

Valerie Senee

List of Publications by Year in descending order

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16
papers

966
citations

687220

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940416

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docs citations

16
times ranked

1711
citing authors

#	ARTICLE	IF	CITATIONS
1	Mutations in GLIS3 are responsible for a rare syndrome with neonatal diabetes mellitus and congenital hypothyroidism. <i>Nature Genetics</i> , 2006, 38, 682-687.	9.4	327
2	Wolcott-Rallison Syndrome: Clinical, Genetic, and Functional Study of EIF2AK3 Mutations and Suggestion of Genetic Heterogeneity. <i>Diabetes</i> , 2004, 53, 1876-1883.	0.3	165
3	A Novel Hypomorphic <i>PDX1</i> Mutation Responsible for Permanent Neonatal Diabetes With Subclinical Exocrine Deficiency. <i>Diabetes</i> , 2010, 59, 733-740.	0.3	85
4	A Missense Mutation in <i>PPP1R15B</i> Causes a Syndrome Including Diabetes, Short Stature, and Microcephaly. <i>Diabetes</i> , 2015, 64, 3951-3962.	0.3	71
5	YIPF5 mutations cause neonatal diabetes and microcephaly through endoplasmic reticulum stress. <i>Journal of Clinical Investigation</i> , 2020, 130, 6338-6353.	3.9	58
6	Wolcott-Rallison syndrome due to the same mutation (W522X) in EIF2AK3 in two unrelated families and review of the literature*. <i>Pediatric Diabetes</i> , 2010, 11, 279-285.	1.2	43
7	Abnormal FGFR 3 Expression in Cartilage of Thanatophoric Dysplasia Fetuses. <i>Human Molecular Genetics</i> , 1997, 6, 1899-1906.	1.4	40
8	DNAJC3 deficiency induces β -cell mitochondrial apoptosis and causes syndromic young-onset diabetes. <i>European Journal of Endocrinology</i> , 2021, 184, 455-468.	1.9	29
9	Severe FOXP3+ and Naïve T Lymphopenia in a Non-IPEX Form of Autoimmune Enteropathy Combined With an Immunodeficiency. <i>Gastroenterology</i> , 2007, 132, 1694-1704.	0.6	26
10	SLC29A3 mutation in a patient with syndromic diabetes with features of pigmented hypertrichotic dermatosis with insulin-dependent diabetes, H syndrome and Faisalabad histiocytosis. <i>Diabetes and Metabolism</i> , 2013, 39, 281-285.	1.4	24
11	Mutations and variants of ONECUT1 in diabetes. <i>Nature Medicine</i> , 2021, 27, 1928-1940.	15.2	24
12	dUTPase (<i>DUT</i>) Is Mutated in a Novel Monogenic Syndrome With Diabetes and Bone Marrow Failure. <i>Diabetes</i> , 2017, 66, 1086-1096.	0.3	22
13	A novel ALMS1 splice mutation in a non-obese juvenile-onset insulin-dependent syndromic diabetic patient. <i>European Journal of Human Genetics</i> , 2014, 22, 140-143.	1.4	19
14	Microscopic and ultrastructural features in Wolcott-Rallison syndrome, a permanent neonatal diabetes mellitus: about two autopsy cases. <i>Pediatric Diabetes</i> , 2015, 16, 510-520.	1.2	13
15	Crohn's disease associated CARD15 (NOD2) variants are not involved in the susceptibility to type 1 diabetes. <i>Molecular Genetics and Metabolism</i> , 2005, 86, 379-383.	0.5	12
16	Juvenile-Onset Diabetes and Congenital Cataract: "Double-Gene" Mutations Mimicking a Syndromic Diabetes Presentation. <i>Genes</i> , 2017, 8, 309.	1.0	8