Maurizio Delvecchio

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	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
2	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
3	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
4	The Italian National Survey for Prader–Willi syndrome: An epidemiologic study. American Journal of Medical Genetics, Part A, 2008, 146A, 861-872.	0.7	81
5	Treatment Options for MODY Patients: A Systematic Review of Literature. Diabetes Therapy, 2020, 11, 1667-1685.	1.2	81
6	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
7	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
8	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
9	Permanent diabetes during the first year of life: multiple gene screening in 54 patients. Diabetologia, 2011, 54, 1693-1701.	2.9	63
10	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
11	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
12	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
13	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
14	Impaired bone remodeling in children with osteogenesis imperfecta treated and untreated with bisphosphonates: the role of DKK1, RANKL, and TNF-α. Osteoporosis International, 2016, 27, 2355-2365.	1.3	52
15	Disorders of glucose metabolism in Prader–Willi syndrome: Results of a multicenter Italian cohort study. Nutrition, Metabolism and Cardiovascular Diseases, 2016, 26, 842-847.	1.1	51
16	Growth and endocrine function in thalassemia major in childhood and adolescence. Journal of Endocrinological Investigation, 2010, 33, 61-68.	1.8	46
17	Metabolic syndrome in children with Prader–Willi syndrome: the effect of obesity. Nutrition, Metabolism and Cardiovascular Diseases, 2010, 21, 269-76.	1.1	46
18	Graves Disease in Children: Thyroid-Stimulating Hormone Receptor Antibodies as Remission Markers. Journal of Pediatrics, 2014, 164, 1189-1194.e1.	0.9	46

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19	The Prevalence of Growth Hormone Deficiency and Celiac Disease in Short Children. Clinical Medicine and Research, 2006, 4, 180-183.	0.4	45
20	Transient neonatal diabetes mellitus is associated with a recurrent (R201H) KCNJ11 (KIR6.2) mutation. Diabetologia, 2005, 48, 2439-2441.	2.9	41
21	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
22	Endothelial dysfunction and cardiovascular risk factors in childhood acute lymphoblastic leukemia survivors. International Journal of Cardiology, 2017, 228, 621-627.	0.8	40
23	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
24	IGF2 Gene Variants and Risk of Hypertension in Obese Children and Adolescents. Pediatric Research, 2010, 67, 340-344.	1.1	36
25	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
26	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
27	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
28	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
29	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
30	Factors predicting final height in early treated congenital hypothyroid patients. Clinical Endocrinology, 2006, 65, 693-697.	1.2	26
31	Potential celiac disease in type 1 diabetes: A multicenter study. Diabetes Research and Clinical Practice, 2011, 92, 53-56.	1.1	26
32	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
33	High frequency of diabetic ketoacidosis at diagnosis of type 1 diabetes in Italian children: a nationwide longitudinal study, 2004–2013. Scientific Reports, 2016, 6, 38844.	1.6	26
34	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
35	Vascular Function and Myocardial Performance Indices in Children Born Small for Gestational Age. Circulation Journal, 2016, 80, 958-963.	0.7	25
36	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

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37	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
38	Low Prevalence of <i>HNF1A</i> 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
39	Clinical Spectrum Associated with Wolfram Syndrome Type 1 and Type 2: A Review on Genotype–Phenotype Correlations. International Journal of Environmental Research and Public Health, 2021, 18, 4796.	1.2	22
40	SODIUM FRACTION EXCRETION RATE IN NOCTURNAL ENURESIS CORRELATES WITH NOCTURNAL POLYURIA AND OSMOLALITY. Journal of Urology, 2004, 171, 2567-2570.	0.2	21
41	Sleep-Disordered Breathing in Obese Children. Chest, 2010, 137, 1085-1090.	0.4	20
42	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
43	Analysis of Circulating Mediators of Bone Remodeling in Prader–Willi Syndrome. Calcified Tissue International, 2018, 102, 635-643.	1.5	19
44	Continuous Subcutaneous Insulin Infusion in Italy: Third National Survey. Diabetes Technology and Therapeutics, 2015, 17, 96-104.	2.4	18
45	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
46	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
47	Final height in short polytransfused thalassemia major patients treated with recombinant growth hormone. Journal of Endocrinological Investigation, 2005, 28, 363-366.	1.8	17
48	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
49	Longâ€ŧerm 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
50	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
51	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
52	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
53	Growth hormone deficiency and antipituitary antibodies in a patient with common variable immunodeficiency. Journal of Endocrinological Investigation, 2009, 32, 637-640.	1.8	14
54	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

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55	Metabolic syndrome in childhood leukemia survivors: a meta-analysis. Endocrine, 2015, 49, 353-360.	1.1	14
56	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
57	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
58	Autoimmune pituitary involvement in Prader–Willi syndrome: new perspective for further research. Endocrine, 2018, 62, 733-736.	1.1	13
59	Alagille Syndrome: A Novel Mutation in JAC1 Gene. Frontiers in Pediatrics, 2019, 7, 199.	0.9	13
60	Caring and living with Prader-Willi syndrome in Italy: integrating children, adults and parents' experiences through a multicentre narrative medicine research. BMJ Open, 2020, 10, e036502.	0.8	13
61	Differences between transient neonatal diabetes mellitus subtypes can guide diagnosis and therapy. European Journal of Endocrinology, 2021, 184, 575-585.	1.9	13
62	Levothyroxine requirement in congenital hypothyroidism: a 12-year longitudinal study. Endocrine, 2015, 50, 674-680.	1.1	12
63	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
64	Evaluation of impact of steroid replacement treatment on bone health in children with 21-hydroxylase deficiency. Endocrine, 2015, 48, 995-1000.	1.1	10
65	Anthropometric characteristics of newborns with Prader–Willi syndrome. American Journal of Medical Genetics, Part A, 2019, 179, 2067-2074.	0.7	10
66	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
67	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
68	Prevalence of positive atopy patch test in an unselected pediatric population. Clinical and Molecular Allergy, 2015, 13, 2.	0.8	8
69	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
70	Cardiovascular Implications in Idiopathic and Syndromic Obesity in Childhood: An Update. Frontiers in Endocrinology, 2020, 11, 330.	1.5	8
71	Cardiovascular dysfunction and vitamin D status in childhood acute lymphoblastic leukemia survivors. World Journal of Pediatrics, 2019, 15, 465-470.	0.8	7
72	Stimulated GH levels during the transition phase in Prader–Willi syndrome. Journal of Endocrinological Investigation, 2021, 44, 1465-1474.	1.8	7

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73	lodine Absorption in Celiac Children: A Longitudinal Pilot Study. Nutrients, 2021, 13, 808.	1.7	7
74	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
75	Coeliac children treated for growth hormone deficiency reach normal final height. Clinical Endocrinology, 2011, 74, 791-792.	1.2	6
76	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
77	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
78	<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
79	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
80	MODY type 2 P59S GCK mutant: founder effect in South of Italy. Clinical Genetics, 2013, 83, 83-87.	1.0	4
81	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
82	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
83	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
84	Clinical heterogeneity of abnormal glucose homeostasis associated with the HNF4A R311H mutation. Italian Journal of Pediatrics, 2014, 40, 58.	1.0	3
85	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
86	EEG Patterns in Patients with Prader–Willi Syndrome. Brain Sciences, 2021, 11, 1045.	1.1	3
87	Cardiometabolic risk in childhood cancer survivors. Minerva Pediatrics, 2021, , .	0.2	2
88	Linear growth and puberty in childhood obesity: what is new?. Minerva Pediatrics, 2022, 73, .	0.2	2
89	Comment on "Real-World Use of a New Hybrid Closed Loop Improves Glycemic Control in Youth with Type 1 Diabetes―by Messer et al Diabetes Technology and Therapeutics, 2022, 24, 455-457. 	2.4	2
90	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

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91	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
92	Editorial: Debates in Clinical Management in Pediatric Endocrinology. Frontiers in Endocrinology, 2021, 12, 663860.	1.5	1
93	Circulating Inhibitory Factor 1 levels in adult patients with Prader–Willi syndrome. Hormone Molecular Biology and Clinical Investigation, 2021, 42, 317-320.	0.3	1
94	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
95	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
96	Polycystic ovary syndrome in pediatric obesity and diabetes. Minerva Pediatrics, 2022, 73, .	0.2	1
97	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
98	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
99	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	Ο
100	Assessment of the hypothalamus-pituitary-adrenal axis with different corticotropin tests in adult patients with Prader-Willi syndrome. Endocrine Abstracts, 0, , .	0.0	0
101	Higher serum levels of the Wnt-signaling antagonist DKK1 in obese respect to Prader-Willi syndrome. Endocrine Abstracts, 0, , .	0.0	Ο
102	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
103	Uric acid and cardiometabolic risk by gender in youth with type 1 diabetes. Scientific Reports, 2022, 12, .	1.6	0