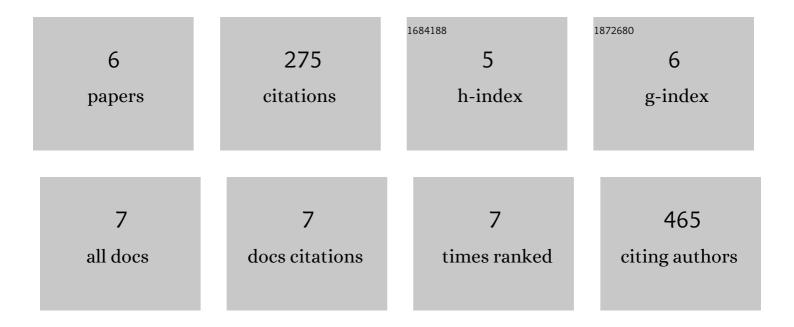
## Geralyn Creadon-Swindell

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/1575040/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	The genetic basis of classic nonketotic hyperglycinemia due to mutations in GLDC and AMT. Genetics in Medicine, 2017, 19, 104-111.	2.4	71
2	Biochemical and molecular predictors for prognosis in nonketotic hyperglycinemia. Annals of Neurology, 2015, 78, 606-618.	5.3	68
3	Mutations in the mitochondrial cysteinyl-tRNA synthase gene, <i>CARS2,</i> lead to a severe epileptic encephalopathy and complex movement disorder. Journal of Medical Genetics, 2015, 52, 532-540.	3.2	62
4	Neurodevelopmental Outcome and Treatment Efficacy of Benzoate andÂDextromethorphan in Siblings with Attenuated NonketoticÂHyperglycinemia. Journal of Pediatrics, 2016, 170, 234-239.	1.8	60
5	Biomarkers of oxidative stress, inflammation, and vascular dysfunction in inherited cystathionine βâ€synthase deficient homocystinuria and the impact of taurine treatment in a phase 1/2 human clinical trial. Journal of Inherited Metabolic Disease, 2019, 42, 424-437.	3.6	11
6	d -Clyceric aciduria does not cause nonketotic hyperglycinemia: A historic co-occurrence. Molecular Genetics and Metabolism, 2017, 121, 80-82.	1.1	2