Gertrud Eckstein

List of Publications by Year
in descending order

9	375	1163117 8	1474206 9
papers	citations	h-index	g-index

9	9	9	1552
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Biallelic Mutations of Methionyl-tRNA Synthetase Cause a Specific Type of Pulmonary Alveolar Proteinosis Prevalent on RÃ©union Island. American Journal of Human Genetics, 2015, 96, 826-831.	6.2	94
2	Bi-allelic Truncating Mutations in TANGO2 Cause Infancy-Onset Recurrent Metabolic Crises with Encephalocardiomyopathy. American Journal of Human Genetics, 2016, 98, 358-362.	6.2	77
3	Mitochondrial DNA mutation analysis from exome sequencingâ€"A more holistic approach in diagnostics of suspected mitochondrial disease. Journal of Inherited Metabolic Disease, 2019, 42, 909-917.	3.6	57
4	Congenital heart disease risk loci identified by genome-wide association study in European patients. Journal of Clinical Investigation, 2021, 131, .	8.2	47
5	Mapping the Genetic Architecture of Gene Regulation in Whole Blood. PLoS ONE, 2014, 9, e93844.	2.5	31
6	Genomic Heterogeneity of Osteosarcoma - Shift from Single Candidates to Functional Modules. PLoS ONE, 2015, 10, e0123082.	2.5	27
7	Human skin-resident host T cells can persist long term after allogeneic stem cell transplantation and maintain recirculation potential. Science Immunology, 2022, 7, eabe2634.	11.9	23
8	Stimulation of soluble guanylyl cyclase (sGC) by riociguat attenuates heart failure and pathological cardiac remodelling. British Journal of Pharmacology, 2022, 179, 2430-2442.	5.4	15
9	Adult-onset variant ataxia-telangiectasia diagnosed by exome and cDNA sequencing. Neurology: Genetics, 2019, 5, e346.	1.9	4