Mohamad Maghnie

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 136
 4,335
 35
 63

 papers
 citations
 h-index
 g-index

 143
 5,168
 5.5
 4.86

 ext. papers
 ext. citations
 avg, IF
 L-index

#	Paper	IF	Citations
136	Expert consensus document: European Consensus Statement on congenital hypogonadotropic hypogonadismpathogenesis, diagnosis and treatment. <i>Nature Reviews Endocrinology</i> , 2015 , 11, 547-64	4 ^{15.2}	462
135	Central diabetes insipidus in children and young adults. New England Journal of Medicine, 2000, 343, 99	18 5 150207	' 314
134	Corticotropin tests for hypothalamic-pituitary- adrenal insufficiency: a metaanalysis. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2008 , 93, 4245-53	5.6	284
133	Diagnosis and management of Silver-Russell syndrome: first international consensus statement. <i>Nature Reviews Endocrinology</i> , 2017 , 13, 105-124	15.2	224
132	Growth hormone (GH) deficiency (GHD) of childhood onset: reassessment of GH status and evaluation of the predictive criteria for permanent GHD in young adults. <i>Journal of Clinical Endocrinology and Metabolism</i> , 1999 , 84, 1324-8	5.6	156
131	Diabetes insipidusdiagnosis and management. Hormone Research in Paediatrics, 2012, 77, 69-84	3.3	154
130	Growth hormone-releasing hormone resistance in pseudohypoparathyroidism type ia: new evidence for imprinting of the Gs alpha gene. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2003 , 88, 4070-4	5.6	121
129	Hypothalamic-pituitary dysfunction in growth hormone-deficient patients with pituitary abnormalities. <i>Journal of Clinical Endocrinology and Metabolism</i> , 1991 , 73, 79-83	5.6	97
128	Cross-sectional reference data for phalangeal quantitative ultrasound from early childhood to young-adulthood according to gender, age, skeletal growth, and pubertal development. <i>Bone</i> , 2006 , 39, 159-73	4.7	94
127	Genetic analysis and evaluation of resistance to thyrotropin and growth hormone-releasing hormone in pseudohypoparathyroidism type Ib. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2007 , 92, 3738-42	5.6	78
126	Diagnosis of GH deficiency in the transition period: accuracy of insulin tolerance test and insulin-like growth factor-I measurement. <i>European Journal of Endocrinology</i> , 2005 , 152, 589-96	6.5	78
125	Central diabetes insipidus in children and young adults: etiological diagnosis and long-term outcome of idiopathic cases. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2014 , 99, 1264-72	5.6	73
124	Thyroid function and structure are affected in childhood obesity. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2008 , 93, 4749-54	5.6	72
123	Results of early reevaluation of growth hormone secretion in short children with apparent growth hormone deficiency. <i>Journal of Pediatrics</i> , 2002 , 140, 445-9	3.6	72
122	Cut-off limits of the GH response to GHRH plus arginine test and IGF-I levels for the diagnosis of GH deficiency in late adolescents and young adults. <i>European Journal of Endocrinology</i> , 2007 , 157, 701-8	6.5	68
121	Novel HESX1 mutations associated with a life-threatening neonatal phenotype, pituitary aplasia, but normally located posterior pituitary and no optic nerve abnormalities. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2006 , 91, 4528-36	5.6	65
120	Idiopathic central diabetes insipidus in children and young adults is commonly associated with vasopressin-cell antibodies and markers of autoimmunity. <i>Clinical Endocrinology</i> , 2006 , 65, 470-8	3.4	61

(2004-2013)

119	Magnetic resonance imaging of CNS in 15,043 children with GH deficiency in KIGS (Pfizer International Growth Database). <i>European Journal of Endocrinology</i> , 2013 , 168, 211-7	6.5	59	
118	Hypopituitarism and stalk agenesis: a congenital syndrome worsened by breech delivery?. <i>Hormone Research</i> , 1991 , 35, 104-8		55	
117	Characteristics of a nationwide cohort of patients presenting with isolated hypogonadotropic hypogonadism (IHH). <i>European Journal of Endocrinology</i> , 2018 , 178, 23-32	6.5	54	
116	TRAIP promotes DNA damage response during genome replication and is mutated in primordial dwarfism. <i>Nature Genetics</i> , 2016 , 48, 36-43	36.3	53	
115	The use of neuroimaging for assessing disorders of pituitary development. <i>Clinical Endocrinology</i> , 2012 , 76, 161-76	3.4	51	
114	Inaccuracy of insulin-like growth factor (IGF) binding protein (IGFBP)-3 assessment in the diagnosis of growth hormone (GH) deficiency from childhood to young adulthood: association to low GH dependency of IGF-II and presence of circulating IGFBP-3 18-kilodalton fragment. <i>Journal of Clinical</i>	5.6	49	
113	Molecular analysis of the GNAS1 gene for the correct diagnosis of Albright hereditary osteodystrophy and pseudohypoparathyroidism. <i>Pediatric Research</i> , 2003 , 53, 749-55	3.2	48	
112	Reassessment of the growth hormone status in young adults with childhood-onset growth hormone deficiency: reappraisal of insulin tolerance testing. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2009 , 94, 4195-204	5.6	47	
111	Recombinant human GH replacement therapy in children with pseudohypoparathyroidism type Ia: first study on the effect on growth. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2010 , 95, 5011-7	5.6	46	
110	The diagnosis of children with central diabetes insipidus. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2007 , 20, 359-75	1.6	45	
109	A longitudinal PRINTO study on growth and puberty in juvenile systemic lupus erythematosus. <i>Annals of the Rheumatic Diseases</i> , 2012 , 71, 511-7	2.4	44	
108	Adult height in patients with permanent growth hormone deficiency with and without multiple pituitary hormone deficiencies. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2006 , 91, 2900-5	5.6	44	
107	The glucagon test in the diagnosis of growth hormone deficiency in children with short stature younger than 6 years. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2009 , 94, 4251-7	5.6	43	
106	Dynamic endocrine testing and magnetic resonance imaging in the long-term follow-up of childhood langerhans cell histiocytosis. <i>Journal of Clinical Endocrinology and Metabolism</i> , 1998 , 83, 3089	9-54	43	
105	GHRH plus arginine in the diagnosis of acquired GH deficiency of childhood-onset. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2002 , 87, 2740-4	5.6	41	
104	Deterioration of growth hormone (GH) response and anterior pituitary function in young adults with childhood-onset GH deficiency and ectopic posterior pituitary: a two-year prospective follow-up study. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2007 , 92, 3875-84	5.6	38	
103	Pitfalls in the diagnosis of central adrenal insufficiency in children. <i>Endocrine Development</i> , 2010 , 17, 96-107		36	
102	Idiopathic central diabetes insipidus is associated with abnormal blood supply to the posterior pituitary gland caused by vascular impairment of the inferior hypophyseal artery system. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2004 , 89, 1891-6	5.6	36	

101	JAG1 Loss-Of-Function Variations as a Novel Predisposing Event in the Pathogenesis of Congenital Thyroid Defects. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2016 , 101, 861-70	5.6	35
100	Relationship of CYP21A2 genotype and serum 17-hydroxyprogesterone and cortisol levels in a large cohort of Italian children with premature pubarche. <i>European Journal of Endocrinology</i> , 2011 , 165, 307-14	6.5	34
99	Management of diabetes insipidus and adipsia in the child. <i>Best Practice and Research in Clinical Endocrinology and Metabolism</i> , 2015 , 29, 415-36	6.5	32
98	Transient beneficial effect of GH replacement therapy and topical GH application on skin ulcers in a boy with prolidase deficiency. <i>Pediatric Dermatology</i> , 2000 , 17, 227-30	1.9	32
97	Cut-off limits of the peak GH response to stimulation tests for the diagnosis of GH deficiency in children and adolescents: study in patients with organic GHD. <i>European Journal of Endocrinology</i> , 2016 , 175, 41-7	6.5	32
96	Classical and non-classical causes of GH deficiency in the paediatric age. <i>Best Practice and Research in Clinical Endocrinology and Metabolism</i> , 2016 , 30, 705-736	6.5	31
95	Pituitary stalk thickening on MRI: when is the best time to re-scan and how long should we continue re-scanning for?. <i>Clinical Endocrinology</i> , 2015 , 83, 449-55	3.4	29
94	The accuracy of the glucagon test compared to the insulin tolerance test in the diagnosis of adrenal insufficiency in young children with growth hormone deficiency. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2010 , 95, 2132-9	5.6	29
93	19q13.11 cryptic deletion: description of two new cases and indication for a role of WTIP haploinsufficiency in hypospadias. <i>European Journal of Human Genetics</i> , 2012 , 20, 852-6	5.3	29
92	Central adrenal insufficiency in children and adolescents. <i>Best Practice and Research in Clinical Endocrinology and Metabolism</i> , 2018 , 32, 425-444	6.5	28
91	Safety Outcomes During Pediatric GH Therapy: Final Results From the Prospective GeNeSIS Observational Program. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2019 , 104, 379-389	5.6	28
90	PREVALENCE AND CORRELATES OF ADHERENCE IN CHILDREN AND ADOLESCENTS TREATED WITH GROWTH HORMONE: A MULTICENTER ITALIAN STUDY. <i>Endocrine Practice</i> , 2017 , 23, 929-941	3.2	27
89	Bridging the gap: metabolic and endocrine care of patients during transition. <i>Endocrine Connections</i> , 2016 , 5, R44-R54	3.5	25
88	Caring for children and adolescents with type 1 diabetes mellitus: Italian Society for Pediatric Endocrinology and Diabetology (ISPED) statements during COVID-19 pandemia. <i>Diabetes Research and Clinical Practice</i> , 2020 , 168, 108372	7.4	24
87	Hypothalamic-pituitary vascularization in pituitary stalk transection syndrome: is the pituitary stalk really transected? The role of gadolinium-DTPA with spin-echo T1 imaging and turbo-FLASH technique. <i>Pediatric Radiology</i> , 1997 , 27, 48-53	2.8	23
86	Early-onset central diabetes insipidus is associated with de novo arginine vasopressin-neurophysin II or Wolfram syndrome 1 gene mutations. <i>European Journal of Endocrinology</i> , 2015 , 172, 461-72	6.5	22
85	Adult height in children with short stature and idiopathic delayed puberty after different management. <i>European Journal of Pediatrics</i> , 2008 , 167, 677-81	4.1	22
84	Current perspective on the pathogenesis of central diabetes insipidus. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2005 , 18, 631-45	1.6	21

(2015-1994)

83	Adrenocorticotrophin stimulation and HLA polymorphisms suggest a high frequency of heterozygosity for steroid 21-hydroxylase deficiency in patients with Turner@syndrome and their families. <i>Clinical Endocrinology</i> , 1994 , 40, 39-45	3.4	21	
82	Posterior pituitary (PP) evaluation in patients with anterior pituitary defect associated with ectopic PP and septo-optic dysplasia. <i>European Journal of Endocrinology</i> , 2011 , 165, 411-20	6.5	20	
81	Φο no harmQmanagement of craniopharyngioma. <i>European Journal of Endocrinology</i> , 2008 , 159 Suppl 1, S95-9	6.5	18	
80	Addressing gaps in care of people with conditions affecting sex development and maturation. <i>Nature Reviews Endocrinology</i> , 2019 , 15, 615-622	15.2	17	
79	Hyponatremia and seizures during desmopressin acetate treatment in hypothyroidism. <i>Journal of Pediatrics</i> , 1990 , 116, 835-6	3.6	17	
78	latrogenic [corrected] extrapontine myelinolysis in central diabetes insipidus: are cyclosporine and 1-desamino-8-D-arginine vasopressin harmful in association?. <i>Journal of Clinical Endocrinology and Metabolism</i> , 1997 , 82, 1749-51	5.6	16	
77	GH response to ghrelin in subjects with congenital GH deficiency: evidence that ghrelin action requires hypothalamic-pituitary connections. <i>European Journal of Endocrinology</i> , 2007 , 156, 449-54	6.5	16	
76	Central diabetes insipidus in children: Diagnosis and management. <i>Best Practice and Research in Clinical Endocrinology and Metabolism</i> , 2020 , 34, 101440	6.5	15	
75	Role of MRI T2-DRIVE in the assessment of pituitary stalk abnormalities without gadolinium in pituitary diseases. <i>European Journal of Endocrinology</i> , 2018 , 178, 613-622	6.5	15	
74	Short Stature Diagnosis and Referral. <i>Frontiers in Endocrinology</i> , 2017 , 8, 374	5.7	15	
73	T2*-based MR imaging (gradient echo or susceptibility-weighted imaging) in midline and off-midline intracranial germ cell tumors: a pilot study. <i>Neuroradiology</i> , 2018 , 60, 89-99	3.2	15	
7 ²	Wolfram syndrome 1 in the Italian population: genotype-phenotype correlations. <i>Pediatric Research</i> , 2020 , 87, 456-462	3.2	14	
71	Contribution of LHX4 Mutations to Pituitary Deficits in a Cohort of 417 Unrelated Patients. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2017 , 102, 290-301	5.6	13	
70	Germline prokineticin receptor 2 (PROKR2) variants associated with central hypogonadism cause differental modulation of distinct intracellular pathways. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2014 , 99, E458-63	5.6	13	
69	Diagnosis of hypochondroplasia: the role of radiological interpretation. Italian Study Group for Hypochondroplasia. <i>Pediatric Radiology</i> , 2001 , 31, 203-8	2.8	13	
68	Individualised growth response optimisation (iGRO) tool: an accessible and easy-to-use growth prediction system to enable treatment optimisation for children treated with growth hormone. Journal of Pediatric Endocrinology and Metabolism, 2017, 30, 1019-1026	1.6	12	
67	Antipituitary antibodies in patients with pituitary abnormalities and hormonal deficiency. <i>Clinical Endocrinology</i> , 1994 , 40, 809-10	3.4	12	
66	GH deficiency status combined with GH receptor polymorphism affects response to GH in children. European Journal of Endocrinology, 2015 , 173, 777-89	6.5	11	

65	Plasma total adiponectin levels in pediatrics: reference intervals calculated as a continuous variable of age. <i>Clinical Biochemistry</i> , 2012 , 45, 1703-5	3.5	11
64	Growth hormone secretory pattern and response to treatment in children with short stature followed to adult height. <i>Clinical Endocrinology</i> , 2003 , 59, 27-33	3.4	11
63	Age- and sex-matched reference curves for serum collagen type I C-telopeptides and bone alkaline phosphatase in children and adolescents: An alternative multivariate statistical analysis approach. <i>Clinical Biochemistry</i> , 2016 , 49, 802-7	3.5	10
62	Failure to increase insulin-like growth factor-I synthesis is involved in the mechanisms of growth retardation of children with inherited liver disorders. <i>Clinical Endocrinology</i> , 1998 , 48, 747-55	3.4	10
61	Impairment of the opioidergic control of luteinizing hormone secretion in Turner@syndrome: lack of effect of gonadal steroid therapy. <i>Journal of Clinical Endocrinology and Metabolism</i> , 1988 , 66, 1024-8	5.6	10
60	Pituitary gland imaging and outcome. <i>Endocrine Development</i> , 2012 , 23, 16-29		9
59	The advantage of measuring spontaneous growth hormone (GH) secretion compared with the insulin tolerance test in the diagnosis of GH deficiency in young adults. <i>Clinical Endocrinology</i> , 2007 , 67, 78-84	3.4	9
58	Antibodies Against Hypothalamus and Pituitary Gland in Childhood-Onset Brain Tumors and Pituitary Dysfunction. <i>Frontiers in Endocrinology</i> , 2020 , 11, 16	5.7	8
57	Quantitative ultrasound detects bone impairment after bone marrow transplantation in children and adolescents affected by hematological diseases. <i>Bone</i> , 2008 , 43, 177-182	4.7	8
56	IGF1 for the diagnosis of growth hormone deficiency in children and adolescents: a reappraisal. <i>Endocrine Connections</i> , 2020 , 9, 1095-1102	3.5	8
55	Familial neurohypophyseal diabetes insipidus in 13 kindreds and 2 novel mutations in the vasopressin gene. <i>European Journal of Endocrinology</i> , 2019 , 181, 233-244	6.5	8
54	Congenital hypogonadotropic hypogonadism/Kallmann syndrome is associated with statural gain in both men and women: a monocentric study. <i>European Journal of Endocrinology</i> , 2020 , 182, 185	6.5	8
53	Clinical Manifestations and Metabolic Outcomes of Seven Adults With Silver-Russell Syndrome. Journal of Clinical Endocrinology and Metabolism, 2018 , 103, 2225-2233	5.6	7
52	Growth hormone response to growth hormone-releasing hormone varies with the hypothalamic-pituitary abnormalities. <i>European Journal of Endocrinology</i> , 1996 , 135, 198-204	6.5	7
51	Highly conserved non-coding sequences and the 18q critical region for short stature: a common mechanism of disease?. <i>PLoS ONE</i> , 2008 , 3, e1460	3.7	7
50	Giant Urticaria and Acral Peeling in a Child with Coronavirus Disease 2019. <i>Journal of Pediatrics</i> , 2021 , 230, 261-263	3.6	7
49	Growth hormone and treatment outcomes: expert review of current clinical practice. <i>Pediatric Endocrinology Reviews</i> , 2011 , 9, 554-65	1.1	7
48	Accuracy and Limitations of the Growth Hormone (GH) Releasing Hormone-Arginine Retesting in Young Adults With Childhood-Onset GH Deficiency. <i>Frontiers in Endocrinology</i> , 2019 , 10, 525	5.7	6

47	Sleep disorders in Prader-Willi syndrome, evidence from animal models and humans. <i>Sleep Medicine Reviews</i> , 2021 , 57, 101432	10.2	6
46	Pretreatment Endocrine Disorders Due to Optic Pathway Gliomas in Pediatric Neurofibromatosis Type 1: Multicenter Study. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020 , 105,	5.6	6
45	Shared Decision-Making in Growth Hormone Therapy-Implications for Patient Care. <i>Frontiers in Endocrinology</i> , 2018 , 9, 688	5.7	6
44	The Effect of Lockdown and Physical Activity on Glycemic Control in Italian Children and Young Patients With Type 1 Diabetes. <i>Frontiers in Endocrinology</i> , 2021 , 12, 690222	5.7	6
43	Hypothalamic-pituitary magnetic resonance imaging in growth hormone deficiency. <i>Expert Review of Endocrinology and Metabolism</i> , 2006 , 1, 413-423	4.1	5
42	Neonatal Hypoglycemia and Brain Vulnerability. Frontiers in Endocrinology, 2021, 12, 634305	5.7	5
41	Primary Adrenal Insufficiency in Childhood: Data From a Large Nationwide Cohort. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2021 , 106, 762-773	5.6	5
40	Reliability of clonidine testing for the diagnosis of growth hormone deficiency in children and adolescents. <i>Clinical Endocrinology</i> , 2018 , 89, 765-770	3.4	5
39	Cost-consequence analysis for human recombinant growth hormone (r-hGH) treatment administered via different devices in children with growth hormone deficiency in Italy. <i>ClinicoEconomics and Outcomes Research</i> , 2019 , 11, 525-537	1.7	4
38	Familial ROBO1 deletion associated with ectopic posterior pituitary, duplication of the pituitary stalk and anterior pituitary hypoplasia. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2019 , 32, 95	-9] .6	4
37	Practical Approach to Using Trend Arrows on Real-Time Continuous Glucose Monitoring System in Type 1 Diabetes Adolescents Living Camp Setting Treated With Multiple Daily Injection or Continuous Subcutaneous Insulin Infusion Insulin Therapy. <i>Journal of Diabetes Science and</i>	4.1	3
36	Technology, 2021, 15, 1098-1103 Predictors of renal complications in pediatric patients with type 1 diabetes mellitus: A prospective cohort study. <i>Journal of Diabetes and Its Complications</i> , 2018, 32, 955-960	3.2	3
35	Motor function improvement after intravenous pamidronate in osteoporosis pseudoglioma syndrome. <i>Journal of Pediatrics</i> , 2006 , 149, 734	3.6	3
34	Growth and Puberty in Juvenile Dermatomyositis: A Longitudinal Cohort Study. <i>Arthritis Care and Research</i> , 2020 , 72, 265-273	4.7	3
33	The phenotypic spectrum associated with OTX2 mutations in humans. <i>European Journal of Endocrinology</i> , 2021 , 185, 121-135	6.5	3
32	Advances in differential diagnosis and management of growth hormone deficiency in children. <i>Nature Reviews Endocrinology</i> , 2021 , 17, 608-624	15.2	3
31	Sleep disturbances in craniopharyngioma: a challenging diagnosis. <i>Journal of Neurology</i> , 2021 , 268, 436	52 5 4 3 69	9 3
30	MEHMO syndrome and the link between brain, pituitary and pancreas. EBioMedicine, 2019, 42, 26-27	8.8	2

29	Management of childhood-onset craniopharyngioma in Italy: A multicenter seven year follow-up study of 145 patients. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2021 ,	5.6	2
28	Cognitive Profiles and Brain Volume Are Affected in Patients with Silver-Russell Syndrome. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020 , 105,	5.6	2
27	Endocrine Outcomes In Central Diabetes Insipidus: the Predictive Value of Neuroimaging "Mismatch Pattern". <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020 , 105,	5.6	2
26	New Horizons in Short Children Born Small for Gestational Age. <i>Frontiers in Pediatrics</i> , 2021 , 9, 655931	3.4	2
25	Endothelial impairment evaluation by peripheral arterial tonometry in pediatric endocrinopathies: A narrative review. <i>World Journal of Diabetes</i> , 2021 , 12, 810-826	4.7	2
24	Patient Satisfaction of Telemedicine in Pediatric and Young Adult Type 1 Diabetes Patients During Covid-19 Pandemic <i>Frontiers in Public Health</i> , 2022 , 10, 857561	6	2
23	Endothelial Dysfunction in Childhood Cancer Survivors: A Narrative Review Life, 2021, 12,	3	2
22	Encefalopatia di Hashimoto. <i>L Endocrinologo</i> , 2019 , 20, 98-99	О	1
21	Urinary Tract Involvement in Wolfram Syndrome: A Narrative Review. <i>International Journal of Environmental Research and Public Health</i> , 2021 , 18,	4.6	1
20	Prognostic factors in children and adolescents with differentiated thyroid carcinoma treated with total thyroidectomy and RAI: a real-life multicentric study. <i>European Journal of Nuclear Medicine and Molecular Imaging</i> , 2021 , 1	8.8	1
19	Characterization of Two Novel Variants of the Steroidogenic Acute Regulatory Protein Identified in a Girl with Classic Lipoid Congenital Adrenal Hyperplasia. <i>International Journal of Molecular Sciences</i> , 2020 , 21,	6.3	1
18	Summary of Expert Opinion on the Management of Children With Chronic Kidney Disease and Growth Failure With Human Growth Hormone. <i>Frontiers in Endocrinology</i> , 2020 , 11, 587	5.7	1
17	Maternal Uniparental Disomy of Chromosome 20 (UPD(20)mat) as Differential Diagnosis of Silver Russell Syndrome: Identification of Three New Cases. <i>Genes</i> , 2021 , 12,	4.2	1
16	Cognitive and White Matter Microstructure Development in Congenital Hypothyroidism and Familial Thyroid Disorders. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2021 , 106, e3990-e4006	5.6	1
15	Cardiometabolic risk in childhood cancer survivors. <i>Minerva Pediatrics</i> , 2021 ,	1.5	1
14	Neuromuscular and Neuroendocrinological Features Associated With -Related Arthrogryposis Multiplex Congenita in a Sicilian Family: A Case Report. <i>Frontiers in Neurology</i> , 2021 , 12, 704747	4.1	1
13	The first European consensus on principles of management for achondroplasia. <i>Orphanet Journal of Rare Diseases</i> , 2021 , 16, 333	4.2	1
12	Important Tools for Use by Pediatric Endocrinologists in the Assessment of Short Stature. <i>JCRPE Journal of Clinical Research in Pediatric Endocrinology</i> , 2021 , 13, 124-135	1.9	O

LIST OF PUBLICATIONS

11	Case Report: Lipoma of the Tuber Cinereum Mimicking a Pituitary Gland Abnormality in a Girl With Central Precocious Puberty. <i>Frontiers in Endocrinology</i> , 2021 , 12, 766253	5.7	О
10	Gene expression signatures predict response to therapy with growth hormone. <i>Pharmacogenomics Journal</i> , 2021 , 21, 594-607	3.5	O
9	A case of interference in testosterone, DHEA-S and progesterone measurements by second generation immunoassays. <i>Clinical Chemistry and Laboratory Medicine</i> , 2021 , 59, e275-e277	5.9	О
8	Growth in Children With Noonan Syndrome and Effects of Growth Hormone Treatment on Adult Height <i>Frontiers in Endocrinology</i> , 2021 , 12, 761171	5.7	О
7	Disorders of Water Balance 2019 , 553-582		
6	Letter to the Editor: "Forty-One Individuals With Mutations in the AVP-NPII Gene Associated With Familial Neurohypophyseal Diabetes Insipidus.". <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020 , 105,	5.6	
5	Is growth hormone treatment in young children safe for the heart?. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2004 , 89, 5271-2	5.6	
4	Important Tools for Use by Pediatric Endocrinologists in the Assessment of Short Stature. <i>JCRPE Journal of Clinical Research in Pediatric Endocrinology</i> , 2021 , 13, 124-135	1.9	
3	Pituitary Gland Imaging 2016 , 123-146		
2	An Atypical Case of Late-Onset Wolfram Syndrome 1 without Diabetes Insipidus <i>International Journal of Environmental Research and Public Health</i> , 2022 , 19,	4.6	
1	Osteogenesis Imperfecta/Ehlers-Danlos Overlap Syndrome and Neuroblastoma-Case Report and Review of Literature <i>Genes</i> , 2022 , 13,	4.2	