Joseph G Vockley

List of Publications by Year in descending order

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1163117 1474206 9 827 8 9 citations g-index h-index papers 9 9 9 2116 docs citations times ranked citing authors all docs

| # | Article | IF | CITATIONS |
|---|---|------|-----------|
| 1 | Parent-of-origin-specific signatures of de novo mutations. Nature Genetics, 2016, 48, 935-939. | 21.4 | 266 |
| 2 | Aberrant splicing induced by the most common <i>EPG5</i> mutation in an individual with Vici syndrome. Brain, 2016, 139, e52-e52. | 7.6 | 14 |
| 3 | New observations on maternal age effect on germline de novo mutations. Nature Communications, 2016, 7, 10486. | 12.8 | 166 |
| 4 | Utility of whole-genome sequencing for detection of newborn screening disorders in a population cohort of 1,696 neonates. Genetics in Medicine, 2016, 18, 221-230. | 2.4 | 101 |
| 5 | Diagnosis of D-Bifunctional Protein Deficiency through Whole-Genome Sequencing: Implications for Cost-Effective Care. Molecular Syndromology, 2015, 6, 141-146. | 0.8 | 5 |
| 6 | Genetic Variants That Predispose to DNA Double-Strand Breaks in Lymphocytes From a Subset of Patients With Familial Colorectal Carcinomas. Gastroenterology, 2015, 149, 1872-1883.e9. | 1.3 | 31 |
| 7 | Systematic evaluation of underlying defects in DNA repair as an approach to case-only assessment of familial prostate cancer. Oncotarget, 2015, 6, 39614-39633. | 1.8 | 13 |
| 8 | Germline Variation in Cancer-Susceptibility Genes in a Healthy, Ancestrally Diverse Cohort: Implications for Individual Genome Sequencing. PLoS ONE, 2014, 9, e94554. | 2.5 | 81 |
| 9 | Mutations in NOTCH1 Cause Adams-Oliver Syndrome. American Journal of Human Genetics, 2014, 95, 275-284. | 6.2 | 150 |