Alessandro Salvatoni

List of Publications by Year
in descending order

33 papers	1,052 citations	471371 17 h-index	454834 30 g-index
35 all docs	35 docs citations	35 times ranked	1757 citing authors

#	Article	IF	Citations
1	Genetic Analysis of Italian Patients with Congenital Hyperinsulinism of Infancy. Hormone Research in Paediatrics, 2013, 79, 236-242.	0.8	382
2	The Italian National Survey for Praderâ€“Willi syndrome: An epidemiologic study. American Journal of Medical Genetics, Part A, 2008, 146A, 861-872.	0.7	81
3	Children with Praderâ€“Willi syndrome exhibit more evident meal-induced responses in plasma ghrelin and peptide YY levels than obese and lean children. European Journal of Endocrinology, 2010, 162, 499-505.	1.9	56
4	Growth Hormone Therapy and Respiratory Disorders: Long-Term Follow-up in PWS Children. Journal of Clinical Endocrinology and Metabolism, 2013, 98, E1516-E1523.	1.8	53
5	Assessment of central adrenal insufficiency in children and adolescents with Praderâ€“Willi syndrome. Clinical Endocrinology, 2012, 76, 843-850.	1.2	42
6	Bone mineral density in diabetic children and adolescents: a follow-up study. Bone, 2004, 34, 900-904.	1.4	38
7	Intrafamilial spread of enterovirus infections at the clinical onset of type 1 diabetes. Pediatric Diabetes, 2013, 14, 407-416.	1.2	32
8	Vitamin D status, enterovirus infection, and type 1 diabetes in Italian children/adolescents. Pediatric Diabetes, 2018, 19, 923-929.	1.2	32
9	Thyroid function in patients with Prader-Willi syndrome: an Italian multicenter study of 339 patients. Journal of Pediatric Endocrinology and Metabolism, 2019, 32, 159-165.	0.4	32
10	The Effect of Alginate in Gastroesophageal Reflux in Infants. Paediatric Drugs, 2018, 20, 575-583.	1.3	31
11	Central adrenal insufficiency in young adults with Praderâ€Willi Syndrome. Clinical Endocrinology, 2013, 79, 371-378.	1.2	29
12	The role of socio-economic and clinical factors on HbA1c in children and adolescents with type 1 diabetes: an Italian multicentre survey. Pediatric Diabetes, 2017, 18, 241-248.	1.2	28
13	Revealing enterovirus infection in chronic human disorders: An integrated diagnostic approach. Scientific Reports, 2017, 7, 5013.	1.6	28
14	Whole Genome SNP Genotyping and Exome Sequencing Reveal Novel Genetic Variants and Putative Causative Genes in Congenital Hyperinsulinism. PLoS ONE, 2013, 8, e68740.	1.1	25
15	Growth hormone secretory pattern in non-obese children and adolescents with Prader-Willi syndrome. Journal of Pediatric Endocrinology and Metabolism, 2011, 24, 477-81.	0.4	24
16	Prevalence of pathogenetic MC4R mutations in Italian children with early Onset obesity, tall stature and familial history of obesity. BMC Medical Genetics, 2009, 10, 25.	2.1	23
17	Hyponatremia and seizures during desmopressin acetate treatment in hypothyroidism. Journal of Pediatrics, 1990, 116, 835-836.	0.9	18
18	Esophageal Impedance Baseline Is Age Dependent. Journal of Pediatric Gastroenterology and Nutrition, 2013, 57, 506-513.	0.9	17

#	Article	IF	Citations
19	Oral ranitidine and duration of gastric pH >4.0 in infants with persisting reflux symptoms. Acta Paediatrica, International Journal of Paediatrics, 2006, 95, 176-181.	0.7	16
20	Ketoacidosis at diagnosis in childhood-onset diabetes and the risk of retinopathy 20years later. Journal of Diabetes and Its Complications, 2016, 30, 55-60.	1.2	11
21	Molecular cytogenetics, RFLP analysis and clinical characterization of a de novo trisomy 10p case. Annales De GÃ©nÃ©tique, 2000, 43, 45-50.	0.4	10
22	Anthropometric characteristics of newborns with Praderâ€“Willi syndrome. American Journal of Medical Genetics, Part A, 2019, 179, 2067-2074.	0.7	10
23	Enteroviruses and causality of type 1 diabetes: how close are we?. Pediatric Diabetes, 2012, 13, 92-99.	1.2	9
24	Benefits of multidisciplinary care in Prader-Willi syndrome. Expert Review of Endocrinology and Metabolism, 2021, 16, 63-71.	1.2	7
25	Hypothyroidism and sodium valproate. Journal of Pediatrics, 1983, 103, 1005-1006.	0.9	4
26	Long-term side effects of growth hormone treatment in children with Praderâ€“Willi syndrome. Expert Review of Endocrinology and Metabolism, 2014, 9, 369-375.	1.2	3
27	Uniparental disomy and pretreatment IGF-1 may predict elevated IGF-1 levels in Prader-Willi patients on GH treatment. Growth Hormone and IGF Research, 2019, 48-49, 9-15.	0.5	3
28	Application of DEXA in body composition assessment in children. Annals of Diagnostic Paediatric Pathology, 1998, 2, 49-51.	0.0	2
29	Enteroviruses in Blood. , 2013, , 143-155.		2
30	Increased parental perception of sleep disordered breathing in a cohort of infants with ALTE/BRUE events. Minerva Pediatrics, 2018, , .	0.2	2
31	A rare genetic disorder causing persistent severe neonatal hypoglycaemia the diagnostic workup. BMJ Case Reports, 2012, 2012, bcr0320125979-bcr0320125979.	0.2	1
32	Decreasing prevalence of retinopathy in childhoodâ€onset type 1 diabetes over the last decade: A comparison of two cohorts diagnosed 10â€‰years apart. Diabetes, Obesity and Metabolism, 2021, 23, 1950-1955.	2.2	1
33	The turner phenotype and the different types of human X isochromosome. Human Genetics, 1982, 62, 93-93.	1.8	0