Valerie Senee

List of Publications by Year
in descending order

16	966	687220 13	940416 16
papers	citations	h-index	g-index

16	16	16	1711
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	DNAJC3 deficiency induces Î²-cell mitochondrial apoptosis and causes syndromic young-onset diabetes. European Journal of Endocrinology, 2021, 184, 455-468.	1.9	29
2	Mutations and variants of ONECUT1 in diabetes. Nature Medicine, 2021, 27, 1928-1940.	15.2	24
3	YIPF5 mutations cause neonatal diabetes and microcephaly through endoplasmic reticulum stress. Journal of Clinical Investigation, 2020, 130, 6338-6353.	3.9	58
4	dUTPase (DUT) Is Mutated in a Novel Monogenic Syndrome With Diabetes and Bone Marrow Failure. Diabetes, 2017, 66, 1086-1096.	0.3	22
5	Juvenile-Onset Diabetes and Congenital Cataract: â€œDouble-Geneâ€•Mutations Mimicking a Syndromic Diabetes Presentation. Genes, 2017, 8, 309.	1.0	8
6	Microscopic and ultrastructural features in Wolcott-Rallison syndrome, a permanent neonatal diabetes mellitus: about two autopsy cases. Pediatric Diabetes, 2015, 16, 510-520.	1.2	13
7	A Missense Mutation in PPP1R15B Causes a Syndrome Including Diabetes, Short Stature, and Microcephaly. Diabetes, 2015, 64, 3951-3962.	0.3	71
8	A novel ALMS1 splice mutation in a non-obese juvenile-onset insulin-dependent syndromic diabetic patient. European Journal of Human Genetics, 2014, 22, 140-143.	1.4	19
9	SLC29A3 mutation in a patient with syndromic diabetes with features of pigmented hypertrichotic dermatosis with insulin-dependent diabetes, H syndrome and Faisalabad histiocytosis. Diabetes and Metabolism, 2013, 39, 281-285.	1.4	24
10	Wolcott-Rallison syndrome due to the same mutation (W522X) in EIF2AK3 in two unrelated families and review of the literature*. Pediatric Diabetes, 2010, 11, 279-285.	1.2	43
11	A Novel Hypomorphic PDX1 Mutation Responsible for Permanent Neonatal Diabetes With Subclinical Exocrine Deficiency. Diabetes, 2010, 59, 733-740.	0.3	85
12	Severe FOXP3+ and NaÃ¯ve T Lymphopenia in a Non-IPEX Form of Autoimmune Enteropathy Combined With an Immunodeficiency. Gastroenterology, 2007, 132, 1694-1704.	0.6	26
13	Mutations in GLIS3 are responsible for a rare syndrome with neonatal diabetes mellitus and congenital hypothyroidism. Nature Genetics, 2006, 38, 682-687.	9.4	327
14	Crohnâ€™s disease associated CARD15 (NOD2) variants are not involved in the susceptibility to type 1 diabetes. Molecular Genetics and Metabolism, 2005, 86, 379-383.	0.5	12
15	Wolcott-Rallison Syndrome: Clinical, Genetic, and Functional Study of EIF2AK3 Mutations and Suggestion of Genetic Heterogeneity. Diabetes, 2004, 53, 1876-1883.	0.3	165
16	Abnormal FGFR 3 Expression in Cartilage of Thanatophoric Dysplasia Fetuses. Human Molecular Genetics, 1997, 6, 1899-1906.	1.4	40