Maurizio Delvecchio

List of Publications by Year
in descending order

103	2,479	218592	243529
papers	citations	26 h-index	44 g-index

108	108	108	3364
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Linear growth and puberty in childhood obesity: what is new?. Minerva Pediatrics, 2022, 73, .	0.2	2
2	Polycystic ovary syndrome in pediatric obesity and diabetes. Minerva Pediatrics, 2022, 73, .	0.2	1
3	Comment on â€œReal-World Use of a New Hybrid Closed Loop Improves Glycemic Control in Youth with Type 1 Diabetesâ€•by Messer et al.. Diabetes Technology and Therapeutics, 2022, 24, 455-457.	2.4	2
4	Isolated childhood growth hormone deficiency: a 30-year experience on final height and a new prediction model. Journal of Endocrinological Investigation, 2022, , 1.	1.8	0
5	Uric acid and cardiometabolic risk by gender in youth with type 1 diabetes. Scientific Reports, 2022, 12, .	1.6	0
6	Stimulated GH levels during the transition phase in Praderâ€“Willi syndrome. Journal of Endocrinological Investigation, 2021, 44, 1465-1474.	1.8	7
7	Differences between transient neonatal diabetes mellitus subtypes can guide diagnosis and therapy. European Journal of Endocrinology, 2021, 184, 575-585.	1.9	13
8	Editorial: Debates in Clinical Management in Pediatric Endocrinology. Frontiers in Endocrinology, 2021, 12, 663860.	1.5	1
9	Inflammatory Status and Glycemic Control Level of Patients with Type 2 Diabetes and Periodontitis: A Randomized Clinical Trial. International Journal of Environmental Research and Public Health, 2021, 18, 3018.	1.2	18
10	Circulating Inhibitory Factor 1 levels in adult patients with Praderâ€“Willi syndrome. Hormone Molecular Biology and Clinical Investigation, 2021, 42, 317-320.	0.3	1
11	Iodine Absorption in Celiac Children: A Longitudinal Pilot Study. Nutrients, 2021, 13, 808.	1.7	7
12	Clinical Spectrum Associated with Wolfram Syndrome Type 1 and Type 2: A Review on Genotypeâ€“Phenotype Correlations. International Journal of Environmental Research and Public Health, 2021, 18, 4796.	1.2	22
13	Albuminuric and non-albuminuric reduced eGFR phenotypes in youth with type 1 diabetes: Factors associated with cardiometabolic risk. Nutrition, Metabolism and Cardiovascular Diseases, 2021, 31, 2033-2041.	1.1	7
14	Beyond the barriers of the use of continuous subcutaneous insulin therapy in type 1 diabetes: a new opportunity from catheter-less insulin pumps. AboutOpen, 2021, 8, 55-70.	0.2	1
15	Cardiometabolic risk in childhood cancer survivors. Minerva Pediatrics, 2021, , .	0.2	2
16	Relationships between HbA1c and continuous glucose monitoring metrics of glycaemic control and glucose variability in a large cohort of children and adolescents with type 1 diabetes. Diabetes Research and Clinical Practice, 2021, 177, 108933.	1.1	12
17	A Novel Genetic Variant in the WFS1 Gene in a Patient with Partial Uniparental Mero-Isodisomy of Chromosome 4. International Journal of Molecular Sciences, 2021, 22, 8082.	1.8	1
18	EEG Patterns in Patients with Praderâ€“Willi Syndrome. Brain Sciences, 2021, 11, 1045.	1.1	3

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19	Effectiveness of a closedâ€loop control system and a virtual educational camp for children and adolescents with type 1 diabetes: A prospective, multicentre, realâ€life study. Diabetes, Obesity and Metabolism, 2021, 23, 2484-2491.	2.2	18
20	Vascular and Myocardial Function in Young People with Type 1 Diabetes Mellitus: Insulin Pump Therapy Versus Multiple Daily Injections Insulin Regimen. Experimental and Clinical Endocrinology and Diabetes, 2021, , .	0.6	1
21	Telemedicine and <scp>COVID</scp> â€19 pandemic: The perfect storm to mark a change in diabetes care. Results from a worldâ€wide crossâ€sectional webâ€based survey. Pediatric Diabetes, 2021, 22, 1115-1119.	1.2	31
22	High Glycemic Variability Is Associated with Worse Continuous Glucose Monitoring Metrics in Children and Adolescents with Type 1 Diabetes. Hormone Research in Paediatrics, 2021, 94, 369-373.	0.8	5
23	Caring and living with Prader-Willi syndrome in Italy: integrating children, adults and parentsâ€™ experiences through a multicentre narrative medicine research. BMJ Open, 2020, 10, e036502.	0.8	13
24	Dietary cholesterol supplementation and inhibitory factor 1 serum levels in two dizygotic Smith-Lemli-Opitz syndrome twins: a case report. Italian Journal of Pediatrics, 2020, 46, 161.	1.0	1
25	The Effect of Gaseous Ozone Therapy in Conjunction with Periodontal Treatment on Glycated Hemoglobin Level in Subjects with Type 2 Diabetes Mellitus: An Unmasked Randomized Controlled Trial. International Journal of Environmental Research and Public Health, 2020, 17, 5467.	1.2	26
26	Has COVID-19 Delayed the Diagnosis and Worsened the Presentation of Type 1 Diabetes in Children?. Diabetes Care, 2020, 43, 2870-2872.	4.3	182
27	Longâ€term glycemic control and glucose variability assessed with continuous glucose monitoring in a pediatric population with type 1 diabetes: Determination of optimal sampling duration. Pediatric Diabetes, 2020, 21, 1485-1492.	1.2	17
28	Cardiovascular Implications in Idiopathic and Syndromic Obesity in Childhood: An Update. Frontiers in Endocrinology, 2020, 11, 330.	1.5	8
29	Treatment Options for MODY Patients: A Systematic Review of Literature. Diabetes Therapy, 2020, 11, 1667-1685.	1.2	81
30	Can Anti-Thyroid Antibodies Influence the Outcome of Primary Chronic Immune Thrombocytopenia in Children?. Endocrine, Metabolic and Immune Disorders - Drug Targets, 2020, 20, 351-355.	0.6	6
31	1264-P: Distinguishing between Obese Patients with Type 1 Diabetes (T1DMob) and Type 2 Diabetes in Adolescence (T2DMad) at Presentation. Diabetes, 2020, 69, .	0.3	1
32	Anthropometric characteristics of newborns with Praderâ€“Willi syndrome. American Journal of Medical Genetics, Part A, 2019, 179, 2067-2074.	0.7	10
33	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
34	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
35	Alagille Syndrome: A Novel Mutation in JAG1 Gene. Frontiers in Pediatrics, 2019, 7, 199.	0.9	13
36	Cardiovascular dysfunction and vitamin D status in childhood acute lymphoblastic leukemia survivors. World Journal of Pediatrics, 2019, 15, 465-470.	0.8	7

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37	Growth hormone treatment improves final height and nutritional status of children with chronic kidney disease and growth deceleration. Journal of Endocrinological Investigation, 2018, 41, 325-331.	1.8	4
38	Analysis of Circulating Mediators of Bone Remodeling in Praderâ€“Willi Syndrome. Calcified Tissue International, 2018, 102, 635-643.	1.5	19
39	Autoimmune pituitary involvement in Praderâ€“Willi syndrome: new perspective for further research. Endocrine, 2018, 62, 733-736.	1.1	13
40	Can HbA1c combined with fasting plasma glucose help to assess priority for GCK-MODY vs HNF1A-MODY genetic testing?. Acta Diabetologica, 2018, 55, 981-983.	1.2	14
41	Effectiveness and safety of long-term treatment with sulfonylureas in patients with neonatal diabetes due to KCNJ11 mutations: an international cohort study. Lancet Diabetes and Endocrinology,the, 2018, 6, 637-646.	5.5	120
42	Metabolic Outcomes, Bone Health, and Risk of Polycystic Ovary Syndrome in Girls with Idiopathic Central Precocious Puberty Treated with Gonadotropin-Releasing Hormone Analogues. Hormone Research in Paediatrics, 2017, 87, 162-169.	0.8	25
43	Non-alcoholic fatty liver disease is associated with early left ventricular dysfunction in childhood acute lymphoblastic leukaemia survivors. European Journal of Endocrinology, 2017, 176, 111-121.	1.9	8
44	Endothelial dysfunction and cardiovascular risk factors in childhood acute lymphoblastic leukemia survivors. International Journal of Cardiology, 2017, 228, 621-627.	0.8	40
45	High Sclerostin and Dickkopf-1 (DKK-1) Serum Levels in Children and Adolescents With Type 1 Diabetes Mellitus. Journal of Clinical Endocrinology and Metabolism, 2017, 102, 1174-1181.	1.8	67
46	Monogenic Diabetes Accounts for 6.3% of Cases Referred to 15 Italian Pediatric Diabetes Centers During 2007 to 2012. Journal of Clinical Endocrinology and Metabolism, 2017, 102, 1826-1834.	1.8	88
47	A Multicenter Retrospective Survey regarding Diabetic Ketoacidosis Management in Italian Children with Type 1 Diabetes. Journal of Diabetes Research, 2016, 2016, 1-6.	1.0	28
48	Incidence of Type 1 Diabetes among Children and Adolescents in Italy between 2009 and 2013: The Role of a Regional Childhood Diabetes Registry. Journal of Diabetes Research, 2016, 2016, 1-7.	1.0	19
49	Disorders of glucose metabolism in Praderâ€“Willi syndrome: Results of a multicenter Italian cohort study. Nutrition, Metabolism and Cardiovascular Diseases, 2016, 26, 842-847.	1.1	51
50	High frequency of diabetic ketoacidosis at diagnosis of type 1 diabetes in Italian children: a nationwide longitudinal study, 2004â€“2013. Scientific Reports, 2016, 6, 38844.	1.6	26
51	A novel OTX2 gene frameshift mutation in a child with microphthalmia, ectopic pituitary and growth hormone deficiency. Journal of Pediatric Endocrinology and Metabolism, 2016, 29, 603-5.	0.4	5
52	Vascular Function and Myocardial Performance Indices in Children Born Small for Gestational Age. Circulation Journal, 2016, 80, 958-963.	0.7	25
53	<scp>CHARGE</scp> syndrome and common variable immunodeficiency: A case report and review of literature. Pediatric Allergy and Immunology, 2016, 27, 546-550.	1.1	5
54	Impaired bone remodeling in children with osteogenesis imperfecta treated and untreated with bisphosphonates: the role of DKK1, RANKL, and TNF-Î±. Osteoporosis International, 2016, 27, 2355-2365.	1.3	52

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55	Final height in Italian patients with congenital hypothyroidism detected by neonatal screening: a 20-year observational study. Italian Journal of Pediatrics, 2015, 41, 82.	1.0	13
56	Prevalence of positive atopy patch test in an unselected pediatric population. Clinical and Molecular Allergy, 2015, 13, 2.	0.8	8
57	Microdeletion of 12q24.31: Report of a girl with intellectual disability, stereotypies, seizures and facial dysmorphisms. American Journal of Medical Genetics, Part A, 2015, 167, 438-444.	0.7	32
58	Continuous Subcutaneous Insulin Infusion in Italy: Third National Survey. Diabetes Technology and Therapeutics, 2015, 17, 96-104.	2.4	18
59	Metabolic syndrome in childhood leukemia survivors: a meta-analysis. Endocrine, 2015, 49, 353-360.	1.1	14
60	Levothyroxine requirement in congenital hypothyroidism: a 12-year longitudinal study. Endocrine, 2015, 50, 674-680.	1.1	12
61	Survey on etiological diagnosis of diabetes in 1244 Italian diabetic children and adolescents: Impact of access to genetic testing. Diabetes Research and Clinical Practice, 2015, 107, e15-e18.	1.1	24
62	Evaluation of impact of steroid replacement treatment on bone health in children with 21-hydroxylase deficiency. Endocrine, 2015, 48, 995-1000.	1.1	10
63	Factors associated with different results of allergy tests in children with dust mite-induced atopic dermatitis. Allergologia Et Immunopathologia, 2015, 43, 238-242.	1.0	1
64	Low Prevalence of HNF1A Mutations After Molecular Screening of Multiple MODY Genes in 58 Italian Families Recruited in the Pediatric or Adult Diabetes Clinic From a Single Italian Hospital. Diabetes Care, 2014, 37, e258-e260.	4.3	23
65	Prolactin May Be Increased in Newly Diagnosed Celiac Children and Adolescents and Decreases after 6 Months of Gluten-Free Diet. Hormone Research in Paediatrics, 2014, 81, 309-313.	0.8	15
66	A novel CISD2 intragenic deletion, optic neuropathy and platelet aggregation defect in Wolfram syndrome type 2. BMC Medical Genetics, 2014, 15, 88.	2.1	59
67	Clinical heterogeneity of abnormal glucose homeostasis associated with the HNF4A R311H mutation. Italian Journal of Pediatrics, 2014, 40, 58.	1.0	3
68	Increasing burden, younger age at onset and worst metabolic control in migrant than in Italian children with type 1 diabetes: an emerging problem in pediatric clinics. Acta Diabetologica, 2014, 51, 263-267.	1.2	14
69	Graves Disease in Children: Thyroid-Stimulating Hormone Receptor Antibodies as Remission Markers. Journal of Pediatrics, 2014, 164, 1189-1194.e1.	0.9	46
70	MODY type 2 P59S GCK mutant: founder effect in South of Italy. Clinical Genetics, 2013, 83, 83-87.	1.0	4
71	Identification of Candidate Children for Maturity-Onset Diabetes of the Young Type 2 (MODY2) Gene Testing: A Seven-Item Clinical Flowchart (7-iF). PLoS ONE, 2013, 8, e79933.	1.1	33
72	Involvement of Hypothalamus Autoimmunity in Patients with Autoimmune Hypopituitarism: Role of Antibodies to Hypothalamic Cells. Journal of Clinical Endocrinology and Metabolism, 2012, 97, 3684-3690.	1.8	61

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73	Emerging Effects of Early Environmental Factors over Genetic Background for Type 1 Diabetes Susceptibility: Evidence from a Nationwide Italian Twin Study. Journal of Clinical Endocrinology and Metabolism, 2012, 97, E1483-E1491.	1.8	39
74	Potential celiac disease in type 1 diabetes: A multicenter study. Diabetes Research and Clinical Practice, 2011, 92, 53-56.	1.1	26
75	Use of Integrated Real-Time Continuous Glucose Monitoring/Insulin Pump System in Children and Adolescents with Type 1 Diabetes: A 3-Year Follow-Up Study. Diabetes Technology and Therapeutics, 2011, 13, 99-103.	2.4	26
76	Coeliac children treated for growth hormone deficiency reach normal final height. Clinical Endocrinology, 2011, 74, 791-792.	1.2	6
77	Detection of antipituitary and antihypothalamus antibodies to investigate the role of pituitary or hypothalamic autoimmunity in patients with selective idiopathic hypopituitarism. Clinical Endocrinology, 2011, 75, 361-366.	1.2	56
78	Metabolic, inflammatory, endothelial and haemostatic markers in a group of Italian obese children and adolescents. European Journal of Pediatrics, 2011, 170, 845-850.	1.3	76
79	Permanent diabetes during the first year of life: multiple gene screening in 54 patients. Diabetologia, 2011, 54, 1693-1701.	2.9	63
80	Anti-Pituitary Antibodies in Children With Newly Diagnosed Celiac Disease: A Novel Finding Contributing to Linear-Growth Impairment. American Journal of Gastroenterology, 2010, 105, 691-696.	0.2	41
81	Sleep-Disordered Breathing in Obese Children. Chest, 2010, 137, 1085-1090.	0.4	20
82	IGF2 Gene Variants and Risk of Hypertension in Obese Children and Adolescents. Pediatric Research, 2010, 67, 340-344.	1.1	36
83	Predictive Role of the Immunostaining Pattern of Immunofluorescence and the Titers of Antipituitary Antibodies at Presentation for the Occurrence of Autoimmune Hypopituitarism in Patients with Autoimmune Polyendocrine Syndromes over a Five-Year Follow-Up. Journal of Clinical Endocrinology and Metabolism, 2010, 95, 3750-3757.	1.8	56
84	Growth and endocrine function in thalassemia major in childhood and adolescence. Journal of Endocrinological Investigation, 2010, 33, 61-68.	1.8	46
85	Thyroid function and thyroid autoimmunity in childhood acute lymphoblastic leukemia off-therapy patients treated only with chemotherapy. Journal of Endocrinological Investigation, 2010, 33, 135-139.	1.8	9
86	Metabolic syndrome in children with Praderâ€“Willi syndrome: the effect of obesity. Nutrition, Metabolism and Cardiovascular Diseases, 2010, 21, 269-76.	1.1	46
87	Growth hormone deficiency and antipituitary antibodies in a patient with common variable immunodeficiency. Journal of Endocrinological Investigation, 2009, 32, 637-640.	1.8	14
88	Effects of moderate-severe exercise on blood glucose in Type 1 diabetic adolescents treated with insulin pump or glargine insulin. Journal of Endocrinological Investigation, 2009, 32, 519-524.	1.8	15
89	Sulfonylurea treatment in a girl with neonatal diabetes (KCNJ11 R201H) and celiac disease: Impact of low compliance to the gluten free diet. Diabetes Research and Clinical Practice, 2009, 84, 332-334.	1.1	3
90	Acute Pancreatitis in a Girl with Panhypopituitarism Due to Craniopharyngioma on Growth Hormone Treatment. A Combination of Risk Factors. Hormone Research in Paediatrics, 2009, 71, 372-375.	0.8	3

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91	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
92	Longitudinal Assessment of Levo-Thyroxine Therapy for Congenital Hypothyroidism: Relationship with Aetiology, Bone Maturation and Biochemical Features. Hormone Research in Paediatrics, 2007, 68, 105-112.	0.8	15
93	Growth Hormone Treatment in Prepubertal Children With Celiac Disease and Growth Hormone Deficiency. Journal of Pediatric Gastroenterology and Nutrition, 2007, 45, 433-437.	0.9	17
94	Factors predicting final height in early treated congenital hypothyroid patients. Clinical Endocrinology, 2006, 65, 693-697.	1.2	26
95	The Prevalence of Growth Hormone Deficiency and Celiac Disease in Short Children. Clinical Medicine and Research, 2006, 4, 180-183.	0.4	45
96	Isolated increased serum TSH response to TRH is prevalent in celiac disease and predicts poor response to treatment. Gastroenterologie Clinique Et Biologique, 2005, 29, 1063-1064.	0.9	0
97	Transient neonatal diabetes mellitus is associated with a recurrent (R201H) KCNJ11 (KIR6.2) mutation. Diabetologia, 2005, 48, 2439-2441.	2.9	41
98	Final height in short polytransfused thalassemia major patients treated with recombinant growth hormone. Journal of Endocrinological Investigation, 2005, 28, 363-366.	1.8	17
99	SODIUM FRACTION EXCRETION RATE IN NOCTURNAL ENURESIS CORRELATES WITH NOCTURNAL POLYURIA AND OSMOLALITY. Journal of Urology, 2004, 171, 2567-2570.	0.2	21
100	In congenital hypothyroidism bone maturation at birth may be a predictive factor of psychomotor development during the first Year of life irrespective of other variables related to treatment. European Journal of Endocrinology, 2003, 149, 1-6.	1.9	65
101	Assessment of the hypothalamus-pituitary-adrenal axis with different corticotropin tests in adult patients with Prader-Willi syndrome. Endocrine Abstracts, 0, , .	0.0	0
102	Higher serum levels of the Wnt-signaling antagonist DKK1 in obese respect to Prader-Willi syndrome. Endocrine Abstracts, 0, , .	0.0	0
103	The Silent Epidemic of Diabetic Ketoacidosis at Diagnosis of Type 1 Diabetes in Children and Adolescents in Italy During the COVID-19 Pandemic in 2020. Frontiers in Endocrinology, 0, 13, .	1.5	9