Hideki Matsumoto

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/5047091/publications.pdf

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		1478505	1372567
10	113	6	10
papers	citations	h-index	g-index

10	10	10	156
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Recent advances in understanding beta-ketothiolase (mitochondrial acetoacetyl-CoA thiolase, T2) deficiency. Journal of Human Genetics, 2019, 64, 99-111.	2.3	30
2	Successful Everolimus Treatment of Kaposiform Hemangioendothelioma With Kasabach-Merritt Phenomenon: Clinical Efficacy and Adverse Effects of mTOR Inhibitor Therapy. Journal of Pediatric Hematology/Oncology, 2016, 38, e322-e325.	0.6	22
3	Deficiency of 3â€hydroxybutyrate dehydrogenase (BDH1) in mice causes low ketone body levels and fatty liver during fasting. Journal of Inherited Metabolic Disease, 2020, 43, 960-968.	3.6	21
4	Prepubertal onset of slipped capital femoral epiphysis associated with hypothyroidism: a case report and literature review. BMC Endocrine Disorders, 2017, 17, 59.	2.2	13
5	Autosomal dominant Hashimotoâ€™s thyroiditis with a mutation in <i>TNFAIP3</i>. Clinical Pediatric Endocrinology, 2019, 28, 91-96.	0.8	8
6	In vitro functional analysis of four variants of human asparagine synthetase. Journal of Inherited Metabolic Disease, 2021, 44, 1226-1234.	3.6	7
7	Microdeletion in Xq28 with a polymorphic inversion in a patient with FLNA-associated progressive lung disease. Respiratory Investigation, 2019, 57, 395-398.	1.8	6
8	Japanese patients with mitochondrial 3â€‘hydroxyâ€‘3â€‘methylglutarylâ€‘CoA synthase deficiency: In vitro functional analysis of five novel HMGCS2 mutations. Experimental and Therapeutic Medicine, 2020, 20, 1-1.	1.8	3
9	Founder genetic variants of ABCC4 and ABCC11 in the Japanese population are not associated with the development of subacute myeloâ€opticoâ€neuropathy (SMON). Molecular Genetics & Genomic Medicine, 2022, 10, e1845.	1.2	2
10	Immediate postnatal central hypothyroidism caused by maternal Graves' disease: Importance of early screening. Clinical Case Reports (discontinued), 2022, 10, .	0.5	1