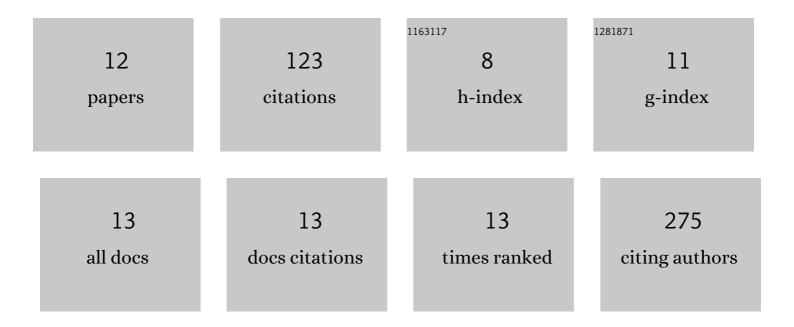
Robert Meyer

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/5121005/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	Targeted Next Generation Sequencing Approach in Patients Referred for Silver-Russell Syndrome Testing Increases the Mutation Detection Rate and Provides Decisive Information for Clinical Management. Journal of Pediatrics, 2017, 187, 206-212.e1.	1.8	22
2	Loss of supervillin causes myopathy with myofibrillar disorganization and autophagic vacuoles. Brain, 2020, 143, 2406-2420.	7.6	15
3	C2orf69 mutations disrupt mitochondrial function and cause a multisystem human disorder with recurring autoinflammation. Journal of Clinical Investigation, 2021, 131, .	8.2	13
4	One test for all: whole exome sequencing significantly improves the diagnostic yield in growth retarded patients referred for molecular testing for Silver–Russell syndrome. Orphanet Journal of Rare Diseases, 2021, 16, 42.	2.7	12
5	NLRP genes and their role in preeclampsia and multi-locus imprinting disorders. Journal of Perinatal Medicine, 2018, 46, 169-173.	1.4	11
6	Next generation sequencing and imprinting disorders: Current applications and future perspectives: Lessons from Silver-Russell syndrome. Molecular and Cellular Probes, 2019, 44, 1-7.	2.1	11
7	<i>HMGA2</i> Variants in Silver-Russell Syndrome: Homozygous and Heterozygous Occurrence. Journal of Clinical Endocrinology and Metabolism, 2020, 105, 2401-2407.	3.6	11
8	Patient with an autosomalâ€recessive <scp><i>MBTPS1</i></scp> â€linked phenotype and clinical features of <scp>Silver–Russell</scp> syndrome. American Journal of Medical Genetics, Part A, 2020, 182, 2727-2730.	1.2	10
9	Inherited cases of <scp><i>CNOT3</i></scp> â€associated intellectual developmental disorder with speech delay, autism, and dysmorphic facies. Clinical Genetics, 2020, 98, 408-412.	2.0	9
10	Germline variants in DNA repair genes, including <i>BRCA1</i> / <i>2</i> , may cause familial myeloproliferative neoplasms. Blood Advances, 2021, 5, 3373-3376.	5.2	7
11	Aggressive infantile myofibromatosis with intestinal involvement. Molecular and Cellular Pediatrics, 2021, 8, 7.	1.8	2
19	Die CfH-lupierAbademie 2020 – virtueller Verreiter in der CfH Medizinische Cenetik, 2020, 32, 347,348	0.2	0

12 Die GfH-JuniorAkademie 2020 – virtueller Vorreiter in der GfH. Medizinische Genetik, 2020, 32, 347-348. 0.2 0