Jose Ballarin

List of Publications by Year
in descending order

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Version: 2024-02-01

66 papers	2,289 citations	201674 27 h-index	223800 46 g-index
67 all docs	67 docs citations	67 times ranked	2895 citing authors

#	Article	IF	Citations
1	Pregnancy in Pateints With Exstrophy-Epispadias Complex: Are Higher Rates of Complications and Spontaneous Abortion Inevitable?. Urology, 2021, 154, 326-332.	1.0	2
2	Genetic Variants Associated with Chronic Kidney Disease in a Spanish Population. Scientific Reports, 2020, 10, 144.	3.3	29
3	The genetic architecture of membranous nephropathy and its potential to improve non-invasive diagnosis. Nature Communications, 2020, 11, 1600.	12.8	120
4	IgA Nephropathy in Elderly Patients. Clinical Journal of the American Society of Nephrology: CJASN, 2019, 14, 1183-1192.	4.5	18
5	Three cases of monoclonal gammopathy of renal significance after kidney transplantation. De novo C3 glomerulopathy. Nefrologia, 2019, 39, 198-201.	0.4	1
6	MYH9 Associated nephropathy. Nefrologia, 2019, 39, 133-140.	0.4	6
7	Mycophenolate mofetil versus cyclophosphamide for remission induction in ANCA-associated vasculitis: a randomised, non-inferiority trial. Annals of the Rheumatic Diseases, 2019, 78, 399-405.	0.9	165
8	NefropatÃa asociada a mutaciÃ³n del gen MYH9. Nefrologia, 2019, 39, 133-140.	0.4	9
9	Autosomal Dominant Polycystic Kidney Disease: Clinical Assessment of Rapid Progression. American Journal of Nephrology, 2018, 48, 308-317.	3.1	15
10	A kidney-disease gene panel allows a comprehensive genetic diagnosis of cystic andÂglomerular inherited kidney diseases. Kidney International, 2018, 94, 363-371.	5.2	109
11	Autosomal Dominant Tubulointerstitial Kidney Disease: Clinical Presentation of Patients With ADTKD-UMOD and ADTKD-MUC1. American Journal of Kidney Diseases, 2018, 72, 411-418.	1.9	42
12	Transethnic, Genome-Wide Analysis Reveals Immune-Related Risk Alleles and Phenotypic Correlates in Pediatric Steroid-Sensitive Nephrotic Syndrome. Journal of the American Society of Nephrology: JASN, 2018, 29, 2000-2013.	6.1	72
13	Contribution of theTTC21Bgene to glomerular and cystic kidney diseases. Nephrology Dialysis Transplantation, 2017, 32, gfv453.	0.7	26
14	Vitamin E-coated dialysis membranes reduce the levels of oxidative genetic damage in hemodialysis patients. Mutation Research - Genetic Toxicology and Environmental Mutagenesis, 2017, 815, 16-21.	1.7	19
15	Efficacy of mycophenolate treatment in adults with steroid-dependent/frequently relapsing idiopathic nephrotic syndrome. CKJ: Clinical Kidney Journal, 2017, 10, 632-638.	2.9	4
16	Rituximab for Steroid-Dependent or Frequently Relapsing Idiopathic Nephrotic Syndrome in Adults: A Retrospective, Multicenter Study in Spain. BioDrugs, 2017, 31, 239-249.	4.6	14
17	Renal angiomyolipoma bleeding in a patient with TSC2/PKD1 contiguous gene syndrome after 17 years of renal replacement therapy. Nefrologia, 2017, 37, 87-92.	0.4	10
18	Markers of endothelial damage in patients with chronic kidney disease on hemodialysis. American Journal of Physiology - Renal Physiology, 2017, 312, F673-F681.	2.7	33

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19	DNA damage in kidney transplant patients. Role of organ origin. Environmental and Molecular Mutagenesis, 2017, 58, 712-718.	2.2	5
20	Sangrado de angiomiolipoma renal en paciente con sÃ¬ndrome de genes contiguos (TSC2/PKD1) tras 17 aÃ±os de tratamiento renal sustitutivo. Nefrologia, 2017, 37, 87-92.	0.4	12
21	Cardiovascular calcifications in chronic kidney disease: Potential therapeutic implications. Nefrologia, 2016, 36, 597-608.	0.4	15
22	Detection of cardiovascular calcifications: Is it a useful tool for nephrologists?. Nefrologia, 2016, 36, 587-596.	0.4	7
23	Calcificaciones cardiovasculares en la enfermedad renal crÃ³nica: Potenciales implicaciones terapÃ©uticas. Nefrologia, 2016, 36, 597-608.	0.4	18
24	Levels of DNA damage in peripheral blood lymphocytes of patients undergoing standard hemodialysis vs on-line hemodiafiltration: A comet assay investigation. Mutation Research - Genetic Toxicology and Environmental Mutagenesis, 2016, 808, 1-7.	1.7	5
25	DetecciÃ³n de las calcificaciones cardiovasculares: Â¿una herramienta Ãºtil para el nefrÃ³logo?. Nefrologia, 2016, 36, 587-596.	0.4	12
26	Unfermented grape juice reduce genomic damage on patients undergoing hemodialysis. Food and Chemical Toxicology, 2016, 92, 1-7.	3.6	22
27	Genetic damage in patients moving from hemodialysis to online hemodiafiltration. Mutagenesis, 2016, 31, 131-135.	2.6	9
28	Clinical and Practical Use of Calcimimetics in Dialysis Patients With Secondary Hyperparathyroidism. Clinical Journal of the American Society of Nephrology: CJASN, 2016, 11, 161-174.	4.5	52
29	Radiosensitivity in patients suffering from chronic kidney disease. International Journal of Radiation Biology, 2015, 91, 172-178.	1.8	11
30	Insight into response to mTOR inhibition when PKD1 and TSC2 are mutated. BMC Medical Genetics, 2015, 16, 39.	2.1	15
31	Targeted next-generation sequencing in steroid-resistant nephrotic syndrome: mutations in multiple glomerular genes may influence disease severity. European Journal of Human Genetics, 2015, 23, 1192-1199.	2.8	72
32	Genomic damage as a biomarker of chronic kidney disease status. Environmental and Molecular Mutagenesis, 2015, 56, 301-312.	2.2	23
33	HLA-DQA1 and PLA2R1 Polymorphisms and Risk of Idiopathic Membranous Nephropathy. Clinical Journal of the American Society of Nephrology: CJASN, 2014, 9, 335-343.	4.5	72
34	Cost-effective PKHD1 genetic testing for autosomal recessive polycystic kidney disease. Pediatric Nephrology, 2014, 29, 223-234.	1.7	14
35	Diagnosis of autosomal dominant polycystic kidney disease using efficient PKD1 and PKD2 targeted nextâ€generation sequencing. Molecular Genetics & Genomic Medicine, 2014, 2, 412-421.	1.2	67
36	Time in hemodialysis modulates the levels of genetic damage in hemodialysis patients. Environmental and Molecular Mutagenesis, 2014, 55, 363-368.	2.2	9

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37	Renal replacement therapy in ADPKD patients: a 25-year survey based on the Catalan registry. BMC Nephrology, 2013, 14, 186.	1.8	33
38	Longâ€term outcome of antineutrophil cytoplasmic antibodyâ€associated small vessel vasculitis after renal transplantation. Clinical Transplantation, 2013, 27, 338-347.	1.6	37
39	Recombinant PTH associated with hypercalcaemia and renal failure: TableÂ1.. CKJ: Clinical Kidney Journal, 2013, 6, 93-95.	2.9	8
40	Lanthanum carbonate for the control of hyperphosphatemia in chronic renal failure patients: a new oral powder formulation – safety, efficacy, and patient adherence. Patient Preference and Adherence, 2013, 7, 1147.	1.8	10
41	Genomic damage as an independent predictor marker of mortality in hemodialysis patients. Clinical Nephrology, 2013, 80, 81-87.	0.7	10
42	Successful Transplantation of Organs From a Donor With Postneurosurgical Meningitis Caused by Escherichia coli. Transplantation, 2012, 93, e11-e13.	1.0	11
43	Assessing the effectiveness of rapamycin on angiomyolipoma in tuberous sclerosis: a two years trial. Orphanet Journal of Rare Diseases, 2012, 7, 87.	2.7	41
44	Genomic instability in chronic renal failure patients. Environmental and Molecular Mutagenesis, 2012, 53, 343-349.	2.2	21
45	Angioplasty and stent treatment of transplant renal artery stenosis. Nefrologia, 2012, 32, 455-8.	0.4	7
46	Acute renal failure associated to paroxysmal nocturnal haemoglobinuria leads to intratubular haemosiderin accumulation and CD163 expression. Nephrology Dialysis Transplantation, 2011, 26, 3408-3411.	0.7	57
47	Low-dose sirolimus combined with angiotensin-converting enzyme inhibitor and statin stabilizes renal function and reduces glomerular proliferation in poor prognosis IgA nephropathy. Nephrology Dialysis Transplantation, 2011, 26, 3596-3602.	0.7	28
48	Clinical Value ofNPHS2Analysis in Early- and Adult-Onset Steroid-Resistant Nephrotic Syndrome. Clinical Journal of the American Society of Nephrology: CJASN, 2011, 6, 344-354.	4.5	65
49	Clinical Utility of Genetic Testing in Children and Adults with Steroid-Resistant Nephrotic Syndrome. Clinical Journal of the American Society of Nephrology: CJASN, 2011, 6, 1139-1148.	4.5	189
50	Genetic damage in chronic renal failure patients is associated with the glomerular filtration rate index. Mutagenesis, 2010, 25, 603-608.	2.6	28
51	Oxidative DNA damage in chronic renal failure patients. Nephrology Dialysis Transplantation, 2010, 25, 879-885.	0.7	51
52	Calcimimetics in the Chronic Kidney Disease-Mineral and Bone Disorder. International Journal of Artificial Organs, 2009, 32, 108-121.	1.4	15
53	Nephrin mutations cause childhood- and adult-onset focal segmental glomerulosclerosis. Kidney International, 2009, 76, 1268-1276.	5.2	111
54	TRPC6 mutational analysis in a large cohort of patients with focal segmental glomerulosclerosis. Nephrology Dialysis Transplantation, 2009, 24, 3089-3096.	0.7	99

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55	UPDATE ON THE TREATMENT OF CHRONIC KIDNEY DISEASEâ€MINERAL AND BONE DISORDER. Journal of Renal Care, 2009, 35, 19-27.	1.2	12
56	Prevalence of Cysts in Seminal Tract and Abnormal Semen Parameters in Patients with Autosomal Dominant Polycystic Kidney Disease. Clinical Journal of the American Society of Nephrology: CJASN, 2008, 3, 790-793.	4.5	57
57	Study of candidate genes affecting the progression of renal disease in autosomal dominant polycystic kidney disease type 1. Nephrology Dialysis Transplantation, 2007, 22, 1567-1577.	0.7	25
58	Treatment of idiopathic membranous nephropathy with the combination of steroids, tacrolimus and mycophenolate mofetil: results of a pilot study. Nephrology Dialysis Transplantation, 2007, 22, 3196-3201.	0.7	40
59	Importancia de la imagen en los sÃndromes renopulmonares. Dialisis Y Trasplante, 2007, 28, 126-127.	0.4	0
60	Genetic Testing for X-Linked Alport Syndrome by Direct Sequencing of COL4A5 cDNA From Hair Root RNA Samples. American Journal of Kidney Diseases, 2007, 50, 257.e1-257.e14.	1.9	27
61	Expression of concentrative nucleoside transporters SLC28 (CNT1, CNT2, and CNT3) along the rat nephron: Effect of diabetes. Kidney International, 2005, 68, 665-672.	5.2	41
62	Sodium retention in cirrhotic rats is associated with increased renal abundance of sodium transporter proteins. Kidney International, 2005, 67, 622-630.	5.2	29
63	Collagen type IV (Â3-Â4) nephropathy: from isolated haematuria to renal failure. Nephrology Dialysis Transplantation, 2004, 19, 2429-2432.	0.7	39
64	Diagnostic, prognostic and pathogenic value of the direct immunofluorescence test in cutaneous leukocytoclastic vasculitis. International Journal of Dermatology, 2004, 43, 19-26.	1.0	42
65	Lack of effect of clinical doses of cyclosporin A on erythrocyte Na+/K+-ATPase activity. Clinical Science, 1999, 97, 283.	4.3	1
66	Effects of cyclosporine A on Na,K-ATPase expression in the renal epithelial cell line NBL-1. Kidney International, 1996, 50, 1483-1489.	5.2	19