Ian Marshall

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/1330641/publications.pdf

Version: 2024-02-01

1163117 940533 19 676 8 16 citations h-index g-index papers 19 19 19 673 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	EXTENSIVE PERSONAL EXPERIENCE: Prenatal Diagnosis for Congenital Adrenal Hyperplasia in 532 Pregnancies. Journal of Clinical Endocrinology and Metabolism, 2001, 86, 5651-5657.	3.6	226
2	A Novel Dominant Negative Mutation of OTX2 Associated with Combined Pituitary Hormone Deficiency. Journal of Clinical Endocrinology and Metabolism, 2008, 93, 4351-4359.	3.6	121
3	Treatment with Growth Hormone and Luteinizing Hormone Releasing Hormone Analog Improves Final Adult Height in Children with Congenital Adrenal Hyperplasia. Journal of Clinical Endocrinology and Metabolism, 2005, 90, 3318-3325.	3.6	97
4	Urinary mycoestrogens, body size and breast development in New Jersey girls. Science of the Total Environment, 2011, 409, 5221-5227.	8.0	82
5	Guidelines for the Development of Comprehensive Care Centers for Congenital Adrenal Hyperplasia: Guidance from the CARES Foundation Initiative. International Journal of Pediatric Endocrinology (Springer), 2010, 2010, 275213.	1.6	52
6	Vitamin D in the maternal–fetal–neonatal interface: clinical implications and requirements for supplementation. Journal of Maternal-Fetal and Neonatal Medicine, 2013, 26, 633-638.	1.5	36
7	Youngest Reported Patient Presenting with an Androgen Producing Sclerosing Stromal Ovarian Tumor. Journal of Pediatric and Adolescent Gynecology, 2014, 27, e121-e124.	0.7	11
8	Management of congenital adrenal hyperplasia: beyond conventional glucocorticoid therapy. Current Opinion in Pediatrics, 2019, 31, 550-554.	2.0	11
9	Bilateral Sclerosing Stromal Ovarian Tumor in an Adolescent. Case Reports in Radiology, 2015, 2015, 1-4.	0.3	9
10	Another Unusual Presentation of McCune Albright Syndrome with Fibrous Dysplasia, Unilateral Testicular Enlargement, and Testicular Microlithiasis. Journal of Pediatric Endocrinology and Metabolism, 2010, 23, 513-5.	0.9	7
11	Heterozygous NPR2 Mutation in Two Family Members with Short Stature and Skeletal Dysplasia. Case Reports in Endocrinology, 2018, 2018, 1-4.	0.4	7
12	Delayed methimazole-induced agranulocytosis in a 6-year old patient with Graves' disease. International Journal of Pediatric Endocrinology (Springer), 2016, 2016, 16.	1.6	4
13	Rare Presentation of Neurofibromatosis and Turner Syndrome in a Pediatric Patient. Mental Illness, 2017, 9, 6810.	0.8	4
14	Profile of the Pediatric Endocrine Clinic at New York–Presbyterian Hospital, New York Weill Cornell Center ¹ . Journal of Clinical Endocrinology and Metabolism, 1999, 84, 4444-4449.	3.6	3
15	Pallister-Hall Syndrome Presenting in Adolescence. Case Reports in Genetics, 2019, 2019, 1-4.	0.2	3
16	Clinical Suspicion of Maturity Onset of Diabetes of the Young in Pediatric Patients Diagnosed with Diabetes Mellitus. Indian Journal of Pediatrics, 2012, 79, 955-958.	0.8	2
17	Recurrent Thyrotoxicosis due to Both Graves' Disease and Hashimoto's Thyroiditis in the Same Three Patients. Case Reports in Endocrinology, 2016, 2016, 1-4.	0.4	1
18	Mineralocorticoid Disorders, Genetic Basis of., 2003,, 679-685.		0

ARTICLE IF CITATIONS

19 Genetic Testing for Pituitary Disease., 2004, , 141-145. o