Dimitrios T Papadimitriou

List of Publications by Year
in descending order

59	752	516215 16	552369 26
papers	citations	h-index	g-index

65	65	65	1125
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Mineralocorticoid receptor mutations are the principal cause of renal type 1 pseudohypoaldosteronism. Human Mutation, 2007, 28, 33-40.	1.1	79
2	Maturational tempo differences in relation to the timing of the onset of puberty in girls. Acta Paediatrica, International Journal of Paediatrics, 2008, 97, 217-220.	0.7	60
3	Deficit in Anterior Pituitary Function and Variable Immune Deficiency (DAVID) in Children Presenting with Adrenocorticotropin Deficiency and Severe Infections. Journal of Clinical Endocrinology and Metabolism, 2012, 97, E121-E128.	1.8	54
4	Novel Twinkle (PEO1) gene mutations in mendelian progressive external ophthalmoplegia. Journal of Neurology, 2008, 255, 1384-1391.	1.8	45
5	Puberty in Subjects with Complete Androgen Insensitivity Syndrome. Hormone Research in Paediatrics, 2006, 65, 126-131.	0.8	41
6	Sex Differences in the Secular Changes in Pubertal Maturation. Pediatrics, 2001, 108, e65-e65.	1.0	33
7	Timing of Pubertal Onset in Girls: Evidence for Non-Gaussian Distribution. Journal of Clinical Endocrinology and Metabolism, 2008, 93, 4422-4425.	1.8	33
8	The Big Vitamin D Mistake. Journal of Preventive Medicine and Public Health, 2017, 50, 278-281.	0.7	32
9	Mineralocorticoid Deficiency in Post-Operative Cerebral Salt Wasting. Journal of Pediatric Endocrinology and Metabolism, 2007, 20, 1145-50.	0.4	30
10	Replacement of Male Mini-Puberty. Journal of the Endocrine Society, 2019, 3, 1275-1282.	0.1	30
11	Low TSH levels are not associated with osteoporosis in childhood. European Journal of Endocrinology, 2007, 157, 221-223.	1.9	23
12	Prevalence of overweight and obesity in young Greek men. Obesity Reviews, 2008, 9, 100-103.	3.1	21
13	Negativation of type 1 diabetesâ€associated autoantibodies to glutamic acid decarboxylase and insulin in children treated with oral calcitriol (å£æœé¨åŒ–ä¸‰é†‡æ²»ç——ç»´æŒå„¿ç«¥1åž‹ç³–å°¿ç—…ç›„å…³çš„è°·æ°¨é…¸è„±ç¾§é…¶å’Žèƒ°	0.8	19
14	Overweight and obesity decreased in Greek schoolchildren from 2009 toÂ2012 during the early phase of the economic crisis. Acta Paediatrica, International Journal of Paediatrics, 2016, 105, 200-205.	0.7	19
15	Thyroid dysfunction associated with increased low-density lipoprotein cholesterol in epileptic children treated with carbamazepine monotherapy: A causal relationship?. European Journal of Paediatric Neurology, 2007, 11, 358-361.	0.7	18
16	Pubertal Maturation of Contemporary Greek Boys: No Evidence of a Secular Trend. Journal of Adolescent Health, 2011, 49, 434-436.	1.2	18
17	Reconsidering the Sex Differences in the Incidence of Pubertal Disorders. Hormone and Metabolic Research, 2005, 37, 708-710.	0.7	14
18	Greek young men grow taller. Acta Paediatrica, International Journal of Paediatrics, 2008, 97, 1105-1107.	0.7	14

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19	Characteristics of the short children referred to an academic paediatric endocrine clinic in Greece. Journal of Paediatrics and Child Health, 2012, 48, 263-267.	0.4	14
20	Association between population vitamin D status and SARS-CoV-2 related serious-critical illness and deaths: An ecological integrative approach. World Journal of Virology, 2021, 10, 111-129.	1.3	14
21	Secular Trend in Body Height of Schoolchildren in Northeast Attica, Greece. Journal of Pediatric Endocrinology and Metabolism, 2009, 22, 13-7.	0.4	10
22	Constitutional Advancement of Growth Is Associated with Early Puberty in Girls. Hormone Research in Paediatrics, 2011, 76, 273-277.	0.8	10
23	Maternal uniparental disomy of chromosomeÂ4 and homozygous novel mutation in the WFS1 gene in a paediatric patient with Wolfram syndrome. Diabetes and Metabolism, 2015, 41, 433-435.	1.4	10
24	The autoimmune hypothesis for acute bilateral cataract in typeÂ1 diabetes. Diabetes and Metabolism, 2016, 42, 386-387.	1.4	10
25	Validation of a Food Frequency Questionnaire Designed for Children 10-12 Years: The Panacea-FFQ. Pediatric Research, 2011, 70, 778-778.	1.1	9
26	Calcium sensing receptor in pregnancies complicated by gestational diabetes mellitus. Placenta, 2014, 35, 632-638.	0.7	8
27	Anastrozole plus leuprorelin in early maturing girls with compromised growth: the â€œGAILâ€•study. Journal of Endocrinological Investigation, 2016, 39, 439-446.	1.8	8
28	Heterozygous mutations in the cholesterol side-chain cleavage enzyme gene (CYP11A1) can cause transient adrenal insufficiency and life-threatening failure to thrive. Hormones, 2018, 17, 419-421.	0.9	8
29	Endocrine-Disrupting Chemicals and Early Puberty in Girls. Children, 2021, 8, 492.	0.6	8
30	<smlcap>L</smlcap>-Dopa Stimulates Cortisol Secretion through Adrenocorticotropic Hormone Release in Short Children. Hormone Research in Paediatrics, 2015, 84, 319-322.	0.8	6
31	Late diagnosis of 3Î²-Hydroxysteroid dehydrogenase deficiency: the pivotal role of gas chromatography-mass spectrometry urinary steroid metabolome analysis and a novel homozygous nonsense mutation in the HSD3B2 gene. Journal of Pediatric Endocrinology and Metabolism, 2021, 34, 131-136.	0.4	6
32	Growth variations with opposite clinical outcomes and the emerging role of IGF-1. Trends in Endocrinology and Metabolism, 2022, 33, 359-370.	3.1	6
33	Cerebral Salt Wasting Complicated by Central Diabetes Insipidus and Growth Hormone Deficiency. Indian Journal of Pediatrics, 2018, 85, 580-581.	0.3	5
34	Rifampicin for COVID-19. World Journal of Virology, 2022, 11, 90-97.	1.3	4
35	<smlcap>L</smlcap>-Dopa Is a Potent Stimulator of Cortisol in Short Children. Hormone Research in Paediatrics, 2014, 81, 386-390.	0.8	3
36	Lessons from Wolfram Syndrome: Initiation of DDAVP Therapy Causes Renal Salt Wasting Due to Elevated ANP/BNP Levels, Rescued by Fludrocortisone Treatment. Indian Journal of Pediatrics, 2021, 88, 582-585.	0.3	3

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37	Treatment of alopecia totalis/universalis/focalis with vitamin D and analogs: Three case reports and a literature review. World Journal of Clinical Pediatrics, 2021, 10, 192-199.	0.6	3
38	A novel heterozygous mutation in the glucokinase gene is responsible for an early-onset mild form of maturity-onset diabetes of the young, type 2. Diabetes and Metabolism, 2015, 41, 342-343.	1.4	2
39	Secular changes in the final height of Greek girls are levelling off. Acta Paediatrica, International Journal of Paediatrics, 2017, 106, 341-343.	0.7	2
40	Autism spectrum disorder, anxiety and severe depression in a male patient with deletion and duplication in the 21q22.3 region: A case report. Biomedical Reports, 2019, 1, 1-5.	0.9	2
41	A novel heterozygous mutation in the SLC5A2 gene causing severe glycosuria, mild failure to thrive, and subclinical hypoglycemia. Journal of Diabetes, 2021, 13, 688-692.	0.8	2
42	MON-541 Successful Treatment of Normocalcemic Hyperparathyroidism in Children. Journal of the Endocrine Society, 2019, 3, .	0.1	2
43	Power Doppler ultrasound: a potentially useful alternative in diagnosing pelvic pathologic conditions. Clinical and Experimental Obstetrics and Gynecology, 1996, 23, 229-32.	0.1	2
44	Combined Growth Hormone-Releasing Hormone and Growth Hormone-Releasing Peptide-6 Test for the Evaluation of Growth Hormone Secretion in Children with Growth Hormone Deficiency and Growth Hormone Neurosecretory Dysfunction. Hormone Research, 2008, 70, 215-223.	1.8	1
45	Successful Treatment of Severe Atopic Dermatitis with Calcitriol and Paricalcitol in an 8-Year-Old Girl. Case Reports in Pediatrics, 2018, 2018, 1-5.	0.2	1
46	A novel detrimental homozygous mutation in the WFS1 gene in two sisters from nonconsanguineous parents with untreated diabetes insipidus. Clinical Case Reports (discontinued), 2019, 7, 2355-2357.	0.2	1
47	A Greek girl with 11Î²-hydroxylase deficiency due to compound heterozygosity for two novel mutations in CYP11B1 gene. Endocrinology, Diabetes and Metabolism Case Reports, 2015, 2015, 150074.	0.2	1
48	Hypoprolactinemia as a Clue to Diagnosis of Mild Central Hypothyroidism due to IGSF1 Deficiency. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2020, 12, 218-222.	0.4	1
49	A Mixed-Longitudinal Study of Height Velocity of Greek Schoolchildren and the Milestones of the Adolescent Growth Spurt. Children, 2022, 9, 790.	0.6	1
50	MON-072 A 2 -Year Old Girl with Turner Syndrome and Neurofibromatosis Type 1. Journal of the Endocrine Society, 2020, 4, .	0.1	0
51	Association Between Population Vitamin D Status and SARS-CoV-2 Related Serious-Critical Illness and Deaths. Journal of the Endocrine Society, 2021, 5, A270-A271.	0.1	0
52	Exaggerated Premature Adrenarche in Boys: Comparative Study of a Therapeutic Intervention With the Aromatase Inhibitor Anastrozole Versus Morning Low Dose Hydrocortisone. Journal of the Endocrine Society, 2021, 5, A666-A667.	0.1	0
53	Anastrozole Improves Near Adult Height in Boys With Compromised Height Potential, as Monotherapy or in Combination With a GnRH Analogue. Journal of the Endocrine Society, 2021, 5, A673-A673.	0.1	0
54	PDB23 Investigating InternistsÃ¢â‚¬â„¢ EndocrinologistsÃ¢â‚¬â„¢ and PaediatriciansÃ¢â‚¬â„¢ Attitudes in Regards to the VALUE of Innovative Diabetes Mellitus Self-Monitoring Technologies in Greece. Value in Health, 2021, 24, S81-S82.	0.1	0

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55	SUN-028 Concomitant Mutations in the POR and AR Genes in a Boy Presenting with Micropenis and Premature Adrenarche. Journal of the Endocrine Society, 2020, 4, .	0.1	0
56	Vitamin D revisited: Individualized Vitamin D normal values according to PTH levels; incidence and treatment of Normocalcemic Hyperparathyroidism in children. Endocrine Abstracts, 0, , .	0.0	0
57	OR15-05 Gain in Near Adult Height Using the Combination of an LHRH Analogue and an Aromatase Inhibitor in Early Maturing Girls with Compromised Growth. The â€œGailâ€•Study ISRCTN11469487. Journal of the Endocrine Society, 2020, 4, .	0.1	0
58	MON-078 WFS1 Related Disorder in A 4-Month Old Girl. Journal of the Endocrine Society, 2020, 4, .	0.1	0
59	Acute Bilateral Cataract in Type 1 Diabetes Mellitus. Annals of Pediatrics & Child Health, 2015, 3, .	0.5	0