Nadia Tinto

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/7693615/publications.pdf

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17	362	840119 11	887659 17
papers	citations	h-index	g-index

17	17	17	576
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Monogenic Diabetes Accounts for 6.3% of Cases Referred to 15 Italian Pediatric Diabetes Centers During 2007 to 2012. Journal of Clinical Endocrinology and Metabolism, 2017, 102, 1826-1834.	1.8	88
2	Glucokinase Gene Mutations: Structural and Genotype-Phenotype Analyses in MODY Children from South Italy. PLoS ONE, 2008, 3, e1870.	1.1	44
3	Glucokinase (GCK) Mutations and Their Characterization in MODY2 Children of Southern Italy. PLoS ONE, 2012, 7, e38906.	1.1	37
4	Identification of Candidate Children for Maturity-Onset Diabetes of the Young Type 2 (MODY2) Gene Testing: A Seven-Item Clinical Flowchart (7-iF). PLoS ONE, 2013, 8, e79933.	1.1	33
5	High Frequency of Haplotype HLA-DQ7 in Celiac Disease Patients from South Italy: Retrospective Evaluation of 5,535 Subjects at Risk of Celiac Disease. PLoS ONE, 2015, 10, e0138324.	1.1	23
6	A ceRNA Circuitry Involving the Long Noncoding RNA Klhl14-AS, Pax8, and Bcl2 Drives Thyroid Carcinogenesis. Cancer Research, 2019, 79, 5746-5757.	0.4	23
7	Dietary Thiols: A Potential Supporting Strategy against Oxidative Stress in Heart Failure and Muscular Damage during Sports Activity. International Journal of Environmental Research and Public Health, 2020, 17, 9424.	1.2	23
8	Molecular Epidemiology of Mitochondrial Cardiomyopathy: A Search among Mitochondrial and Nuclear Genes. International Journal of Molecular Sciences, 2021, 22, 5742.	1.8	17
9	Can HbA1c combined with fasting plasma glucose help to assess priority for GCK-MODY vs HNF1A-MODY genetic testing?. Acta Diabetologica, 2018, 55, 981-983.	1.2	14
10	The Hidden Fragility in the Heart of the Athletes: A Review of Genetic Biomarkers. International Journal of Molecular Sciences, 2020, 21, 6682.	1.8	14
11	Cystic Fibrosis-Related Diabetes (CFRD): Overview of Associated Genetic Factors. Diagnostics, 2021, 11, 572.	1.3	14
12	NGS Analysis Revealed Digenic Heterozygous GCK and HNF1A Variants in a Child with Mild Hyperglycemia: A Case Report. Diagnostics, 2021, 11, 1164.	1.3	8
13	Metabolic Treatment of Wolfram Syndrome. International Journal of Environmental Research and Public Health, 2022, 19, 2755.	1.2	7
14	Prenatal diagnosis of HNF1b mutation allows recognition of neonatal dysglycemia. Acta Diabetologica, 2021, 58, 393-395.	1.2	5
15	Molecular diagnosis of MODY3 permitted to reveal a de novo 12q24.31 deletion and to explain a complex phenotype in a young diabetic patient. Clinical Chemistry and Laboratory Medicine, 2019, 57, e306-e310.	1.4	4
16	Generation of an iPSC line (UNINAi001-A) from a girl with neonatal-onset epilepsy and non-syndromic intellectual disability carrying the homozygous KCNQ3 p.PHE534ILEfs*15 variant and of an iPSC line (UNINAi002-A) from a non-carrier, unaffected brother. Stem Cell Research, 2021, 53, 102311.	0.3	4
17	Congenital diabetes mellitus. Minerva Pediatrica, 2020, 72, 240-249.	2.6	4