

Karel Pacak

List of Publications by Year in descending order

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497
papers

36,219
citations

2795

94
h-index

4750

169
g-index

509
all docs

509
docs citations

509
times ranked

19871
citing authors

#	ARTICLE	IF	CITATIONS
1	Pheochromocytoma and Paraganglioma: An Endocrine Society Clinical Practice Guideline. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2014, 99, 1915-1942.	1.8	2,031
2	Phaeochromocytoma. <i>Lancet</i> , The, 2005, 366, 665-675.	6.3	1,462
3	Biochemical Diagnosis of Pheochromocytoma. <i>JAMA - Journal of the American Medical Association</i> , 2002, 287, 1427-34.	3.8	994
4	Irisin and FGF21 Are Cold-Induced Endocrine Activators of Brown Fat Function in Humans. <i>Cell Metabolism</i> , 2014, 19, 302-309.	7.2	643
5	Cushing's Syndrome Due to Ectopic Corticotropin Secretion: Twenty Years' Experience at the National Institutes of Health. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2005, 90, 4955-4962.	1.8	588
6	Pheochromocytoma: recommendations for clinical practice from the First International Symposium. <i>Nature Clinical Practice Endocrinology and Metabolism</i> , 2007, 3, 92-102.	2.9	581
7	Comprehensive Molecular Characterization of Pheochromocytoma and Paraganglioma. <i>Cancer Cell</i> , 2017, 31, 181-193.	7.7	532
8	Recent Advances in Genetics, Diagnosis, Localization, and Treatment of Pheochromocytoma. <i>Annals of Internal Medicine</i> , 2001, 134, 315.	2.0	512
9	Preoperative Management of the Pheochromocytoma Patient. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2007, 92, 4069-4079.	1.8	497
10	The North American Neuroendocrine Tumor Society Consensus Guideline for the Diagnosis and Management of Neuroendocrine Tumors. <i>Pancreas</i> , 2010, 39, 775-783.	0.5	493
11	An immunohistochemical procedure to detect patients with paraganglioma and pheochromocytoma with germline SDHB, SDHC, or SDHD gene mutations: a retrospective and prospective analysis. <i>Lancet Oncology</i> , The, 2009, 10, 764-771.	5.1	477
12	Biochemical Diagnosis of Pheochromocytoma: How to Distinguish True- from False-Positive Test Results. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2003, 88, 2656-2666.	1.8	447
13	Clinical and molecular genetics of patients with the Carney-Stratakis syndrome and germline mutations of the genes coding for the succinate dehydrogenase subunits SDHB, SDHC, and SDHD. <i>European Journal of Human Genetics</i> , 2008, 16, 79-88.	1.4	446
14	Mapping of human brown adipose tissue in lean and obese young men. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2017, 114, 8649-8654.	3.3	370
15	Comparison of 18F-Fluoro-L-DOPA, 18F-Fluoro-Deoxyglucose, and 18F-Fluorodopamine PET and 123I-MIBG Scintigraphy in the Localization of Pheochromocytoma and Paraganglioma. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2009, 94, 4757-4767.	1.8	361
16	Stress-Induced Norepinephrine Release in the Hypothalamic Paraventricular Nucleus and Pituitary-Adrenocortical and Sympathoadrenal Activity: In Vivo Microdialysis Studies. <i>Frontiers in Neuroendocrinology</i> , 1995, 16, 89-150.	2.5	348
17	Somatic <i>HIF2A</i> Gain-of-Function Mutations in Paraganglioma with Polycythemia. <i>New England Journal of Medicine</i> , 2012, 367, 922-930.	13.9	338
18	Superiority of Fluorodeoxyglucose Positron Emission Tomography to Other Functional Imaging Techniques in the Evaluation of Metastatic SDHB-Associated Pheochromocytoma and Paraganglioma. <i>Journal of Clinical Oncology</i> , 2007, 25, 2262-2269.	0.8	316

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19	Plasma methoxytyramine: A novel biomarker of metastatic pheochromocytoma and paraganglioma in relation to established risk factors of tumour size, location and SDHB mutation status. <i>European Journal of Cancer</i> , 2012, 48, 1739-1749.	1.3	304
20	Malignant pheochromocytoma: current status and initiatives for future progress. <i>Endocrine-Related Cancer</i> , 2004, 11, 423-436.	1.6	299
21	High Frequency ofSDHBGermline Mutations in Patients with Malignant Catecholamine-Producing Paragangliomas: Implications for Genetic Testing. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2006, 91, 4505-4509.	1.8	299
22	Current Approaches and Recommended Algorithm for the Diagnostic Localization of Pheochromocytoma. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2004, 89, 479-491.	1.8	296
23	Molecular Subtypes of <i>KIT/PDGFR</i> Wild-Type Gastrointestinal Stromal Tumors. <i>JAMA Oncology</i> , 2016, 2, 922.	3.4	291
24	Succinate Dehydrogenase Mutation Underlies Global Epigenomic Divergence in Gastrointestinal Stromal Tumor. <i>Cancer Discovery</i> , 2013, 3, 648-657.	7.7	288
25	Prospective Study of ⁶⁸ Ga-DOTATATE Positron Emission Tomography/Computed Tomography for Detecting Gastro-Entero-Pancreatic Neuroendocrine Tumors and Unknown Primary Sites. <i>Journal of Clinical Oncology</i> , 2016, 34, 588-596.	0.8	287
26	Measurements of Plasma Methoxytyramine, Normetanephrine, and Metanephrine as Discriminators of Different Hereditary Forms of Pheochromocytoma. <i>Clinical Chemistry</i> , 2011, 57, 411-420.	1.5	282
27	<i>MAX</i> Mutations Cause Hereditary and Sporadic Pheochromocytoma and Paraganglioma. <i>Clinical Cancer Research</i> , 2012, 18, 2828-2837.	3.2	277
28	Pheochromocytomas in von Hippel-Lindau Syndrome and Multiple Endocrine Neoplasia Type 2 Display Distinct Biochemical and Clinical Phenotypes. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2001, 86, 1999-2008.	1.8	262
29	Clinical Presentations, Biochemical Phenotypes, and Genotype-Phenotype Correlations in Patients with Succinate Dehydrogenase Subunit B-Associated Pheochromocytomas and Paragangliomas. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2007, 92, 779-786.	1.8	262
30	New Perspectives on Pheochromocytoma and Paraganglioma: Toward a Molecular Classification. <i>Endocrine Reviews</i> , 2017, 38, 489-515.	8.9	241
31	Staging and Functional Characterization of Pheochromocytoma and Paraganglioma by ¹⁸ F-Fluorodeoxyglucose (¹⁸ F-FDG) Positron Emission Tomography. <i>Journal of the National Cancer Institute</i> , 2012, 104, 700-708.	3.0	240
32	Superiority of 6-[¹⁸ F]-Fluorodopamine Positron Emission Tomography Versus [¹³¹ I]-Metaiodobenzylguanidine Scintigraphy in the Localization of Metastatic Pheochromocytoma. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2003, 88, 4083-4087.	1.8	237
33	Treatment of malignant pheochromocytoma/paraganglioma with cyclophosphamide, vincristine, and dacarbazine. <i>Cancer</i> , 2008, 113, 2020-2028.	2.0	234
34	EANM 2012 guidelines for radionuclide imaging of phaeochromocytoma and paraganglioma. <i>European Journal of Nuclear Medicine and Molecular Imaging</i> , 2012, 39, 1977-1995.	3.3	223
35	Superiority of [⁶⁸ Ga]-DOTATATE PET/CT to Other Functional Imaging Modalities in the Localization of <i>SDHB</i> -Associated Metastatic Pheochromocytoma and Paraganglioma. <i>Clinical Cancer Research</i> , 2015, 21, 3888-3895.	3.2	223
36	The optimal imaging of adrenal tumours: a comparison of different methods. <i>Endocrine-Related Cancer</i> , 2007, 14, 587-599.	1.6	218

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37	6-[¹⁸ F]Fluorodopamine Positron Emission Tomographic (PET) Scanning for Diagnostic Localization of Pheochromocytoma. <i>Hypertension</i> , 2001, 38, 6-8.	1.3	215
38	Biochemical and Clinical Manifestations of Dopamine-Producing Paragangliomas: Utility of Plasma Methoxytyramine. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2005, 90, 2068-2075.	1.8	213
39	Succinate Dehydrogenase Kidney Cancer: An Aggressive Example of the Warburg Effect in Cancer. <i>Journal of Urology</i> , 2012, 188, 2063-2071.	0.2	211
40	European Association of Nuclear Medicine Practice Guideline/Society of Nuclear Medicine and Molecular Imaging Procedure Standard 2019 for radionuclide imaging of pheochromocytoma and paraganglioma. <i>European Journal of Nuclear Medicine and Molecular Imaging</i> , 2019, 46, 2112-2137.	3.3	208
41	Mitochondrial Complex II: At the Crossroads. <i>Trends in Biochemical Sciences</i> , 2017, 42, 312-325.	3.7	192
42	Reactivation of Dihydroorotate Dehydrogenase-Driven Pyrimidine Biosynthesis Restores Tumor Growth of Respiration-Deficient Cancer Cells. <i>Cell Metabolism</i> , 2019, 29, 399-416.e10.	7.2	190
43	Genetics, diagnosis, management and future directions of research of pheochromocytoma and paraganglioma: a position statement and consensus of the Working Group on Endocrine Hypertension of the European Society of Hypertension. <i>Journal of Hypertension</i> , 2020, 38, 1443-1456.	0.3	190
44	Monoamine Oxidase A-mediated Enhanced Catabolism of Norepinephrine Contributes to Adverse Remodeling and Pump Failure in Hearts With Pressure Overload. <i>Circulation Research</i> , 2010, 106, 193-202.	2.0	187
45	Current Treatment of Malignant Pheochromocytoma. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2007, 92, 1217-1225.	1.8	180
46	Pheochromocytoma Catecholamine Phenotypes and Prediction of Tumor Size and Location by Use of Plasma Free Metanephrines. <i>Clinical Chemistry</i> , 2005, 51, 735-744.	1.5	177
47	SDHB/SDHA immunohistochemistry in pheochromocytomas and paragangliomas: a multicenter interobserver variation analysis using virtual microscopy: a Multinational Study of the European Network for the Study of Adrenal Tumors (ENS@T). <i>Modern Pathology</i> , 2015, 28, 807-821.	2.9	176
48	The Molecular Pathogenesis of Hereditary and Sporadic Adrenocortical and Adrenomedullary Tumors. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2002, 87, 5367-5384.	1.8	174
49	Metastatic Pheochromocytoma/Paraganglioma Related to Primary Tumor Development in Childhood or Adolescence: Significant Link to SDHB Mutations. <i>Journal of Clinical Oncology</i> , 2011, 29, 4137-4142.	0.8	170
50	Catecholamine metabolomic and secretory phenotypes in pheochromocytoma. <i>Endocrine-Related Cancer</i> , 2010, 18, 97-111.	1.6	169
51	Pheochromocytoma and paraganglioma: Diagnosis, genetics, management, and treatment. <i>Current Problems in Cancer</i> , 2014, 38, 7-41.	1.0	163
52	Understanding catecholamine metabolism as a guide to the biochemical diagnosis of pheochromocytoma. <i>Reviews in Endocrine and Metabolic Disorders</i> , 2001, 2, 297-311.	2.6	156
53	A novel EPAS1/HIF2A germline mutation in a congenital polycythemia with paraganglioma. <i>Journal of Molecular Medicine</i> , 2013, 91, 507-512.	1.7	155
54	Hypertension in Pheochromocytoma: Characteristics and Treatment. <i>Endocrinology and Metabolism Clinics of North America</i> , 2011, 40, 295-311.	1.2	153

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73	Adverse Drug Reactions in Patients with Pheochromocytoma. <i>Drug Safety</i> , 2007, 30, 1031-1062.	1.4	122
74	Pituitary Adenoma With Paraganglioma/Pheochromocytoma (3PAs) and Succinate Dehydrogenase Defects in Humans and Mice. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2015, 100, E710-E719.	1.8	121
75	Characteristics of Pediatric vs Adult Pheochromocytomas and Paragangliomas. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2017, 102, 1122-1132.	1.8	120
76	Current Management of Pheochromocytoma/Paraganglioma: A Guide for the Practicing Clinician in the Era of Precision Medicine. <i>Cancers</i> , 2019, 11, 1505.	1.7	120
77	Heterogeneous neurochemical responses to different stressors: a test of Selye's doctrine of nonspecificity. <i>American Journal of Physiology - Regulatory Integrative and Comparative Physiology</i> , 1998, 275, R1247-R1255.	0.9	119
78	Molecular Imaging of Gastroenteropancreatic Neuroendocrine Tumors: Current Status and Future Directions. <i>Journal of Nuclear Medicine</i> , 2016, 57, 1949-1956.	2.8	119
79	Central Nervous System Imprinting of the G Protein Gs α and Its Role in Metabolic Regulation. <i>Cell Metabolism</i> , 2009, 9, 548-555.	7.2	118
80	Germ-line PHD1 and PHD2 mutations detected in patients with pheochromocytoma/paraganglioma-polycythemia. <i>Journal of Molecular Medicine</i> , 2015, 93, 93-104.	1.7	118
81	Clinical aspects of SDHx-related pheochromocytoma and paraganglioma. <i>Endocrine-Related Cancer</i> , 2009, 16, 391-400.	1.6	117
82	Biochemical Diagnosis and Localization of Pheochromocytoma: Can We Reach a Consensus?. <i>Annals of the New York Academy of Sciences</i> , 2006, 1073, 332-347.	1.8	115
83	New imaging approaches to pheochromocytomas and paragangliomas. <i>Clinical Endocrinology</i> , 2010, 72, 137-145.	1.2	112
84	Biochemically Silent Abdominal Paragangliomas in Patients with Mutations in the Succinate Dehydrogenase Subunit B Gene. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2008, 93, 4826-4832.	1.8	111
85	Krebs Cycle Metabolite Profiling for Identification and Stratification of Pheochromocytomas/Paragangliomas due to Succinate Dehydrogenase Deficiency. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2014, 99, 3903-3911.	1.8	111
86	Noradrenergic activation in the paraventricular nucleus during acute and chronic immobilization stress in rats: an in vivo microdialysis study. <i>Brain Research</i> , 1992, 589, 91-96.	1.1	110
87	Is Supine Rest Necessary before Blood Sampling for Plasma Metanephrines?. <i>Clinical Chemistry</i> , 2007, 53, 352-354.	1.5	110
88	Characteristics And Outcomes Of Metastatic Sdhb And Sporadic Pheochromocytoma/Paraganglioma: An National Institutes Of Health Study. <i>Endocrine Practice</i> , 2016, 22, 302-314.	1.1	110
89	False-negative 123I-MIBG SPECT is most commonly found in SDHB-related pheochromocytoma or paraganglioma with high frequency to develop metastatic disease. <i>Endocrine-Related Cancer</i> , 2012, 19, 83-93.	1.6	106
90	Plasma Metadrenalines: Do they Provide Useful Information about Sympatho-Adrenal Function and Catecholamine Metabolism?. <i>Clinical Science</i> , 1995, 88, 533-542.	1.8	105

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91	Utility of Plasma Free Metanephrines for Detecting Childhood Pheochromocytoma. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2002, 87, 1955-1960.	1.8	104
92	Update on pediatric pheochromocytoma. <i>Pediatric Nephrology</i> , 2009, 24, 943-950.	0.9	102
93	The Effects of Carbidopa on Uptake of 6-18F-Fluoro-L-DOPA in PET of Pheochromocytoma and Extraadrenal Abdominal Paraganglioma. <i>Journal of Nuclear Medicine</i> , 2007, 48, 1599-1606.	2.8	101
94	Reference intervals for plasma free metanephrines with an age adjustment for normetanephrine for optimized laboratory testing of pheochromocytoma. <i>Annals of Clinical Biochemistry</i> , 2013, 50, 62-69.	0.8	98
95	Prospective comparison of 68Ga-DOTATATE and 18F-FDOPA PET/CT in patients with various pheochromocytomas and paragangliomas with emphasis on sporadic cases. <i>European Journal of Nuclear Medicine and Molecular Imaging</i> , 2016, 43, 1248-1257.	3.3	96
96	Drugs and Pheochromocytoma – Don't Be Fooled by Every Elevated Metanephrine. <i>New England Journal of Medicine</i> , 2011, 364, 2268-2270.	13.9	95
97	New Roles of Carboxypeptidase E in Endocrine and Neural Function and Cancer. <i>Endocrine Reviews</i> , 2012, 33, 216-253.	8.9	95
98	Effects of immobilization on in vivo release of norepinephrine in the bed nucleus of the stria terminalis in conscious rats. <i>Brain Research</i> , 1995, 688, 242-246.	1.1	92
99	Downregulation of metastasis suppressor genes in malignant pheochromocytoma. <i>International Journal of Cancer</i> , 2005, 114, 139-143.	2.3	92
100	Brown Fat Imaging with 18F-6-Fluorodopamine PET/CT, 18F-FDG PET/CT, and 123I-MIBG SPECT: A Study of Patients Being Evaluated for Pheochromocytoma. <i>Journal of Nuclear Medicine</i> , 2007, 48, 1077-1083.	2.8	92
101	Metastatic Paraganglioma. <i>Seminars in Oncology</i> , 2010, 37, 627-637.	0.8	91
102	Age at Diagnosis of Pheochromocytoma Differs According to Catecholamine Phenotype and Tumor Location. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2011, 96, 375-384.	1.8	90
103	Cardiac sympathetic denervation preceding motor signs in Parkinson disease. <i>Clinical Autonomic Research</i> , 2007, 17, 118-121.	1.4	88
104	Pheochromocytoma as an endocrine emergency. <i>Reviews in Endocrine and Metabolic Disorders</i> , 2003, 4, 121-128.	2.6	87
105	Novel HIF2A mutations disrupt oxygen sensing, leading to polycythemia, paragangliomas, and somatostatinomas. <i>Blood</i> , 2013, 121, 2563-2566.	0.6	87
106	Genotype-phenotype correlations in pheochromocytoma and paraganglioma: a systematic review and individual patient meta-analysis. <i>Endocrine-Related Cancer</i> , 2019, 26, 539-550.	1.6	87
107	Pheochromocytoma crisis induced by glucocorticoids: a report of four cases and review of the literature. <i>European Journal of Endocrinology</i> , 2008, 158, 423-429.	1.9	86
108	Discordant localization of 2-[18F]-fluoro-2-deoxy-D-glucose in 6-[18F]-fluorodopamine- and [123I]-metaiodobenzylguanidine-negative metastatic pheochromocytoma sites. <i>Nuclear Medicine Communications</i> , 2006, 27, 31-36.	0.5	85

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109	Partial Adrenalectomy: Underused First Line Therapy for Small Adrenal Tumors. <i>Journal of Urology</i> , 2010, 184, 18-25.	0.2	85
110	15 YEARS OF PARAGANGLIOMA: Imaging and imaging-based treatment of pheochromocytoma and paraganglioma. <i>Endocrine-Related Cancer</i> , 2015, 22, T135-T145.	1.6	84
111	Radiofrequency Ablation: a Novel Approach for Treatment of Metastatic Pheochromocytoma. <i>Journal of the National Cancer Institute</i> , 2001, 93, 648-649.	3.0	83
112	Update of Pheochromocytoma Syndromes: Genetics, Biochemical Evaluation, and Imaging. <i>Frontiers in Endocrinology</i> , 2018, 9, 515.	1.5	82
113	Minimal changes in environmental temperature result in a significant increase in energy expenditure and changes in the hormonal homeostasis in healthy adults. <i>European Journal of Endocrinology</i> , 2010, 163, 863-872.	1.9	80
114	International consensus on initial screening and follow-up of asymptomatic SDHx mutation carriers. <i>Nature Reviews Endocrinology</i> , 2021, 17, 435-444.	4.3	80
115	Accuracy of recommended sampling and assay methods for the determination of plasma-free and urinary fractionated metanephrines in the diagnosis of pheochromocytoma and paraganglioma: a systematic review. <i>Endocrine</i> , 2017, 56, 495-503.	1.1	79
116	Impaired Basal and Restraint-Induced Epinephrine Secretion in Corticotropin-Releasing Hormone-Deficient Mice ¹ . <i>Endocrinology</i> , 2000, 141, 1142-1150.	1.4	78
117	The role of 6-[¹⁸ F]fluorodopamine positron emission tomography in the localization of adrenal pheochromocytoma associated with von Hippel-Lindau syndrome. <i>European Journal of Endocrinology</i> , 2007, 156, 483-487.	1.9	78
118	SDH-related pheochromocytoma and paraganglioma. <i>Best Practice and Research in Clinical Endocrinology and Metabolism</i> , 2010, 24, 415-424.	2.2	78
119	Subclinical phaeochromocytoma. <i>Best Practice and Research in Clinical Endocrinology and Metabolism</i> , 2012, 26, 507-515.	2.2	76
120	Pheochromocytoma in von Hippel-Lindau Disease: Distinct Histopathologic Phenotype Compared to Pheochromocytoma in Multiple Endocrine Neoplasia Type 2. <i>Endocrine Pathology</i> , 2002, 13, 17-28.	5.2	74
121	Use of ¹⁸ F-fluorodopamine positron emission tomography (PET) as first-line investigation for the diagnosis and localization of non-metastatic and metastatic phaeochromocytoma (PHEO). <i>Clinical Endocrinology</i> , 2009, 71, 11-17.	1.2	74
122	Plasma metanephrines in renal failure. <i>Kidney International</i> , 2005, 67, 668-677.	2.6	73
123	Opposing effects of HIF1 [±] and HIF2 [±] on chromaffin cell phenotypic features and tumor cell proliferation: Insights from MYC-associated factor X. <i>International Journal of Cancer</i> , 2014, 135, 2054-2064.	2.3	72
124	Results of ⁶⁸ Gallium-DOTATATE PET/CT Scanning in Patients with Multiple Endocrine Neoplasia Type 1. <i>Journal of the American College of Surgeons</i> , 2015, 221, 509-517.	0.2	72
125	Molecular imaging and radionuclide therapy of pheochromocytoma and paraganglioma in the era of genomic characterization of disease subgroups. <i>Endocrine-Related Cancer</i> , 2019, 26, R627-R652.	1.6	72
126	Genomic Landscape of Pheochromocytoma and Paraganglioma. <i>Trends in Cancer</i> , 2018, 4, 6-9.	3.8	71

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127	CRH Deficiency Impairs but Does Not Block Pituitary-Adrenal Responses to Diverse Stressors. <i>Neuroendocrinology</i> , 2000, 71, 79-87.	1.2	68
128	New Advances in the Biochemical Diagnosis of Pheochromocytoma. <i>Annals of the New York Academy of Sciences</i> , 2002, 970, 29-40.	1.8	68
129	Correlation Between In Vivo ¹⁸ F-FDG PET and Immunohistochemical Markers of Glucose Uptake and Metabolism in Pheochromocytoma and Paraganglioma. <i>Journal of Nuclear Medicine</i> , 2014, 55, 1253-1259.	2.8	67
130	Differential expression of the regulated catecholamine secretory pathway in different hereditary forms of pheochromocytoma. <i>American Journal of Physiology - Endocrinology and Metabolism</i> , 2008, 295, E1223-E1233.	1.8	66
131	Functional and Oncologic Outcomes of Partial Adrenalectomy for Pheochromocytoma in Patients With von Hippel-Lindau Syndrome After at Least 5 Years of Followup. <i>Journal of Urology</i> , 2010, 184, 1855-1859.	0.2	66
132	Histone deacetylase inhibitors increase glucocerebrosidase activity in Gaucher disease by modulation of molecular chaperones. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013, 110, 966-971.	3.3	66
133	New functional imaging modalities for chromaffin tumors, neuroblastomas and ganglioneuromas. <i>Trends in Endocrinology and Metabolism</i> , 2005, 16, 66-72.	3.1	65
134	Emergencies Caused by Pheochromocytoma, Neuroblastoma, or Ganglioneuroma. <i>Endocrinology and Metabolism Clinics of North America</i> , 2006, 35, 699-724.	1.2	65
135	Effects of Single or Repeated Immobilization on Release of Norepinephrine and Its Metabolites in the Central Nucleus of the Amygdala in Conscious Rats. <i>Neuroendocrinology</i> , 1993, 57, 626-633.	1.2	64
136	Interrelations between Sympathoadrenal System and Hypothalamo-Pituitary-Adrenocortical/Thyroid Systems in Rats Exposed to Cold Stress. <i>Journal of Neuroendocrinology</i> , 1996, 8, 533-541.	1.2	64
137	INCREASED PLASMA NOREPINEPHRINE CONCENTRATION IN CATS WITH INTERSTITIAL CYSTITIS. <i>Journal of Urology</i> , 2001, 165, 2051-2054.	0.2	64
138	Localization of Medullary Thyroid Carcinoma Metastasis in a Multiple Endocrine Neoplasia Type 2A Patient by 6-[¹⁸ F]-Fluorodopamine Positron Emission Tomography. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2003, 88, 637-641.	1.8	64
139	A clinical overview of pheochromocytomas/paragangliomas and carcinoid tumors. <i>Nuclear Medicine and Biology</i> , 2008, 35, S27-S34.	0.3	64
140	Superiority of ⁶⁸ Ga-DOTATATE over ¹⁸ F-FDG and anatomic imaging in the detection of succinate dehydrogenase mutation (SDHx)-related pheochromocytoma and paraganglioma in the pediatric population. <i>European Journal of Nuclear Medicine and Molecular Imaging</i> , 2018, 45, 787-797.	3.3	64
141	Vorinostat suppresses hypoxia signaling by modulating nuclear translocation of hypoxia inducible factor 1 alpha. <i>Oncotarget</i> , 2017, 8, 56110-56125.	0.8	64
142	Succinate-to-Fumarate Ratio as a New Metabolic Marker to Detect the Presence of SDHB/D-related Paraganglioma: Initial Experimental and Ex Vivo Findings. <i>Endocrinology</i> , 2014, 155, 27-32.	1.4	63
143	SDHB-related pheochromocytoma and paraganglioma penetrance and genotype-phenotype correlations. <i>Journal of Cancer Research and Clinical Oncology</i> , 2017, 143, 1421-1435.	1.2	63
144	PheoSeq. <i>Journal of Molecular Diagnostics</i> , 2017, 19, 575-588.	1.2	63

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145	Stress-Induced Norepinephrine Release in the Paraventricular Nucleus of Rats with Brainstem Hemisections: A Microdialysis Study. <i>Neuroendocrinology</i> , 1993, 58, 196-201.	1.2	62
146	Characterization of an animal model of aggressive metastatic pheochromocytoma linked to a specific gene signature. <i>Clinical and Experimental Metastasis</i> , 2009, 26, 239-250.	1.7	62
147	Novel insights into the polycythemiaâ€“paragangliomaâ€“somatostatinoma syndrome. <i>Endocrine-Related Cancer</i> , 2016, 23, 899-908.	1.6	62
148	Prognostic Utility of Total 68Ga-DOTATATE-Avid Tumor Volume in Patients With Neuroendocrine Tumors. <i>Gastroenterology</i> , 2018, 154, 998-1008.e1.	0.6	62
149	Current Role of Metaiodobenzylguanidine in the Diagnosis of Pheochromocytoma and Medullary Thyroid Cancer. <i>Seminars in Nuclear Medicine</i> , 2011, 41, 364-368.	2.5	61
150	An N-terminal truncated carboxypeptidase E splice isoform induces tumor growth and is a biomarker for predicting future metastasis in human cancers. <i>Journal of Clinical Investigation</i> , 2011, 121, 880-892.	3.9	61
151	Diagnostic Localization of Pheochromocytoma. <i>Annals of the New York Academy of Sciences</i> , 2002, 970, 170-176.	1.8	60
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