Aude Servais

List of Publications by Year
in descending order

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46 papers	1,794 citations	430442 18 h-index	276539 41 g-index
46 all docs	46 docs citations	46 times ranked	1942 citing authors

#	Article	IF	Citations
1	Acquired and genetic complement abnormalities play a critical role in dense deposit disease and other C3 glomerulopathies. Kidney International, 2012, 82, 454-464.	2.6	454
2	Primary glomerulonephritis with isolated C3 deposits: a new entity which shares common genetic risk factors with haemolytic uraemic syndrome. Journal of Medical Genetics, 2006, 44, 193-199.	1.5	259
3	Nephropathic cystinosis: an international consensus document. Nephrology Dialysis Transplantation, 2014, 29, iv87-iv94.	0.4	164
4	Update on Lysinuric Protein Intolerance, a Multi-faceted Disease Retrospective cohort analysis from birth to adulthood. Orphanet Journal of Rare Diseases, 2017, 12, 3.	1.2	78
5	Late-Onset Nephropathic Cystinosis. Clinical Journal of the American Society of Nephrology: CJASN, 2008, 3, 27-35.	2.2	68
6	Increased risk of solid renal tumors in lithium-treated patients. Kidney International, 2014, 86, 184-190.	2.6	62
7	Cystinuria: clinical practice recommendation. Kidney International, 2021, 99, 48-58.	2.6	58
8	Phenotype, treatment practice and outcome in the cobalaminâ€dependent remethylation disorders and MTHFR deficiency: Data from the Eâ€HOD registry. Journal of Inherited Metabolic Disease, 2019, 42, 333-352.	1.7	53
9	Controversies and research agenda in nephropathic cystinosis: conclusions from a â€œKidney Disease: Improving Global Outcomesâ€•(KDIGO) Controversies Conference. Kidney International, 2016, 89, 1192-1203.	2.6	52
10	Identification of genetic causes for sporadic steroid-resistant nephrotic syndrome in adults. Kidney International, 2018, 94, 1013-1022.	2.6	51
11	Quantification of Interstitial Fibrosis by Image Analysis on Routine Renal Biopsy in Patients Receiving Cyclosporine. Transplantation, 2007, 84, 1595-1601.	0.5	44
12	Heterogeneous pattern of renal disease associated with homozygous Factor H deficiency. Human Pathology, 2011, 42, 1305-1311.	1.1	41
13	Management of bone disease in cystinosis: Statement from an international conference. Journal of Inherited Metabolic Disease, 2019, 42, 1019-1029.	1.7	39
14	An international cohort study spanning five decades assessed outcomes of nephropathic cystinosis. Kidney International, 2021, 100, 1112-1123.	2.6	31
15	C3 Glomerulopathy. Contributions To Nephrology, 2013, 181, 185-193.	1.1	28
16	Excellent long-term outcome of renal transplantation in cystinosis patients. Orphanet Journal of Rare Diseases, 2015, 10, 90.	1.2	27
17	The European Rare Kidney Disease Registry (ERKReg): objectives, design and initial results. Orphanet Journal of Rare Diseases, 2021, 16, 251.	1.2	26
18	Longâ€term metabolic followâ€up and clinical outcome of 35 patients with maple syrup urine disease. Journal of Inherited Metabolic Disease, 2017, 40, 783-792.	1.7	25

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19	Longâ€term outcome of methylmalonic aciduria after kidney, liver, or combined liverâ€kidney transplantation: The French experience. Journal of Inherited Metabolic Disease, 2020, 43, 234-243.	1.7	20
20	Cystathionine Î²â€synthase deficiency in the <scp>Eâ€HOD registryâ€part</scp> I: pyridoxine responsiveness as a determinant of biochemical and clinical phenotype at diagnosis. Journal of Inherited Metabolic Disease, 2021, 44, 677-692.	1.7	20
21	Clinical and histological differences between adults and children in new onset IgA nephropathy. Pediatric Nephrology, 2020, 35, 1897-1905.	0.9	20
22	Statistical color texture descriptors for histological images analysis. , 2011, , .		19
23	Renal involvement in lysinuric protein intolerance: contribution of pathology to assessment of heterogeneity of renal lesions. Human Pathology, 2017, 62, 160-169.	1.1	18
24	COVID-19 outbreak in vaccinated patients from a haemodialysis unit: antibody titres as a marker of protection from infection. Nephrology Dialysis Transplantation, 2022, 37, 1357-1365.	0.4	17
25	Heterogeneous histologic and clinical evolution in 3 cases of dense deposit disease with long-term follow-up. Human Pathology, 2014, 45, 2326-2333.	1.1	15
26	Central nervous system complications in adult cystinosis patients. Journal of Inherited Metabolic Disease, 2020, 43, 348-356.	1.7	14
27	Infectious and digestive complications in glycogen storage disease type Ib: Study of a French cohort. Molecular Genetics and Metabolism Reports, 2020, 23, 100581.	0.4	12
28	Expert guidance on the multidisciplinary management of cystinosis in adolescent and adult patients. CKJ: Clinical Kidney Journal, 2022, 15, 1675-1684.	1.4	9
29	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.	1.4	8
30	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.	0.4	7
31	Long-term renal outcome in methylmalonic acidemia in adolescents and adults. Orphanet Journal of Rare Diseases, 2021, 16, 220.	1.2	7
32	Idiopathic nephrotic syndrome relapse following COVID-19 vaccination: a series of 25 cases. CKJ: Clinical Kidney Journal, 2022, 15, 1574-1582.	1.4	7
33	Comparison of Postdonation Kidney Function Between Caucasian Donors and Low-risk APOL1 Genotype Living Kidney Donors of African Ancestry. Transplantation, 2018, 102, e462-e463.	0.5	5
34	Postauthorization safety study of betaine anhydrous. Journal of Inherited Metabolic Disease, 2022, 45, 719-733.	1.7	5
35	The Authors Reply:. Kidney International, 2014, 86, 857-858.	2.6	4
36	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.	0.5	4

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37	FDX2 and ISCU Gene Variations Lead to Rhabdomyolysis With Distinct Severity and Iron Regulation. Neurology: Genetics, 2022, 8, e648.	0.9	4
38	Biopsy-proven kidney involvement in hypocomplementemic urticarial vasculitis. BMC Nephrology, 2022, 23, 67.	0.8	3
39	Central Nervous System Complications in Cystinosis: The Role of Neuroimaging. Cells, 2022, 11, 682.	1.8	3
40	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, Volume 18, 277-287.	1.0	3
41	Pregnancy in cystinosis patients with chronic kidney disease: A European case series. Journal of Inherited Metabolic Disease, 2022, 45, 963-968.	1.7	3
42	APOL1 risk genotype in Europe: Data in patients with focal segmental glomerulosclerosis and after renal transplantation. Nephrologie Et Therapeutique, 2019, 15, S85-S89.	0.2	2
43	The Case | Atrophic kidney and ocular abnormalities. Kidney International, 2020, 98, 1059-1060.	2.6	2
44	Very longâ€term outcomes in 23 patients with <scp>cblA</scp> type methylmalonic acidemia. Journal of Inherited Metabolic Disease, 0, , .	1.7	2
45	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.	1.2	1
46	The Case | A 69-year-old man with purpura and acute renal failure. Kidney International, 2018, 94, 435-436.	2.6	0