

# Zoran Gucev

## List of Publications by Year in descending order

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Version: 2024-02-01

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	The copy number variation landscape of congenital anomalies of the kidney and urinary tract. <i>Nature Genetics</i> , 2019, 51, 117-127.	21.4	144
2	Whole exome sequencing frequently detects a monogenic cause in early onset nephrolithiasis and nephrocalcinosis. <i>Kidney International</i> , 2018, 93, 204-213.	5.2	133
3	Prevalence of Monogenic Causes in Pediatric Patients with Nephrolithiasis or Nephrocalcinosis. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2016, 11, 664-672.	4.5	105
4	Treatment and long-term outcome in primary distal renal tubular acidosis. <i>Nephrology Dialysis Transplantation</i> , 2019, 34, 981-991.	0.7	75
5	Mutations in SLC26A1 Cause Nephrolithiasis. <i>American Journal of Human Genetics</i> , 2016, 98, 1228-1234.	6.2	41
6	Somapacitan, a once-weekly reversible albumin-binding GH derivative, in children with GH deficiency: A randomized dose-escalation trial. <i>Clinical Endocrinology</i> , 2017, 87, 350-358.	2.4	38
7	Exome Sequencing and Identification of Phenocopies in Patients With Clinically Presumed Hereditary Nephropathies. <i>American Journal of Kidney Diseases</i> , 2020, 76, 460-470.	1.9	33
8	Long-term renal outcome in children with OCRL mutations: retrospective analysis of a large international cohort. <i>Nephrology Dialysis Transplantation</i> , 2016, 33, gfw350.	0.7	27
9	Metabolic Profiles in Obese Children and Adolescents with Insulin Resistance. <i>Open Access Macedonian Journal of Medical Sciences</i> , 2018, 6, 511-518.	0.2	22
10	Obesity in Childhood and Adolescence, Genetic Factors. <i>Prilozi - Makedonska Akademija Na Naukite I Umetnostite Oddelenie Za Medicinski Nauki</i> , 2017, 38, 121-133.	0.5	21
11	Copy Number Variant Analysis and Genome-wide Association Study Identify Loci with Large Effect for Vesicoureteral Reflux. <i>Journal of the American Society of Nephrology: JASN</i> , 2021, 32, 805-820.	6.1	17
12	A Comprehensive Cohort Analysis Comparing Growth and GH Therapy Response in IGF1R Mutation Carriers and SGA Children. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020, 105, e1705-e1717.	3.6	12
13	Cystinuria AA (B): digenic inheritance with three mutations in two cystinuria genes. <i>Journal of Genetics</i> , 2011, 90, 157-159.	0.7	9
14	First-line therapy in atypical hemolytic uremic syndrome: consideration on infants with a poor prognosis. <i>Italian Journal of Pediatrics</i> , 2014, 40, 101.	2.6	8
15	A Novel GH1 Mutation in a Family with Isolated Growth Hormone Deficiency Type II. <i>Hormone Research in Paediatrics</i> , 2012, 77, 200-204.	1.8	7
16	Rare heterozygous GDF6 variants in patients with renal anomalies. <i>European Journal of Human Genetics</i> , 2020, 28, 1681-1693.	2.8	7
17	McCune-Albright Syndrome (MAS): Early and Extensive Bone Fibrous Dysplasia Involvement and "Mistaken Identity" Oophorectomy. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2010, 23, 837-42.	0.9	6
18	Bilateral Polycystic Kidneys in a Girl with WAGR Syndrome. <i>Indian Journal of Pediatrics</i> , 2011, 78, 1290-1292.	0.8	6

#	ARTICLE	IF	CITATIONS
19	IGF1R Gene Alterations in Children Born Small for Gestational Age (SGA). Open Access Macedonian Journal of Medical Sciences, 2018, 6, 2040-2044.	0.2	6
20	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
21	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
22	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
23	Pathogenic variants in RNPC3 are associated with hypopituitarism and primary ovarian insufficiency. Genetics in Medicine, 2022, 24, 384-397.	2.4	4
24	The Child Health Care System of Macedonia. Journal of Pediatrics, 2016, 177, S127-S137.	1.8	3
25	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
26	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
27	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
28	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
29	Poststreptococcal glomerulonephritis in children with congenital anomalies of the kidney and urinary tract. Renal Failure, 2015, 37, 1440-1443.	2.1	2
30	5<sup>th</sup> Rare Disease South Eastern Europe (SEE) Meeting, Skopje, Macedonia (November) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 5 Medicinski Nauki, 2017, 38, 119-123.	0.5	2
31	Cover Image, Volume 179A, Number 7, July 2019. , 2019, 179, .		2
32	Friedreich's ataxia (FA) associated with diabetes mellitus type 1 and hypertrophic cardiomyopathy: analysis of a FA family. , 2009, 63, 110-1.		2
33	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
34	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
35	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
36	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

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37	Response to "CLOVE(S) Syndrome: Expanding the Acronym". American Journal of Medical Genetics, Part A, 2009, 149A, 295-295.	1.2	0
38	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
39	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
40	IGF1R Gene Alterations in Small for Gestational Age (SGA) Children. Open Access Macedonian Journal of Medical Sciences, 2018, 6, 790-793.	0.2	0
41	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