Monica Furlano

List of Publications by Year
in descending order

15 papers	392 citations	1040056 9 h-index	996975 15 g-index
15 all docs	15 docs citations	15 times ranked	509 citing authors

#	Article	IF	Citations
1	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
2	Clinical and Genetic Features of Autosomal Dominant Alport Syndrome: A Cohort Study. American Journal of Kidney Diseases, 2021, 78, 560-570.e1.	1.9	48
3	Clinical utility of genetic testing in early-onset kidney disease: seven genes are the main players. Nephrology Dialysis Transplantation, 2022, 37, 687-696.	0.7	44
4	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
5	New therapeutic options for Alport syndrome. Nephrology Dialysis Transplantation, 2019, 34, 1272-1279.	0.7	37
6	Genetic kidney diseases as an underrecognized cause of chronic kidney disease: the key role of international registry reports. CKJ: Clinical Kidney Journal, 2021, 14, 1879-1885.	2.9	36
7	Novel homozygous OSGEP gene pathogenic variants in two unrelated patients with Galloway-Mowat syndrome: case report and review of the literature. BMC Nephrology, 2019, 20, 126.	1.8	16
8	Autosomal Dominant Polycystic Kidney Disease: Clinical Assessment of Rapid Progression. American Journal of Nephrology, 2018, 48, 308-317.	3.1	15
9	Generation of integration-free induced pluripotent stem cell lines derived from two patients with X-linked Alport syndrome (XLAS). Stem Cell Research, 2017, 25, 291-295.	0.7	13
10	Treatment and long-term outcome in primary nephrogenic diabetes insipidus. Nephrology Dialysis Transplantation, 2023, 38, 2120-2130.	0.7	9
11	Integration-free induced pluripotent stem cells derived from a patient with autosomal recessive Alport syndrome (ARAS). Stem Cell Research, 2017, 25, 1-5.	0.7	8
12	Comparative analysis of tools to predict rapid progression in autosomal dominant polycystic kidney disease. CKJ: Clinical Kidney Journal, 2022, 15, 912-921.	2.9	5
13	How genomics reclassifies diseases: the case of Alport syndrome. CKJ: Clinical Kidney Journal, 2020, 13, 933-935.	2.9	4
14	Autosomal dominant polycystic kidney disease: possibly the least silent cause of chronic kidney disease. CKJ: Clinical Kidney Journal, 2021, 14, 2281-2284.	2.9	3
15	Flank pain has a significant adverse impact on quality of life in ADPKD: the CYSTic-QoL study. CKJ: Clinical Kidney Journal, 2022, 15, 2063-2071.	2.9	3