

# Tony Huynh

## List of Publications by Year in descending order

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Version: 2024-02-01

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	Long-Term Efficacy of T3 Analogue Triac in Children and Adults With MCT8 Deficiency: A Real-Life Retrospective Cohort Study. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2022, 107, e1136-e1147.	1.8	15
2	Challenges of managing congenital hyperinsulinism in remote Aboriginal Australian communities. <i>Journal of Paediatrics and Child Health</i> , 2021, 57, 727-731.	0.4	0
3	Pre-analytical mysteries: A case of severe hypervitaminosis D and mild hypercalcaemia. <i>Biochimica Medica</i> , 2021, 31, 149-155.	1.2	1
4	Disease characteristics of MCT8 deficiency: an international, retrospective, multicentre cohort study. <i>Lancet Diabetes and Endocrinology</i> , 2020, 8, 594-605.	5.5	50
5	A Rare and Unusual Cause of Unilateral Ureteric Obstruction in a Child. <i>Clinical Chemistry</i> , 2020, 66, 1006-1009.	1.5	0
6	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
7	CT can stratify patients as low risk for tibial neuropathy following a talus fracture. <i>Emergency Radiology</i> , 2019, 26, 541-548.	1.0	0
8	An Unusual Cause of Metabolic Alkalosis and Hypocalcemia in Childhood. <i>Clinical Chemistry</i> , 2019, 65, 514-517.	1.5	1
9	Immunoassay interference secondary to therapeutic high-dose biotin: A paediatric case report. <i>Journal of Paediatrics and Child Health</i> , 2018, 54, 572-575.	0.4	4
10	Diabetes in a child on growth hormone therapy: Questions. <i>Pediatric Nephrology</i> , 2018, 33, 77-78.	0.9	1
11	Therapeutic plasma exchange normalizes insulin-mediated response in a child with type 1 diabetes and insulin autoimmune syndrome. <i>Pediatric Diabetes</i> , 2018, 19, 171-179.	1.2	8
12	Diabetes in a child on growth hormone therapy: Answers. <i>Pediatric Nephrology</i> , 2018, 33, 79-80.	0.9	0
13	A novel mutation in the TG gene (G2322S) causing congenital hypothyroidism in a Sudanese family: a case report. <i>BMC Medical Genetics</i> , 2018, 19, 69.	2.1	13
14	Virilisation in siblings secondary to transdermal bioidentical testosterone exposure. <i>Journal of Paediatrics and Child Health</i> , 2017, 53, 301-305.	0.4	3
15	Disorders of sex development: insights from targeted gene sequencing of a large international patient cohort. <i>Genome Biology</i> , 2016, 17, 243.	3.8	241
16	Urine metabolomic profiling of a female adolescent with PIT-1 mutation before and during growth hormone therapy: Insights into the metabolic effects of growth hormone. <i>Growth Hormone and IGF Research</i> , 2013, 23, 29-36.	0.5	12
17	Selective modulation through the glucocorticoid receptor ameliorates muscle pathology in mdx mice. <i>Journal of Pathology</i> , 2013, 231, 223-235.	2.1	31
18	The effects of MyD88 deficiency on disease phenotype in dysferlin-deficient A/J mice: role of endogenous TLR ligands. <i>Journal of Pathology</i> , 2013, 231, 199-209.	2.1	22

#	ARTICLE	IF	CITATIONS
19	VBP15, a novel anti-inflammatory and membrane stabilizer, improves muscular dystrophy without side effects. <i>EMBO Molecular Medicine</i> , 2013, 5, 1569-1585.	3.3	148
20	VBP15, a Glucocorticoid Analogue, Is Effective at Reducing Allergic Lung Inflammation in Mice. <i>PLoS ONE</i> , 2013, 8, e63871.	1.1	24
21	Omigapil Treatment Decreases Fibrosis and Improves Respiratory Rate in dy2J Mouse Model of Congenital Muscular Dystrophy. <i>PLoS ONE</i> , 2013, 8, e65468.	1.1	37
22	Effects of Dantrolene Therapy on Disease Phenotype in Dystrophin Deficient mdx Mice. <i>PLOS Currents</i> , 2013, 5, .	1.4	7
23	A Novel V185DfsX4 Mutation of the AAAS Gene in a 2-year-old Boy with Triple A Syndrome. <i>Clinical Pediatric Endocrinology</i> , 2009, 18, 73-75.	0.4	3
24	Benefit of Early Commencement of Growth Hormone Therapy in Children with Prader-Willi Syndrome. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2009, 22, 1151-8.	0.4	13
25	The association between ketoacidosis and 25(OH)-vitamin D <sub>3</sub> levels at presentation in children with type 1 diabetes mellitus. <i>Pediatric Diabetes</i> , 2009, 10, 38-43.	1.2	35