Amrit Bhangoo

List of Publications by Year
in descending order

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27 papers	552 citations	516710 16 h-index	642732 23 g-index
28 all docs	28 docs citations	28 times ranked	977 citing authors

#	Article	IF	Citations
1	Fasting Serum IGFBP-1 as a Marker of Insulin Resistance in Diverse School Age Groups. Frontiers in Endocrinology, 2022, 13, 840361.	3.5	1
2	Response to Letter to the Editor from Rosenfield: â€œGlucocorticoid Resistance in Premature Adrenarche and PCOS: From Childhood to Adulthoodâ€. Journal of the Endocrine Society, 2021, 5, bvaa166.	0.2	0
3	Glucocorticoid Resistance in Premature Adrenarche and PCOS: From Childhood to Adulthood. Journal of the Endocrine Society, 2020, 4, bvaa111.	0.2	6
4	Phenotypic spectrum and responses to recombinant human IGF1 (rhIGF1) therapy in patients with homozygous intronic pseudoexon growth hormone receptor mutation. European Journal of Endocrinology, 2018, 178, 481-489.	3.7	17
5	LHX3 deficiency presenting in the United States with severe developmental delay in a child of Syrian refugee parents. Endocrinology, Diabetes and Metabolism Case Reports, 2018, 2018, .	0.5	2
6	Case 16: Hypoglycemia in a Term Appropriate for Gestational Age Infant. , 2018, , 111-116.		0
7	Endocrinopathies in HIV, AIDS and HAART. Reviews in Endocrine and Metabolic Disorders, 2013, 14, 101-103.	5.7	3
8	Pathophysiology of GHRH-growth hormone-IGF1 axis in HIV/AIDS. Reviews in Endocrine and Metabolic Disorders, 2013, 14, 113-118.	5.7	17
9	Bone and vitamin D metabolism in HIV. Reviews in Endocrine and Metabolic Disorders, 2013, 14, 119-125.	5.7	17
10	HIV and thyroid dysfunction. Reviews in Endocrine and Metabolic Disorders, 2013, 14, 127-131.	5.7	35
11	Insulin resistance, lipodystrophy and cardiometabolic syndrome in HIV/AIDS. Reviews in Endocrine and Metabolic Disorders, 2013, 14, 133-140.	5.7	43
12	Human immune deficiency virus (HIV) infection and the hypothalamic pituitary adrenal axis. Reviews in Endocrine and Metabolic Disorders, 2013, 14, 105-112.	5.7	21
13	Role of 11Î²HSD Type 2 Enzyme Activity in Essential Hypertension and Children with Chronic Kidney Disease (CKD). Journal of Clinical Endocrinology and Metabolism, 2012, 97, 3622-3629.	3.6	23
14	Endothelial Function as Measured by Peripheral Arterial Tonometry Increases during Pubertal Advancement. Hormone Research in Paediatrics, 2011, 76, 226-233.	1.8	40
15	Isolated mild clitoral hypertrophy may reveal 46,XY disorders of sex development in infancy due to17Î²HSD-3defect confirmed by molecular analysis. Gynecological Endocrinology, 2011, 27, 890-894.	1.7	9
16	The Clinical and Molecular Heterogeneity of 17Î²HSD-3 Enzyme Deficiency. Hormone Research in Paediatrics, 2010, 74, 229-240.	1.8	93
17	Isolated micropenis reveals partial androgen insensitivity syndrome confirmed by molecular analysis. Asian Journal of Andrology, 2010, 12, 561-566.	1.6	33
18	The genetics of idiopathic hypogonadotropic hypogonadism:unraveling the biology of human sexual development. Pediatric Endocrinology Reviews, 2009, 6, 395-404.	1.2	19

#	Article	IF	Citations
19	Novel Mutation in Cytochrome P450c17 Causes Complete Combined 17Î±-Hydroxylase/17,20-Lyase Deficiency. Journal of Pediatric Endocrinology and Metabolism, 2008, 21, 185-90.	0.9	12
20	Resting Energy Expenditure in Insulin Resistance Falls with Decompensation of Insulin Secretion in Obese Children. Journal of Pediatric Endocrinology and Metabolism, 2008, 21, 359-67.	0.9	4
21	Effect of Metformin and Rosiglitazone in a Prepubertal Boy with AlstrÃ¶m Syndrome. Journal of Pediatric Endocrinology and Metabolism, 2007, 20, 1045-52.	0.9	16
22	Characterization of Insulin Resistance Syndrome in Children and Young Adults. When to Screen for Prediabetes?. Journal of Pediatric Endocrinology and Metabolism, 2007, 20, 989-99.	0.9	3
23	Role of a Founder c.201_202delCT Mutation and New Phenotypic Features of Congenital Lipoid Adrenal Hyperplasia in Palestinians. Journal of Clinical Endocrinology and Metabolism, 2007, 92, 4000-4008.	3.6	30
24	Phenotypic features of 46, XX females with StAR protein mutations. Pediatric Endocrinology Reviews, 2007, 5, 633-41.	1.2	20
25	Donor Splice Mutation in the 11Î²-Hydroxylase (CYP11B1) Gene Resulting in Sex Reversal: A Case Report and Review of the Literature. Journal of Pediatric Endocrinology and Metabolism, 2006, 19, 1267-82.	0.9	18
26	Phenotypic variations in lipoid congenital adrenal hyperplasia. Pediatric Endocrinology Reviews, 2006, 3, 258-71.	1.2	28
27	Phenotypic Features Associated with Mutations in Steroidogenic Acute Regulatory Protein. Journal of Clinical Endocrinology and Metabolism, 2005, 90, 6303-6309.	3.6	42