

Imre F Schene

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/1682416/publications.pdf>

Version: 2024-02-01

10
papers

402
citations

1307594

7
h-index

1281871

11
g-index

12
all docs

12
docs citations

12
times ranked

827
citing authors

#	ARTICLE	IF	CITATIONS
1	The potential and limitations of intrahepatic cholangiocyte organoids to study inborn errors of metabolism. <i>Journal of Inherited Metabolic Disease</i> , 2022, 45, 353-365.	3.6	4
2	Mutation-specific reporter for optimization and enrichment of prime editing. <i>Nature Communications</i> , 2022, 13, 1028.	12.8	16
3	Treatment of ARS deficiencies with specific amino acids. <i>Genetics in Medicine</i> , 2021, 23, 2202-2207.	2.4	18
4	Large-scale Production of LGR5-Positive Bipotential Human Liver Stem Cells. <i>Hepatology</i> , 2020, 72, 257-270.	7.3	89
5	Prime editing for functional repair in patient-derived disease models. <i>Nature Communications</i> , 2020, 11, 5352.	12.8	134
6	Misdiagnosis of CTX due to propofol: The interference of total intravenous propofol anaesthesia with bile acid profiling. <i>Journal of Inherited Metabolic Disease</i> , 2020, 43, 843-851.	3.6	5
7	Human extrahepatic and intrahepatic cholangiocyte organoids show region-specific differentiation potential and model cystic fibrosis-related bile duct disease. <i>Scientific Reports</i> , 2020, 10, 21900.	3.3	43
8	Aminoacyl-tRNA synthetase deficiencies in search of common themes. <i>Genetics in Medicine</i> , 2019, 21, 319-330.	2.4	70
9	Glycogen Storage Disease Type IV: A Rare Cause for Neuromuscular Disorders or Often Missed?. <i>JIMD Reports</i> , 2018, 45, 99-104.	1.5	7
10	Pitfalls in Diagnosing Neuraminidase Deficiency: Psychosomatics and Normal Sialic Acid Excretion. <i>JIMD Reports</i> , 2015, 25, 9-13.	1.5	15