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List of Publications by Citations

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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.

11
papers

266
citations

7
h-index

12
g-index

12
ext. papers

291
ext. citations

3.8
avg, IF

2.56
L-index

#	Paper	IF	Citations
11	The genetic basis of congenital hyperinsulinism. <i>Journal of Medical Genetics</i> , 2009 , 46, 289-99	5.8	115
10	The predictive value of preoperative fluorine-18-L-3,4-dihydroxyphenylalanine positron emission tomography-computed tomography scans in children with congenital hyperinsulinism of infancy. <i>Journal of Pediatric Surgery</i> , 2011 , 46, 204-8	2.6	42
9	Familial focal congenital hyperinsulinism. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2011 , 96, 24-8	5.6	26
8	The heterogeneity of focal forms of congenital hyperinsulinism. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2012 , 97, E94-9	5.6	25
7	Role of 18F-DOPA PET/CT imaging in congenital hyperinsulinism. <i>Reviews in Endocrine and Metabolic Disorders</i> , 2010 , 11, 165-9	10.5	23
6	Persistent hyperinsulinaemic hypoglycaemia of infancy: 15 years experience at the Royal Children's Hospital (RCH), Melbourne. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2005 , 18, 1103-9	1.6	11
5	The heterogeneity of hyperinsulinaemic hypoglycaemia in 19 patients with Beckwith-Wiedemann syndrome due to KvDMR1 hypomethylation. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2015 , 28, 83-6	1.6	10
4	Mutational analysis of the GYS2 gene in patients diagnosed with ketotic hypoglycaemia. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2012 , 25, 963-7	1.6	6
3	Satisfaction of care in a tertiary level diabetes clinic: correlations with diabetes knowledge, clinical outcome and health-related quality of life. <i>Journal of Paediatrics and Child Health</i> , 2008 , 44, 432-7	1.3	4
2	Galactokinase deficiency in a patient with congenital hyperinsulinism. <i>JIMD Reports</i> , 2012 , 5, 7-11	1.9	3
1	Refinement of the critical genomic region for hypoglycaemia in the Chromosome 9p deletion syndrome. <i>Wellcome Open Research</i> , 2019 , 4, 149	4.8	1