# Lauren Fishbein

List of Publications by Year
in descending order



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|---|---|---|---|
| 35 papers | 2,180 citations | 471509 17 h-index | 395702 33 g-index |
| 35 all docs | 35 docs citations | 35 times ranked | 3397 citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Comprehensive Molecular Characterization of Pheochromocytoma and Paraganglioma. Cancer Cell, 2017, 31, 181-193. | 16.8 | 532 |
| 2 | Genetic Analysis of 779 Advanced Differentiated and Anaplastic Thyroid Cancers. Clinical Cancer Research, 2018, 24, 3059-3068. | 7.0 | 366 |
| 3 | Inherited Mutations in Pheochromocytoma and Paraganglioma: Why All Patients Should Be Offered Genetic Testing. Annals of Surgical Oncology, 2013, 20, 1444-1450. | 1.5 | 182 |
| 4 | Pheochromocytoma and paraganglioma: understanding the complexities of the genetic background. Cancer Genetics, 2012, 205, 1-11. | 0.4 | 177 |
| 5 | Whole-exome sequencing identifies somatic ATRX mutations in pheochromocytomas and paragangliomas. Nature Communications, 2015, 6, 6140. | 12.8 | 143 |
| 6 | Pheochromocytoma and Paraganglioma. Hematology/Oncology Clinics of North America, 2016, 30, 135-150. | 2.2 | 127 |
| 7 | The North American Neuroendocrine Tumor Society Consensus Guidelines for Surveillance and Medical Management of Pancreatic Neuroendocrine Tumors. Pancreas, 2020, 49, 863-881. | 1.1 | 88 |
| 8 | Pheochromocytoma/Paraganglioma: Review of Perioperative Management of Blood Pressure and Update on Genetic Mutations Associated With Pheochromocytoma. Journal of Clinical Hypertension, 2013, 15, 428-434. | 2.0 | 62 |
| 9 | The North American Neuroendocrine Tumor Society Consensus Guidelines for Surveillance and Management of Metastatic and/or Unresectable Pheochromocytoma and Paraganglioma. Pancreas, 2021, 50, 469-493. | 1.1 | 55 |
| 10 | Development of new preclinical models to advance adrenocortical carcinoma research. Endocrine-Related Cancer, 2018, 25, 437-451. | 3.1 | 45 |
| 11 | Predicting Metastatic Potential in Pheochromocytoma and Paraganglioma: A Comparison of PASS and GAPP Scoring Systems. Journal of Clinical Endocrinology and Metabolism, 2020, 105, e4661-e4670. | 3.6 | 40 |
| 12 | In vitro studies of steroid hormones in neurofibromatosis 1 tumors and schwann cells. Molecular Carcinogenesis, 2007, 46, 512-523. | 2.7 | 39 |
| 13 | Muscle oxidative phosphorylation quantitation using creatine chemical exchange saturation transfer (CrCEST) MRI in mitochondrial disorders. JCI Insight, 2016, 1, e88207. | 5.0 | 38 |
| 14 | Intricacies of the Molecular Machinery of Catecholamine Biosynthesis and Secretion by Chromaffin Cells of the Normal Adrenal Medulla and in Pheochromocytoma and Paraganglioma. Cancers, 2019, 11, 1121. | 3.7 | 36 |
| 15 | Analysis of somatic NF1 promoter methylation in plexiform neurofibromas and Schwann cells. Cancer Genetics and Cytogenetics, 2005, 157, 181-186. | 1.0 | 26 |
| 16 | Genetics of pheochromocytoma and paraganglioma. Current Opinion in Endocrinology, Diabetes and Obesity, 2021, 28, 283-290. | 2.3 | 22 |
| 17 | Tumor detection rates in screening of individuals with SDHx-related hereditary paragangliomaâ€“pheochromocytoma syndrome. Genetics in Medicine, 2020, 22, 2101-2107. | 2.4 | 20 |
| 18 | The Changing Paradigm of Head and Neck Paragangliomas: What Every Otolaryngologist Needs to Know. Annals of Otology, Rhinology and Laryngology, 2020, 129, 1135-1143. | 1.1 | 20 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | Elucidating the Role of the Maternal Embryonic Leucine Zipper Kinase in Adrenocortical Carcinoma. Endocrinology, 2018, 159, 2532-2544. | 2.8 | 19 |
| 20 | Chromaffin cell biology: inferences from The Cancer Genome Atlas. Cell and Tissue Research, 2018, 372, 339-346. | 2.9 | 17 |
| 21 | Head and Neck Paragangliomas: An Update on the Molecular Classification, State-of-the-Art Imaging, and Management Recommendations. Radiology Imaging Cancer, 2022, 4, e210088. | 1.6 | 17 |
| 22 | Pheochromocytoma/Paraganglioma: Is This a Genetic Disorder?. Current Cardiology Reports, 2019, 21, 104. | 2.9 | 16 |
| 23 | SDHB knockout and succinate accumulation are insufficient for tumorigenesis but dual SDHB/NF1 loss yields SDHx-like pheochromocytomas. Cell Reports, 2022, 38, 110453. | 6.4 | 16 |
| 24 | Correlation Between Plasma Catecholamines, Weight, and Diabetes in Pheochromocytoma and Paraganglioma. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e4028-e4038. | 3.6 | 13 |
| 25 | Targeted genomic analysis of 364 adrenocortical carcinomas. Endocrine-Related Cancer, 2021, 28, 671-681. | 3.1 | 13 |
| 26 | Mastermind Like Transcriptional Coactivator 3 (MAML3) Drives Neuroendocrine Tumor Progression. Molecular Cancer Research, 2021, 19, 1476-1485. | 3.4 | 11 |
| 27 | The Jewel in the Crown: Specific Aims Section of Investigator-Initiated Grant Proposals. Journal of the Endocrine Society, 2017, 1, 1194-1202. | 0.2 | 8 |
| 28 | Genotype-Phenotype Features of Germline Variants of the TMEM127 Pheochromocytoma Susceptibility Gene: A 10-Year Update. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e350-e364. | 3.6 | 8 |
| 29 | Pathological and Genetic Stratification for Management of Adrenocortical Carcinoma. Journal of Clinical Endocrinology and Metabolism, 2022, 107, 1159-1169. | 3.6 | 7 |
| 30 | Paclitaxel is necessary for improved survival in epithelial ovarian cancers with homologous recombination gene mutations. Oncotarget, 2016, 7, 48577-48585. | 1.8 | 6 |
| 31 | Pheochromocytoma and Paraganglioma Susceptibility Genes. JAMA Oncology, 2017, 3, 1212. | 7.1 | 4 |
| 32 | Discovery of new susceptibility genes: proceed cautiously. Genetics in Medicine, 2018, 20, 1512-1514. | 2.4 | 3 |
| 33 | Case report of a paraganglioma arising from a mature cystic teratoma of the ovary. Gynecologic Oncology Reports, 2020, 32, 100537. | 0.6 | 3 |
| 34 | Case of Metastatic Pheochromocytoma and Meningiomas in a Patient With Lynch Syndrome. JCO Precision Oncology, 2022, 6, e2100251. | 3.0 | 1 |
| 35 | Impact of COVID-19 on individuals with paraganglioma/pheochromocytoma history and/or hereditary risk.. Journal of Clinical Oncology, 2022, 40, 10613-10613. | 1.6 | 0 |