Stefania Pedicelli

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/8957049/publications.pdf

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15	410	932766 10	996533 15
papers	citations	h-index	g-index

15	15	15	679
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Routine Screening by Brain Magnetic Resonance Imaging Is Not Indicated in Every Girl With Onset of Puberty Between the Ages of 6 and 8 Years. Journal of Clinical Endocrinology and Metabolism, 2014, 99, 4455-4461.	1.8	77
2	Controversies in the Definition and Treatment of Idiopathic Short Stature (ISS). JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2011, 1, 105-115.	0.4	72
3	Genetic homogeneity but IgG subclassâ€“dependent clinical variability of alloimmune membranous nephropathy with anti-neutral endopeptidase antibodies. Kidney International, 2015, 87, 602-609.	2.6	57
4	Serum insulin-like growth factor-I (IGF-I) reference ranges for chemiluminescence assay in childhood and adolescence. Data from a population of in- and out-patients. Growth Hormone and IGF Research, 2012, 22, 134-138.	0.5	31
5	Water Balance and â€˜Salt Wasting' in the First Year of Life: The Role of Aldosterone-Signaling Defects. Hormone Research in Paediatrics, 2016, 86, 143-153.	0.8	31
6	Brain Magnetic Resonance Imaging as First-Line Investigation for Growth Hormone Deficiency Diagnosis in Early Childhood. Hormone Research in Paediatrics, 2015, 84, 323-330.	0.8	26
7	Glucose tolerance affects pubertal growth and final height of children with cystic fibrosis. Pediatric Pulmonology, 2015, 50, 144-149.	1.0	23
8	Prevalence of prediabetes in children and adolescents by class of obesity. Pediatric Obesity, 2022, 17, e12900.	1.4	20
9	Iatrogenic Kwashiorkor in three infants on a diet of rice beverages. Pediatric Allergy and Immunology, 2011, 22, 878-879.	1.1	18
10	Congenital primary adrenal insufficiency and selective aldosterone defects presenting as salt-wasting in infancy: a single center 10-year experience. Italian Journal of Pediatrics, 2016, 42, 73.	1.0	17
11	Early retesting by GHRHÂ+Âarginine test shows normal GH response in most children with idiopathic GH deficiency. Journal of Endocrinological Investigation, 2015, 38, 429-436.	1.8	10
12	Lipoid congenital adrenal hyperplasia by steroidogenic acute regulatory protein (STAR) gene mutation in an Italian infant: an uncommon cause of adrenal insufficiency. Italian Journal of Pediatrics, 2017, 43, 57.	1.0	9
13	25OH vitamin D levels in pediatric patients affected by Praderâ€“Willi syndrome. Journal of Endocrinological Investigation, 2018, 41, 739-742.	1.8	9
14	Responses to GHRH plus arginine test are more concordant with IGF-I circulating levels than responses to arginine and clonidine provocative tests. Journal of Endocrinological Investigation, 2012, 35, 742-7.	1.8	7
15	Treatment of boric acid overdose in two infants with Continuous Venovenous Hemodialysis. Clinical Toxicology, 2015, 53, 920-922.	0.8	3