Li-Ting Li

List of Publications by Year
in descending order

16	384	933264 10	940416 16
papers	citations	h-index	g-index

17	17	17	566
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Defining pathogenicity of <scp>NOTCH2</scp> variants for diagnosis of Alagille syndrome type 2 using a large cohort of patients. Liver International, 2022, 42, 1836-1848.	1.9	3
2	Impact of Genotype, Serum Bile Acids, and Surgical Biliary Diversion on Native Liver Survival in FIC1 Deficiency. Hepatology, 2021, 74, 892-906.	3.6	25
3	Whole-Genome Sequencing Reveals Large ATP8B1 Deletion/Duplications as Second Mutations Missed by Exome-Based Sequencing. Journal of Molecular Diagnostics, 2021, 23, 1491-1499.	1.2	2
4	ABCB11 deficiency presenting as transient neonatal cholestasis: Correlation with genotypes and BSEP expression. Liver International, 2020, 40, 2788-2796.	1.9	10
5	Lowâ€GGT intrahepatic cholestasis associated with biallelic USP53 variants: Clinical, histological and ultrastructural characterization. Liver International, 2020, 40, 1142-1150.	1.9	34
6	Genotype correlates with the natural history of severe bile salt export pump deficiency. Journal of Hepatology, 2020, 73, 84-93.	1.8	61
7	Novel missense mutation in VPS33B is associated with isolated low gammaâ€glutamyltransferase cholestasis: Attenuated, incomplete phenotype of arthrogryposis, renal dysfunction, and cholestasis syndrome. Human Mutation, 2019, 40, 2247-2257.	1.1	13
8	Comprehensive bile acid profiling in hereditary intrahepatic cholestasis: Genetic and clinical correlations. Liver International, 2018, 38, 1676-1685.	1.9	14
9	UGT1A1 genotypes and unconjugated hyperbilirubinemia phenotypes in post-neonatal Chinese children. Medicine (United States), 2018, 97, e13576.	0.4	17
10	Defects in myosin VB are associated with a spectrum of previously undiagnosed low Î³â€glutamyltransferase cholestasis. Hepatology, 2017, 65, 1655-1669.	3.6	107
11	The Features of GGT in Patients with ATP8B1 or ABCB11 Deficiency Improve the Diagnostic Efficiency. PLoS ONE, 2016, 11, e0153114.	1.1	12
12	Wilson disease with hepatic presentation in an eight-month-old boy. World Journal of Gastroenterology, 2015, 21, 8981.	1.4	25
13	JAG1 Mutation Spectrum and Origin in Chinese Children with Clinical Features of Alagille Syndrome. PLoS ONE, 2015, 10, e0130355.	1.1	30
14	Hypothyroidism Associated with ATP8B1 Deficiency. Journal of Pediatrics, 2015, 167, 1334-1339.e1.	0.9	15
15	Two novel VPS33B mutations in a patient with arthrogryposis, renal dysfunction and cholestasis syndrome in mainland China. World Journal of Gastroenterology, 2014, 20, 326.	1.4	6
16	ARC syndrome with high GGT cholestasis caused by VPS33B mutations. World Journal of Gastroenterology, 2014, 20, 4830.	1.4	10