

# Alessandro Salvatoni

## List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

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	Benefits of multidisciplinary care in Prader-Willi syndrome. <i>Expert Review of Endocrinology and Metabolism</i> , <b>2021</b> , 16, 63-71	4.1	3
32	Decreasing prevalence of retinopathy in childhood-onset type 1 diabetes over the last decade: A comparison of two cohorts diagnosed 10 years apart. <i>Diabetes, Obesity and Metabolism</i> , <b>2021</b> , 23, 1950-1955	6.7	67
31	Uniparental disomy and pretreatment IGF-1 may predict elevated IGF-1 levels in Prader-Willi patients on GH treatment. <i>Growth Hormone and IGF Research</i> , <b>2019</b> , 48-49, 9-15	2	3
30	Thyroid function in patients with Prader-Willi syndrome: an Italian multicenter study of 339 patients. <i>Journal of Pediatric Endocrinology and Metabolism</i> , <b>2019</b> , 32, 159-165	1.6	18
29	Anthropometric characteristics of newborns with Prader-Willi syndrome. <i>American Journal of Medical Genetics, Part A</i> , <b>2019</b> , 179, 2067-2074	2.5	6
28	Vitamin D status, enterovirus infection, and type 1 diabetes in Italian children/adolescents. <i>Pediatric Diabetes</i> , <b>2018</b> , 19, 923-929	3.6	22
27	Increased parental perception of sleep disordered breathing in a cohort of infants with ALTE/BRUE events. <i>Minerva Pediatrics</i> , <b>2018</b> ,	1.5	2
26	The Effect of Alginate in Gastroesophageal Reflux in Infants. <i>Paediatric Drugs</i> , <b>2018</b> , 20, 575-583	4.2	19
25	The role of socio-economic and clinical factors on HbA1c in children and adolescents with type 1 diabetes: an Italian multicentre survey. <i>Pediatric Diabetes</i> , <b>2017</b> , 18, 241-248	3.6	23
24	Revealing enterovirus infection in chronic human disorders: An integrated diagnostic approach. <i>Scientific Reports</i> , <b>2017</b> , 7, 5013	4.9	20
23	Ketoacidosis at diagnosis in childhood-onset diabetes and the risk of retinopathy 20years later. <i>Journal of Diabetes and Its Complications</i> , <b>2016</b> , 30, 55-60	3.2	9
22	Long-term side effects of growth hormone treatment in children with Prader-Willi syndrome. <i>Expert Review of Endocrinology and Metabolism</i> , <b>2014</b> , 9, 369-375	4.1	2
21	Enteroviruses in Blood <b>2013</b> , 143-155		1
20	Intrafamilial spread of enterovirus infections at the clinical onset of type 1 diabetes. <i>Pediatric Diabetes</i> , <b>2013</b> , 14, 407-16	3.6	30
19	Esophageal impedance baseline is age dependent. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , <b>2013</b> , 57, 506-13	2.8	16
18	Growth hormone therapy and respiratory disorders: long-term follow-up in PWS children. <i>Journal of Clinical Endocrinology and Metabolism</i> , <b>2013</b> , 98, E1516-23	5.6	40
17	Genetic analysis of Italian patients with congenital hyperinsulinism of infancy. <i>Hormone Research in Paediatrics</i> , <b>2013</b> , 79, 236-42	3.3	14

16	Central adrenal insufficiency in young adults with Prader-Willi syndrome. <i>Clinical Endocrinology</i> , <b>2013</b> , 79, 371-8	3.4	25
15	Whole genome SNP genotyping and exome sequencing reveal novel genetic variants and putative causative genes in congenital hyperinsulinism. <i>PLoS ONE</i> , <b>2013</b> , 8, e68740	3.7	21
14	Assessment of central adrenal insufficiency in children and adolescents with Prader-Willi syndrome. <i>Clinical Endocrinology</i> , <b>2012</b> , 76, 843-50	3.4	36
13	A rare genetic disorder causing persistent severe neonatal hypoglycaemia the diagnostic workup. <i>BMJ Case Reports</i> , <b>2012</b> , 2012,	0.9	1
12	Enteroviruses and causality of type 1 diabetes: how close are we?. <i>Pediatric Diabetes</i> , <b>2012</b> , 13, 92-9	3.6	9
11	Growth hormone secretory pattern in non-obese children and adolescents with Prader-Willi syndrome. <i>Journal of Pediatric Endocrinology and Metabolism</i> , <b>2011</b> , 24, 477-81	1.6	18
10	Children with Prader-Willi syndrome exhibit more evident meal-induced responses in plasma ghrelin and peptide YY levels than obese and lean children. <i>European Journal of Endocrinology</i> , <b>2010</b> , 162, 499-505	6.5	48
9	Prevalence of pathogenetic MC4R mutations in Italian children with early onset obesity, tall stature and familial history of obesity. <i>BMC Medical Genetics</i> , <b>2009</b> , 10, 25	2.1	19
8	The Italian National Survey for Prader-Willi syndrome: an epidemiologic study. <i>American Journal of Medical Genetics, Part A</i> , <b>2008</b> , 146A, 861-72	2.5	65
7	Oral ranitidine and duration of gastric pH >4.0 in infants with persisting reflux symptoms. <i>Acta Paediatrica, International Journal of Paediatrics</i> , <b>2006</b> , 95, 176-81	3.1	13
6	Bone mineral density in diabetic children and adolescents: a follow-up study. <i>Bone</i> , <b>2004</b> , 34, 900-4	4.7	36
5	Molecular cytogenetics, RFLP analysis and clinical characterization of a de novo trisomy 10p case. <i>Annales De Génétique</i> , <b>2000</b> , 43, 45-50		9
4	Application of DEXA in body composition assessment in children <b>1998</b> , 2, 49-51		2
3	Hyponatremia and seizures during desmopressin acetate treatment in hypothyroidism. <i>Journal of Pediatrics</i> , <b>1990</b> , 116, 835-6	3.6	17
2	Hypothyroidism and sodium valproate. <i>Journal of Pediatrics</i> , <b>1983</b> , 103, 1005-6	3.6	4
1	The Turner phenotype and the different types of human X isochromosome. <i>Human Genetics</i> , <b>1982</b> , 62, 93	6.3	