Min-Sun Kim

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/10246852/publications.pdf

Version: 2024-02-01

1684188 1588992 19 69 5 8 citations h-index g-index papers 55 20 20 20 times ranked docs citations citing authors all docs

#	Article	IF	CITATIONS
1	Clinical Utility of [18F]FDG-PET /CT in Pericardial Disease. Current Cardiology Reports, 2019, 21, 107.	2.9	18
2	Clinical Presentation and Treatment Outcomes of Children and Adolescents With Pheochromocytoma and Paraganglioma in a Single Center in Korea. Frontiers in Endocrinology, 2020, 11, 610746.	3.5	13
3	Long-Term Antithyroid Drug Treatment of Graves' Disease in Children and Adolescents: A 20-Year Single-Center Experience. Frontiers in Endocrinology, 2021, 12, 687834.	3.5	10
4	Clinical and molecular characterization of Korean children with infantile and late-onset Pompe disease: 10 years of experience with enzyme replacement therapy at a single center. Korean Journal of Pediatrics, 2019, 62, 224-234.	1.9	5
5	Tailored management of life-threatening complications related to severe obesity in a young adult with Prader-Willi syndrome. Annals of Pediatric Endocrinology and Metabolism, 2022, 27, 148-152.	2.3	5
6	Clinical characteristics, treatment outcomes, and occurrence of diabetes mellitus after pancreatic resection of solid pseudopapillary tumor in children and adolescents: A single institution experience with 51 cases. Pancreatology, 2021, 21, 509-514.	1.1	3
7	Endocrine and Metabolic Illnesses in Young Adults with Prader–Willi Syndrome. Journal of Personalized Medicine, 2022, 12, 858.	2.5	3
8	Hypertriglyceridemia with acute pancreatitis in a 14-year-old girl with diabetic ketoacidosis. Annals of Pediatric Endocrinology and Metabolism, 2022, 27, 73-77.	2.3	2
9	Nonclassic congenital lipoid adrenal hyperplasia diagnosed at 17 months in a Korean boy with normal male genitalia: emphasis on pigmentation as a diagnostic clue. Annals of Pediatric Endocrinology and Metabolism, 2020, 25, 46-51.	2.3	2
10	First female Korean child with Coffin-Lowry syndrome: a novel variant in RPS6KA3 diagnosed by exome sequencing and a literature review. Annals of Pediatric Endocrinology and Metabolism, 2023, 28, 67-72.	2.3	2
11	Natural History and Molecular Characteristics of Korean Patients with Mucopolysaccharidosis Type III. Journal of Personalized Medicine, 2022, 12, 665.	2.5	2
12	Efficacy and safety of the recombinant human growth hormone in short children born small for gestational age. Medicine (United States), 2021, 100, e26711.	1.0	1
13	Wilson disease diagnosed incidentally by targeted gene panel sequencing in a Korean boy with severe obesity. Annals of Pediatric Endocrinology and Metabolism, 2022, 27, 229-235.	2.3	1
14	Appropriate Age for Height Control Treatment in Patients With Marfan Syndrome. Frontiers in Endocrinology, 2021, 12, 708931.	3.5	1
15	Multiple endocrine neoplasia type 2 and autoimmune polyendocrine syndromes (type 1 diabetes) Tj ETQq1 1 0.	784314 rg 1.6	gBT /Overlock 1 1
16	Development and validation of the Pediatric-Youth Hyperphagia Assessment for Prader-Willi syndrome. Epidemiology and Health, 2022, 44, e2022014.	1.9	0
17	The First Korean Case of Baraitser-Winter Cerebro-Fronto-Facial Syndrome with a Novel Mutation in Diagnosed Via Targeted Gene Panel Sequencing and Literature Review. Annals of Clinical and Laboratory Science, 2020, 50, 818-824.	0.2	0
18	A boy with Coffin-Siris syndrome with a novel frameshift mutation in ARID1B. Neuroendocrinology Letters, 2021, 41, 285-289.	0.2	0

#	Article	IF	CITATIONS
19	Recombinant growth hormone therapy in children with Turner Syndrome in Korea: a phase III Randomized Trial. BMC Endocrine Disorders, 2021, 21, 243.	2.2	O