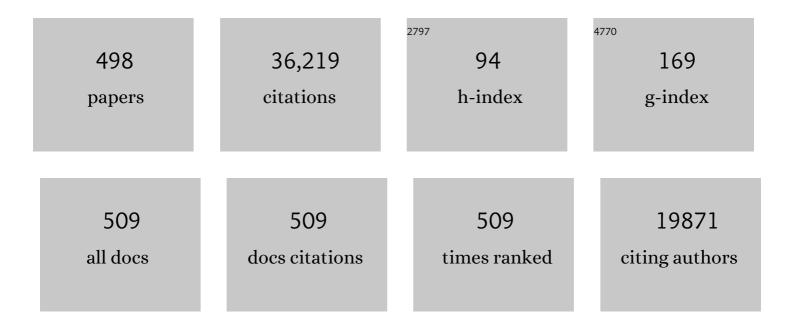
List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Pheochromocytoma and Paraganglioma: An Endocrine Society Clinical Practice Guideline. Journal of Clinical Endocrinology and Metabolism, 2014, 99, 1915-1942.	1.8	2,031
2	Phaeochromocytoma. Lancet, The, 2005, 366, 665-675.	6.3	1,462
3	Biochemical Diagnosis of Pheochromocytoma. JAMA - Journal of the American Medical Association, 2002, 287, 1427-34.	3.8	994
4	Irisin and FGF21 Are Cold-Induced Endocrine Activators of Brown Fat Function in Humans. Cell Metabolism, 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. Journal of Clinical Endocrinology and Metabolism, 2005, 90, 4955-4962.	1.8	588
6	Pheochromocytoma: recommendations for clinical practice from the First International Symposium. Nature Clinical Practice Endocrinology and Metabolism, 2007, 3, 92-102.	2.9	581
7	Comprehensive Molecular Characterization of Pheochromocytoma and Paraganglioma. Cancer Cell, 2017, 31, 181-193.	7.7	532
8	Recent Advances in Genetics, Diagnosis, Localization, and Treatment of Pheochromocytoma. Annals of Internal Medicine, 2001, 134, 315.	2.0	512
9	Preoperative Management of the Pheochromocytoma Patient. Journal of Clinical Endocrinology and Metabolism, 2007, 92, 4069-4079.	1.8	497
10	The North American Neuroendocrine Tumor Society Consensus Guideline for the Diagnosis and Management of Neuroendocrine Tumors. Pancreas, 2010, 39, 775-783.	0.5	493
11	An immunohistochemical procedure to detect patients with paraganglioma and phaeochromocytoma with germline SDHB, SDHC, or SDHD gene mutations: a retrospective and prospective analysis. Lancet Oncology, The, 2009, 10, 764-771.	5.1	477
12	Biochemical Diagnosis of Pheochromocytoma: How to Distinguish True- from False-Positive Test Results. Journal of Clinical Endocrinology and Metabolism, 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. European Journal of Human Genetics, 2008, 16, 79-88.	1.4	446
14	Mapping of human brown adipose tissue in lean and obese young men. Proceedings of the National Academy of Sciences of the United States of America, 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. Journal of Clinical Endocrinology and Metabolism, 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. Frontiers in Neuroendocrinology, 1995, 16, 89-150.	2.5	348
17	Somatic <i>HIF2A</i> Gain-of-Function Mutations in Paraganglioma with Polycythemia. New England Journal of Medicine, 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. Journal of Clinical Oncology, 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. European Journal of Cancer, 2012, 48, 1739-1749.	1.3	304
20	Malignant pheochromocytoma: current status and initiatives for future progress. Endocrine-Related Cancer, 2004, 11, 423-436.	1.6	299
21	High Frequency ofSDHBGermline Mutations in Patients with Malignant Catecholamine-Producing Paragangliomas: Implications for Genetic Testing. Journal of Clinical Endocrinology and Metabolism, 2006, 91, 4505-4509.	1.8	299
22	Current Approaches and Recommended Algorithm for the Diagnostic Localization of Pheochromocytoma. Journal of Clinical Endocrinology and Metabolism, 2004, 89, 479-491.	1.8	296
23	Molecular Subtypes of <i>KIT/PDGFRA</i> Wild-Type Gastrointestinal Stromal Tumors. JAMA Oncology, 2016, 2, 922.	3.4	291
24	Succinate Dehydrogenase Mutation Underlies Clobal Epigenomic Divergence in Gastrointestinal Stromal Tumor. Cancer Discovery, 2013, 3, 648-657.	7.7	288
25	Prospective Study of <sup>68</sup> Ga-DOTATATE Positron Emission Tomography/Computed Tomography for Detecting Gastro-Entero-Pancreatic Neuroendocrine Tumors and Unknown Primary Sites. Journal of Clinical Oncology, 2016, 34, 588-596.	0.8	287
26	Measurements of Plasma Methoxytyramine, Normetanephrine, and Metanephrine as Discriminators of Different Hereditary Forms of Pheochromocytoma. Clinical Chemistry, 2011, 57, 411-420.	1.5	282
27	<i>MAX</i> Mutations Cause Hereditary and Sporadic Pheochromocytoma and Paraganglioma. Clinical Cancer Research, 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. Journal of Clinical Endocrinology and Metabolism, 2001, 86, 1999-2008.	1.8	262
29	Clinical Presentations, Biochemical Phenotypes, and Genotype-Phenotype Correlations in Patients withSuccinate Dehydrogenase Subunit B-Associated Pheochromocytomas and Paragangliomas. Journal of Clinical Endocrinology and Metabolism, 2007, 92, 779-786.	1.8	262
30	New Perspectives on Pheochromocytoma and Paraganglioma: Toward a Molecular Classification. Endocrine Reviews, 2017, 38, 489-515.	8.9	241
31	Staging and Functional Characterization of Pheochromocytoma and Paraganglioma by 18F-Fluorodeoxyglucose (18F-FDG) Positron Emission Tomography. Journal of the National Cancer Institute, 2012, 104, 700-708.	3.0	240
32	Superiority of 6-[18F]-Fluorodopamine Positron Emission TomographyVersus[131I]-Metaiodobenzylguanidine Scintigraphy in the Localization of Metastatic Pheochromocytoma. Journal of Clinical Endocrinology and Metabolism, 2003, 88, 4083-4087.	1.8	237
33	Treatment of malignant pheochromocytoma/paraganglioma with cyclophosphamide, vincristine, and dacarbazine. Cancer, 2008, 113, 2020-2028.	2.0	234
34	EANM 2012 guidelines for radionuclide imaging of phaeochromocytoma and paraganglioma. European Journal of Nuclear Medicine and Molecular Imaging, 2012, 39, 1977-1995.	3.3	223
35	Superiority of [68Ga]-DOTATATE PET/CT to Other Functional Imaging Modalities in the Localization of <i>SDHB</i> -Associated Metastatic Pheochromocytoma and Paraganglioma. Clinical Cancer Research, 2015, 21, 3888-3895.	3.2	223
36	The optimal imaging of adrenal tumours: a comparison of different methods. Endocrine-Related Cancer, 2007, 14, 587-599.	1.6	218

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

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55	Paternal versus maternal transmission of a stimulatory G-protein $\hat{I}\pm$ subunit knockout produces opposite effects on energy metabolism. Journal of Clinical Investigation, 2000, 105, 615-623.	3.9	151
56	<sup>68</sup> Ga-DOTATATE PET/CT in the Localization of Head and Neck Paragangliomas Compared with Other Functional Imaging Modalities and CT/MRI. Journal of Nuclear Medicine, 2016, 57, 186-191.	2.8	148
57	Hypoxia-Inducible Factor Signaling in Pheochromocytoma: Turning the Rudder in the Right Direction. Journal of the National Cancer Institute, 2013, 105, 1270-1283.	3.0	146
58	Functional Imaging of Endocrine Tumors: Role of Positron Emission Tomography. Endocrine Reviews, 2004, 25, 568-580.	8.9	145
59	Endocrine Withdrawal Syndromes. Endocrine Reviews, 2003, 24, 523-538.	8.9	144
60	Loss of Meal-Induced Decrease in Plasma Ghrelin Levels in Patients with Anorexia Nervosa. Journal of Clinical Endocrinology and Metabolism, 2003, 88, 1678-1682.	1.8	142
61	Effects of Various Stressors on In Vivo Norepinephrine Release in the Hypothalamic Paraventricular Nucleus and on the Pituitary-Adrenocortical Axis. Annals of the New York Academy of Sciences, 1995, 771, 115-130.	1.8	141
62	Usefulness of <sup>123</sup> I-MIBG Scintigraphy in the Evaluation of Patients with Known or Suspected Primary or Metastatic Pheochromocytoma or Paraganglioma: Results from a Prospective Multicenter Trial. Journal of Nuclear Medicine, 2009, 50, 1448-1454.	2.8	139
63	PET/CT comparing 68Ga-DOTATATE and other radiopharmaceuticals and in comparison with CT/MRI for the localization of sporadic metastatic pheochromocytoma and paraganglioma. European Journal of Nuclear Medicine and Molecular Imaging, 2016, 43, 1784-1791.	3.3	138
64	Comparison of 6- <sup>18</sup> F-Fluorodopamine PET with <sup>123</sup> I-Metaiodobenzylguanidine and <sup>111</sup> In-Pentetreotide Scintigraphy in Localization of Nonmetastatic and Metastatic Pheochromocytoma. Journal of Nuclear Medicine, 2008, 49, 1613-1619.	2.8	137
65	Landscape of the mitochondrial Hsp90 metabolome in tumours. Nature Communications, 2013, 4, 2139.	5.8	135
66	Monoamine Oxidase B Prompts Mitochondrial and Cardiac Dysfunction in Pressure Overloaded Hearts. Antioxidants and Redox Signaling, 2014, 20, 267-280.	2.5	135
67	Succinate Dehydrogenase (SDH) D Subunit (SDHD) Inactivation in a Growth-Hormone-Producing Pituitary Tumor: A New Association for SDH?. Journal of Clinical Endocrinology and Metabolism, 2012, 97, E357-E366.	1.8	134
68	The Role of [18F]Fluorodeoxyglucose Positron Emission Tomography and [111In]-Diethylenetriaminepentaacetate-d-Phe-Pentetreotide Scintigraphy in the Localization of Ectopic Adrenocorticotropin-Secreting Tumors Causing Cushing's Syndrome. Journal of Clinical Endocrinology and Metabolism, 2004, 89, 2214-2221.	1.8	133
69	Functional Imaging of <i>&gt;SDHx </i> >Related Head and Neck Paragangliomas: Comparison of < sup > 18  F-Fluorodihydroxyphenylalanine, < sup > 18  F-Fluorodopamine, < sup > 18  F-Fluoro-2-Dec PET, < sup > 123  I-Metaiodobenzylguanidine Scintigraphy, and < sup > 111  In-Pentetreotide Scintigraphy. Journal of Clinical Endocrinology and Metabolism. 2011, 96, 2779-2785.	oxy- <scp> 1.8</scp>	d-Glu
70	New Syndrome of Paraganglioma and Somatostatinoma Associated With Polycythemia. Journal of Clinical Oncology, 2013, 31, 1690-1698.	0.8	129
71	Personalized Management of Pheochromocytoma and Paraganglioma. Endocrine Reviews, 2022, 43, 199-239.	8.9	127
72	Current Approaches and Recent Developments in the Management of Head and Neck Paragangliomas. Endocrine Reviews, 2014, 35, 795-819.	8.9	124

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

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91	Utility of Plasma Free Metanephrines for Detecting Childhood Pheochromocytoma. Journal of Clinical Endocrinology and Metabolism, 2002, 87, 1955-1960.	1.8	104
92	Update on pediatric pheochromocytoma. Pediatric Nephrology, 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. Journal of Nuclear Medicine, 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 phaeochromocytoma. Annals of Clinical Biochemistry, 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. European Journal of Nuclear Medicine and Molecular Imaging, 2016, 43, 1248-1257.	3.3	96
96	Drugs and Pheochromocytoma — Don't Be Fooled by Every Elevated Metanephrine. New England Journal of Medicine, 2011, 364, 2268-2270.	13.9	95
97	New Roles of Carboxypeptidase E in Endocrine and Neural Function and Cancer. Endocrine Reviews, 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. Brain Research, 1995, 688, 242-246.	1.1	92
99	Downregulation of metastasis suppressor genes in malignant pheochromocytoma. International Journal of Cancer, 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. Journal of Nuclear Medicine, 2007, 48, 1077-1083.	2.8	92
101	Metastatic Paraganglioma. Seminars in Oncology, 2010, 37, 627-637.	0.8	91
102	Age at Diagnosis of Pheochromocytoma Differs According to Catecholamine Phenotype and Tumor Location. Journal of Clinical Endocrinology and Metabolism, 2011, 96, 375-384.	1.8	90
103	Cardiac sympathetic denervation preceding motor signs in Parkinson disease. Clinical Autonomic Research, 2007, 17, 118-121.	1.4	88
104	Pheochromocytoma as an endocrine emergency. Reviews in Endocrine and Metabolic Disorders, 2003, 4, 121-128.	2.6	87
105	Novel HIF2A mutations disrupt oxygen sensing, leading to polycythemia, paragangliomas, and somatostatinomas. Blood, 2013, 121, 2563-2566.	0.6	87
106	Genotype–phenotype correlations in pheochromocytoma and paraganglioma: a systematic review and individual patient meta-analysis. Endocrine-Related Cancer, 2019, 26, 539-550.	1.6	87
107	Pheochromocytoma crisis induced by glucocorticoids: a report of four cases and review of the literature. European Journal of Endocrinology, 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. Nuclear Medicine Communications, 2006, 27, 31-36.	0.5	85

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109	Partial Adrenalectomy: Underused First Line Therapy for Small Adrenal Tumors. Journal of Urology, 2010, 184, 18-25.	0.2	85
110	15 YEARS OF PARAGANGLIOMA: Imaging and imaging-based treatment of pheochromocytoma and paraganglioma. Endocrine-Related Cancer, 2015, 22, T135-T145.	1.6	84
111	Radiofrequency Ablation: a Novel Approach for Treatment of Metastatic Pheochromocytoma. Journal of the National Cancer Institute, 2001, 93, 648-649.	3.0	83
112	Update of Pheochromocytoma Syndromes: Genetics, Biochemical Evaluation, and Imaging. Frontiers in Endocrinology, 2018, 9, 515.	1.5	82
113	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.	1.8	81
114	Minimal changes in environmental temperature result in a significant increase in energy expenditure and changes in the hormonal homeostasis in healthy adults. European Journal of Endocrinology, 2010, 163, 863-872.	1.9	80
115	International consensus on initial screening and follow-up of asymptomatic SDHx mutation carriers. Nature Reviews Endocrinology, 2021, 17, 435-444.	4.3	80
116	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. Endocrine, 2017, 56, 495-503.	1.1	79
117	Impaired Basal and Restraint-Induced Epinephrine Secretion in Corticotropin-Releasing Hormone- Deficient Mice <sup>1</sup> . Endocrinology, 2000, 141, 1142-1150.	1.4	78
118	The role of 6-[18F]fluorodopamine positron emission tomography in the localization of adrenal pheochromocytoma associated with von Hippel–Lindau syndrome. European Journal of Endocrinology, 2007, 156, 483-487.	1.9	78
119	SDH-related pheochromocytoma and paraganglioma. Best Practice and Research in Clinical Endocrinology and Metabolism, 2010, 24, 415-424.	2.2	78
120	Subclinical phaeochromocytoma. Best Practice and Research in Clinical Endocrinology and Metabolism, 2012, 26, 507-515.	2.2	76
121	Pheochromocytoma in von Hippel-Lindau Disease: Distinct Histopathologic Phenotype Compared to Pheochromocytoma in Multiple Endocrine Neoplasia Type 2. Endocrine Pathology, 2002, 13, 17-28.	5.2	74
122	Use of 6â€[ <sup>18</sup> F]â€fluorodopamine positron emission tomography (PET) as firstâ€line investigation for the diagnosis and localization of nonâ€metastatic and metastatic phaeochromocytoma (PHEO). Clinical Endocrinology, 2009, 71, 11-17.	1.2	74
123	Plasma metanephrines in renal failure. Kidney International, 2005, 67, 668-677.	2.6	73
124	Opposing effects of HIF1α and HIF2α on chromaffin cell phenotypic features and tumor cell proliferation: Insights from MYCâ€associated factor X. International Journal of Cancer, 2014, 135, 2054-2064.	2.3	72
125	Results of 68Gallium-DOTATATE PET/CT Scanning in Patients with Multiple Endocrine Neoplasia Type 1. Journal of the American College of Surgeons, 2015, 221, 509-517.	0.2	72
126	Molecular imaging and radionuclide therapy of pheochromocytoma and paraganglioma in the era of genomic characterization of disease subgroups. Endocrine-Related Cancer, 2019, 26, R627-R652.	1.6	72

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127	Genomic Landscape of Pheochromocytoma and Paraganglioma. Trends in Cancer, 2018, 4, 6-9.	3.8	71
128	CRH Deficiency Impairs but Does Not Block Pituitary-Adrenal Responses to Diverse Stressors. Neuroendocrinology, 2000, 71, 79-87.	1.2	68
129	New Advances in the Biochemical Diagnosis of Pheochromocytoma. Annals of the New York Academy of Sciences, 2002, 970, 29-40.	1.8	68
130	Correlation Between In Vivo <sup>18</sup> F-FDG PET and Immunohistochemical Markers of Glucose Uptake and Metabolism in Pheochromocytoma and Paraganglioma. Journal of Nuclear Medicine, 2014, 55, 1253-1259.	2.8	67
131	Differential expression of the regulated catecholamine secretory pathway in different hereditary forms of pheochromocytoma. American Journal of Physiology - Endocrinology and Metabolism, 2008, 295, E1223-E1233.	1.8	66
132	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.	0.2	66
133	Histone deacetylase inhibitors increase glucocerebrosidase activity in Gaucher disease by modulation of molecular chaperones. Proceedings of the National Academy of Sciences of the United States of America, 2013, 110, 966-971.	3.3	66
134	New functional imaging modalities for chromaffin tumors, neuroblastomas and ganglioneuromas. Trends in Endocrinology and Metabolism, 2005, 16, 66-72.	3.1	65
135	Emergencies Caused by Pheochromocytoma, Neuroblastoma, or Ganglioneuroma. Endocrinology and Metabolism Clinics of North America, 2006, 35, 699-724.	1.2	65
136	Effects of Single or Repeated Immobilization on Release of Norepinephrine and Its Metabolites in the Central Nucleus of the Amygdala in Conscious Rats. Neuroendocrinology, 1993, 57, 626-633.	1.2	64
137	Interrelations between Sympathoadrenal System andHypothalamo-Pituitary-Adrenocortical/Thyroid Systemsin Rats Exposed to Cold Stress. Journal of Neuroendocrinology, 1996, 8, 533-541.	1.2	64
138	INCREASED PLASMA NOREPINEPHRINE CONCENTRATION IN CATS WITH INTERSTITIAL CYSTITIS. Journal of Urology, 2001, 165, 2051-2054.	0.2	64
139	Localization of Medullary Thyroid Carcinoma Metastasis in a Multiple Endocrine Neoplasia Type 2A Patient by 6-[18F]-Fluorodopamine Positron Emission Tomography. Journal of Clinical Endocrinology and Metabolism, 2003, 88, 637-641.	1.8	64
140	A clinical overview of pheochromocytomas/paragangliomas and carcinoid tumors. Nuclear Medicine and Biology, 2008, 35, S27-S34.	0.3	64
141	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.	3.3	64
142	Vorinostat suppresses hypoxia signaling by modulating nuclear translocation of hypoxia inducible factor 1 alpha. Oncotarget, 2017, 8, 56110-56125.	0.8	64
143	Succinate-to-Fumarate Ratio as a New Metabolic Marker to Detect the Presence of SDHB/D-related Paraganglioma: Initial Experimental and Ex Vivo Findings. Endocrinology, 2014, 155, 27-32.	1.4	63
144	SDHB-related pheochromocytoma and paraganglioma penetrance and genotype–phenotype correlations. Journal of Cancer Research and Clinical Oncology, 2017, 143, 1421-1435.	1.2	63

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