Maturity-Onset Diabetes of the Young in Children With

Diabetes Care

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Citation Report

#	Article	IF	Citations
1	Opposite Clinical Phenotypes of Glucokinase Disease: Description of a Novel Activating Mutation and Contiguous Inactivating Mutations in Human Glucokinase (GCK) Gene. Molecular Endocrinology, 2009, 23, 1983-1989.	3.7	30
2	Update on mutations in glucokinase (GCK), which cause maturity-onset diabetes of the young, permanent neonatal diabetes, and hyperinsulinemic hypoglycemia. Human Mutation, 2009, 30, 1512-1526.	1.1	403
3	Current literature in diabetes. Diabetes/Metabolism Research and Reviews, 2010, 26, i-xi.	1.7	0
4	Clinical application of best practice guidelines for the genetic diagnosis of MODY2 and MODY3. Diabetic Medicine, 2010, 27, 1331-1333.	1.2	6
5	Glucokinase diabetes in 103 families from a country-based study in the Czech Republic: geographically restricted distribution of two prevalent GCK mutations. Pediatric Diabetes, 2010, 11, 529-535.	1.2	50
6	A novel synonymous substitution in the GCK gene causes aberrant splicing in an Italian patient with GCK-MODY phenotype. Diabetes Research and Clinical Practice, 2011, 92, e23-e26.	1.1	5
7	Diagnosis and management of maturity onset diabetes of the young (MODY). BMJ: British Medical Journal, 2011, 343, d6044-d6044.	2.4	154
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10	MODY. Diabetes Care, 2011, 34, 1878-1884.	4.3	265
11	Review on monogenic diabetes. Current Opinion in Endocrinology, Diabetes and Obesity, 2011, 18, 252-258.	1.2	45
12	Stress Hyperglycemia: A Sign of Familial Diabetes in Children. Pediatrics, 2011, 128, e1614-e1617.	1.0	5
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17	Incidental mild hyperglycemia in children: two MODY 2 families identified in Brazilian subjects. Arquivos Brasileiros De Endocrinologia E Metabologia, 2012, 56, 519-524.	1.3	8
18	Novel glucokinase mutations in patients with monogenic diabetes â€“ clinical outline of GCK â€MD and potential for founder effect in Slavic population. Clinical Genetics, 2012, 81, 278-283.	1.0	13
19	Onset of type 1 diabetes mellitus in two patients with maturity onset diabetes of the young. Pediatric Diabetes, 2012, 13, 208-212.	1.2	15

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20	The glucokinase mutation p.T206P is common among MODY patients of Jewish Ashkenazi descent. Pediatric Diabetes, 2012, 13, e14-e21.	1.2	7
21	Hyperglycaemia and Î²-cell antibodies: Is it always pre-type 1 diabetes?. Diabetes Research and Clinical Practice, 2013, 100, e20-e22.	1.1	4
22	MODY type 2 P59S GCK mutant: founder effect in South of Italy. Clinical Genetics, 2013, 83, 83-87.	1.0	4
23	The spectrum ofHNF1Agene mutations in Greek patients with MODY3: relative frequency and identification of seven novel germline mutations. Pediatric Diabetes, 2013, 14, 526-534.	1.2	12
24	Prognostic Accuracy of Continuous Glucose Monitoring in the Prediction of Diabetes Mellitus in Children with Incidental Hyperglycemia: Receiver Operating Characteristic Analysis. Diabetes Technology and Therapeutics, 2013, 15, 580-585.	2.4	9
25	Low Prevalence of HNF1A Mutations After Molecular Screening of Multiple MODY Genes in 58 Italian Families Recruited in the Pediatric or Adult Diabetes Clinic From a Single Italian Hospital. Diabetes Care, 2014, 37, e258-e260.	4.3	23
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34	Molecular diagnosis of maturity-onset diabetes of the young (MODY) in Turkish children by using targeted next-generation sequencing. Journal of Pediatric Endocrinology and Metabolism, 2015, 28, 1265-71.	0.4	28
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36	Survey on etiological diagnosis of diabetes in 1244 Italian diabetic children and adolescents: Impact of access to genetic testing. Diabetes Research and Clinical Practice, 2015, 107, e15-e18.	1.1	24
37	Adolescent nonâ€adherence reveals a genetic cause forÂdiabetes. Diabetic Medicine, 2015, 32, e20-3.	1.2	5

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45	Abnormalities in alternative splicing in diabetes: therapeutic targets. Journal of Molecular Endocrinology, 2017, 59, R93-R107.	1.1	33
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60	Screening for monogenic diabetes in primary care. Primary Care Diabetes, 2020, 14, 1-11.	0.9	10
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