Tony Huynh

List of Publications by Year
in descending order

25 papers	676 citations	840119 11 h-index	752256 20 g-index
25 all docs	25 docs citations	25 times ranked	1216 citing authors

#	Article	IF	Citations
1	Disorders of sex development: insights from targeted gene sequencing of a large international patient cohort. Genome Biology, 2016, 17, 243.	3.8	241
2	VBP15, a novel antiâ€inflammatory and membraneâ€stabilizer, improves muscular dystrophy without side effects. EMBO Molecular Medicine, 2013, 5, 1569-1585.	3.3	148
3	Disease characteristics of MCT8 deficiency: an international, retrospective, multicentre cohort study. Lancet Diabetes and Endocrinology,the, 2020, 8, 594-605.	5.5	50
4	Omigapil Treatment Decreases Fibrosis and Improves Respiratory Rate in dy2J Mouse Model of Congenital Muscular Dystrophy. PLoS ONE, 2013, 8, e65468.	1.1	37
5	The association between ketoacidosis and 25(OH)-vitamin D₃levels at presentation in children with type 1 diabetes mellitus. Pediatric Diabetes, 2009, 10, 38-43.	1.2	35
6	Selective modulation through the glucocorticoid receptor ameliorates muscle pathology in mdx mice. Journal of Pathology, 2013, 231, 223-235.	2.1	31
7	VBP15, a Glucocorticoid Analogue, Is Effective at Reducing Allergic Lung Inflammation in Mice. PLoS ONE, 2013, 8, e63871.	1.1	24
8	The effects of MyD88 deficiency on disease phenotype in dysferlin-deficient A/J mice: role of endogenous TLR ligands. Journal of Pathology, 2013, 231, 199-209.	2.1	22
9	Long-Term Efficacy of T3 Analogue Triac in Children and Adults With MCT8 Deficiency: A Real-Life Retrospective Cohort Study. Journal of Clinical Endocrinology and Metabolism, 2022, 107, e1136-e1147.	1.8	15
10	Benefit of Early Commencement of Growth Hormone Therapy in Children with Prader-Willi Syndrome. Journal of Pediatric Endocrinology and Metabolism, 2009, 22, 1151-8.	0.4	13
11	A novel mutation in the TG gene (G2322S) causing congenital hypothyroidism in a Sudanese family: a case report. BMC Medical Genetics, 2018, 19, 69.	2.1	13
12	Urine metabonomic profiling of a female adolescent with PIT-1 mutation before and during growth hormone therapy: Insights into the metabolic effects of growth hormone. Growth Hormone and IGF Research, 2013, 23, 29-36.	0.5	12
13	Therapeutic plasma exchange normalizes insulin-mediated response in a child with type 1 diabetes and insulin autoimmune syndrome. Pediatric Diabetes, 2018, 19, 171-179.	1.2	8
14	Effects of Dantrolene Therapy on Disease Phenotype in Dystrophin Deficient mdx Mice. PLOS Currents, 2013, 5, .	1.4	7
15	Clinical and Laboratory Aspects of Insulin Autoantibody-Mediated Glycaemic Dysregulation and Hyperinsulinaemic Hypoglycaemia: Insulin Autoimmune Syndrome and Exogenous Insulin Antibody Syndrome. , 2020, 41, 93-102.		7
16	Immunoassay interference secondary to therapeutic highâ€dose biotin: A paediatric case report. Journal of Paediatrics and Child Health, 2018, 54, 572-575.	0.4	4
17	A Novel V185DfsX4 Mutation of the AAAS Gene in a 2-year-old Boy with Triple A Syndrome. Clinical Pediatric Endocrinology, 2009, 18, 73-75.	0.4	3
18	Virilisation in siblings secondary to transdermal â€˜bioidenticalâ€™ testosterone exposure. Journal of Paediatrics and Child Health, 2017, 53, 301-305.	0.4	3

#	Article	IF	Citations
19	Diabetes in a child on growth hormone therapy: Questions. Pediatric Nephrology, 2018, 33, 77-78.	0.9	1
20	An Unusual Cause of Metabolic Alkalosis and Hypocalcemia in Childhood. Clinical Chemistry, 2019, 65, 514-517.	1.5	1
21	Pre-analytical mysteries: A case of severe hypervitaminosis D and mild hypercalcaemia. Biochemia Medica, 2021, 31, 149-155.	1.2	1
22	Diabetes in a child on growth hormone therapy: Answers. Pediatric Nephrology, 2018, 33, 79-80.	0.9	0
23	CT can stratify patients as low risk for tibial neuropathy following a talus fracture. Emergency Radiology, 2019, 26, 541-548.	1.0	0
24	A Rare and Unusual Cause of Unilateral Ureteric Obstruction in a Child. Clinical Chemistry, 2020, 66, 1006-1009.	1.5	0
25	Challenges of managing congenital hyperinsulinism in remote Aboriginal Australian communities. Journal of Paediatrics and Child Health, 2021, 57, 727-731.	0.4	0