

# Kapoor Rr

## List of Publications by Year in descending order

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Version: 2024-02-01

39  
papers

1,524  
citations

394390

19  
h-index

345203

36  
g-index

40  
all docs

40  
docs citations

40  
times ranked

1494  
citing authors

#	ARTICLE	IF	CITATIONS
1	Insulin Mutation Screening in 1,044 Patients With Diabetes. <i>Diabetes</i> , 2008, 57, 1034-1042.	0.6	347
2	Clinical and molecular characterisation of 300 patients with congenital hyperinsulinism. <i>European Journal of Endocrinology</i> , 2013, 168, 557-564.	3.7	190
3	Persistent Hyperinsulinemic Hypoglycemia and Maturity-Onset Diabetes of the Young Due to Heterozygous <i>HNF4A</i> Mutations. <i>Diabetes</i> , 2008, 57, 1659-1663.	0.6	133
4	Advances in the diagnosis and management of hyperinsulinemic hypoglycemia. <i>Nature Clinical Practice Endocrinology and Metabolism</i> , 2009, 5, 101-112.	2.8	84
5	Hyperinsulinism—hyperammonaemia syndrome: novel mutations in the <i>GLUD1</i> gene and genotype—phenotype correlations. <i>European Journal of Endocrinology</i> , 2009, 161, 731-735.	3.7	81
6	3-Hydroxyacyl-Coenzyme A Dehydrogenase Deficiency and Hyperinsulinemic Hypoglycemia: Characterization of a Novel Mutation and Severe Dietary Protein Sensitivity. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2009, 94, 2221-2225.	3.6	72
7	Measuring the Impact of Diagnostic Decision Support on the Quality of Clinical Decision Making: Development of a Reliable and Valid Composite Score. <i>Journal of the American Medical Informatics Association: JAMIA</i> , 2003, 10, 563-572.	4.4	58
8	Hyperinsulinism in Developmental Syndromes. <i>Endocrine Development</i> , 2009, 14, 95-113.	1.3	56
9	Genetics of congenital hyperinsulinemic hypoglycemia. <i>Seminars in Pediatric Surgery</i> , 2011, 20, 13-17.	1.1	55
10	Genome-Wide Homozygosity Analysis Reveals <i>HADH</i> Mutations as a Common Cause of Diazoxide-Responsive Hyperinsulinemic-Hypoglycemia in Consanguineous Pedigrees. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2011, 96, E498-E502.	3.6	51
11	Hyperinsulinemic hypoglycemia in children and adolescents: Recent advances in understanding of pathophysiology and management. <i>Reviews in Endocrine and Metabolic Disorders</i> , 2020, 21, 577-597.	5.7	45
12	Clinical and molecular characterisation of hyperinsulinaemic hypoglycaemia in infants born small-for-gestational age. <i>Archives of Disease in Childhood: Fetal and Neonatal Edition</i> , 2013, 98, F356-F358.	2.8	44
13	Leucine-sensitive hyperinsulinaemic hypoglycaemia in patients with loss of function mutations in 3-Hydroxyacyl-CoA Dehydrogenase. <i>Orphanet Journal of Rare Diseases</i> , 2012, 7, 25.	2.7	41
14	Adjacent mutations in the gating loop of Kir6.2 produce neonatal diabetes and hyperinsulinism. <i>EMBO Molecular Medicine</i> , 2009, 1, 166-177.	6.9	36
15	Using referral rates for genetic testing to determine the incidence of a rare disease: The minimal incidence of congenital hyperinsulinism in the UK is 1 in 28,389. <i>PLoS ONE</i> , 2020, 15, e0228417.	2.5	29
16	The Heterogeneity of Focal Forms of Congenital Hyperinsulinism. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2012, 97, E94-E99.	3.6	26
17	Diabetes mellitus, exocrine pancreatic deficiency, hypertrichosis, hyperpigmentation, and chronic inflammation: confirmation of a syndrome. <i>Pediatric Diabetes</i> , 2009, 10, 193-197.	2.9	25
18	The Clinical Problem of Hyperinsulinemic Hypoglycemia and Resultant Infantile Spasms. <i>Pediatrics</i> , 2010, 126, e1231-e1236.	2.1	23

#	ARTICLE	IF	CITATIONS
19	Congenital Hyperinsulinism due to a Compound Heterozygous <i>ABCC8</i> Mutation with Spontaneous Resolution at Eight Weeks. <i>Hormone Research in Paediatrics</i> , 2010, 73, 287-292.	1.8	23
20	Congenital Hyperinsulinism due to mutations in HNF4A and HADH. <i>Reviews in Endocrine and Metabolic Disorders</i> , 2010, 11, 185-191.	5.7	21
21	Xq27.1 Duplication Encompassing <i>SOX3</i> : Variable Phenotype and Smallest Duplication Associated with Hypopituitarism to Date – A Large Case Series of Unrelated Patients and a Literature Review. <i>Hormone Research in Paediatrics</i> , 2019, 92, 382-389.	1.8	15
22	Prolactinoma in childhood and adolescence – Tumour size at presentation predicts management strategy: Single centre series and a systematic review and meta-analysis. <i>Clinical Endocrinology</i> , 2021, 94, 413-423.	2.4	11
23	Transient neonatal hyperinsulinaemic hypoglycaemia: perinatal predictors of length and cost of stay. <i>European Journal of Pediatrics</i> , 2018, 177, 1823-1829.	2.7	10
24	Autoimmune Encephalitis and Autism Spectrum Disorder. <i>Frontiers in Psychiatry</i> , 2021, 12, 775017.	2.6	10
25	Congenital hyperinsulinism: marked clinical heterogeneity in siblings with identical mutations in the <i>ABCC8</i> gene. <i>Clinical Endocrinology</i> , 2012, 76, 312-313.	2.4	8
26	Focal congenital hyperinsulinism in a patient with septo-optic dysplasia. <i>Nature Reviews Endocrinology</i> , 2010, 6, 646-650.	9.6	5
27	Hyperinsulinaemic hypoglycaemia in deoxyguanosine kinase deficiency. <i>Clinical Endocrinology</i> , 2019, 91, 900-903.	2.4	4
28	Acromesomelic Dysplasia, Type Maroteaux: Impact of Long-Term (8 Years) High-Dose Growth Hormone Treatment on Growth Velocity and Final Height in 2 Siblings. <i>Hormone Research in Paediatrics</i> , 2020, 93, 335-342.	1.8	4
29	IgG4-related hypophysitis in adolescence. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2021, 34, 395-399.	0.9	4
30	Exceptional diazoxide sensitivity in hyperinsulinaemic hypoglycaemia due to a novel HNF4A mutation. <i>Endocrinology, Diabetes and Metabolism Case Reports</i> , 2019, 2019, .	0.5	3
31	Central Diabetes Insipidus in Children and Adolescents: Twenty-Six Year Experience from a Single Centre. <i>International Journal of Endocrinology</i> , 2022, 2022, 1-6.	1.5	3
32	<i>HNF4A</i> and Hyperinsulinemic Hypoglycemia. <i>Frontiers in Diabetes</i> , 2012, , 182-190.	0.4	2
33	Pituitary apoplexy in an adolescent male with macroprolactinoma presenting as middle cerebral artery territory infarction. <i>Annals of Pediatric Endocrinology and Metabolism</i> , 2022, 27, 320-324.	2.3	2
34	New-onset diabetes mellitus following pediatric liver transplantation. <i>Pediatric Transplantation</i> , 2013, 17, 5-7.	1.0	1
35	A novel heterozygous mutation in the insulin receptor gene presenting with type A severe insulin resistance syndrome. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2020, 33, 809-812.	0.9	1
36	Title is missing!. , 2020, 15, e0228417.		0

#	ARTICLE	IF	CITATIONS
37	Title is missing!. , 2020, 15, e0228417.		0
38	Title is missing!. , 2020, 15, e0228417.		0
39	Title is missing!. , 2020, 15, e0228417.		0