MagdalÃ©na NeÅoldovÃ¡

List of Publications by Year
in descending order

12 papers	99 citations	1478505 6 h-index	1372567 10 g-index
12 all docs	12 docs citations	12 times ranked	277 citing authors

#	Article	IF	Citations
1	NKD1 marks intestinal and liver tumors linked to aberrant Wnt signaling. Cellular Signalling, 2015, 27, 245-256.	3.6	19
2	Rare variants in known and novel candidate genes predisposing to statin-associated myopathy. Pharmacogenomics, 2016, 17, 1405-1414.	1.3	17
3	Donor PNPLA3 and TM6SF2 Variant Alleles Confer Additive Risks for Graft Steatosis After Liver Transplantation. Transplantation, 2020, 104, 526-534.	1.0	16
4	Variable X-chromosome inactivation and enlargement of pericentral glutamine synthetase zones in the liver of heterozygous females with OTC deficiency. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2018, 472, 1029-1039.	2.8	12
5	Alpha-1 Antitrypsin and Hepatocellular Carcinoma in Liver Cirrhosis: SERPINA1 MZ or MS Genotype Carriage Decreases the Risk. International Journal of Molecular Sciences, 2021, 22, 10560.	4.1	8
6	Relevance of low viral load in haemodialysed patients with chronic hepatitis C virus infection. World Journal of Gastroenterology, 2015, 21, 5496.	3.3	8
7	Liver stiffness measured by two-dimensional shear-wave elastography predicts hepatic vein pressure gradient at high values in liver transplant candidates with advanced liver cirrhosis. PLoS ONE, 2021, 16, e0244934.	2.5	6
8	IL28B rs12979860 T allele protects against CMV disease in liver transplant recipients in the postâ€prophylaxis and late period. Transplant Infectious Disease, 2019, 21, e13124.	1.7	5
9	PNPLA3 rs738409 G allele carriers with genotype 1b HCV cirrhosis have lower viral load but develop liver failure at younger age. PLoS ONE, 2019, 14, e0222609.	2.5	3
10	USP18 downregulation in peripheral blood mononuclear cells predicts nonresponse to interferon-based triple therapy in patients with chronic hepatitis C, genotype 1: a pilot study. Therapeutics and Clinical Risk Management, 2015, 11, 1853.	2.0	2
11	ABCB4 disease mimicking morbus Wilson: A potential diagnostic pitfall. Biomedical Papers of the Medical Faculty of the University Palacký, Olomouc, Czechoslovakia, 2020, 164, 121-125.	0.6	2
12	Hereditary haemochromatosis caused by homozygous <scp>HJV</scp> mutation evolved through paternal disomy. Clinical Genetics, 2015, 87, 96-98.	2.0	1