Alessandro Salina

List of Publications by Year
in descending order

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14 papers	272 citations	1478505 6 h-index	1125743 13 g-index
14 all docs	14 docs citations	14 times ranked	548 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.	3.6	88
2	A novel CISD2 intragenic deletion, optic neuropathy and platelet aggregation defect in Wolfram syndrome type 2. BMC Medical Genetics, 2014, 15, 88.	2.1	59
3	Wolfram Syndrome: New Mutations, Different Phenotype. PLoS ONE, 2012, 7, e29150.	2.5	55
4	Wolfram syndrome 1 in the Italian population: genotypeâ€“phenotype correlations. Pediatric Research, 2020, 87, 456-462.	2.3	20
5	The coexistence of type 1 diabetes, MODY2 and metabolic syndrome in a young girl. Acta Diabetologica, 2012, 49, 401-404.	2.5	14
6	Glucokinase mutations in pediatric patients with impaired fasting glucose. Acta Diabetologica, 2017, 54, 913-923.	2.5	11
7	Mother and daughter carrying the same KCNJ11 mutation but with a different response to switching from insulin to sulfonylurea. Diabetes Research and Clinical Practice, 2011, 94, e50-e52.	2.8	6
8	Diabetes Mellitus Diagnosed in Childhood and Adolescence With Negative Autoimmunity: Results of Genetic Investigation. Frontiers in Endocrinology, 0, 13, .	3.5	6
9	Hyperglycaemia and Î²-cell antibodies: Is it always pre-type 1 diabetes?. Diabetes Research and Clinical Practice, 2013, 100, e20-e22.	2.8	4
10	Novel homozygous mutation in exon 5 of <scp>WFS1</scp> gene in an Apulian family with mild phenotypic expression of Wolfram syndrome. Clinical Genetics, 2014, 86, 197-198.	2.0	3
11	Comment on: Clinical application of best practice guidelines for genetic diagnosis of MODY2. Diabetes Research and Clinical Practice, 2012, 95, e29-e30.	2.8	2
12	A mild impairment of K+ATP channel function caused by two different ABCC8 defects in an Italian newborn. Acta Diabetologica, 2018, 55, 201-203.	2.5	2
13	Estimation of genetic risk for Type 1 diabetes mellitus in newborns on dried blood spot. Journal of Endocrinological Investigation, 2010, 33, 406-408.	3.3	1
14	A Novel Genetic Variant in the WFS1 Gene in a Patient with Partial Uniparental Mero-Isodisomy of Chromosome 4. International Journal of Molecular Sciences, 2021, 22, 8082.	4.1	1