

Riccardo Magistroni

List of Publications by Year in descending order

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Version: 2024-02-01

73
papers

5,720
citations

172207

29
h-index

114278

63
g-index

83
all docs

83
docs citations

83
times ranked

6378
citing authors

#	ARTICLE	IF	CITATIONS
1	Tolvaptan in Patients with Autosomal Dominant Polycystic Kidney Disease. <i>New England Journal of Medicine</i> , 2012, 367, 2407-2418.	13.9	1,267
2	Unified Criteria for Ultrasonographic Diagnosis of ADPKD. <i>Journal of the American Society of Nephrology: JASN</i> , 2009, 20, 205-212.	3.0	590
3	Genome-wide association study identifies susceptibility loci for IgA nephropathy. <i>Nature Genetics</i> , 2011, 43, 321-327.	9.4	528
4	Discovery of new risk loci for IgA nephropathy implicates genes involved in immunity against intestinal pathogens. <i>Nature Genetics</i> , 2014, 46, 1187-1196.	9.4	505
5	Validation of the Oxford classification of IgA nephropathy in cohorts with different presentations and treatments. <i>Kidney International</i> , 2014, 86, 828-836.	2.6	373
6	New developments in the genetics, pathogenesis, and therapy of IgA nephropathy. <i>Kidney International</i> , 2015, 88, 974-989.	2.6	211
7	The MEST score provides earlier risk prediction in IgA nephropathy. <i>Kidney International</i> , 2016, 89, 167-175.	2.6	190
8	Corticosteroids in IgA Nephropathy. <i>Journal of the American Society of Nephrology: JASN</i> , 2015, 26, 2248-2258.	3.0	187
9	Genotype-Renal Function Correlation in Type 2 Autosomal Dominant Polycystic Kidney Disease. <i>Journal of the American Society of Nephrology: JASN</i> , 2003, 14, 1164-1174.	3.0	129
10	Coexistence of Different Circulating Anti-Podocyte Antibodies in Membranous Nephropathy. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2012, 7, 1394-1400.	2.2	123
11	Genome-Wide Linkage Scan of a Large Family with IgA Nephropathy Localizes a Novel Susceptibility Locus to Chromosome 2q36. <i>Journal of the American Society of Nephrology: JASN</i> , 2007, 18, 2408-2415.	3.0	112
12	Rituximab or Cyclophosphamide in the Treatment of Membranous Nephropathy: The RI-CYCLO Randomized Trial. <i>Journal of the American Society of Nephrology: JASN</i> , 2021, 32, 972-982.	3.0	103
13	Direct characterization of target podocyte antigens and auto-antibodies in human membranous glomerulonephritis: Alfa-enolase and borderline antigens. <i>Journal of Proteomics</i> , 2011, 74, 2008-2017.	1.2	101
14	Progressive Loss of Renal Function Is an Age-Dependent Heritable Trait in Type 1 Autosomal Dominant Polycystic Kidney Disease. <i>Journal of the American Society of Nephrology: JASN</i> , 2005, 16, 755-762.	3.0	84
15	Urine Proteome Analysis May Allow Noninvasive Differential Diagnosis of Diabetic Nephropathy. <i>Diabetes Care</i> , 2010, 33, 2409-2415.	4.3	83
16	COVID-19 pneumonia in a kidney transplant recipient successfully treated with tocilizumab and hydroxychloroquine. <i>American Journal of Transplantation</i> , 2020, 20, 1902-1906.	2.6	81
17	Hypokalemia in Patients with COVID-19. <i>Clinical and Experimental Nephrology</i> , 2021, 25, 401-409.	0.7	78
18	Tonsillectomy in a European Cohort of 1,147 Patients with IgA Nephropathy. <i>Nephron</i> , 2016, 132, 15-24.	0.9	60

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19	Genetic Variation of DKK3 May Modify Renal Disease Severity in ADPKD. <i>Journal of the American Society of Nephrology: JASN</i> , 2010, 21, 1510-1520.	3.0	59
20	The prevalence of autosomal dominant polycystic kidney disease (ADPKD): A meta-analysis of European literature and prevalence evaluation in the Italian province of Modena suggest that ADPKD is a rare and underdiagnosed condition. <i>PLoS ONE</i> , 2018, 13, e0190430.	1.1	57
21	Precocious activation of genes of the renin-angiotensin system and the fibrogenic cascade in IgA glomerulonephritis. <i>Kidney International</i> , 2003, 64, 149-159.	2.6	56
22	Pregnancy and Progression of IgA Nephropathy: Results of an Italian Multicenter Study. <i>American Journal of Kidney Diseases</i> , 2010, 56, 506-512.	2.1	53
23	A Review of the Imaging Techniques for Measuring Kidney and Cyst Volume in Establishing Autosomal Dominant Polycystic Kidney Disease Progression. <i>American Journal of Nephrology</i> , 2018, 48, 67-78.	1.4	51
24	Oxalate Nephropathy Caused by Excessive Vitamin C Administration in 2 Patients With COVID-19. <i>Kidney International Reports</i> , 2020, 5, 1815-1822.	0.4	45
25	Influence of ACE I/D gene polymorphism in the progression of renal failure in autosomal dominant polycystic kidney disease: a meta-analysis. <i>Nephrology Dialysis Transplantation</i> , 2006, 21, 3155-3163.	0.4	35
26	Improving treatment decisions using personalized risk assessment from the International IgA Nephropathy Prediction Tool. <i>Kidney International</i> , 2020, 98, 1009-1019.	2.6	35
27	Evaluation of the Classification Accuracy of the Kidney Biopsy Direct Immunofluorescence through Convolutional Neural Networks. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2020, 15, 1445-1454.	2.2	34
28	Lipoprotein glomerulopathy treated with LDL-apheresis (Heparin-induced Extracorporeal Lipoprotein) Tj ETQq0 0 0 ggBT /Overlock 10 Tf 0,4 32	0.4	32
29	Proteomic analysis of urine from proteinuric patients shows a proteolytic activity directed against albumin. <i>Nephrology Dialysis Transplantation</i> , 2009, 24, 1672-1681.	0.4	30
30	Deciphering Variability of PKD1 and PKD2 in an Italian Cohort of 643 Patients with Autosomal Dominant Polycystic Kidney Disease (ADPKD). <i>Scientific Reports</i> , 2016, 6, 30850.	1.6	28
31	Defective glycolysis and the use of 2-deoxy-d-glucose in polycystic kidney disease: from animal models to humans. <i>Journal of Nephrology</i> , 2017, 30, 511-519.	0.9	28
32	Incidence, risk factors and outcome of acute kidney injury (AKI) in patients with COVID-19. <i>Clinical and Experimental Nephrology</i> , 2021, 25, 1203-1214.	0.7	27
33	Acid base disorders in patients with COVID-19. <i>International Urology and Nephrology</i> , 2022, 54, 405-410.	0.6	26
34	The frail world of haemodialysis patients in the COVID-19 pandemic era: a systematic scoping review. <i>Journal of Nephrology</i> , 2021, 34, 1387-1403.	0.9	24
35	Can tonsillectomy modify the innate and adaptive immunity pathways involved in IgA nephropathy?. <i>Journal of Nephrology</i> , 2015, 28, 51-58.	0.9	23
36	Severe acute respiratory SARS-CoV-2 infection in dialysis patients in northern Italy: a single-centre experience. <i>CKJ: Clinical Kidney Journal</i> , 2020, 13, 334-339.	1.4	19

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37	The Role of the Renin-Angiotensin System in Severe Acute Respiratory Syndrome-CoV-2 Infection. <i>Blood Purification</i> , 2021, 50, 263-267.	0.9	17
38	ADPKD current management and ongoing trials. <i>Journal of Nephrology</i> , 2020, 33, 223-237.	0.9	16
39	OUP accepted manuscript. CKJ: <i>Clinical Kidney Journal</i> , 2020, 13, 265-268.	1.4	16
40	Identification and Characterization of a New Autoimmune Protein in Membranous Nephropathy by Immunoscreening of a Renal cDNA Library. <i>PLoS ONE</i> , 2012, 7, e48845.	1.1	14
41	Twenty-four-hour serum creatinine variation is associated with poor outcome in the novel coronavirus disease 2019 (COVID-19) patients. <i>Kidney Research and Clinical Practice</i> , 2021, 40, 231-240.	0.9	14
42	A pilot study to evaluate tolerability and safety of a modified Atkins diet in ADPKD patients. <i>PharmaNutrition</i> , 2019, 9, 100154.	0.8	13
43	Lipoprotein Glomerulopathy Associated with a Mutation in Apolipoprotein E. <i>Clinical Medicine Insights: Case Reports</i> , 2013, 6, CCRRep.S12209.	0.3	12
44	Reliability of Total Renal Volume Computation in Polycystic Kidney Disease From Magnetic Resonance Imaging. <i>Academic Radiology</i> , 2015, 22, 1376-1384.	1.3	12
45	Comparison of Total Kidney Volume Quantification Methods in Autosomal Dominant Polycystic Disease for a Comprehensive Disease Assessment. <i>American Journal of Nephrology</i> , 2017, 45, 373-379.	1.4	12
46	TRPP2 dysfunction decreases ATP-evoked calcium, induces cell aggregation and stimulates proliferation in T lymphocytes. <i>BMC Nephrology</i> , 2019, 20, 355.	0.8	12
47	Rituximab versus steroids and cyclophosphamide for the treatment of primary membranous nephropathy: protocol of a pilot randomised controlled trial. <i>BMJ Open</i> , 2019, 9, e029232.	0.8	11
48	GREASE II. A phase II randomized, 12-month, parallel-group, superiority study to evaluate the efficacy of a Modified Atkins Diet in Autosomal Dominant Polycystic Kidney Disease patients. <i>PharmaNutrition</i> , 2020, 13, 100206.	0.8	10
49	Long-term effects of COVID-19 in a patient on maintenance dialysis. <i>Hemodialysis International</i> , 2020, 24, E50-E54.	0.4	9
50	One-year persistence of neutralizing SARS-CoV-2 antibodies in dialysis patients recovered from COVID-19. <i>Hemodialysis International</i> , 2021, 25, E53-E56.	0.4	8
51	Clinical Predictors of Nondiabetic Kidney Disease in Patients with Diabetes: A Single-Center Study. <i>International Journal of Nephrology</i> , 2021, 2021, 1-7.	0.7	7
52	A validated model of disease progression in IgA nephropathy. <i>Journal of Nephrology</i> , 2006, 19, 32-40.	0.9	7
53	Artificial intelligence in glomerular diseases. <i>Pediatric Nephrology</i> , 2022, 37, 2533-2545.	0.9	7
54	Confidence Calibration for Deep Renal Biopsy Immunofluorescence Image Classification. , 2021, , .		5

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55	Interstitial fluid obtained from kidney biopsy as new source of renal biomarkers. Journal of Nephrology, 2011, 24, 329-337.	0.9	4
56	Seroconversion after COVID-19 vaccine in a dialysis patient on immunosuppressants. CKJ: Clinical Kidney Journal, 2021, 14, 1983-1984.	1.4	4
57	Acid base disorders in patients with COVID-19. International Urology and Nephrology, 2021, , .	0.6	4
58	Ethical challenges in managing unvaccinated patients receiving chronic in-centre haemodialysis. CKJ: Clinical Kidney Journal, 2022, 15, 615-617.	1.4	4
59	Which criteria should we use to end isolation in hemodialysis patients with COVID-19?. CKJ: Clinical Kidney Journal, 0, , .	1.4	3
60	Comment on the Paper: 'Novel Approach to Estimate Kidney and Cyst Volumes Using Mid-Slice Magnetic Resonance Images in Polycystic Kidney Disease'. American Journal of Nephrology, 2014, 39, 163-164.	1.4	2
61	Geometry-independent assessment of renal volume in polycystic kidney disease from magnetic resonance imaging. , 2015, 2015, 3081-4.		2
62	Tolvaptan: Clinical Evidence for Slowing the Progression of Autosomal Dominant Polycystic Kidney Disease. Giornale De Tecniche Nefrologiche & Dialitiche, 2017, 29, 80-84.	0.1	2
63	Clinical Presentation, Renal Histopathological Findings, and Outcome in Patients with Monoclonal Gammopathy and Kidney Disease. International Journal of Nephrology, 2021, 2021, 1-9.	0.7	2
64	Response letter to the Editorial: "Ketogenic diet in ADPKD patients". PharmaNutrition, 2021, 16, 100268.	0.8	2
65	Epidermal growth factor receptor polymorphism and autosomal dominant polycystic kidney disease. Journal of Nephrology, 2003, 16, 110-5.	0.9	2
66	Hybrid dialysis: a promising strategy to reduce hospital access during the SARS-CoV-2 pandemic. BMJ Case Reports, 2020, 13, e236411.	0.2	1
67	Methicillin-Resistant <i>Staphylococcus aureus</i> Peritonitis due to Hematogenous Dissemination from Central Venous Catheter in a Maintenance Dialysis Patient. Case Reports in Nephrology and Dialysis, 2021, 11, 281-285.	0.3	1
68	Monoclonal B lymphocytosis in a kidney transplant recipient. BMJ Case Reports, 2021, 14, e242889.	0.2	0
69	Advances in Genetics of Immunoglobulin A Nephropathy. , 2016, , 19-42.		0
70	AKI in hospitalized patients with COVID-19: a single-center experience. Giornale Italiano Di Nefrologia: Organo Ufficiale Della Società Italiana Di Nefrologia, 2021, 38, .	0.3	0
71	Immunosuppressive therapy reduction and early post-infection graft function in kidney transplant recipients with COVID-19.. Giornale Italiano Di Nefrologia: Organo Ufficiale Della Società Italiana Di Nefrologia, 2021, 38, .	0.3	0
72	Reactogenicity of COVID-19 vaccine in hemodialysis patients: a single-center retrospective study.. Giornale Italiano Di Nefrologia: Organo Ufficiale Della Società Italiana Di Nefrologia, 2022, 39, .	0.3	0

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73	MO137: Sarcoidosis in a Living Kidney Donor Candidate: Case Report and Review of the Literature. Nephrology Dialysis Transplantation, 2022, 37, .	0.4	0