tract obstruction in utero. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2001, 439, 560-570.	2.8	34
155	Glomerular crescents in renal amyloidosis: An epiphenomenon or distinct pathology?. Pathology International, 2001, 51, 179-186.	1.3	38
156	Three autopsied cases of cystic fibrosis in Japan. Pathology International, 2001, 51, 467-472.	1.3	2
157	Induction of Collecting Duct Morphogenesis In Vitro by Heparin-Binding Epidermal Growth Factor-Like Growth Factor. Journal of the American Society of Nephrology: JASN, 2001, 12, 964-972.	6.1	21
158	Podocyte cell cycle regulation and proliferation in collapsing glomerulopathies. Kidney International, 2000, 58, 137-143.	5.2	171
159	Phenotypic Characteristics and Cyclin-Dependent Kinase Inhibitors Repression in Hyperplastic Epithelial Pathology in Idiopathic Focal Segmental Glomerulosclerosis. Laboratory Investigation, 2000, 80, 869-880.	3.7	63
160	Carcinosarcoma with rhabdoid features of the urinary bladder in a 2â€yearâ€old girl: Possible histogenesis of stem cell origin. Pathology International, 2000, 50, 973-978.	1.3	20
161	Corresponding distributions of increased endothelinâ€B receptor expression and increased endothelinâ€1 expression in the aorta of apolipoprotein Eâ€deficient mice with advanced atherosclerosis. Pathology International, 2000, 50, 929-936.	1.3	44
162	Pathogenesis of glomerulosclerosis: role of epithelial interactions. Clinical and Experimental Nephrology, 2000, 4, 173-181.	1.6	4

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163	Heterotopic thyroid tissue at the porta hepatis in a fetus with trisomy 18. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2000, 436, 498.	2.8	13
164	A NEW SPECIES OF ALPHEUS (DECAPODA, CARIDEA, ALPHEIDAE) FROM KAGOSHIMA BAY, JAPAN. Crustaceana, 2000, 73, 1109-1120.	0.3	4
165	Prune-belly syndrome diagnosed at 14 weeks' gestation with severe urethral obstruction but normal kidneys. Pediatric Nephrology, 1999, 13, 135-137.	1.7	22
166	Glomerular expression of cell-cycle-regulatory proteins in human crescentic glomerulonephritis. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 1999, 435, 422-427.	2.8	45
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