

Kapoor Rr

List of Publications by Year in Descending Order

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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.

37
papers

1,213
citations

19
h-index

34
g-index

40
ext. papers

1,380
ext. citations

4.5
avg, IF

3.85
L-index

#	Paper	IF	Citations
37	Central Diabetes Insipidus in Children and Adolescents: Twenty-Six Year Experience from a Single Centre.. <i>International Journal of Endocrinology</i> , 2022 , 2022, 9397130	2.7	0
36	IgG4-related hypophysitis in adolescence. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2021 , 34, 395-399	1.6	1
35	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	3.4	6
34	Autoimmune Encephalitis and Autism Spectrum Disorder.. <i>Frontiers in Psychiatry</i> , 2021 , 12, 775017	5	1
33	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	10.5	19
32	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	3.7	11
31	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	1.6	1
30	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	3.3	4
29	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 2020 , 15, e0228417		
28	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 2020 , 15, e0228417		
27	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 2020 , 15, e0228417		
26	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 2020 , 15, e0228417		
25	Hyperinsulinaemic hypoglycaemia in deoxyguanosine kinase deficiency. <i>Clinical Endocrinology</i> , 2019 , 91, 900-903	3.4	1
24	Xq27.1 Duplication Encompassing SOX3: 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	3.3	9
23	Exceptional diazoxide sensitivity in hyperinsulinaemic hypoglycaemia due to a novel HNF4A mutation. <i>Endocrinology, Diabetes and Metabolism Case Reports</i> , 2019 , 2019,	1.4	1
22	Transient neonatal hyperinsulinaemic hypoglycaemia: perinatal predictors of length and cost of stay. <i>European Journal of Pediatrics</i> , 2018 , 177, 1823-1829	4.1	8
21	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, F356-8	4.7	30

20	Clinical and molecular characterisation of 300 patients with congenital hyperinsulinism. <i>European Journal of Endocrinology</i> , 2013 , 168, 557-64	6.5	147
19	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	4.2	26
18	HNF4A and Hyperinsulinemic Hypoglycemia. <i>Frontiers in Diabetes</i> , 2012 , 182-190	0.6	1
17	Congenital hyperinsulinism: marked clinical heterogeneity in siblings with identical mutations in the ABCC8 gene. <i>Clinical Endocrinology</i> , 2012 , 76, 312-3	3.4	8
16	The heterogeneity of focal forms of congenital hyperinsulinism. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2012 , 97, E94-9	5.6	25
15	Genetics of congenital hyperinsulinemic hypoglycemia. <i>Seminars in Pediatric Surgery</i> , 2011 , 20, 13-7	2.1	48
14	Genome-wide homozygosity analysis reveals HADH mutations as a common cause of diazoxide-responsive hyperinsulinemic-hypoglycemia in consanguineous pedigrees. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2011 , 96, E498-502	5.6	44
13	The clinical problem of hyperinsulinemic hypoglycemia and resultant infantile spasms. <i>Pediatrics</i> , 2010 , 126, e1231-6	7.4	20
12	Congenital hyperinsulinism due to a compound heterozygous ABCC8 mutation with spontaneous resolution at eight weeks. <i>Hormone Research in Paediatrics</i> , 2010 , 73, 287-92	3.3	19
11	Focal congenital hyperinsulinism in a patient with septo-optic dysplasia. <i>Nature Reviews Endocrinology</i> , 2010 , 6, 646-50	15.2	4
10	Congenital hyperinsulinism due to mutations in HNF4A and HADH. <i>Reviews in Endocrine and Metabolic Disorders</i> , 2010 , 11, 185-91	10.5	19
9	Hyperinsulinism-hyperammonaemia syndrome: novel mutations in the GLUD1 gene and genotype-phenotype correlations. <i>European Journal of Endocrinology</i> , 2009 , 161, 731-5	6.5	65
8	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, 2221-5	5.6	64
7	Adjacent mutations in the gating loop of Kir6.2 produce neonatal diabetes and hyperinsulinism. <i>EMBO Molecular Medicine</i> , 2009 , 1, 166-77	12	34
6	Diabetes mellitus, exocrine pancreatic deficiency, hypertrichosis, hyperpigmentation, and chronic inflammation: confirmation of a syndrome. <i>Pediatric Diabetes</i> , 2009 , 10, 193-7	3.6	21
5	Hyperinsulinism in developmental syndromes. <i>Endocrine Development</i> , 2009 , 14, 95-113		45
4	Advances in the diagnosis and management of hyperinsulinemic hypoglycemia. <i>Nature Clinical Practice Endocrinology and Metabolism</i> , 2009 , 5, 101-12		67
3	Persistent hyperinsulinemic hypoglycemia and maturity-onset diabetes of the young due to heterozygous HNF4A mutations. <i>Diabetes</i> , 2008 , 57, 1659-63	0.9	113

2	Insulin mutation screening in 1,044 patients with diabetes: mutations in the INS gene are a common cause of neonatal diabetes but a rare cause of diabetes diagnosed in childhood or adulthood. <i>Diabetes</i> , 2008 , 57, 1034-42	0.9	299
1	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, 563-72	8.6	51