Nicos Skordis

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/2556794/publications.pdf

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15	178	1163117 8	1125743 13
papers	citations	h-index	g-index

15	15	15	235
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Central Precocious Puberty Caused by Novel Mutations in the Promoter and 5â€²-UTR Region of the Imprinted MKRN3 Gene. Frontiers in Endocrinology, 2019, 10, 677.	3.5	23
2	Molecular Defects of the CYP21A2 Gene in Greek-Cypriot Patients with Congenital Adrenal Hyperplasia. Hormone Research in Paediatrics, 2011, 75, 180-186.	1.8	22
3	GnRH Deficient Patients With Congenital Hypogonadotropic Hypogonadism: Novel Genetic Findings in ANOS1, RNF216, WDR11, FGFR1, CHD7, and POLR3A Genes in a Case Series and Review of the Literature. Frontiers in Endocrinology, 2020, 11, 626.	3.5	20
4	Endocrine profile and phenotype-genotype correlation in unrelated patients with non-classical congenital adrenal hyperplasia. Clinical Biochemistry, 2011, 44, 959-963.	1.9	18
5	Phenotypic variability of hyperandrogenemia in females heterozygous for CYP21A2 mutations. Indian Journal of Endocrinology and Metabolism, 2014, 18, 72.	0.4	17
6	Variations in the 3â€²UTR of theCYP21A2Gene in Heterozygous Females with Hyperandrogenaemia. International Journal of Endocrinology, 2017, 2017, 1-8.	1.5	12
7	Rare mutations in the CYP21A2 gene detected in congenital adrenal hyperplasia. Clinical Biochemistry, 2009, 42, 1363-1367.	1.9	11
8	Late diagnosis of 5alpha steroid-reductase deficiency due to IVS12A>G mutation of the SRD5a2 gene in an adolescent girl presented with primary amenorrhea. Hormones, 2011, 10, 230-235.	1.9	11
9	Genotype Is Associated to the Degree of Virilization in Patients With Classic Congenital Adrenal Hyperplasia. Frontiers in Endocrinology, 2018, 9, 733.	3.5	9
10	The Spectrum of Genetic Defects in Congenital Adrenal Hyperplasia in the Population of Cyprus: A Retrospective Analysis. Hormone and Metabolic Research, 2019, 51, 586-594.	1.5	8
11	46,XY complete gonadal dysgenesis in a familial case with a rare mutation in the desert hedgehog (DHH) gene. Hormones, 2019, 18, 315-320.	1.9	7
12	Pathogenic and Low-Frequency Variants in Children With Central Precocious Puberty. Frontiers in Endocrinology, 2021, 12, 745048.	3.5	7
13	Molecular modelling of novel ADCY3 variantÂpredicts a molecular target for tackling obesity. International Journal of Molecular Medicine, 2021, 49, .	4.0	7
14	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.	0.9	6
15	Polycystic ovarian syndrome in adolescents: From diagnostic criteria to therapeutic management. Acta Biomedica, 2020, 91, e2020085.	0.3	0