Aude Servais

List of Publications by Citations

39 papers	1,230 citations	17 h-index	35 g-index
46 ext. papers	1,505 ext. citations	5.1 avg, IF	3.46 L-index

#	Paper	IF	Citations
39	Acquired and genetic complement abnormalities play a critical role in dense deposit disease and other C3 glomerulopathies. *Kidney International*, **2012**, 82, 454-64	9.9	360
38	Primary glomerulonephritis with isolated C3 deposits: a new entity which shares common genetic risk factors with haemolytic uraemic syndrome. *Journal of Medical Genetics*, **2007**, 44, 193-9	5.8	223
37	Nephropathic cystinosis: an international consensus document. *Nephrology Dialysis Transplantation*, **2014**, 29 Suppl 4, iv87-94	4.3	116
36	Update on Lysinuric Protein Intolerance, a Multi-faceted Disease Retrospective cohort analysis from birth to adulthood. *Orphanet Journal of Rare Diseases*, **2017**, 12, 3	4.2	55
35	Late-onset nephropathic cystinosis: clinical presentation, outcome, and genotyping. *Clinical Journal of the American Society of Nephrology: CJASN*, **2008**, 3, 27-35	6.9	52
34	Increased risk of solid renal tumors in lithium-treated patients. *Kidney International*, **2014**, 86, 184-90	9.9	45
33	Controversies and research agenda in nephropathic cystinosis: conclusions from a "Kidney Disease: Improving Global Outcomes" (KDIGO) Controversies Conference. *Kidney International*, **2016**, 89, 1192-203	9.9	37
32	Heterogeneous pattern of renal disease associated with homozygous factor H deficiency. *Human Pathology*, **2011**, 42, 1305-11	3.7	36
31	Quantification of interstitial fibrosis by image analysis on routine renal biopsy in patients receiving cyclosporine. *Transplantation*, **2007**, 84, 1595-601	1.8	36
30	Identification of genetic causes for sporadic steroid-resistant nephrotic syndrome in adults. *Kidney International*, **2018**, 94, 1013-1022	9.9	31
29	Management of bone disease in cystinosis: Statement from an international conference. *Journal of Inherited Metabolic Disease*, **2019**, 42, 1019-1029	5.4	28
28	Phenotype, treatment practice and outcome in the cobalamin-dependent remethylation disorders and MTHFR deficiency: Data from the E-HOD registry. *Journal of Inherited Metabolic Disease*, **2019**, 42, 333-352	5.4	28
27	C3 glomerulopathy. *Contributions To Nephrology*, **2013**, 181, 185-93	1.6	23
26	Cystinuria: clinical practice recommendation. *Kidney International*, **2021**, 99, 48-58	9.9	21
25	Long-term metabolic follow-up and clinical outcome of 35 patients with maple syrup urine disease. *Journal of Inherited Metabolic Disease*, **2017**, 40, 783-792	5.4	17
24	Excellent long-term outcome of renal transplantation in cystinosis patients. *Orphanet Journal of Rare Diseases*, **2015**, 10, 90	4.2	17
23	Statistical color texture descriptors for histological images analysis **2011**,		17

22	Heterogeneous histologic and clinical evolution in 3 cases of dense deposit disease with long-term follow-up. *Human Pathology*, **2014**, 45, 2326-33	3.7	12
21	Renal involvement in lysinuric protein intolerance: contribution of pathology to assessment of heterogeneity of renal lesions. *Human Pathology*, **2017**, 62, 160-169	3.7	10
20	Long-term outcome of methylmalonic aciduria after kidney, liver, or combined liver-kidney transplantation: The French experience. *Journal of Inherited Metabolic Disease*, **2020**, 43, 234-243	5.4	10
19	The European Rare Kidney Disease Registry (ERKReg): objectives, design and initial results. *Orphanet Journal of Rare Diseases*, **2021**, 16, 251	4.2	8
18	An international cohort study spanning five decades assessed outcomes of nephropathic cystinosis. *Kidney International*, **2021**, 100, 1112-1123	9.9	7
17	Central nervous system complications in adult cystinosis patients. *Journal of Inherited Metabolic Disease*, **2020**, 43, 348-356	5.4	7
16	Cystathionine β-synthase deficiency in the E-HOD registry-part I: pyridoxine responsiveness as a determinant of biochemical and clinical phenotype at diagnosis. *Journal of Inherited Metabolic Disease*, **2021**, 44, 677-692	5.4	7
15	Clinical and histological differences between adults and children in new onset IgA nephropathy. *Pediatric Nephrology*, **2020**, 35, 1897-1905	3.2	5
14	Infectious and digestive complications in glycogen storage disease type Ib: Study of a French cohort. *Molecular Genetics and Metabolism Reports*, **2020**, 23, 100581	1.8	4
13	Comparison of Postdonation Kidney Function Between Caucasian Donors and Low-risk APOL1 Genotype Living Kidney Donors of African Ancestry. *Transplantation*, **2018**, 102, e462-e463	1.8	4
12	The authors reply. *Kidney International*, **2014**, 86, 857-8	9.9	3
11	Enteral tube feeding in patients receiving dietary treatment for metabolic diseases: A retrospective analysis in a large French cohort. *Molecular Genetics and Metabolism Reports*, **2021**, 26, 100655	1.8	3
10	Neonatal factors related to survival and intellectual and developmental outcome of patients with early-onset urea cycle disorders. *Molecular Genetics and Metabolism*, **2020**, 130, 110-117	3.7	2
9	Effects of L-Carnitine on Mineral Metabolism in the Multicentre, Randomized, Double Blind, Placebo-Controlled CARNIDIAL Trial. *American Journal of Nephrology*, **2018**, 48, 349-356	4.6	2
8	APOL1 risk genotype in Europe: Data in patients with focal segmental glomerulosclerosis and after renal transplantation. *Nephrologie Et Therapeutique*, **2019**, 15 Suppl 1, S85-S89	0.6	1
7	The Case | Atrophic kidney and ocular abnormalities. *Kidney International*, **2020**, 98, 1059-1060	9.9	1
6	Long-term renal outcome in methylmalonic acidemia in adolescents and adults. *Orphanet Journal of Rare Diseases*, **2021**, 16, 220	4.2	1
5	FDX2 and ISCU Gene Variations Lead to Rhabdomyolysis With Distinct Severity and Iron Regulation.. *Neurology: Genetics*, **2022**, 8, e648	3.8	0

4	Biopsy-proven kidney involvement in hypocomplementemic urticarial vasculitis.. *BMC Nephrology*, **2022**, 23, 67	2.7	0
3	Home Blood Pressure Measurement and Self-Interpretation of Blood Pressure Readings During Pregnancy: Hy-Result e-Health Prospective Study.. *Vascular Health and Risk Management*, **2022**, 18, 277-287	4.4	0
2	Intravenous administration of a branched-chain amino-acid-free solution in children and adults with acute decompensation of maple syrup urine disease: a prospective multicentre observational study.. *Orphanet Journal of Rare Diseases*, **2022**, 17, 202	4.2	0
1	The Case | A 69-year-old man with purpura and acute renal failure. *Kidney International*, **2018**, 94, 435-436	9.9	