

# Maria F Faienza

## List of Publications by Citations

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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.

129  
papers

2,268  
citations

27  
h-index

39  
g-index

140  
ext. papers

2,857  
ext. citations

4.4  
avg, IF

4.74  
L-index

#	Paper	IF	Citations
129	SOX2 plays a critical role in the pituitary, forebrain, and eye during human embryonic development. <i>Journal of Clinical Endocrinology and Metabolism</i> , <b>2008</b> , 93, 1865-73	5.6	130
128	Postmenopausal osteoporosis: the role of immune system cells. <i>Clinical and Developmental Immunology</i> , <b>2013</b> , 2013, 575936		96
127	SOS1 mutations in Noonan syndrome: molecular spectrum, structural insights on pathogenic effects, and genotype-phenotype correlations. <i>Human Mutation</i> , <b>2011</b> , 32, 760-72	4.7	82
126	The dangerous link between childhood and adulthood predictors of obesity and metabolic syndrome. <i>Internal and Emergency Medicine</i> , <b>2016</b> , 11, 175-82	3.7	66
125	Oxidative stress in obesity and metabolic syndrome in children and adolescents. <i>Hormone Research in Paediatrics</i> , <b>2012</b> , 78, 158-64	3.3	65
124	Metabolic, inflammatory, endothelial and haemostatic markers in a group of Italian obese children and adolescents. <i>European Journal of Pediatrics</i> , <b>2011</b> , 170, 845-50	4.1	61
123	Analysis of cyclin-dependent kinase inhibitor genes (CDKN2A, CDKN2B, and CDKN2C) in childhood rhabdomyosarcoma. <i>Genes Chromosomes and Cancer</i> , <b>1996</b> , 15, 217-22	5	54
122	Osteoporosis and obesity: Role of Wnt pathway in human and murine models. <i>World Journal of Orthopedics</i> , <b>2014</b> , 5, 242-6	2.2	48
121	LIGHT/TNFSF14 increases osteoclastogenesis and decreases osteoblastogenesis in multiple myeloma-bone disease. <i>Oncotarget</i> , <b>2014</b> , 5, 12950-67	3.3	47
120	High Sclerostin and Dickkopf-1 (DKK-1) Serum Levels in Children and Adolescents With Type 1 Diabetes Mellitus. <i>Journal of Clinical Endocrinology and Metabolism</i> , <b>2017</b> , 102, 1174-1181	5.6	45
119	Nonalcoholic fatty liver disease in prepubertal children born small for gestational age: influence of rapid weight catch-up growth. <i>Hormone Research in Paediatrics</i> , <b>2013</b> , 79, 103-9	3.3	43
118	(TA)8 allele in the UGT1A1 gene promoter of a Caucasian with Gilbert's syndrome. <i>Haematologica</i> , <b>1999</b> , 84, 106-9	6.6	43
117	Impaired bone remodeling in children with osteogenesis imperfecta treated and untreated with bisphosphonates: the role of DKK1, RANKL, and TNF- $\alpha$ . <i>Osteoporosis International</i> , <b>2016</b> , 27, 2355-2365	5.3	40
116	Osteoclastogenesis in children with 21-hydroxylase deficiency on long-term glucocorticoid therapy: the role of receptor activator of nuclear factor-kappaB ligand/osteoprotegerin imbalance. <i>Journal of Clinical Endocrinology and Metabolism</i> , <b>2009</b> , 94, 2269-76	5.6	40
115	Transient neonatal diabetes mellitus is associated with a recurrent (R201H) KCNJ11 (KIR6.2) mutation. <i>Diabetologia</i> , <b>2005</b> , 48, 2439-41	10.3	39
114	17beta-Hydroxysteroid dehydrogenase-3 deficiency: from pregnancy to adolescence. <i>Journal of Endocrinological Investigation</i> , <b>2009</b> , 32, 666-70	5.2	36
113	Clinical, endocrine, and molecular findings in 17beta-hydroxysteroid dehydrogenase type 3 deficiency. <i>Journal of Endocrinological Investigation</i> , <b>2008</b> , 31, 85-91	5.2	36

112	Metabolic Bone Disease of Prematurity: Diagnosis and Management. <i>Frontiers in Pediatrics</i> , <b>2019</b> , 7, 143-148	3.4	35
111	High irisin levels are associated with better glycemic control and bone health in children with Type 1 diabetes. <i>Diabetes Research and Clinical Practice</i> , <b>2018</b> , 141, 10-17	7.4	35
110	Monoclonal antibodies for treating osteoporosis. <i>Expert Opinion on Biological Therapy</i> , <b>2018</b> , 18, 149-157	5.4	35
109	High dickkopf-1 levels in sera and leukocytes from children with 21-hydroxylase deficiency on chronic glucocorticoid treatment. <i>American Journal of Physiology - Endocrinology and Metabolism</i> , <b>2013</b> , 304, E546-54	6	33
108	Glucocorticoid-induced osteoporosis in children with 21-hydroxylase deficiency. <i>BioMed Research International</i> , <b>2013</b> , 2013, 250462	3	32
107	Irisin serum levels are positively correlated with bone mineral status in a population of healthy children. <i>Pediatric Research</i> , <b>2019</b> , 85, 484-488	3.2	30
106	Risk factors for subclinical atherosclerosis in diabetic and obese children. <i>International Journal of Medical Sciences</i> , <b>2013</b> , 10, 338-43	3.7	29
105	Uric acid: from a biological advantage to a potential danger. A focus on cardiovascular effects. <i>Vascular Pharmacology</i> , <b>2019</b> , 120, 106565	5.9	28
104	The association with Turner syndrome significantly affects the course of Hashimoto's thyroiditis in children, irrespective of karyotype. <i>Endocrine</i> , <b>2015</b> , 50, 777-82	4	27
103	IGF2 gene variants and risk of hypertension in obese children and adolescents. <i>Pediatric Research</i> , <b>2010</b> , 67, 340-4	3.2	27
102	Homozygous deletions of cyclin-dependent kinase inhibitor genes, p16(INK4A) and p18, in childhood T cell lineage acute lymphoblastic leukemias. <i>Leukemia</i> , <b>1996</b> , 10, 255-60	10.7	26
101	Bone Fragility in Turner Syndrome: Mechanisms and Prevention Strategies. <i>Frontiers in Endocrinology</i> , <b>2016</b> , 7, 34	5.7	25
100	Insulin resistance and endothelial function in children and adolescents. <i>International Journal of Cardiology</i> , <b>2014</b> , 174, 343-7	3.2	24
99	PTPN11 gene mutation and severe neonatal hypertrophic cardiomyopathy: what is the link?. <i>Pediatric Cardiology</i> , <b>2009</b> , 30, 1012-5	2.1	23
98	Mechanisms of enhanced osteoclastogenesis in girls and young women with Turner Syndrome. <i>Bone</i> , <b>2015</b> , 81, 228-236	4.7	22
97	A novel heterozygous SOX2 mutation causing congenital bilateral anophthalmia, hypogonadotropic hypogonadism and growth hormone deficiency. <i>Gene</i> , <b>2014</b> , 534, 282-5	3.8	22
96	Endothelial dysfunction and cardiovascular risk factors in childhood acute lymphoblastic leukemia survivors. <i>International Journal of Cardiology</i> , <b>2017</b> , 228, 621-627	3.2	22
95	Skeleton and glucose metabolism: a bone-pancreas loop. <i>International Journal of Endocrinology</i> , <b>2015</b> , 2015, 758148	2.7	22

94	Peculiarities of presentation and evolution over time of Hashimoto's thyroiditis in children and adolescents with Down syndrome. <i>Hormones</i> , <b>2015</b> , 14, 410-6	3.1	22
93	Neonatal hyperbilirubinemia and Gilbert syndrome. <i>Journal of Perinatal Medicine</i> , <b>2002</b> , 30, 166-9	2.7	22
92	Genotype-phenotype correlation in juvenile Paget disease: role of molecular alterations of the TNFRSF11B gene. <i>Endocrine</i> , <b>2012</b> , 42, 266-71	4	21
91	Mechanisms Involved in Childhood Obesity-Related Bone Fragility. <i>Frontiers in Endocrinology</i> , <b>2019</b> , 10, 269	5.7	20
90	Role of antithyroid autoimmunity as a predictive biomarker of chronic immune thrombocytopenia. <i>Pediatric Blood and Cancer</i> , <b>2019</b> , 66, e27452	3	20
89	How Physical Activity across the Lifespan Can Reduce the Impact of Bone Ageing: A Literature Review. <i>International Journal of Environmental Research and Public Health</i> , <b>2020</b> , 17,	4.6	19
88	Novel insights in health-promoting properties of sweet cherries. <i>Journal of Functional Foods</i> , <b>2020</b> , 69, 103945-103945	5.1	19
87	Issues in pediatric haemophilia care. <i>Italian Journal of Pediatrics</i> , <b>2013</b> , 39, 24	3.2	19
86	Osteoblasts display different responsiveness to TRAIL-induced apoptosis during their differentiation process. <i>Cell Biochemistry and Biophysics</i> , <b>2013</b> , 67, 1127-36	3.2	19
85	LIGHT/TNFSF14 Promotes Osteolytic Bone Metastases in Non-small Cell Lung Cancer Patients. <i>Journal of Bone and Mineral Research</i> , <b>2020</b> , 35, 671-680	6.3	19
84	Bone health in children and adolescents with steroid-sensitive nephrotic syndrome assessed by DXA and QUS. <i>Pediatric Nephrology</i> , <b>2014</b> , 29, 2147-55	3.2	17
83	Elevated endothelin-1 (ET-1) levels may contribute to hypoadiponectinemia in childhood obesity. <i>Journal of Clinical Endocrinology and Metabolism</i> , <b>2013</b> , 98, E683-93	5.6	17
82	Interstitial and large chromosome 1p deletion occurs in localized and disseminated neuroblastomas and predicts an unfavourable outcome. <i>Cancer Letters</i> , <b>1998</b> , 130, 83-92	9.9	17
81	Factors predicting final height in early treated congenital hypothyroid patients. <i>Clinical Endocrinology</i> , <b>2006</b> , 65, 693-7	3.4	17
80	An update on the role of RANKL-RANK/osteoprotegerin and WNT- $\beta$ -catenin signaling pathways in pediatric diseases. <i>World Journal of Pediatrics</i> , <b>2019</b> , 15, 4-11	4.6	17
79	Pilot study on circulating miRNA signature in children with obesity born small for gestational age and appropriate for gestational age. <i>Pediatric Obesity</i> , <b>2018</b> , 13, 803-811	4.6	17
78	Mechanisms of Enhanced Osteoclastogenesis in Alkaptonuria. <i>American Journal of Pathology</i> , <b>2018</b> , 188, 1059-1068	5.8	16
77	Vascular Function and Myocardial Performance Indices in Children Born Small for Gestational Age. <i>Circulation Journal</i> , <b>2016</b> , 80, 958-63	2.9	16

76	High serum sclerostin levels in children with haemophilia A. <i>British Journal of Haematology</i> , <b>2016</b> , 172, 293-5	4.5	16
75	GilbertQ syndrome and jaundice in glucose-6-phosphate dehydrogenase deficient neonates. <i>Haematologica</i> , <b>1999</b> , 84, 99-102	6.6	16
74	Autoimmune haematological disorders in two Italian children with Kabuki syndrome. <i>Italian Journal of Pediatrics</i> , <b>2014</b> , 40, 10	3.2	15
73	Prolactin may be increased in newly diagnosed celiac children and adolescents and decreases after 6 months of gluten-free diet. <i>Hormone Research in Paediatrics</i> , <b>2014</b> , 81, 309-13	3.3	15
72	Effects of moderate-severe exercise on blood glucose in Type 1 diabetic adolescents treated with insulin pump or glargine insulin. <i>Journal of Endocrinological Investigation</i> , <b>2009</b> , 32, 519-24	5.2	15
71	Metabolic Outcomes, Bone Health, and Risk of Polycystic Ovary Syndrome in Girls with Idiopathic Central Precocious Puberty Treated with Gonadotropin-Releasing Hormone Analogues. <i>Hormone Research in Paediatrics</i> , <b>2017</b> , 87, 162-169	3.3	14
70	High frequency of homozygous deletions of CDK4I gene in childhood acute lymphoblastic leukaemia. <i>British Journal of Haematology</i> , <b>1995</b> , 91, 647-51	4.5	14
69	Metabolic syndrome in childhood leukemia survivors: a meta-analysis. <i>Endocrine</i> , <b>2015</b> , 49, 353-60	4	13
68	Childhood obesity, cardiovascular and liver health: a growing epidemic with age. <i>World Journal of Pediatrics</i> , <b>2020</b> , 16, 438-445	4.6	13
67	Endothelial and Metabolic Function Interactions in Overweight/Obese Children. <i>Journal of Atherosclerosis and Thrombosis</i> , <b>2016</b> , 23, 950-9	4	13
66	Bilirubin levels in the acute hemolytic crisis of G6PD deficiency are related to GilbertQ syndrome. <i>European Journal of Haematology</i> , <b>1999</b> , 62, 307-10	3.8	12
65	Frequent clonal loss of heterozygosity (LOH) in the chromosomal region 1p32 occurs in childhood T cell acute lymphoblastic leukemia (T-ALL) carrying rearrangements of the TAL1 gene. <i>Leukemia</i> , <b>1997</b> , 11, 359-63	10.7	12
64	High expression of TRAIL by osteoblastic differentiated dental pulp stem cells affects myeloma cell viability. <i>Oncology Reports</i> , <b>2018</b> , 39, 2031-2039	3.5	12
63	LIGHT/TNFSF14 as a New Biomarker of Bone Disease in Multiple Myeloma Patients Experiencing Therapeutic Regimens. <i>Frontiers in Immunology</i> , <b>2018</b> , 9, 2459	8.4	12
62	Analysis of Circulating Mediators of Bone Remodeling in Prader-Willi Syndrome. <i>Calcified Tissue International</i> , <b>2018</b> , 102, 635-643	3.9	12
61	Expression of cell cycle regulatory genes in chronic myelogenous leukemia. <i>Haematologica</i> , <b>1998</b> , 83, 771-7	6.6	12
60	Effects of Sweet Cherry Polyphenols on Enhanced Osteoclastogenesis Associated With Childhood Obesity. <i>Frontiers in Immunology</i> , <b>2019</b> , 10, 1001	8.4	11
59	Evidence that fibrinolytic changes in paediatric obesity translate into a hypofibrinolytic state: relative contribution of TAFI and PAI-1. <i>Thrombosis and Haemostasis</i> , <b>2012</b> , 108, 311-7	7	11

58	A novel HESX1 splice mutation causes isolated GH deficiency by interfering with mRNA processing. <i>European Journal of Endocrinology</i> , <b>2011</b> , 164, 705-13	6.5	11
57	Longitudinal assessment of levo-thyroxine therapy for congenital hypothyroidism: relationship with aetiology, bone maturation and biochemical features. <i>Hormone Research in Paediatrics</i> , <b>2007</b> , 68, 105-12	3.3	11
56	Polyphenols and obesity prevention: critical insights on molecular regulation, bioavailability and dose in preclinical and clinical settings. <i>Critical Reviews in Food Science and Nutrition</i> , <b>2021</b> , 61, 1804-1826	11.5	11
55	Analysis of endothelial protein C receptor gene and metabolic profile in Prader-Willi syndrome and obese subjects. <i>Obesity</i> , <b>2012</b> , 20, 1866-70	8	10
54	Impairment of Bone Remodeling in LIGHT/TNFSF14-Deficient Mice. <i>Journal of Bone and Mineral Research</i> , <b>2018</b> , 33, 704-719	6.3	10
53	P16INK4A gene homozygous deletions in human acute leukaemias with alterations of chromosome 9. <i>British Journal of Haematology</i> , <b>1996</b> , 93, 632-6	4.5	9
52	Deletion of the Transcription Factor PGC-1 $\beta$ in Mice Negatively Regulates Bone Mass. <i>Calcified Tissue International</i> , <b>2018</b> , 103, 638-652	3.9	8
51	Caring and living with Prader-Willi syndrome in Italy: integrating children, adults and parents' experiences through a multicentre narrative medicine research. <i>BMJ Open</i> , <b>2020</b> , 10, e036502	3	8
50	Evaluation of impact of steroid replacement treatment on bone health in children with 21-hydroxylase deficiency. <i>Endocrine</i> , <b>2015</b> , 48, 995-1000	4	7
49	LIGHT/TNFSF14 regulates estrogen deficiency-induced bone loss. <i>Journal of Pathology</i> , <b>2020</b> , 250, 440-451	4.1	7
48	Thyroid function and thyroid autoimmunity in childhood acute lymphoblastic leukemia off-therapy patients treated only with chemotherapy. <i>Journal of Endocrinological Investigation</i> , <b>2010</b> , 33, 135-9	5.2	7
47	Prospective evaluation of autoimmune and non-autoimmune subclinical hypothyroidism in Down syndrome children. <i>European Journal of Endocrinology</i> , <b>2020</b> , 182, 385-392	6.5	7
46	UGT1 promoter polymorphism accounts for increased neonatal appearance of hereditary spherocytosis. <i>Blood</i> , <b>1998</b> , 91, 1093	2.2	7
45	Levothyroxine requirement in congenital hypothyroidism: a 12-year longitudinal study. <i>Endocrine</i> , <b>2015</b> , 50, 674-80	4	6
44	Final height in Italian patients with congenital hypothyroidism detected by neonatal screening: a 20-year observational study. <i>Italian Journal of Pediatrics</i> , <b>2015</b> , 41, 82	3.2	6
43	Multiplying effects of COVID-19 lockdown on metabolic risk and fatty liver. <i>European Journal of Clinical Investigation</i> , <b>2021</b> , 51, e13597	4.6	6
42	17 $\beta$ hydroxysteroid dehydrogenase type 3 deficiency: female sex assignment and follow-up. <i>Journal of Endocrinological Investigation</i> , <b>2020</b> , 43, 1711-1716	5.2	6
41	Effect of recombinant insulin-like growth factor-1 treatment on short-term linear growth in a child with Majewski osteodysplastic primordial dwarfism type II and hepatic insufficiency. <i>Journal of Pediatric Endocrinology and Metabolism</i> , <b>2013</b> , 26, 771-4	1.6	5

40	The p53 family member p73 modulates the proliferative role of IGFBP3 in short children born small for gestational age. <i>Molecular Biology of the Cell</i> , <b>2015</b> , 26, 2733-41	3.5	5
39	Osteoclastogenic potential of peripheral blood mononuclear cells in cleidocranial dysplasia. <i>International Journal of Medical Sciences</i> , <b>2014</b> , 11, 356-64	3.7	5
38	Can Anti-Thyroid Antibodies Influence the Outcome of Primary Chronic Immune Thrombocytopenia in Children?. <i>Endocrine, Metabolic and Immune Disorders - Drug Targets</i> , <b>2020</b> , 20, 351-355	2.2	5
37	Venous Thromboembolism in Children: From Diagnosis to Management. <i>International Journal of Environmental Research and Public Health</i> , <b>2020</b> , 17,	4.6	5
36	Regulation of IGFBP3 gene expression in short children born small for gestational age. <i>Growth Hormone and IGF Research</i> , <b>2011</b> , 21, 349-55	2	4
35	Growth Trajectory and Adult Height in Children with Nonclassical Congenital Adrenal Hyperplasia. <i>Hormone Research in Paediatrics</i> , <b>2020</b> , 93, 173-181	3.3	4
34	The genetic background and vitamin D supplementation can affect irisin levels in Prader-Willi syndrome. <i>Journal of Endocrinological Investigation</i> , <b>2021</b> , 44, 2261-2271	5.2	4
33	A novel OTX2 gene frameshift mutation in a child with microphthalmia, ectopic pituitary and growth hormone deficiency. <i>Journal of Pediatric Endocrinology and Metabolism</i> , <b>2016</b> , 29, 603-5	1.6	4
32	NR5A1 Gene Variants: Variable Phenotypes, New Variants, Different Outcomes. <i>Sexual Development</i> , <b>2019</b> , 13, 258-263	1.6	4
31	Non-alcoholic fatty liver disease is associated with early left ventricular dysfunction in childhood acute lymphoblastic leukaemia survivors. <i>European Journal of Endocrinology</i> , <b>2017</b> , 176, 111-121	6.5	3
30	Growth hormone treatment improves final height and nutritional status of children with chronic kidney disease and growth deceleration. <i>Journal of Endocrinological Investigation</i> , <b>2018</b> , 41, 325-331	5.2	3
29	Osteotropic Cancers: From Primary Tumor to Bone. <i>Clinical Reviews in Bone and Mineral Metabolism</i> , <b>2013</b> , 11, 94-102	2.5	3
28	Treatment of osteoporosis in children with glucocorticoid-treated diseases. <i>Expert Review of Endocrinology and Metabolism</i> , <b>2014</b> , 9, 525-534	4.1	3
27	Growth plate gene involvement and isolated short stature. <i>Endocrine</i> , <b>2021</b> , 71, 28-34	4	3
26	Cardiovascular dysfunction and vitamin D status in childhood acute lymphoblastic leukemia survivors. <i>World Journal of Pediatrics</i> , <b>2019</b> , 15, 465-470	4.6	2
25	Ductal size indexed to weight and body surface area correlates with morbidities in preterm infants B2 weeks. <i>Journal of Maternal-Fetal and Neonatal Medicine</i> , <b>2021</b> , 34, 3133-3139	2	2
24	Sulfonylurea treatment in a girl with neonatal diabetes (KCNJ11 R201H) and celiac disease: impact of low compliance to the gluten free diet. <i>Diabetes Research and Clinical Practice</i> , <b>2009</b> , 84, 332-4	7.4	2
23	Physical Activity Modulating Lipid Metabolism in Gallbladder Diseases. <i>Journal of Gastrointestinal and Liver Diseases</i> , <b>2020</b> , 29, 99-110	1.4	2



22	Red blood cell transfusions and potentially related morbidities in neonates under 32 weeksQ gestation. <i>Blood Transfusion</i> , <b>2021</b> , 19, 113-119	3.6	2
21	Monitoring and maintaining bone health in patients with Turner syndrome. <i>Expert Review of Endocrinology and Metabolism</i> , <b>2020</b> , 15, 431-438	4.1	2
20	Extra Virgin Olive Oil Extracts Modulate the Inflammatory Ability of Murine Dendritic Cells Based on Their Polyphenols Pattern: Correlation between Chemical Composition and Biological Function. <i>Antioxidants</i> , <b>2021</b> , 10,	7.1	2
19	Mechanisms of altered bone remodeling in children with type 1 diabetes. <i>World Journal of Diabetes</i> , <b>2021</b> , 12, 997-1009	4.7	2
18	Bone mineral density surveillance for childhood, adolescent, and young adult cancer survivors: evidence-based recommendations from the International Late Effects of Childhood Cancer Guideline Harmonization Group. <i>Lancet Diabetes and Endocrinology</i> , <b>2021</b> , 9, 622-637	18.1	2
17	Mechanisms of Altered Bone Remodeling in Multiple Myeloma. <i>Clinical Reviews in Bone and Mineral Metabolism</i> , <b>2017</b> , 15, 151-161	2.5	1
16	Best determinants of nonalcoholic fatty liver disease and intra-abdominal fat in prepubertal children born small for gestational age: ultrasound technique versus anthropometric data. <i>Hormone Research in Paediatrics</i> , <b>2013</b> , 80, 135-6	3.3	1
15	Acute pancreatitis in a girl with panhypopituitarism due to craniopharyngioma on growth hormone treatment. A combination of risk factors. <i>Hormone Research in Paediatrics</i> , <b>2009</b> , 71, 372-5	3.3	1
14	Cardiovascular, Brain, and Lung Involvement in a Newborn With a Novel FLNA Mutation: A Case Report and Literature Review. <i>Advances in Neonatal Care</i> , <b>2021</b> ,	2	1
13	Mechanisms of Bone Impairment in Sickle Bone Disease. <i>International Journal of Environmental Research and Public Health</i> , <b>2021</b> , 18,	4.6	1
12	The Hyperphagia Questionnaire: Insights From a Multicentric Validation Study in Individuals With Prader Willi Syndrome.. <i>Frontiers in Pediatrics</i> , <b>2022</b> , 10, 829486	3.4	1
11	Thrombopoietin Receptor Agonists in Children with Immune Thrombocytopenia: A New Therapeutic Era. <i>Endocrine, Metabolic and Immune Disorders - Drug Targets</i> , <b>2021</b> , 21, 397-406	2.2	0
10	Role of Wnt-signaling inhibitors DKK-1 and sclerostin in bone fragility associated with Turner syndrome.. <i>Journal of Endocrinological Investigation</i> , <b>2022</b> , 1	5.2	0
9	Growth in Children With Noonan Syndrome and Effects of Growth Hormone Treatment on Adult Height.. <i>Frontiers in Endocrinology</i> , <b>2021</b> , 12, 761171	5.7	0
8	Unusual ultrasonographic finding of ovarian hyperstimulation syndrome in a preterm newborn with severe intra-uterine growth retardation. <i>Journal of Paediatrics and Child Health</i> , <b>2019</b> , 55, 1139-1141	1.3	
7	To test or not to test—this is the problem. <i>Journal of Pediatrics</i> , <b>2011</b> , 159, 168-9; author reply 169	3.6	
6	The use of quantitative ultrasound in a tertiary-level children hospital: role in the follow-up of chronically ill patients.. <i>Journal of Ultrasound</i> , <b>2022</b> , 1	3.4	
5	The role of N-terminal pro-B type natriuretic peptide in subjects with idiopathic sudden sensorineural hearing loss: a paradigm of its vascular function. <i>Journal of Cardiovascular Medicine</i> , <b>2020</b> , 21, 620-621	1.9	



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|---|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----|
| 4 | Antithyroid treatment improves thrombocytopenia in a young patient with Graves Disease. <i>Acta Biomedica</i> , <b>2020</b> , 91, e2020194                                         | 3.2 |
| 3 | Alteration of cell division cycle regulation in human cancers: The role of CDKN2A gene <b>1998</b> , 129-138                                                                       |     |
| 2 | Detection of PittHopkins Syndrome Based on Morphological Facial Features. <i>Applied Sciences (Switzerland)</i> , <b>2021</b> , 11, 12086                                          | 2.6 |
| 1 | Isolated childhood growth hormone deficiency: a 30-year experience on final height and a new prediction model.. <i>Journal of Endocrinological Investigation</i> , <b>2022</b> , 1 | 5.2 |