## W Marston Linehan

## List of Publications by Year in descending order

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323 papers 40,210 citations

<sup>2544</sup> 96 h-index

193 g-index

332 all docs 332 docs citations

times ranked

332

28199 citing authors

#	Article	IF	CITATIONS
1	Identification of the von Hippel-Lindau Disease Tumor Suppressor Gene. Science, 1993, 260, 1317-1320.	12.6	2,723
2	Germline and somatic mutations in the tyrosine kinase domain of the MET proto-oncogene in papillary renal carcinomas. Nature Genetics, $1997$ , $16$ , $68-73$ .	21.4	1,461
3	von Hippel-Lindau disease. Lancet, The, 2003, 361, 2059-2067.	13.7	1,322
4	Comparison of MR/Ultrasound Fusion–Guided Biopsy With Ultrasound-Guided Biopsy for the Diagnosis of Prostate Cancer. JAMA - Journal of the American Medical Association, 2015, 313, 390.	7.4	1,267
5	Reductive carboxylation supports growth in tumour cells with defective mitochondria. Nature, 2012, 481, 385-388.	27.8	1,074
6	Comprehensive Molecular Characterization of Papillary Renal-Cell Carcinoma. New England Journal of Medicine, 2016, 374, 135-145.	27.0	1,040
7	Biochemical Diagnosis of Pheochromocytoma. JAMA - Journal of the American Medical Association, 2002, 287, 1427-34.	7.4	994
8	HIF overexpression correlates with biallelic loss of fumarate hydratase in renal cancer: Novel role of fumarate in regulation of HIF stability. Cancer Cell, 2005, 8, 143-153.	16.8	843
9	Mutations in a novel gene lead to kidney tumors, lung wall defects, and benign tumors of the hair follicle in patients with the Birt-Hogg-Dubé syndrome. Cancer Cell, 2002, 2, 157-164.	16.8	833
10	The Somatic Genomic Landscape of Chromophobe Renal Cell Carcinoma. Cancer Cell, 2014, 26, 319-330.	16.8	665
11	The genetic basis of kidney cancer: a metabolic disease. Nature Reviews Urology, 2010, 7, 277-285.	3.8	634
12	Mutations in the Fumarate Hydratase Gene Cause Hereditary Leiomyomatosis and Renal Cell Cancer in Families in North America. American Journal of Human Genetics, 2003, 73, 95-106.	6.2	563
13	Renal Tumors in the Birt-Hogg-Dubé Syndrome. American Journal of Surgical Pathology, 2002, 26, 1542-1552.	3.7	544
14	Germline mutations in the von Hippel-Lindau disease tumor suppressor gene: Correlations with phenotype. Human Mutation, 1995, 5, 66-75.	2.5	526
15	The Cancer Genome Atlas Comprehensive Molecular Characterization of Renal Cell Carcinoma. Cell Reports, 2018, 23, 313-326.e5.	6.4	523
16	Recent Advances in Genetics, Diagnosis, Localization, and Treatment of Pheochromocytoma. Annals of Internal Medicine, 2001, 134, 315.	3.9	512
17	Improved Identification of von Hippel-Lindau Gene Alterations in Clear Cell Renal Tumors. Clinical Cancer Research, 2008, 14, 4726-4734.	7.0	503
18	Novel mutations of the MET proto-oncogene in papillary renal carcinomas. Oncogene, 1999, 18, 2343-2350.	5.9	487

#	Article	IF	CITATIONS
19	Improved detection of germline mutations in the von Hippel-Lindau disease tumor suppressor gene. Human Mutation, 1998, 12, 417-423.	2.5	452
20	The Genetic Basis of Cancer of the Kidney. Journal of Urology, 2003, 170, 2163-2172.	0.4	447
21	Germline mutations in the Von Hippel-Lindau disease (VHL) gene in families from North America, Europe, and Japan. Human Mutation, 1996, 8, 348-357.	2.5	436
22	Integrated Proteogenomic Characterization of Clear Cell Renal Cell Carcinoma. Cell, 2019, 179, 964-983.e31.	28.9	430
23	Folliculin encoded by the <i>BHD</i> gene interacts with a binding protein, FNIP1, and AMPK, and is involved in AMPK and mTOR signaling. Proceedings of the National Academy of Sciences of the United States of America, 2006, 103, 15552-15557.	7.1	427
24	The contribution of VHL substrate binding and HIF1- $\hat{l}$ ± to the phenotype of VHL loss in renal cell carcinoma. Cancer Cell, 2002, 1, 247-255.	16.8	421
25	Phase II and Biomarker Study of the Dual MET/VEGFR2 Inhibitor Foretinib in Patients With Papillary Renal Cell Carcinoma. Journal of Clinical Oncology, 2013, 31, 181-186.	1.6	401
26	Germline BHD-Mutation Spectrum and Phenotype Analysis of a Large Cohort of Families with Birt-Hogg-Dubé Syndrome. American Journal of Human Genetics, 2005, 76, 1023-1033.	6.2	363
27	The Cancer Genome Atlas of renal cell carcinoma: findings and clinical implications. Nature Reviews Urology, 2019, 16, 539-552.	3.8	357
28	Birt-Hogg-Dub $\tilde{A}$ © Syndrome, a Genodermatosis Associated with Spontaneous Pneumothorax and Kidney Neoplasia, Maps to Chromosome 17p11.2. American Journal of Human Genetics, 2001, 69, 876-882.	6.2	355
29	Lung Cysts, Spontaneous Pneumothorax, and Genetic Associations in 89 Families with Birt-Hogg-Dubé Syndrome. American Journal of Respiratory and Critical Care Medicine, 2007, 175, 1044-1053.	5.6	318
30	High Frequency of <i>SDHB </i> Germline Mutations in Patients with Malignant Catecholamine-Producing Paragangliomas: Implications for Genetic Testing. Journal of Clinical Endocrinology and Metabolism, 2006, 91, 4505-4509.	3.6	299
31	Fusion of splicing factor genes PSF and NonO (p54nrb) to the TFE3 gene in papillary renal cell carcinoma. Oncogene, 1997, 15, 2233-2239.	5.9	298
32	Trisomy 7-harbouring non-random duplication of the mutant MET allele in hereditary papillary renal carcinomas. Nature Genetics, 1998, 20, 66-69.	21.4	291
33	Hereditary Papillary Renal Cell Carcinoma. Journal of Urology, 1994, 151, 561-566.	0.4	289
34	Measurements of Plasma Methoxytyramine, Normetanephrine, and Metanephrine as Discriminators of Different Hereditary Forms of Pheochromocytoma. Clinical Chemistry, 2011, 57, 411-420.	3.2	282
35	Oxidation of Alpha-Ketoglutarate Is Required for Reductive Carboxylation in Cancer Cells with Mitochondrial Defects. Cell Reports, 2014, 7, 1679-1690.	6.4	281
36	Pheochromocytomas in von Hippel-Lindau Syndrome and Multiple Endocrine Neoplasia Type 2 Display Distinct Biochemical and Clinical Phenotypes. Journal of Clinical Endocrinology and Metabolism, 2001, 86, 1999-2008.	3.6	262

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37	EVALUATION AND MANAGEMENT OF RENAL TUMORS IN THE BIRT-HOGG-DUBÉ SYNDROME. Journal of Urology, 2005, 173, 1482-1486.	0.4	260
38	Hereditary leiomyomatosis and renal cell cancer (HLRCC): renal cancer risk, surveillance and treatment. Familial Cancer, 2014, 13, 637-644.	1.9	251
39	Proteomic analysis of laser capture microdissected human prostate cancer andin vitro prostate cell lines. Electrophoresis, 2000, 21, 2235-2242.	2.4	246
40	Multiple Neuroendocrine Tumors of the Pancreas in von Hippel-Lindau Disease Patients. American Journal of Pathology, 1998, 153, 223-231.	3.8	243
41	Hereditary and Sporadic Papillary Renal Carcinomas with c-met Mutations Share a Distinct Morphological Phenotype. American Journal of Pathology, 1999, 155, 517-526.	3.8	243
42	Hereditary Leiomyomatosis and Renal Cell Cancer: A Syndrome Associated With an Aggressive Form of Inherited Renal Cancer. Journal of Urology, 2007, 177, 2074-2080.	0.4	235
43	CLINICAL AND GENETIC CHARACTERIZATION OF PHEOCHROMOCYTOMA IN VON HIPPEL-LINDAU FAMILIES: COMPARISON WITH SPORADIC PHEOCHROMOCYTOMA GIVES INSIGHT INTO NATURAL HISTORY OF PHEOCHROMOCYTOMA. Journal of Urology, 1999, 162, 659-664.	0.4	233
44	Clinical, genetic and radiographic analysis of 108 patients with von Hippel-Lindau disease (VHL) manifested by pancreatic neuroendocrine tumors (PNETs). Surgery, 2007, 142, 814-818.e2.	1.9	232
45	RENAL CANCER IN FAMILIES WITH HEREDITARY RENAL CANCER: PROSPECTIVE ANALYSIS OF A TUMOR SIZE THRESHOLD FOR RENAL PARENCHYMAL SPARING SURGERY. Journal of Urology, 1999, 161, 1475-1479.	0.4	229
46	Multiparametric Magnetic Resonance Imaging and Ultrasound Fusion Biopsy Detect Prostate Cancer in Patients with Prior Negative Transrectal Ultrasound Biopsies. Journal of Urology, 2012, 188, 2152-2157.	0.4	227
47	Molecular genetics and cellular features of TFE3 and TFEB fusion kidney cancers. Nature Reviews Urology, 2014, 11, 465-475.	3.8	227
48	Kidney-Targeted Birt-Hogg-Dube Gene Inactivation in a Mouse Model: Erk1/2 and Akt-mTOR Activation, Cell Hyperproliferation, and Polycystic Kidneys. Journal of the National Cancer Institute, 2008, 100, 140-154.	6.3	223
49	High Frequency of Somatic Frameshift BHD Gene Mutations in Birt-Hogg-Dubé–Associated Renal Tumors. Journal of the National Cancer Institute, 2005, 97, 931-935.	6.3	213
50	Fumarate Hydratase Deficiency in Renal Cancer Induces Glycolytic Addiction and Hypoxia-Inducible Transcription Factor $1^1$ ± Stabilization by Glucose-Dependent Generation of Reactive Oxygen Species. Molecular and Cellular Biology, 2009, 29, 4080-4090.	2.3	212
51	Succinate Dehydrogenase Kidney Cancer: An Aggressive Example of the Warburg Effect in Cancer. Journal of Urology, 2012, 188, 2063-2071.	0.4	211
52	Hereditary Renal Cancers. Radiology, 2003, 226, 33-46.	7.3	210
53	Robotic Partial Nephrectomy for Complex Renal Tumors: Surgical Technique. European Urology, 2008, 53, 514-523.	1.9	210
54	Homozygous loss of <i>BHD</i> causes early embryonic lethality and kidney tumor development with activation of mTORC1 and mTORC2. Proceedings of the National Academy of Sciences of the United States of America, 2009, 106, 18722-18727.	7.1	203

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55	Genetic basis of kidney cancer: Role of genomics for the development of disease-based therapeutics. Genome Research, 2012, 22, 2089-2100.	5.5	202
56	PARENCHYMAL SPARING SURGERY IN PATIENTS WITH HEREDITARY RENAL CELL CARCINOMA: 10-YEAR EXPERIENCE. Journal of Urology, 2001, 165, 777-781.	0.4	198
57	The Glycolytic Shift in Fumarate-Hydratase-Deficient Kidney Cancer Lowers AMPK Levels, Increases Anabolic Propensities and Lowers Cellular Iron Levels. Cancer Cell, 2011, 20, 315-327.	16.8	190
58	Correlation of Magnetic Resonance Imaging Tumor Volume with Histopathology. Journal of Urology, 2012, 188, 1157-1163.	0.4	188
59	Genetic Basis of Cancer of the Kidney. Clinical Cancer Research, 2004, 10, 6282S-6289S.	7.0	187
60	THE RELATIONSHIP BETWEEN RENAL TUMOR SIZE AND METASTASES IN PATIENTS WITH VON HIPPEL-LINDAU DISEASE. Journal of Urology, 2004, 172, 63-65.	0.4	181
61	Sarcomatoid Renal Cell Carcinoma: A Comprehensive Review of the Biology and Current Treatment Strategies. Oncologist, 2012, 17, 46-54.	3.7	177
62	Risk of renal and colonic neoplasms and spontaneous pneumothorax in the Birt-Hogg-Dub $\tilde{A}$ © syndrome. Cancer Epidemiology Biomarkers and Prevention, 2002, 11, 393-400.	2.5	177
63	Original Articles: Kidney Cancer: Hereditary Papillary Renal Cell Carcinoma: Clinical Studies in 10 Families. Journal of Urology, 1995, 153, 907-912.	0.4	176
64	Molecular genetics and clinical features of Birt–Hogg–Dubé syndrome. Nature Reviews Urology, 2015, 12, 558-569.	3.8	175
65	Molecular Pathways: <i>Fumarate Hydratase</i> Deficient Kidney Cancerâ€"Targeting the Warburg Effect in Cancer. Clinical Cancer Research, 2013, 19, 3345-3352.	7.0	172
66	Prevalence of Microscopic lesions in Grossly Normal Renal Parenchyma from Patients with von Hippel-Lindau Disease, Sporadic Renal Cell Carcinoma and No Renal Disease: Clinical Implications. Journal of Urology, 1995, 154, 2010-2015.	0.4	170
67	Catecholamine metabolomic and secretory phenotypes in phaeochromocytoma. Endocrine-Related Cancer, 2010, 18, 97-111.	3.1	169
68	Von Hippel-Lindau (VHL) Inactivation in Sporadic Clear Cell Renal Cancer: Associations with Germline VHL Polymorphisms and Etiologic Risk Factors. PLoS Genetics, 2011, 7, e1002312.	3.5	168
69	Identification and characterization of a novel folliculin-interacting protein FNIP2. Gene, 2008, 415, 60-67.	2.2	163
70	The Metabolic Basis of Kidney Cancer. Cancer Discovery, 2019, 9, 1006-1021.	9.4	163
71	LACK OF RETROPERITONEAL LYMPHADENOPATHY PREDICTS SURVIVAL OF PATIENTS WITH METASTATIC RENAL CELL CARCINOMA. Journal of Urology, 2001, 166, 68-72.	0.4	159
72	Molecular Diagnosis and Therapy of Kidney Cancer. Annual Review of Medicine, 2010, 61, 329-343.	12.2	154

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73	Robotic Partial Nephrectomy for Renal Hilar Tumors: A Multi-Institutional Analysis. Journal of Urology, 2008, 180, 2353-2356.	0.4	147
74	Inactivation of the FLCN Tumor Suppressor Gene Induces TFE3 Transcriptional Activity by Increasing Its Nuclear Localization. PLoS ONE, 2010, 5, e15793.	2.5	146
<b>7</b> 5	Rapid protein display profiling of cancer progression directly from human tissue using a protein biochip. Drug Development Research, 2000, 49, 34-42.	2.9	144
76	Characterization of the Renal Pathology of a Familial Form of Renal Cell Carcinoma Associated With Von Hippel-Lindau Disease: Clinical and Molecular Genetic Implications. Journal of Urology, 1995, 153, 22-26.	0.4	143
77	A Novel Germline Mutation in <i>BAP1</i> Predisposes to Familial Clear-Cell Renal Cell Carcinoma. Molecular Cancer Research, 2013, 11, 1061-1071.	3.4	135
78	Defining Early-Onset Kidney Cancer: Implications for Germline and Somatic Mutation Testing and Clinical Management. Journal of Clinical Oncology, 2014, 32, 431-437.	1.6	135
79	FLCN : The causative gene for Birt-Hogg-Dubé syndrome. Gene, 2018, 640, 28-42.	2.2	133
80	Regression of Metastatic Renal Cell Carcinoma After Cytoreductive Nephrectomy. Journal of Urology, 1993, 150, 463-466.	0.4	132
81	The metabolic basis of kidney cancer. Seminars in Cancer Biology, 2013, 23, 46-55.	9.6	132
82	Identification of the Genes for Kidney Cancer: Opportunity for Disease-Specific Targeted Therapeutics. Clinical Cancer Research, 2007, 13, 671s-679s.	7.0	131
83	UOK 262 cell line, fumarate hydratase deficient (FHâ^'/FHâ^') hereditary leiomyomatosis renal cell carcinoma: in vitro and in vivo model of an aberrant energy metabolic pathway in human cancer. Cancer Genetics and Cytogenetics, 2010, 196, 45-55.	1.0	131
84	FAMILIAL RENAL ONCOCYTOMA: CLINICOPATHOLOGICAL STUDY OF 5 FAMILIES. Journal of Urology, 1998, 160, 335-340.	0.4	127
85	Expression of Birt–Hogg–Dubé gene mRNA in normal and neoplastic human tissues. Modern Pathology, 2004, 17, 998-1011.	5.5	124
86	CYTOREDUCTIVE SURGERY BEFORE HIGH DOSE INTERLEUKIN-2 BASED THERAPY IN PATIENTS WITH METASTATIC RENAL CELL CARCINOMA. Journal of Urology, 1997, 158, 1675-1678.	0.4	120
87	Focus on kidney cancer. Cancer Cell, 2004, 6, 223-228.	16.8	119
88	EARLY ONSET HEREDITARY PAPILLARY RENAL CARCINOMA: GERMLINE MISSENSE MUTATIONS IN THE TYROSINE KINASE DOMAIN OF THE MET PROTO-ONCOGENE. Journal of Urology, 2004, 172, 1256-1261.	0.4	115
89	Hereditary leiomyomatosis and renal cell carcinoma. International Journal of Nephrology and Renovascular Disease, 2014, 7, 253.	1.8	112
90	Dual-color, Break-apart FISH Assay on Paraffin-embedded Tissues as an Adjunct to Diagnosis of Xp11 Translocation Renal Cell Carcinoma and Alveolar Soft Part Sarcoma. American Journal of Surgical Pathology, 2010, 34, 757-766.	3.7	111

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91	Decreased expression of the pro-apoptotic protein Par-4 in renal cell carcinoma. Oncogene, 1999, 18, 1205-1208.	5.9	108
92	Genetic predisposition to kidney cancer. Seminars in Oncology, 2016, 43, 566-574.	2.2	107
93	INTERLEUKIN-2 BASED IMMUNOTHERAPY FOR METASTATIC RENAL CELL CARCINOMA WITH THE KIDNEY IN PLACE. Journal of Urology, 1999, 162, 43-45.	0.4	102
94	Surgical Management of Pheochromocytoma with the Use of Metyrosine. Annals of Surgery, 1990, 212, 621-628.	4.2	101
95	Hereditary kidney cancer. Cancer, 2009, 115, 2252-2261.	4.1	101
96	Translocation Renal Cell Carcinomas in Adults. American Journal of Surgical Pathology, 2012, 36, 654-662.	3.7	98
97	MANAGEMENT OF HEREDITARY PHEOCHROMOCYTOMA IN VON HIPPEL-LINDAU KINDREDS WITH PARTIAL ADRENALECTOMY. Journal of Urology, 1999, 161, 395-398.	0.4	97
98	Epididymal cystadenomas in von Hippel-Lindau disease. Urology, 1997, 49, 926-931.	1.0	96
99	SDHB-Deficient Cancers: The Role of Mutations That Impair Iron Sulfur Cluster Delivery. Journal of the National Cancer Institute, 2016, 108, djv287.	6.3	92
100	PREVALENCE OF MICROSCOPIC TUMORS IN NORMAL APPEARING RENAL PARENCHYMA OF PATIENTS WITH HEREDITARY PAPILLARY RENAL CANCER. Journal of Urology, 2000, 163, 431-433.	0.4	91
101	Alternative splicing of the cell fate determinant Numb in hepatocellular carcinoma. Hepatology, 2015, 62, 1122-1131.	7.3	91
102	Preparative Cytoreductive Surgery in Patients with Metastatic Renal Cell Carcinoma Treated with Adoptive Immunotherapy with Interleukin-2 or Interleukin-2 Plus Lymphokine Activated Killer Cells. Journal of Urology, 1990, 144, 614-617.	0.4	90
103	Partial adrenalectomy: The National Cancer Institute experience. Urology, 2005, 66, 19-23.	1.0	89
104	New Strategies in Renal Cell Carcinoma: Targeting the Genetic and Metabolic Basis of Disease. Clinical Cancer Research, 2015, 21, 10-17.	7.0	88
105	Original Articles: Kidney Cancer: Parenchymal Sparing Surgery in Patients With Hereditary Renal Cell Carcinoma. Journal of Urology, 1995, 153, 913-916.	0.4	87
106	Targeting ABL1-Mediated Oxidative Stress Adaptation in Fumarate Hydratase-Deficient Cancer. Cancer Cell, 2014, 26, 840-850.	16.8	87
107	Detection of an Immunogenic HERV-E Envelope with Selective Expression in Clear Cell Kidney Cancer. Cancer Research, 2016, 76, 2177-2185.	0.9	86
108	Solid renal tumor severity in von Hippel Lindau disease is related to germline deletion length and location. Human Mutation, 2004, 23, 40-46.	2.5	85

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109	Germline PTEN Mutation Cowden Syndrome: An Underappreciated Form of Hereditary Kidney Cancer. Journal of Urology, 2013, 190, 1990-1998.	0.4	85
110	Diagnosis and management of BHD-associated kidney cancer. Familial Cancer, 2013, 12, 397-402.	1.9	85
111	Folliculin Controls Lung Alveolar Enlargement and Epithelial Cell Survival through E-Cadherin, LKB1, and AMPK. Cell Reports, 2014, 7, 412-423.	6.4	84
112	Integrative molecular characterization of sarcomatoid and rhabdoid renal cell carcinoma. Nature Communications, 2021, 12, 808.	12.8	84
113	Endolymphatic sac tumors in von Hippel—Lindau disease. Journal of Neurosurgery, 2004, 100, 480-487.	1.6	83
114	Salvage Partial Nephrectomy for Hereditary Renal Cancer: Feasibility and Outcomes. Journal of Urology, 2008, 179, 67-70.	0.4	83
115	Association of Germline Mutations in the Fumarate Hydratase Gene and Uterine Fibroids in Women With Hereditary Leiomyomatosis and Renal Cell Cancer. Archives of Dermatology, 2008, 144, 1584-92.	1.4	83
116	Regulation of Mitochondrial Oxidative Metabolism by Tumor Suppressor FLCN. Journal of the National Cancer Institute, 2012, 104, 1750-1764.	6.3	82
117	Tumor suppressor FLCN inhibits tumorigenesis of a FLCN-null renal cancer cell line and regulates expression of key molecules in TGF- $\hat{l}^2$ signaling. Molecular Cancer, 2010, 9, 160.	19.2	81
118	Development of a prostate cDNA microarray and statistical gene expression analysis package. , 2000, 28, 12-22.		80
119	Familial Kidney Cancer: Implications of New Syndromes and Molecular Insights. European Urology, 2019, 76, 754-764.	1.9	80
120	Metabolic Reprogramming for Producing Energy and Reducing Power in Fumarate Hydratase Null Cells from Hereditary Leiomyomatosis Renal Cell Carcinoma. PLoS ONE, 2013, 8, e72179.	2.5	80
121	EXPRESSION STUDIES AND MUTATIONAL ANALYSIS OF THE ANDROGEN REGULATED HOMEOBOX GENE NKX3.1 IN BENIGN AND MALIGNANT PROSTATE EPITHELIUM. Journal of Urology, 2001, 165, 1329-1334.	0.4	79
122	Discovering Targets of Non-enzymatic Acylation by Thioester Reactivity Profiling. Cell Chemical Biology, 2017, 24, 231-242.	5.2	79
123	ONC201 kills breast cancer cells <i>i) in vitro</i> by targeting mitochondria. Oncotarget, 2018, 9, 18454-18479.	1.8	77
124	A chemoproteomic portrait of the oncometabolite fumarate. Nature Chemical Biology, 2019, 15, 391-400.	8.0	77
125	Englerin A Stimulates PKCÎ, to Inhibit Insulin Signaling and to Simultaneously Activate HSF1: Pharmacologically Induced Synthetic Lethality. Cancer Cell, 2013, 23, 228-237.	16.8	74
126	Folliculin-interacting proteins Fnip1 and Fnip2 play critical roles in kidney tumor suppression in cooperation with Flcn. Proceedings of the National Academy of Sciences of the United States of America, 2015, 112, E1624-31.	7.1	74

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127	Dynamic Imaging of LDH Inhibition in Tumors Reveals Rapid InÂVivo Metabolic Rewiring and Vulnerability to Combination Therapy. Cell Reports, 2020, 30, 1798-1810.e4.	6.4	73
128	Fe-S cofactors in the SARS-CoV-2 RNA-dependent RNA polymerase are potential antiviral targets. Science, 2021, 373, 236-241.	12.6	71
129	Genotype-Phenotype Correlation in von Hippel-Lindau Disease With Retinal Angiomatosis. JAMA Ophthalmology, 2007, 125, 239.	2.4	70
130	Acute loss of iron–sulfur clusters results in metabolic reprogramming and generation of lipid droplets in mammalian cells. Journal of Biological Chemistry, 2018, 293, 8297-8311.	3.4	70
131	Regulatory Effects of microRNA-92 (miR-92) on <i>VHL</i> Gene Expression and the Hypoxic Activation of miR-210 in Clear Cell Renal Cell Carcinoma. Journal of Cancer, 2011, 2, 515-526.	2.5	69
132	Studying Cancer Families to Identify Kidney Cancer Genes. Annual Review of Medicine, 2003, 54, 217-233.	12.2	66
133	Targeting the Met signaling pathway in renal cancer. Expert Review of Anticancer Therapy, 2009, 9, 785-793.	2.4	66
134	Initial Experience With Robot Assisted Partial Nephrectomy for Multiple Renal Masses. Journal of Urology, 2009, 182, 1280-1286.	0.4	66
135	Functional and Oncologic Outcomes of Partial Adrenalectomy for Pheochromocytoma in Patients With von Hippel-Lindau Syndrome After at Least 5 Years of Followup. Journal of Urology, 2010, 184, 1855-1859.	0.4	66
136	Superiority of 68Ga-DOTATATE over 18F-FDG and anatomic imaging in the detection of succinate dehydrogenase mutation (SDHx )-related pheochromocytoma and paraganglioma in the pediatric population. European Journal of Nuclear Medicine and Molecular Imaging, 2018, 45, 787-797.	6.4	64
137	Mitochondrial DNA alterations underlie an irreversible shift to aerobic glycolysis in fumarate hydratase–deficient renal cancer. Science Signaling, 2021, 14, .	3.6	64
138	Imaging Features of Hereditary Papillary Renal Cancers. Journal of Computer Assisted Tomography, 1997, 21, 737-741.	0.9	63
139	Metabolism of Kidney Cancer: From the Lab to Clinical Practice. European Urology, 2013, 63, 244-251.	1.9	61
140	Updated Recommendations on the Diagnosis, Management, and Clinical Trial Eligibility Criteria for Patients With Renal Medullary Carcinoma. Clinical Genitourinary Cancer, 2019, 17, 1-6.	1.9	60
141	Robot-Assisted Laparoscopic Partial Adrenalectomy for Pheochromocytoma: The National Cancer Institute Technique. European Urology, 2011, 60, 118-124.	1.9	58
142	Therapeutic Targeting of TFE3/IRS-1/PI3K/mTOR Axis in Translocation Renal Cell Carcinoma. Clinical Cancer Research, 2018, 24, 5977-5989.	7.0	58
143	Tumor-Specific Hypermethylation of Epigenetic Biomarkers, Including SFRP1, Predicts for Poorer Survival in Patients from the TCGA Kidney Renal Clear Cell Carcinoma (KIRC) Project. PLoS ONE, 2014, 9, e85621.	2.5	58
144	The folliculin-FNIP1 pathway deleted in human Birt-Hogg-Dubé syndrome is required for murine B-cell development. Blood, 2012, 120, 1254-1261.	1.4	57

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145	The FNIP co-chaperones decelerate the Hsp90 chaperone cycle and enhance drug binding. Nature Communications, 2016, 7, 12037.	12.8	56
146	Gender Specific Mutation Incidence and Survival Associations in Clear Cell Renal Cell Carcinoma (CCRCC). PLoS ONE, 2015, 10, e0140257.	2.5	56
147	The UOK 257 cell line: a novel model for studies of the human Birt–Hogg–Dubé gene pathway. Cancer Genetics and Cytogenetics, 2008, 180, 100-109.	1.0	55
148	A novel fumarate hydratase-deficient HLRCC kidney cancer cell line, UOK268: a model of the Warburg effect in cancer. Cancer Genetics, 2012, 205, 377-390.	0.4	55
149	MicroRNAâ€204â€5p: A novel candidate urinary biomarker of Xp11.2 translocation renal cell carcinoma. Cancer Science, 2019, 110, 1897-1908.	3.9	55
150	Folliculin (Flcn) inactivation leads to murine cardiac hypertrophy through mTORC1 deregulation. Human Molecular Genetics, 2014, 23, 5706-5719.	2.9	54
151	Differential expression of the mismatch repair genehMSH2 in malignant prostate tissue is associated with cancer recurrence. Cancer, 2002, 94, 690-699.	4.1	53
152	Results from a phase II study of bevacizumab and erlotinib in subjects with advanced hereditary leiomyomatosis and renal cell cancer (HLRCC) or sporadic papillary renal cell cancer Journal of Clinical Oncology, 2020, 38, 5004-5004.	1.6	53
153	Genetic basis for kidney cancer: opportunity for disease-specific approaches to therapy. Expert Opinion on Biological Therapy, 2008, 8, 779-790.	3.1	51
154	Identification of intragenic deletions and duplication in the ⟨i⟩FLCN⟨/i⟩ gene in Birtâ€Hoggâ€Dubé syndrome. Genes Chromosomes and Cancer, 2011, 50, 466-477.	2.8	50
155	VHL loss of function and its impact on oncogenic signaling networks in clear cell renal cell carcinoma. International Journal of Biochemistry and Cell Biology, 2009, 41, 753-756.	2.8	49
156	Loss of heterozygosity on the short arm of chromosome 3 in sporadic, von hippel-lindau disease-associated, and familial pheochromocytoma. Genes Chromosomes and Cancer, 1995, 13, 151-156.	2.8	48
157	Management of von Hippel–Lindau-associated kidney cancer. Nature Reviews Urology, 2005, 2, 248-255.	1.4	48
158	Robot-assisted Laparoscopic Partial Adrenalectomy: Initial Experience. Urology, 2011, 77, 775-780.	1.0	47
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