## Aurelia Poujois

List of Publications by Year in descending order

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623734 552781 36 722 14 26 citations g-index h-index papers 41 41 41 579 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Wilson's disease: A 2017 update. Clinics and Research in Hepatology and Gastroenterology, 2018, 42, 512-520.	1.5	92
2	Exchangeable copper: a reflection of the neurological severity in <scp>W</scp> ilson's disease. European Journal of Neurology, 2017, 24, 154-160.	3.3	74
3	Characteristics and prevalence of Wilson's disease: A 2013 observational population-based study in France. Clinics and Research in Hepatology and Gastroenterology, 2018, 42, 57-63.	1.5	56
4	New tools for Wilson's disease diagnosis: exchangeable copper fraction. Annals of Translational Medicine, 2019, 7, S70-S70.	1.7	53
5	High genetic carrier frequency of Wilson's disease in France: discrepancies with clinical prevalence. BMC Medical Genetics, 2018, 19, 143.	2.1	47
6	Wilson's disease: update on pathogenesis, biomarkers and treatments. Journal of Neurology, Neurosurgery and Psychiatry, 2021, 92, 1053-1061.	1.9	44
7	Semiquantitative Scale for Assessing Brain MRI Abnormalities in Wilson Disease: A Validation Study. Movement Disorders, 2020, 35, 994-1001.	3.9	43
8	Wilson disease. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2017, 142, 77-89.	1.8	39
9	Liver transplantation as a rescue therapy for severe neurologic forms of Wilson disease. Neurology, 2020, 94, e2189-e2202.	1.1	36
10	Challenges in the diagnosis of Wilson disease. Annals of Translational Medicine, 2019, 7, S67-S67.	1.7	31
11	Liver Transplantation in Wilson's Disease with Neurological Impairment: Evaluation in 4 Patients. European Neurology, 2017, 77, 5-15.	1.4	23
12	Bioavailable Trace Metals in Neurological Diseases. Current Treatment Options in Neurology, 2016, 18, 46.	1.8	21
13	CCDC115-CDG: A new rare and misleading inherited cause of liver disease. Molecular Genetics and Metabolism, 2018, 124, 228-235.	1.1	20
14	Adherence to treatment, a challenge even in treatable metabolic rare diseases: A cross sectional study of Wilson's disease. Journal of Inherited Metabolic Disease, 2021, 44, 1481-1488.	3.6	19
15	Neurological presentations revealing acquired copper deficiency: diagnosis features, aetiologies and evolution in seven patients. Internal Medicine Journal, 2018, 48, 535-540.	0.8	11
16	Sleep Abnormalities in Wilson's Disease. Current Treatment Options in Neurology, 2018, 20, 46.	1.8	11
17	The hidden face of Wilson's disease. Revue Neurologique, 2018, 174, 589-596.	1.5	9
18	Direct Determination of Non-Ceruloplasmin-Bound Copper in Plasma., 2019,, 249-255.		9

#	Article	IF	Citations
19	Inhibitory rTMS applied on somatosensory cortex in Wilson's disease patients with hand dystonia. Journal of Neural Transmission, 2017, 124, 1161-1170.	2.8	8
20	Sleep Disorders in Wilson's Disease. Current Neurology and Neuroscience Reports, 2019, 19, 84.	4.2	7
21	Sleep disorders in Wilson's disease. Sleep Medicine, 2021, 83, 299-303.	1.6	7
22	Non-invasive diagnosis and follow-up of rare genetic liver diseases. Clinics and Research in Hepatology and Gastroenterology, 2022, 46, 101768.	1.5	7
23	Dystonic Dysarthria in Wilson Disease: Efficacy of Zolpidem. Frontiers in Neurology, 2017, 8, 559.	2.4	6
24	ATP7B variant spectrum in a French pediatric Wilson disease cohort. European Journal of Medical Genetics, 2021, 64, 104305.	1.3	6
25	Eye Involvement in Wilson's Disease: A Review of the Literature. Journal of Clinical Medicine, 2022, 11, 2528.	2.4	6
26	Efficacy and Safety of Two Salts of Trientine in the Treatment of Wilson's Disease. Journal of Clinical Medicine, 2022, 11, 3975.	2.4	6
27	Comprehensive and comparative exploration of the <i>Atp7bâ^'/â^'</i> mouse plasma proteome. Metallomics, 2020, 12, 249-258.	2.4	5
28	Maintenance therapy simplification using a single daily dose: A preliminary real-life feasibility study in patients with Wilson disease. Clinics and Research in Hepatology and Gastroenterology, 2022, 46, 101978.	1.5	4
29	Cardiac involvement in Wilson disease: Review of the literature and description of three cases of sudden death. Journal of Inherited Metabolic Disease, 2021, 44, 1099-1112.	3.6	3
30	Biochemical Markers., 2019, , 115-124.		2
31	Exceptional involvement of medulla oblongata in Wilson disease. Neurology, 2019, 92, 770-771.	1.1	2
32	Validation of a non-motor fluctuations questionnaire in Parkinson's disease. Revue Neurologique, 2022, 178, 347-354.	1.5	2
33	La maladie de Wilson en 2018. Pratique Neurologique - FMC, 2018, 9, 173-185.	0.1	O
34	Wilson's Disease. , 2018, , 147-168.		0
35	Monitoring of Medical Therapy and Copper End Points. , 2019, , 223-232.		0
36	Answer to challenging issues in the management of Wilson's disease. Clinics and Research in Hepatology and Gastroenterology, 2019, 43, e7-e8.	1.5	0