Alessandro Salvatoni

List of Publications by Citations

33 papers	551 citations	16 h-index	23 g-index
35 ext. papers	623 ext. citations	3.4 avg, IF	2.89 L-index

#	Paper	IF	Citations
33	The Italian National Survey for Prader-Willi syndrome: an epidemiologic study. *American Journal of Medical Genetics, Part A*, **2008**, 146A, 861-72	2.5	65
32	Children with Prader-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	6.5	48
31	Growth hormone therapy and respiratory disorders: long-term follow-up in PWS children. *Journal of Clinical Endocrinology and Metabolism*, **2013**, 98, E1516-23	5.6	40
30	Assessment of central adrenal insufficiency in children and adolescents with Prader-Willi syndrome. *Clinical Endocrinology*, **2012**, 76, 843-50	3.4	36
29	Bone mineral density in diabetic children and adolescents: a follow-up study. *Bone*, **2004**, 34, 900-4	4.7	36
28	Intrafamilial spread of enterovirus infections at the clinical onset of type 1 diabetes. *Pediatric Diabetes*, **2013**, 14, 407-16	3.6	30
27	Central adrenal insufficiency in young adults with Prader-Willi syndrome. *Clinical Endocrinology*, **2013**, 79, 371-8	3.4	25
26	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	3.6	23
25	Vitamin D status, enterovirus infection, and type 1 diabetes in Italian children/adolescents. *Pediatric Diabetes*, **2018**, 19, 923-929	3.6	22
24	Whole genome SNP genotyping and exome sequencing reveal novel genetic variants and putative causative genes in congenital hyperinsulinism. *PLoS ONE*, **2013**, 8, e68740	3.7	21
23	Revealing enterovirus infection in chronic human disorders: An integrated diagnostic approach. *Scientific Reports*, **2017**, 7, 5013	4.9	20
22	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	19
21	The Effect of Alginate in Gastroesophageal Reflux in Infants. *Paediatric Drugs*, **2018**, 20, 575-583	4.2	19
20	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	1.6	18
19	Growth hormone secretory pattern in non-obese children and adolescents with Prader-Willi syndrome. *Journal of Pediatric Endocrinology and Metabolism*, **2011**, 24, 477-81	1.6	18
18	Hyponatremia and seizures during desmopressin acetate treatment in hypothyroidism. *Journal of Pediatrics*, **1990**, 116, 835-6	3.6	17
17	Esophageal impedance baseline is age dependent. *Journal of Pediatric Gastroenterology and Nutrition*, **2013**, 57, 506-13	2.8	16

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16	Genetic analysis of Italian patients with congenital hyperinsulinism of infancy. *Hormone Research in Paediatrics*, **2013**, 79, 236-42	3.3	14
15	Oral ranitidine and duration of gastric pH >4.0 in infants with persisting reflux symptoms. *Acta Paediatrica, International Journal of Paediatrics*, **2006**, 95, 176-81	3.1	13
14	Ketoacidosis at diagnosis in childhood-onset diabetes and the risk of retinopathy 20years later. *Journal of Diabetes and Its Complications*, **2016**, 30, 55-60	3.2	9
13	Enteroviruses and causality of type 1 diabetes: how close are we?. *Pediatric Diabetes*, **2012**, 13, 92-9	3.6	9
12	Molecular cytogenetics, RFLP analysis and clinical characterization of a de novo trisomy 10p case. *Annales De G□□ique*, **2000**, 43, 45-50		9
11	Anthropometric characteristics of newborns with Prader-Willi syndrome. *American Journal of Medical Genetics, Part A*, **2019**, 179, 2067-2074	2.5	6
10	Hypothyroidism and sodium valproate. *Journal of Pediatrics*, **1983**, 103, 1005-6	3.6	4
9	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	2	3
8	Benefits of multidisciplinary care in Prader-Willi syndrome. *Expert Review of Endocrinology and Metabolism*, **2021**, 16, 63-71	4.1	3
7	Long-term side effects of growth hormone treatment in children with Prader-Willi syndrome. *Expert Review of Endocrinology and Metabolism*, **2014**, 9, 369-375	4.1	2
6	Application of DEXA in body composition assessment in children **1998**, 2, 49-51		2
5	Increased parental perception of sleep disordered breathing in a cohort of infants with ALTE/BRUE events. *Minerva Pediatrics*, **2018**,	1.5	2
4	Enteroviruses in Blood **2013**, 143-155		1
3	A rare genetic disorder causing persistent severe neonatal hypoglycaemia the diagnostic workup. *BMJ Case Reports*, **2012**, 2012,	0.9	1
2	The Turner phenotype and the different types of human X isochromosome. *Human Genetics*, **1982**, 62, 93	6.3	
1	Decreasing prevalence of retinopathy in childhood-onset type 1 diabetes over the last decade: A comparison of two cohorts diagnosed 10 years apart. *Diabetes, Obesity and Metabolism*, **2021**, 23, 1950-1955	6.7	