Zoran Gucev

List of Publications by Year
in descending order

41 papers	762 citations	840776 11 h-index	552781 26 g-index
42 all docs	42 docs citations	42 times ranked	1210 citing authors

#	Article	IF	Citations
1	Pathogenic variants in RNPC3 are associated with hypopituitarism and primary ovarian insufficiency. Genetics in Medicine, 2022, 24, 384-397.	2.4	4
2	The landscape of Mucopolysaccharidosis in Southern and Eastern European countries: a survey from 19 specialistic centers. Orphanet Journal of Rare Diseases, 2022, 17, 136.	2.7	3
3	Consensus statement on enzyme replacement therapy for mucopolysaccharidosis IVA in Central and South-Eastern European countries. Orphanet Journal of Rare Diseases, 2022, 17, 190.	2.7	2
4	Copy Number Variant Analysis and Genome-wide Association Study Identify Loci with Large Effect for Vesicoureteral Reflux. Journal of the American Society of Nephrology: JASN, 2021, 32, 805-820.	6.1	17
5	Posterior Urethral Valve and Prenataly Resolved Multicystic Dysplastic Kidney. Prilozi - Makedonska Akademija Na Naukite I Umetnostite Oddelenie Za Medicinski Nauki, 2021, 42, 77-81.	0.5	0
6	In Memoriam Academic Momir Polenakovic, Pediatric Nephrology, Rare Diseases and Publishing in Macedonia. Prilozi - Makedonska Akademija Na Naukite I Umetnostite Oddelenie Za Medicinski Nauki, 2021, 42, 27-29.	0.5	0
7	Heterotopic ossifications and Charcot joints: Congenital insensitivity to pain with anhidrosis (CIPA) and a novel NTRK1 gene mutation. European Journal of Medical Genetics, 2020, 63, 103613.	1.3	3
8	A Comprehensive Cohort Analysis Comparing Growth and GH Therapy Response in IGF1R Mutation Carriers and SGA Children. Journal of Clinical Endocrinology and Metabolism, 2020, 105, e1705-e1717.	3.6	12
9	Rare heterozygous GDF6 variants in patients with renal anomalies. European Journal of Human Genetics, 2020, 28, 1681-1693.	2.8	7
10	Exome Sequencing and Identification of Phenocopies in Patients With Clinically Presumed Hereditary Nephropathies. American Journal of Kidney Diseases, 2020, 76, 460-470.	1.9	33
11	Severe digital malformations in a rare variant of fibrodysplasia ossificans progressiva. American Journal of Medical Genetics, Part A, 2019, 179, 1310-1314.	1.2	5
12	Treatment and long-term outcome in primary distal renal tubular acidosis. Nephrology Dialysis Transplantation, 2019, 34, 981-991.	0.7	75
13	Cover Image, Volume 179A, Number 7, July 2019. , 2019, 179, .		2
14	The copy number variation landscape of congenital anomalies of the kidney and urinary tract. Nature Genetics, 2019, 51, 117-127.	21.4	144
15	Whole exome sequencing frequently detects a monogenic cause in early onset nephrolithiasis andÂnephrocalcinosis. Kidney International, 2018, 93, 204-213.	5.2	133
16	Metabolic Profiles in Obese Children and Adolescents with Insulin Resistance. Open Access Macedonian Journal of Medical Sciences, 2018, 6, 511-518.	0.2	22
17	Homeostasis Model Assessment - Insulin Resistance and Sensitivity (HOMA-IR and IS) Index in Overweight Children Born Small for Gestational Age (SGA). Prilozi - Makedonska Akademija Na Naukite I Umetnostite Oddelenie Za Medicinski Nauki, 2018, 39, 83-89.	0.5	4
18	The Spectrum of Kidney Diseases in Children Associated with Low Molecular Weight Proteinuria. Open Access Macedonian Journal of Medical Sciences, 2018, 6, 814-819.	0.2	3

#	Article	IF	Citations
19	IGF1R Gene Alterations in Children Born Small for Gestitional Age (SGA). Open Access Macedonian Journal of Medical Sciences, 2018, 6, 2040-2044.	0.2	6
20	IGF1R Gene Alterations in Small for Gestational Age (SGA) Children. Open Access Macedonian Journal of Medical Sciences, 2018, 6, 790-793.	0.2	0
21	Growth Hormone Treatment in Children Born Small for Gestational Age (SGA). Prilozi - Makedonska Akademija Na Naukite I Umetnostite Oddelenie Za Medicinski Nauki, 2018, 39, 143-149.	0.5	0
22	Low Molecular Weight Proteinuria in Children with Distal Renal Tubular Acidosis. Prilozi - Makedonska Akademija Na Naukite I Umetnostite Oddelenie Za Medicinski Nauki, 2018, 39, 91-95.	0.5	1
23	A 4-Year-Old Boy with Beckwith Wiedemann Syndrome (BWS). Prilozi - Makedonska Akademija Na Naukite I Umetnostite Oddelenie Za Medicinski Nauki, 2018, 39, 131-135.	0.5	1
24	Congenital Anomalies of the Kidney and Urinary Tract in Children Born Small for Gestational Age. Prilozi - Makedonska Akademija Na Naukite I Umetnostite Oddelenie Za Medicinski Nauki, 2017, 38, 53-57.	0.5	3
25	5th Rare Disease South Eastern Europe (SEE) Meeting, Skopje, Macedonia (November) Tj ETQq1 1 0.784314 rgBT /Overl Medicinski Nauki, 2017, 38, 119-123.	0.5	2
26	Somapacitan, a onceâ€weekly reversible albuminâ€binding <scp>GH</scp> derivative, in children with <scp>GH</scp> deficiency: A randomized doseâ€escalation trial. Clinical Endocrinology, 2017, 87, 350-358.	2.4	38
27	Obesity in Childhood and Adolescence, Genetic Factors. Prilozi - Makedonska Akademija Na Naukite I Umetnostite Oddelenie Za Medicinski Nauki, 2017, 38, 121-133.	0.5	21
28	50 Years of the Macedonian Academy of Sciences and Arts 1967-2017 and 48 Years of Publishing the Journal Prilozi (Contributions) of MASA. Prilozi - Makedonska Akademija Na Naukite I Umetnostite Oddelenie Za Medicinski Nauki, 2017, 38, 5-8.	0.5	1
29	Mutations in SLC26A1 Cause Nephrolithiasis. American Journal of Human Genetics, 2016, 98, 1228-1234.	6.2	41
30	Long-term renal outcome in children withOCRLmutations: retrospective analysis of a large international cohort. Nephrology Dialysis Transplantation, 2016, 33, gfw350.	0.7	27
31	The Child Health Care System of Macedonia. Journal of Pediatrics, 2016, 177, S127-S137.	1.8	3
32	Prevalence of Monogenic Causes in Pediatric Patients with Nephrolithiasis or Nephrocalcinosis. Clinical Journal of the American Society of Nephrology: CJASN, 2016, 11, 664-672.	4.5	105
33	Poststreptococcal glomerulonephritis in children with congenital anomalies of the kidney and urinary tract. Renal Failure, 2015, 37, 1440-1443.	2.1	2
34	First-line therapy in atypical hemolytic uremic syndrome: consideration on infants with a poor prognosis. Italian Journal of Pediatrics, 2014, 40, 101.	2.6	8
35	Two Siblings with Niemann-Pick Disease (NPD) Type B: Clinical Findings and Novel Mutations of the Acid Sphingomyelinase Gene. Indian Journal of Pediatrics, 2013, 80, 163-164.	0.8	4
36	A Novel GH1 Mutation in a Family with Isolated Growth Hormone Deficiency Type II. Hormone Research in Paediatrics, 2012, 77, 200-204.	1.8	7

#	Article	IF	Citations
37	Cystinuria AA (B): digenic inheritance with three mutations in two cystinuria genes. Journal of Genetics, 2011, 90, 157-159.	0.7	9
38	Billateral Polycystic Kidneys in a Girl with WAGR Syndrome. Indian Journal of Pediatrics, 2011, 78, 1290-1292.	0.8	6
39	McCune-Albright Syndrome (MAS): Early and Extensive Bone Fibrous Dysplasia Involvement and â€œMistaken Identityâ€•Oophorectomy. Journal of Pediatric Endocrinology and Metabolism, 2010, 23, 837-42.	0.9	6
40	Response to â€œCLOVE(S) Syndrome: Expanding the Acronymâ€• American Journal of Medical Genetics, Part A, 2009, 149A, 295-295.	1.2	0
41	Friedreich's ataxia (FA) associated with diabetes mellitus type 1 and hypertrophic cardiomyopathy: analysis of a FA family. , 2009, 63, 110-1.		2