

# ValÃ©rie SenÃ©e

## List of Publications by Year in descending order

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12  
papers

940  
citations

700390

12  
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1003665

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times ranked

1887  
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.	20.4	335
2	A Novel Hypomorphic <i>PDX1</i> Mutation Responsible for Permanent Neonatal Diabetes With Subclinical Exocrine Deficiency. <i>Diabetes</i> , 2010, 59, 733-740.	0.9	86
3	A Missense Mutation in <i>PPP1R15B</i> Causes a Syndrome Including Diabetes, Short Stature, and Microcephaly. <i>Diabetes</i> , 2015, 64, 3951-3962.	0.9	80
4	YIPF5 mutations cause neonatal diabetes and microcephaly through endoplasmic reticulum stress. <i>Journal of Clinical Investigation</i> , 2020, 130, 6338-6353.	8.2	63
5	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.	3.0	44
6	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.	1.4	28
7	Mutations and variants of <i>ONECUT1</i> in diabetes. <i>Nature Medicine</i> , 2021, 27, 1928-1940.	30.1	27
8	dUTPase ( <i>DUT</i> ) Is Mutated in a Novel Monogenic Syndrome With Diabetes and Bone Marrow Failure. <i>Diabetes</i> , 2017, 66, 1086-1096.	0.9	23
9	A novel <i>ALMS1</i> splice mutation in a non-obese juvenile-onset insulin-dependent syndromic diabetic patient. <i>European Journal of Human Genetics</i> , 2014, 22, 140-143.	2.9	20
10	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.	3.0	13
11	Crohn's disease associated <i>CARD15</i> ( <i>NOD2</i> ) variants are not involved in the susceptibility to type 1 diabetes. <i>Molecular Genetics and Metabolism</i> , 2005, 86, 379-383.	2.2	12
12	Juvenile-Onset Diabetes and Congenital Cataract: "Double-Gene" Mutations Mimicking a Syndromic Diabetes Presentation. <i>Genes</i> , 2017, 8, 309.	2.4	8