

Aude Servais

List of Publications by Year in descending order

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Version: 2024-02-01

46
papers

1,794
citations

430874

18
h-index

276875

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. <i>Kidney International</i> , 2012, 82, 454-464.	5.2	454
2	Primary glomerulonephritis with isolated C3 deposits: a new entity which shares common genetic risk factors with haemolytic uraemic syndrome. <i>Journal of Medical Genetics</i> , 2006, 44, 193-199.	3.2	259
3	Nephropathic cystinosis: an international consensus document. <i>Nephrology Dialysis Transplantation</i> , 2014, 29, iv87-iv94.	0.7	164
4	Update on Lysinuric Protein Intolerance, a Multi-faceted Disease Retrospective cohort analysis from birth to adulthood. <i>Orphanet Journal of Rare Diseases</i> , 2017, 12, 3.	2.7	78
5	Late-Onset Nephropathic Cystinosis. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2008, 3, 27-35.	4.5	68
6	Increased risk of solid renal tumors in lithium-treated patients. <i>Kidney International</i> , 2014, 86, 184-190.	5.2	62
7	Cystinuria: clinical practice recommendation. <i>Kidney International</i> , 2021, 99, 48-58.	5.2	58
8	Phenotype, treatment practice and outcome in the cobalamin-dependent remethylation disorders and MTHFR deficiency: Data from the E&HOD registry. <i>Journal of Inherited Metabolic Disease</i> , 2019, 42, 333-352.	3.6	53
9	Controversies and research agenda in nephropathic cystinosis: conclusions from a "Kidney Disease: Improving Global Outcomes" (KDIGO) Controversies Conference. <i>Kidney International</i> , 2016, 89, 1192-1203.	5.2	52
10	Identification of genetic causes for sporadic steroid-resistant nephrotic syndrome in adults. <i>Kidney International</i> , 2018, 94, 1013-1022.	5.2	51
11	Quantification of Interstitial Fibrosis by Image Analysis on Routine Renal Biopsy in Patients Receiving Cyclosporine. <i>Transplantation</i> , 2007, 84, 1595-1601.	1.0	44
12	Heterogeneous pattern of renal disease associated with homozygous Factor H deficiency. <i>Human Pathology</i> , 2011, 42, 1305-1311.	2.0	41
13	Management of bone disease in cystinosis: Statement from an international conference. <i>Journal of Inherited Metabolic Disease</i> , 2019, 42, 1019-1029.	3.6	39
14	An international cohort study spanning five decades assessed outcomes of nephropathic cystinosis. <i>Kidney International</i> , 2021, 100, 1112-1123.	5.2	31
15	C3 Glomerulopathy. <i>Contributions To Nephrology</i> , 2013, 181, 185-193.	1.1	28
16	Excellent long-term outcome of renal transplantation in cystinosis patients. <i>Orphanet Journal of Rare Diseases</i> , 2015, 10, 90.	2.7	27
17	The European Rare Kidney Disease Registry (ERKReg): objectives, design and initial results. <i>Orphanet Journal of Rare Diseases</i> , 2021, 16, 251.	2.7	26
18	Long-term metabolic follow-up and clinical outcome of 35 patients with maple syrup urine disease. <i>Journal of Inherited Metabolic Disease</i> , 2017, 40, 783-792.	3.6	25

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19	Long-term outcome of methylmalonic aciduria after kidney, liver, or combined liver-kidney transplantation: The French experience. <i>Journal of Inherited Metabolic Disease</i> , 2020, 43, 234-243.	3.6	20
20	Cystathionine Î²-synthase deficiency in the <sc>Eâ€HOD registryâ€part</sc> I: pyridoxine responsiveness as a determinant of biochemical and clinical phenotype at diagnosis. <i>Journal of Inherited Metabolic Disease</i> , 2021, 44, 677-692.	3.6	20
21	Clinical and histological differences between adults and children in new onset IgA nephropathy. <i>Pediatric Nephrology</i> , 2020, 35, 1897-1905.	1.7	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. <i>Human Pathology</i> , 2017, 62, 160-169.	2.0	18
24	COVID-19 outbreak in vaccinated patients from a haemodialysis unit: antibody titres as a marker of protection from infection. <i>Nephrology Dialysis Transplantation</i> , 2022, 37, 1357-1365.	0.7	17
25	Heterogeneous histologic and clinical evolution in 3 cases of dense deposit disease with long-term follow-up. <i>Human Pathology</i> , 2014, 45, 2326-2333.	2.0	15
26	Central nervous system complications in adult cystinosis patients. <i>Journal of Inherited Metabolic Disease</i> , 2020, 43, 348-356.	3.6	14
27	Infectious and digestive complications in glycogen storage disease type Ib: Study of a French cohort. <i>Molecular Genetics and Metabolism Reports</i> , 2020, 23, 100581.	1.1	12
28	Expert guidance on the multidisciplinary management of cystinosis in adolescent and adult patients. <i>CKJ: Clinical Kidney Journal</i> , 2022, 15, 1675-1684.	2.9	9
29	Effects of L-Carnitine on Mineral Metabolism in the Multicentre, Randomized, Double Blind, Placebo-Controlled CARNIDIAL Trial. <i>American Journal of Nephrology</i> , 2018, 48, 349-356.	3.1	8
30	Enteral tube feeding in patients receiving dietary treatment for metabolic diseases: A retrospective analysis in a large French cohort. <i>Molecular Genetics and Metabolism Reports</i> , 2021, 26, 100655.	1.1	7
31	Long-term renal outcome in methylmalonic acidemia in adolescents and adults. <i>Orphanet Journal of Rare Diseases</i> , 2021, 16, 220.	2.7	7
32	Idiopathic nephrotic syndrome relapse following COVID-19 vaccination: a series of 25 cases. <i>CKJ: Clinical Kidney Journal</i> , 2022, 15, 1574-1582.	2.9	7
33	Comparison of Postdonation Kidney Function Between Caucasian Donors and Low-risk APOL1 Genotype Living Kidney Donors of African Ancestry. <i>Transplantation</i> , 2018, 102, e462-e463.	1.0	5
34	Postauthorization safety study of betaine anhydrous. <i>Journal of Inherited Metabolic Disease</i> , 2022, 45, 719-733.	3.6	5
35	The Authors Reply:. <i>Kidney International</i> , 2014, 86, 857-858.	5.2	4
36	Neonatal factors related to survival and intellectual and developmental outcome of patients with early-onset urea cycle disorders. <i>Molecular Genetics and Metabolism</i> , 2020, 130, 110-117.	1.1	4

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37	FDX2 and ISCU Gene Variations Lead to Rhabdomyolysis With Distinct Severity and Iron Regulation. <i>Neurology: Genetics</i> , 2022, 8, e648.	1.9	4
38	Biopsy-proven kidney involvement in hypocomplementemic urticarial vasculitis. <i>BMC Nephrology</i> , 2022, 23, 67.	1.8	3
39	Central Nervous System Complications in Cystinosis: The Role of Neuroimaging. <i>Cells</i> , 2022, 11, 682.	4.1	3
40	Home Blood Pressure Measurement and Self-Interpretation of Blood Pressure Readings During Pregnancy: Hy-Result e-Health Prospective Study. <i>Vascular Health and Risk Management</i> , 2022, Volume 18, 277-287.	2.3	3
41	Pregnancy in cystinosis patients with chronic kidney disease: A European case series. <i>Journal of Inherited Metabolic Disease</i> , 2022, 45, 963-968.	3.6	3
42	APOL1 risk genotype in Europe: Data in patients with focal segmental glomerulosclerosis and after renal transplantation. <i>Nephrologie Et Therapeutique</i> , 2019, 15, S85-S89.	0.5	2
43	The Case Atrophic kidney and ocular abnormalities. <i>Kidney International</i> , 2020, 98, 1059-1060.	5.2	2
44	Very long-term outcomes in 23 patients with cblA type methylmalonic acidemia. <i>Journal of Inherited Metabolic Disease</i> , 0, , .	3.6	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. <i>Orphanet Journal of Rare Diseases</i> , 2022, 17, 202.	2.7	1
46	The Case A 69-year-old man with purpura and acute renal failure. <i>Kidney International</i> , 2018, 94, 435-436.	5.2	0