# Gerald Goh

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/4230924/publications.pdf

Version: 2024-02-01

| | | | |
|---|---|---|---|
| 28 papers | 6,784 citations | 304743 22 h-index | 501196 28 g-index |
| 29 all docs | 29 docs citations | 29 times ranked | 14252 citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Computational Methods for Analysis of Tumor Clonality and Evolutionary History. Methods in Molecular Biology, 2019, 1878, 217-226. | 0.9 | 5 |
| 2 | Fc-Optimized Anti-CD25 Depletes Tumor-Infiltrating Regulatory T Cells and Synergizes with PD-1 Blockade to Eradicate Established Tumors. Immunity, 2017, 46, 577-586. | 14.3 | 323 |
| 3 | Phylogenetic ctDNA analysis depicts early-stage lung cancer evolution. Nature, 2017, 545, 446-451. | 27.8 | 1,287 |
| 4 | Intratumoural evolutionary landscape of high-risk prostate cancer: the PROGENY study of genomic and immune parameters. Annals of Oncology, 2017, 28, 2472-2480. | 1.2 | 45 |
| 5 | Mutational landscape of MCPyV-positive and MCPyV-negative Merkel cell carcinomas with implications for immunotherapy. Oncotarget, 2016, 7, 3403-3415. | 1.8 | 306 |
| 6 | Shifting patterns of genomic variation in the somatic evolution of papillary thyroid carcinoma. BMC Cancer, 2016, 16, 646. | 2.6 | 5 |
| 7 | Clonal Evolutionary Analysis during HER2 Blockade in HER2-Positive Inflammatory Breast Cancer: A Phase II Open-Label Clinical Trial of Afatinib +/- Vinorelbine. PLoS Medicine, 2016, 13, e1002136. | 8.4 | 28 |
| 8 | Identification of a gain-of-function STAT3 mutation (p.Y640F) in lymphocytic variant hypereosinophilic syndrome. Blood, 2016, 127, 948-951. | 1.4 | 52 |
| 9 | DNA replication stress mediates APOBEC3 family mutagenesis in breast cancer. Genome Biology, 2016, 17, 185. | 8.8 | 140 |
| 10 | Absence of <scp>KMT</scp>2D/<scp>MLL</scp>2 mutations in abdominal paraganglioma. Clinical Endocrinology, 2016, 84, 632-634. | 2.4 | 3 |
| 11 | Chromosome 19 amplification correlates with advanced disease in adrenocortical carcinoma. Surgery, 2016, 159, 296-301. | 1.9 | 9 |
| 12 | Wholeâ€exome sequencing defines the mutational landscape of pheochromocytoma and identifies KMT 2 D as a recurrently mutated gene. Genes Chromosomes and Cancer, 2015, 54, 542-554. | 2.8 | 57 |
| 13 | Novel somatic mutations in primary hyperaldosteronism are related to the clinical, radiological and pathological phenotype. Clinical Endocrinology, 2015, 83, 779-789. | 2.4 | 115 |
| 14 | Recurrent gain of function mutation in calcium channel CACNA1H causes early-onset hypertension with primary aldosteronism. ELife, 2015, 4, e06315. | 6.0 | 271 |
| 15 | Increased Levels of Macrophage Inflammatory Proteins Result in Resistance to R5-Tropic HIV-1 in a Subset of Elite Controllers. Journal of Virology, 2015, 89, 5502-5514. | 3.4 | 68 |
| 16 | Characterization of the mutational landscape of anaplastic thyroid cancer via whole-exome sequencing. Human Molecular Genetics, 2015, 24, 2318-2329. | 2.9 | 290 |
| 17 | PRKACA mutations in cortisol-producing adenomas and adrenal hyperplasia: a single-center study of 60 cases. European Journal of Endocrinology, 2015, 172, 677-685. | 3.7 | 64 |
| 18 | Genomic landscape of cutaneous T cell lymphoma. Nature Genetics, 2015, 47, 1011-1019. | 21.4 | 347 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | Whole-Exome Sequencing Characterizes the Landscape of Somatic Mutations and Copy Number Alterations in Adrenocortical Carcinoma. Journal of Clinical Endocrinology and Metabolism, 2015, 100, E493-E502. | 3.6 | 131 |
| 20 | Genetic landscape of metastatic and recurrent head and neck squamous cell carcinoma. Journal of Clinical Investigation, 2015, 126, 169-180. | 8.2 | 156 |
| 21 | Recurrent activating mutation in PRKACA in cortisol-producing adrenal tumors. Nature Genetics, 2014, 46, 613-617. | 21.4 | 211 |
| 22 | Abstract 985: The mutational landscape of LN metastasis and recurrence in HNSCC. , 2014, , . | | 0 |
| 23 | Somatic and germline CACNA1D calcium channel mutations in aldosterone-producing adenomas and primary aldosteronism. Nature Genetics, 2013, 45, 1050-1054. | 21.4 | 519 |
| 24 | Exome sequencing identifies recurrent somatic RAC1 mutations in melanoma. Nature Genetics, 2012, 44, 1006-1014. | 21.4 | 1,052 |
| 25 | Effector CD4+ T Cell Expression Signatures and Immune-Mediated Disease Associated Genes. PLoS ONE, 2012, 7, e38510. | 2.5 | 16 |
| 26 | Application of Whole Exome Sequencing to Identify Disease-Causing Variants in Inherited Human Diseases. Genomics and Informatics, 2012, 10, 214. | 0.8 | 56 |
| 27 | Multiple Recurrent De Novo CNVs, Including Duplications of the 7q11.23 Williams Syndrome Region, Are Strongly Associated with Autism. Neuron, 2011, 70, 863-885. | 8.1 | 1,146 |
| 28 | Association and Mutation Analyses of 16p11.2 Autism Candidate Genes. PLoS ONE, 2009, 4, e4582. | 2.5 | 80 |