# Robert Meyer

List of Publications by Year
in descending order



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|---|---|---|---|
| 12 papers | 123 citations | 1163117 8 h-index | 1281871 11 g-index |
| 13 all docs | 13 docs citations | 13 times ranked | 275 citing authors |

| # | 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 Silverâ€“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 | HMGA2 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>MBTPS1</scp>â€linked phenotype and clinical features of <scp>Silverâ€“Russell</scp> syndrome. American Journal of Medical Genetics, Part A, 2020, 182, 2727-2730. | 1.2 | 10 |
| 9 | Inherited cases of <scp>CNOT3</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 BRCA1/2, 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 |
| 12 | Die GfH-JuniorAkademie 2020 â€“ virtueller Vorreiter in der GfH. Medizinische Genetik, 2020, 32, 347-348. | 0.2 | 0 |