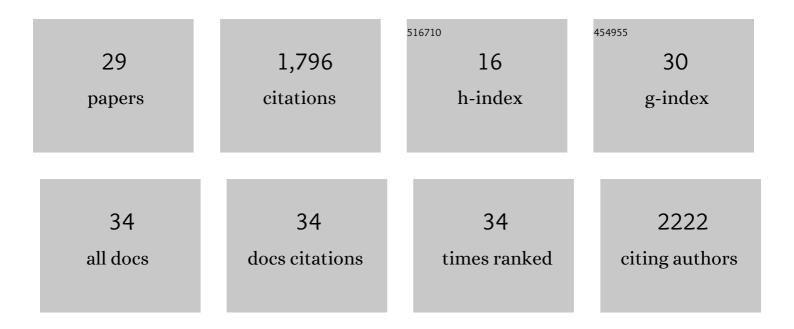
Kanetee Busiah

List of Publications by Year in descending order

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KANFTEF RUSIAH

#	Article	IF	CITATIONS
1	High Prevalence of Early Endocrine Disorders After Childhood Brain Tumors in a Large Cohort. Journal of Clinical Endocrinology and Metabolism, 2022, 107, e2156-e2166.	3.6	6
2	Presenting features and molecular genetics of primary hyperparathyroidism in the paediatric population. European Journal of Endocrinology, 2021, 184, 343-351.	3.7	9
3	Congenital hyperinsulinism: 2 case reports with different rare variants in ABCC8. Annals of Pediatric Endocrinology and Metabolism, 2021, 26, 60-65.	2.3	2
4	The 2021 European Training Requirements in Paediatric Endocrinology and Diabetes. Hormone Research in Paediatrics, 2021, , .	1.8	0
5	Neonatal Diabetes Mellitus. Frontiers in Pediatrics, 2020, 8, 540718.	1.9	37
6	Long-term Metabolic and Socioeducational Outcomes of Transient Neonatal Diabetes: A Longitudinal and Cross-sectional Study. Diabetes Care, 2020, 43, 1191-1199.	8.6	5
7	Glibenclamide oral suspension: Suitable and effective in patients with neonatal diabetes. Pediatric Diabetes, 2019, 20, 246-254.	2.9	16
8	Improved General and Height-Specific Quality of Life in Children With Short Stature After 1 Year on Growth Hormone. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 2103-2111.	3.6	25
9	Successful off-label sulfonylurea treatment of neonatal diabetes mellitus due to chromosome 6 abnormalities. Pediatric Diabetes, 2018, 19, 663-669.	2.9	19
10	A novel <i> <scp>NEUROG3</scp> </i> mutation in neonatal diabetes associated with a neuroâ€intestinal syndrome. Pediatric Diabetes, 2018, 19, 381-387.	2.9	17
11	The transition from pediatric to adult care for youth with epilepsy: Basic biological, sociological, and psychological issues. Epilepsy and Behavior, 2017, 69, 170-176.	1.7	45
12	High Prevalence of Polycystic Ovary Syndrome in Type 1 Diabetes Mellitus Adolescents: Is There a Difference Depending on the NIH and Rotterdam Criteria?. Hormone Research in Paediatrics, 2017, 87, 333-341.	1.8	15
13	Biallelic Mutations in LIPT2 Cause a Mitochondrial Lipoylation Defect Associated with Severe Neonatal Encephalopathy. American Journal of Human Genetics, 2017, 101, 283-290.	6.2	55
14	Hyperglycémies syndromiques néonatales. , 2017, , 463-476.		0
15	Evaluation of the pharmacokinetics of glibenclamide tablet given, off label, orally to children suffering from neonatal syndromic hyperglycemia. European Journal of Clinical Pharmacology, 2016, 72, 1373-1379.	1.9	3
16	Differentiating Transient Idiopathic Hyperglycaemia and Neonatal Diabetes Mellitus in Preterm Infants. Hormone Research in Paediatrics, 2015, 84, 68-72.	1.8	8
17	Sulfonylurea Therapy Benefits Neurological and Psychomotor Functions in Patients With Neonatal Diabetes Owing to Potassium Channel Mutations. Diabetes Care, 2015, 38, 2033-2041.	8.6	75
18	RFX6 Regulates Insulin Secretion by Modulating Ca2+ Homeostasis in Human β Cells. Cell Reports, 2014, 9, 2206-2218.	6.4	73

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#	Article	IF	CITATIONS
19	Human Pancreas Endocrine Cell Populations and Activating <i>ABCC8</i> Mutations. Hormone Research in Paediatrics, 2014, 82, 59-64.	1.8	11
20	Small-molecule inhibitors of the cystic fibrosis transmembrane conductance regulator increase pancreatic endocrine cell development in rat and mouse. Diabetologia, 2013, 56, 330-339.	6.3	6
21	Transcription factor gene MNX1 is a novel cause of permanent neonatal diabetes in a consanguineous family. Diabetes and Metabolism, 2013, 39, 276-280.	2.9	48
22	Neuropsychological dysfunction and developmental defects associated with genetic changes in infants with neonatal diabetes mellitus: a prospective cohort study. Lancet Diabetes and Endocrinology,the, 2013, 1, 199-207.	11.4	87
23	Childhood Craniopharyngioma: Hypothalamus-Sparing Surgery Decreases the Risk of Obesity. Journal of Clinical Endocrinology and Metabolism, 2013, 98, 2376-2382.	3.6	170
24	Disruption of a Novel Krüppel-like Transcription Factor p300-regulated Pathway for Insulin Biosynthesis Revealed by Studies of the c331 INS Mutation Found in Neonatal Diabetes Mellitus. Journal of Biological Chemistry, 2011, 286, 28414-28424.	3.4	72
25	Molecular Diagnosis of Neonatal Diabetes Mellitus Using Next-Generation Sequencing of the Whole Exome. PLoS ONE, 2010, 5, e13630.	2.5	102
26	Childhood craniopharyngioma: greater hypothalamic involvement before surgery is associated with higher homeostasis model insulin resistance index. BMC Pediatrics, 2009, 9, 24.	1.7	34
27	New <i>ABCC8</i> Mutations in Relapsing Neonatal Diabetes and Clinical Features. Diabetes, 2007, 56, 1737-1741.	0.6	83
28	Activating Mutations in the <i>ABCC8</i> Gene in Neonatal Diabetes Mellitus. New England Journal of Medicine, 2006, 355, 456-466.	27.0	591
29	Kir6.2 Mutations Are a Common Cause of Permanent Neonatal Diabetes in a Large Cohort of French Patients. Diabetes, 2004, 53, 2719-2722.	0.6	171