

# Pheochromocytoma and Paraganglioma: An Endocrine

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Citation Report

#	ARTICLE	IF	CITATIONS
1	A rare cause of Cushing's syndrome: an ACTH-secreting pheochromocytoma. <i>BMJ Case Reports</i> , 2014, 2014, bcr2014205487-bcr2014205487.	0.2	16
3	SDHB-Associated Paraganglioma in a Pediatric Patient and Literature Review on Hereditary Pheochromocytoma-Paraganglioma Syndromes. <i>Case Reports in Endocrinology</i> , 2014, 2014, 1-5.	0.2	13
4	Laboratory Evaluation of Pheochromocytoma and Paraganglioma. <i>Clinical Chemistry</i> , 2014, 60, 1486-1499.	1.5	161
5	The genetics of endocrine neoplasia. <i>Current Problems in Cancer</i> , 2014, 38, 262-273.	1.0	0
7	MANAGEMENT OF ENDOCRINE DISEASE: Clinical management of paragangliomas. <i>European Journal of Endocrinology</i> , 2014, 171, R231-R243.	1.9	51
8	Current Approaches and Recent Developments in the Management of Head and Neck Paragangliomas. <i>Endocrine Reviews</i> , 2014, 35, 795-819.	8.9	124
9	Paraganglioma and cyanotic congenital heart disease: The role of tislular hipoxia. <i>EndocrinologÃa Y NutriciÃ³n (English Edition)</i> , 2015, 62, 413-414.	0.5	0
11	Hypertension presenting early in pregnancy. <i>Clinical Case Reports (discontinued)</i> , 2015, 3, 1056-1057.	0.2	1
13	2015 Meet-The-Professor: Endocrine Case Management. , 2015, , .		0
15	Minimally invasive parathyroidectomy in patients with previous thyroid surgery. <i>EndocrinologÃa Y NutriciÃ³n (English Edition)</i> , 2015, 62, 414-416.	0.5	0
16	Het feochromocytoom. <i>Bijblijven (Amsterdam, Netherlands)</i> , 2015, 31, 290-298.	0.0	0
17	Urinary clonidine suppression testing for the diagnosis of pheochromocytoma. <i>Journal of Hypertension</i> , 2015, 33, 2286-2293.	0.3	3
18	Adrenal Pheochromocytoma Incidentally Discovered in a Patient With Parkinsonism. <i>Medicine (United Tj ETQq0 0 0 rgBT /Overlock 10 T</i>	0.4	1
19	Supine or sitting plasma metanephrine screening? A unifying solution for patients and doctors. <i>Clinical Endocrinology</i> , 2015, 82, 776-777.	1.2	4
20	Pheochromocytoma diagnosed pathologically with previous negative serum markers. <i>Journal of Surgical Oncology</i> , 2015, 112, 492-495.	0.8	18
21	Laparoscopic transperitoneal resection of left paraÃeortic retroperitoneal paraganglioma. <i>Surgical Practice</i> , 2015, 19, 189-190.	0.1	0
22	Metastatic pheochromocytoma in a 23ÃeyearÃeold woman with an unclassified variant in the von Hippel Lindau disease gene: how can the pathogenicity of this variant be determined?. <i>Clinical Endocrinology</i> , 2015, 83, 15-19.	1.2	3
23	Risk of catecholaminergic crisis following glucocorticoid administration in patients with an adrenal mass: a literature review. <i>Clinical Endocrinology</i> , 2015, 83, 622-628.	1.2	18

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24	Recurrent pheochromocytoma along the laparoscopic portal sites. <i>Internal Medicine Journal</i> , 2015, 45, 359-361.	0.5	0
25	Clinical course and prognostic factors in patients with malignant pheochromocytoma and paraganglioma: A single institution experience. <i>Journal of Surgical Oncology</i> , 2015, 112, 815-821.	0.8	29
26	Robotic lateral transabdominal adrenalectomy. <i>Journal of Surgical Oncology</i> , 2015, 112, 305-309.	0.8	13
27	Catecholamine Crisis Precipitated by Intra-Articular Glucocorticoid Administration in a Patient with Paraganglioma. <i>AACE Clinical Case Reports</i> , 2015, 1, e265-e268.	0.4	0
28	Endocrine hypertension: An overview on the current etiopathogenesis and management options. <i>World Journal of Hypertension</i> , 2015, 5, 14.	0.8	27
29	Incidentally Discovered Aldosterone and Cortisol Cosecreting Adrenal Cortical Adenoma. <i>The Ewha Medical Journal</i> , 2015, 38, 129.	0.1	0
30	Pheochromocytomas and Paragangliomas: Clinical and Genetic Approaches. <i>Frontiers in Endocrinology</i> , 2015, 6, 126.	1.5	18
31	Ipertensione arteriosa: il ruolo dell'€™internista. <i>Italian Journal of Medicine</i> , 2015, 3, 1.	0.2	0
32	A large mesenteric paraganglioma with lymphovascular invasion. <i>BMJ Case Reports</i> , 2015, 2015, bcr2015209601-bcr2015209601.	0.2	3
33	Next-generation sequencing for the diagnosis of hereditary pheochromocytoma and paraganglioma syndromes. <i>Current Opinion in Endocrinology, Diabetes and Obesity</i> , 2015, 22, 169-179.	1.2	35
34	Endocrine Hypertension and Chronic Kidney Disease. , 2015, , 185-231.		0
35	Adrenal Incidentalomas. <i>Endocrinology and Metabolism Clinics of North America</i> , 2015, 44, 335-354.	1.2	23
36	Lack of utility of SDHB mutation testing in adrenergic metastatic pheochromocytoma. <i>European Journal of Endocrinology</i> , 2015, 172, 89-95.	1.9	17
37	Spatiotemporal Heterogeneity Characterizes the Genetic Landscape of Pheochromocytoma and Defines Early Events in Tumorigenesis. <i>Clinical Cancer Research</i> , 2015, 21, 4451-4460.	3.2	25
38	15 YEARS OF PARAGANGLIOMA: Pathology of pheochromocytoma and paraganglioma. <i>Endocrine-Related Cancer</i> , 2015, 22, T123-T133.	1.6	54
39	Strategic Chemotherapy for Pheochromocytoma?. <i>Endocrinology</i> , 2015, 156, 3880-3881.	1.4	1
40	The Importance of Exclusion of Obstructive Sleep Apnea During Screening for Adrenal Adenoma and Diagnosis of Pheochromocytoma. <i>Journal of Investigative Medicine High Impact Case Reports</i> , 2015, 3, 232470961560706.	0.3	2
42	LABORATORIO DE HORMONAS: ASPECTOS PRÁCTICOS. <i>Revista Médica Clínica Las Condes</i> , 2015, 26, 776-787.0.2		1

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43	Usefulness of Somatostatin Receptor Scintigraphy ( <sup>99m</sup> Tc-[HYNIC, <sub>3</sub> ] Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 752 T with <sup>18</sup> F-FDG PET/CT in SDHx Gene-Related Pheochromocytomas and Paragangliomas Detected by Computed Tomography. <i>Neuroendocrinology</i> , 2015, 101, 321-330.	1.2	13
44	Adrenal disease: The GP's role. <i>InnovAiT</i> , 2015, 8, 83-88.	0.0	0
45	A review of the management of positive biochemical screening for pheochromocytoma and paraganglioma: a salutary tale. <i>International Journal of Clinical Practice</i> , 2015, 69, 802-809.	0.8	7
46	Per-operative Hemodynamic Instability in Normotensive Patients With Incidentally Discovered Pheochromocytomas. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2015, 100, 417-421.	1.8	61
47	Pheochromocytoma and Paraganglioma in Cyanotic Congenital Heart Disease. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2015, 100, 1325-1334.	1.8	77
48	Penetrance and Clinical Features of Pheochromocytoma in a Six-Generation Family Carrying a Germline TMEM127 Mutation. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2015, 100, E308-E318.	1.8	44
49	Diagnosing pheochromocytoma/paraganglioma in a patient presenting with critical illness: biochemistry versus imaging. <i>Clinical Endocrinology</i> , 2015, 83, 298-302.	1.2	35
50	Semiquantitative <sup>123</sup> I-Metaiodobenzylguanidine Scintigraphy to Distinguish Pheochromocytoma and Paraganglioma from Physiologic Adrenal Uptake and Its Correlation with Genotype-Dependent Expression of Catecholamine Transporters. <i>Journal of Nuclear Medicine</i> , 2015, 56, 839-846.	2.8	30
51	Resection of a large carotid paraganglioma in Carney-Stratakis syndrome: a multidisciplinary feat. <i>BMJ Case Reports</i> , 2015, 2015, bcr2014208271-bcr2014208271.	0.2	2
52	The Challenges of Arterial Hypertension. <i>Frontiers in Cardiovascular Medicine</i> , 2015, 2, 2.	1.1	2
53	Menopause. <i>Nature Reviews Disease Primers</i> , 2015, 1, 15004.	18.1	288
54	Screening in asymptomatic SDHx mutation carriers: added value of <sup>18</sup> F-FDG PET/CT at initial diagnosis and 1-year follow-up. <i>European Journal of Nuclear Medicine and Molecular Imaging</i> , 2015, 42, 868-876.	3.3	23
55	Management of metastatic pheochromocytoma and paraganglioma: use of iodine-131-meta-iodobenzylguanidine therapy in a tertiary referral centre. <i>QJM - Monthly Journal of the Association of Physicians</i> , 2015, 108, 361-368.	0.2	24
56	Supine or sitting? Economic considerations regarding patient position during plasma metanephrine analysis for the exclusion of chromaffin tumours. <i>Clinical Endocrinology</i> , 2015, 82, 462-463.	1.2	18
57	Malignant phenotype and two SDHD mutations in a family with paraganglioma syndrome type 1. <i>Genetical Research</i> , 2015, 97, e3.	0.3	4
58	Superiority of [ <sup>68</sup> Ga]-DOTATATE PET/CT to Other Functional Imaging Modalities in the Localization of SDHB-Associated Metastatic Pheochromocytoma and Paraganglioma. <i>Clinical Cancer Research</i> , 2015, 21, 3888-3895.	3.2	223
59	Current trends in partial adrenalectomy. <i>Current Opinion in Urology</i> , 2015, 25, 89-94.	0.9	19
60	Preoperative Metyrosine Improves Cardiovascular Outcomes for Patients Undergoing Surgery for Pheochromocytoma and Paraganglioma. <i>Annals of Surgical Oncology</i> , 2015, 22, 646-654.	0.7	42

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61	Metastatic sympathetic paraganglioma in a patient with loss of the SDHC gene. <i>Familial Cancer</i> , 2015, 14, 615-619.	0.9	9
62	Loss of succinate dehydrogenase activity results in dependency on pyruvate carboxylation for cellular anabolism. <i>Nature Communications</i> , 2015, 6, 8784.	5.8	169
63	15 YEARS OF PARAGANGLIOMA: Clinical manifestations of paraganglioma syndromes types 1â€“5. <i>Endocrine-Related Cancer</i> , 2015, 22, T91-T103.	1.6	167
64	Is the endocrine research pipeline broken? A systematic evaluation of the Endocrine Society clinical practice guidelines and trial registration. <i>BMC Medicine</i> , 2015, 13, 187.	2.3	19
65	Pheochromocytomas in Multiple Endocrine Neoplasia Type 2. <i>Recent Results in Cancer Research</i> , 2015, 204, 157-178.	1.8	10
66	Medullary Thyroid Carcinoma. <i>Recent Results in Cancer Research</i> , 2015, , .	1.8	4
67	Supine or sitting? the voice of the patient stakeholders: economics <i>vs</i> clinically and medically sound. <i>Clinical Endocrinology</i> , 2015, 82, 464-465.	1.2	10
68	Biochemical Testing After Pheochromocytoma Removal: How Early?. <i>Hormone and Metabolic Research</i> , 2015, 47, 633-636.	0.7	1
69	Komplikationen in der Nebennierenchirurgie. , 2015, , 131-146.		0
70	Failure of metyrosine therapy for preoperative management of pheochromocytoma: a case report. <i>Canadian Journal of Anaesthesia</i> , 2015, 62, 1303-1307.	0.7	11
71	Multi-omics analysis defines core genomic alterations in pheochromocytomas and paragangliomas. <i>Nature Communications</i> , 2015, 6, 6044.	5.8	153
73	Genetics of Apparently Sporadic Pheochromocytoma and Paraganglioma in a Chinese Population. <i>Hormone and Metabolic Research</i> , 2015, 47, 833-838.	0.7	11
74	Functional imaging for pheochromocytomaâ€“paraganglioma: a step closer to understanding its place in clinical practice. <i>Endocrine</i> , 2015, 50, 6-8.	1.1	9
75	Life-threatening events in patients with pheochromocytoma. <i>European Journal of Endocrinology</i> , 2015, 173, 757-764.	1.9	84
76	Structural and functional consequences of succinate dehydrogenase subunit B mutations. <i>Endocrine-Related Cancer</i> , 2015, 22, 387-397.	1.6	19
77	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
78	Gasless Single-Port RoboSurgeon Surgery in Urology. , 2015, , .		5
79	Recommendations for somatic and germline genetic testing of single pheochromocytoma and paraganglioma based on findings from a series of 329 patients. <i>Journal of Medical Genetics</i> , 2015, 52, 647-656.	1.5	102

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80	Schildklier carcinoom. Bijblijven (Amsterdam, Netherlands), 2015, 31, 238-249.	0.0	0
81	Paraganglioma asociado a cardiopatía congénita cianótica: papel de la hipoxia tisular. Endocrinología Y Nutrición: Organó De La Sociedad Española De Endocrinología Y Nutrición, 2015, 62, 413-414.	0.8	0
82	Rare Cause of Severe Hypertension in a Young Woman. Hypertension, 2015, 65, 21-24.	1.3	1
83	Paraganglioma and pheochromocytoma: from genetics to personalized medicine. Nature Reviews Endocrinology, 2015, 11, 101-111.	4.3	396
84	Chronic Kidney Disease and Hypertension. , 2015, , .		0
85	Biochemically Silent Pheochromocytoma Presenting with Hypertensive Crisis During Surgery. AACE Clinical Case Reports, 2016, 2, e333-e336.	0.4	3
86	Pheochromocytoma in Pregnancy. Journal of Women's Health Care, 2016, 5, .	0.2	2
87	Structured assessment and followup for patients with hereditary kidney tumour syndromes. Canadian Urological Association Journal, 2016, 10, 214.	0.3	12
89	Mediastinal paragangliomas related to SDHx gene mutations. Kardiochirurgia I Torakochirurgia Polska, 2016, 3, 276-282.	0.1	12
90	Endocrine Hypertension. , 2016, , 556-588.		8
91	Is Biochemical Assessment of Pheochromocytoma Necessary in Adrenal Incidentalomas with Magnetic Resonance Imaging Features not Suggestive of Pheochromocytoma?. Endocrine Practice, 2016, 22, 533-539.	1.1	0
92	Commentary. Clinical Chemistry, 2016, 62, 928-928.	1.5	0
93	±-Methyldopa Interference in Urinary Normetanephrine Measurement by LC-MS/MS?. journal of applied laboratory medicine, The, 2016, 1, 321-324.	0.6	1
94	Hipertensi3n secundaria a paraganglioma: presentaci3n de un caso y revisi3n de la literatura. Iatreia, 2016, 29, .	0.1	0
95	Disentangling of Malignancy from Benign Pheochromocytomas/Paragangliomas. PLoS ONE, 2016, 11, e0168413.	1.1	42
96	Pheochromocytoma Masked by Mutation in the TH Gene. Clinical Chemistry, 2016, 62, 924-928.	1.5	1
97	A Case of Asymptomatic Multiple Paragangliomas. Nihon Rinsho Geka Gakkai Zasshi (Journal of Japan) Tj ETQq0 0 0 rgBT /Overlock 10 T	0.6	0
98	Stress, catecholaminergic system and cancer. Stress, 2016, 19, 419-428.	0.8	83

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99	Robot-assisted laparoscopic resection of large retroperitoneal paraganglioma – initial experience from China. <i>International Journal of Medical Robotics and Computer Assisted Surgery</i> , 2016, 12, 686-693.	1.2	10
100	Utility of <sup>FDG</sup>-PET imaging in screening for succinate dehydrogenase B and D mutation-related lesions. <i>Clinical Endocrinology</i> , 2016, 85, 172-179.	1.2	13
101	Getting personal: Head and neck cancer management in the era of genomic medicine. <i>Head and Neck</i> , 2016, 38, E2250-8.	0.9	19
102	Pheochromocytoma and paraganglioma. <i>Current Opinion in Oncology</i> , 2016, 28, 5-10.	1.1	40
103	Lesson of the month 2: Catecholamine-induced cardiomyopathy – pitfalls in diagnosis and medical management. <i>Clinical Medicine</i> , 2016, 16, 201-203.	0.8	5
104	Management of adrenal incidentalomas: European Society of Endocrinology Clinical Practice Guideline in collaboration with the European Network for the Study of Adrenal Tumors. <i>European Journal of Endocrinology</i> , 2016, 175, G1-G34.	1.9	1,173
105	Resection of Pheochromocytoma in a Patient Requiring Coronary Artery Bypass Grafting: First Things First. <i>AACE Clinical Case Reports</i> , 2016, 2, e25-e29.	0.4	2
106	Somatostatin-secreting Pheochromocytoma Mimicking Insulin-dependent Diabetes Mellitus. <i>Internal Medicine</i> , 2016, 55, 2985-2991.	0.3	9
107	Metoclopramide unmasks potentially misleading contralateral suppression in patients undergoing adrenal vein sampling for primary aldosteronism. <i>Journal of Hypertension</i> , 2016, 34, 2258-2265.	0.3	17
108	Management Of Catecholamine-Secreting Tumors In Pregnancy: A Review. <i>Endocrine Practice</i> , 2016, 22, 357-370.	1.1	25
110	An old retrocardiac mass fortuitously reclassified as paraganglioma. <i>Annales D'Endocrinologie</i> , 2016, 77, 668-669.	0.6	0
111	Venlafaxine drug interaction in the diagnosis of pheochromocytoma. <i>Endocrinología Y Nutrición (English Edition)</i> , 2016, 63, 569-570.	0.5	0
112	Predictors of malignancy in patients with pheochromocytomas/paragangliomas: Asian Indian experience. <i>Endocrine Connections</i> , 2016, 5, 89-97.	0.8	15
113	Interacción farmacológica de venlafaxina en el diagnóstico de feocromocitoma. <i>Endocrinología Y Nutrición: Organo De La Sociedad Espanola De Endocrinología Y Nutricion</i> , 2016, 63, 569-570.	0.8	1
115	Pheochromocytoma-paraganglioma: Biochemical and genetic diagnosis. <i>Nefrologia</i> , 2016, 36, 481-488.	0.2	5
116	Paraganglioma of the Seminal Vesicle Case Report and Review of the Literature. <i>Journal of Endourology Case Reports</i> , 2016, 2, 227-231.	0.3	5
119	Low specificity of urinary 3-methoxytyramine in screening of dopamine-secreting pheochromocytomas and paragangliomas. <i>Clinical Biochemistry</i> , 2016, 49, 1205-1208.	0.8	9
120	European Society of Endocrinology Clinical Practice Guideline for long-term follow-up of patients operated on for a phaeochromocytoma or a paraganglioma. <i>European Journal of Endocrinology</i> , 2016, 174, G1-G10.	1.9	352

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121	Diagnosis and Management of Hereditary Pheochromocytoma and Paraganglioma. Recent Results in Cancer Research, 2016, 205, 105-124.	1.8	6
122	Adrenal Incidentalomas: Clinical Controversies and Modified Recommendations. American Journal of Roentgenology, 2016, 206, 1170-1178.	1.0	31
123	Novel SDHB and TMEM127 Mutations in Patients with Pheochromocytoma/Paraganglioma Syndrome. Pathology and Oncology Research, 2016, 22, 673-679.	0.9	13
124	MANAGEMENT OF ENDOCRINE DISEASE: Recurrence or new tumors after complete resection of pheochromocytomas and paragangliomas: a systematic review and meta-analysis. European Journal of Endocrinology, 2016, 175, R135-R145.	1.9	52
125	Attention Deficit Hyperactivity Disorder in Pediatric Patients with Pheochromocytoma and Paraganglioma. Hormone and Metabolic Research, 2016, 48, 509-513.	0.7	6
126	Feocromocitoma-paraganglioma: del diagn�stico bioqu�mico al gen�tico. Nefrologia, 2016, 36, 481-488.	0.2	14
127	A case report of malignant hypertension in a young woman. BMC Nephrology, 2016, 17, 65.	0.8	2
128	Advances in our understanding of the prognosis of adrenal incidentaloma. Expert Review of Endocrinology and Metabolism, 2016, 11, 529-541.	1.2	1
129	Feocromocitoma y paraganglioma. Medicine, 2016, 12, 795-801.	0.0	0
130	A call to action and a lifecourse strategy to address the global burden of raised blood pressure on current and future generations: the Lancet Commission on hypertension. Lancet, The, 2016, 388, 2665-2712.	6.3	670
131	Genetic predisposition to endocrine tumors: Diagnosis, surveillance and challenges in care. Seminars in Oncology, 2016, 43, 582-590.	0.8	28
133	Adrenal incidentalomas: A guide to assessment, treatment and follow-up. Maturitas, 2016, 92, 79-85.	1.0	16
134	Interest of systematic screening of pheochromocytoma in patients with neurofibromatosis type 1. European Journal of Endocrinology, 2016, 175, 335-344.	1.9	53
135	Resistant Hypertension. Practical Case Studies in Hypertension Management, 2016, , .	0.0	0
136	Precision medicine in pheochromocytoma and paraganglioma: current and future concepts. Journal of Internal Medicine, 2016, 280, 559-573.	2.7	49
137	Symptom-dependent cut-offs of urine metanephrines improve diagnostic accuracy for detecting pheochromocytomas in two separate cohorts, compared to symptom-independent cut-offs. Endocrine, 2016, 54, 206-216.	1.1	7
139	Germline mutations and genotype�phenotype correlation in Asian Indian patients with pheochromocytoma and paraganglioma. European Journal of Endocrinology, 2016, 175, 311-323.	1.9	27
140	Feocromocitoma: estudo retrospectivo multic�ntrico. Revista Portuguesa De Endocrinologia Diabetes E Metabolismo, 2016, 11, 156-162.	0.1	0



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141	A 6-Base Pair in Frame Germline Deletion in Exon 7 Of <i>RET</i> Leads to Increased RET Phosphorylation, ERK Activation, and MEN2A. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2016, 101, 1016-1022.	1.8	14
142	Phaeochromocytoma and Paraganglioma. <i>Advances in Experimental Medicine and Biology</i> , 2016, 956, 239-259.	0.8	49
143	Flushing in (neuro)endocrinology. <i>Reviews in Endocrine and Metabolic Disorders</i> , 2016, 17, 373-380.	2.6	52
144	The Author's Reply: inappropriate adrenoreceptor blockade prior to pheochromocytoma removal "A timely reappraisal". <i>Clinical Endocrinology</i> , 2016, 85, 990-991.	1.2	0
145	No influence of antihypertensive agents on plasma free metanephrines. <i>Clinical Biochemistry</i> , 2016, 49, 1368-1371.	0.8	10
147	Treatment of pheochromocytomas and paragangliomas: genetic approach?. <i>International Journal of Endocrine Oncology</i> , 2016, 3, 325-331.	0.4	0
148	Endocrine Hypertension: A Practical Approach. <i>Advances in Experimental Medicine and Biology</i> , 2016, 956, 215-237.	0.8	23
149	Durable response to lenvatinib in progressive, therapy-refractory, metastatic paraganglioma. <i>International Journal of Endocrine Oncology</i> , 2016, 3, 285-289.	0.4	7
150	Seasonal variations in plasma free metanephrine concentrations are not evident in the West of Ireland. <i>Clinical Chemistry and Laboratory Medicine</i> , 2016, 54, e289-e292.	1.4	4
151	Contemporary Perioperative and Anesthetic Management of Pheochromocytoma and Paraganglioma. <i>Advances in Anesthesia</i> , 2016, 34, 181-196.	0.5	2
153	Nuclear Imaging in Metastatic Paraganglioma. <i>Journal of Nuclear Medicine Technology</i> , 2016, 44, 251-252.	0.4	2
155	[PP.25.14] RECURRENCE OR NEW TUMOURS AFTER COMPLETE RESECTION OF PHAEOCHROMOCYTOMAS AND PARAGANGLIOMAS. <i>Journal of Hypertension</i> , 2016, 34, e269.	0.3	2
156	Are patients with hormonally functional pheochromocytoma and paraganglioma initially receiving a proper adrenoceptor blockade? A retrospective cohort study. <i>Clinical Endocrinology</i> , 2016, 85, 62-69.	1.2	14
157	National Society Of Genetic Counselors Natalie Weissberger Paul National Leadership Award Address: "Patients and Research: Paths to Personal and Professional Growth". <i>Journal of Genetic Counseling</i> , 2016, 25, 617-620.	0.9	1
158	Rapidly Growing Chest Wall Mass in a Case of Sporadic Metastatic Paraganglioma. <i>Clinical Nuclear Medicine</i> , 2016, 41, 399-400.	0.7	2
159	Evaluating the optimum rest period prior to blood collection for fractionated plasma free metanephrines analysis. <i>Practical Laboratory Medicine</i> , 2016, 5, 39-46.	0.6	5
160	Pheochromocytoma in Urologic Practice. <i>European Urology Focus</i> , 2016, 1, 231-240.	1.6	7
161	Catecholamines Facilitate Fuel Expenditure and Protect Against Obesity via a Novel Network of the Gut-Brain Axis in Transcription Factor Skn-1 -deficient Mice. <i>EBioMedicine</i> , 2016, 8, 60-71.	2.7	24

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162	Failed pneumoperitoneum for laparoscopic surgery following autologous Deep Inferior Epigastric Perforator (DIEP) flap breast reconstruction: a case report. <i>BMC Surgery</i> , 2016, 16, 28.	0.6	2
163	Radiopharmaceuticals in paraganglioma imaging: too many members on board?. <i>European Journal of Nuclear Medicine and Molecular Imaging</i> , 2016, 43, 391-393.	3.3	6
165	Diagnosis of pheochromocytoma in a hemodialysis patient through measurement of plasma catecholamines. <i>Hemodialysis International</i> , 2016, 20, E6-9.	0.4	2
166	Rethinking pheochromocytomas and paragangliomas from a genomic perspective. <i>Oncogene</i> , 2016, 35, 1080-1089.	2.6	50
167	Complex MAX Rearrangement in a Family With Malignant Pheochromocytoma, Renal Oncocytoma, and Erythrocytosis. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2016, 101, 453-460.	1.8	47
168	Dopamine concentration in blood platelets is elevated in patients with head and neck paragangliomas. <i>Clinical Chemistry and Laboratory Medicine</i> , 2016, 54, 1395-401.	1.4	9
169	US-guided Biopsy of Neck Lesions: The Head and Neck Neuroradiologist's Perspective. <i>Radiographics</i> , 2016, 36, 226-243.	1.4	15
170	A simple and rapid analytical method based on solid-phase extraction and liquid chromatography-tandem mass spectrometry for the simultaneous determination of free catecholamines and metanephrines in urine and its application to routine clinical analysis. <i>Clinical Biochemistry</i> , 2016, 49, 573-579.	0.8	36
171	MANAGEMENT OF ENDOCRINE DISEASE: Outcome of adrenal sparing surgery in heritable pheochromocytoma. <i>European Journal of Endocrinology</i> , 2016, 174, R9-R18.	1.9	54
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173	PET/CT comparing 68Ga-DOTATATE and other radiopharmaceuticals and in comparison with CT/MRI for the localization of sporadic metastatic pheochromocytoma and paraganglioma. <i>European Journal of Nuclear Medicine and Molecular Imaging</i> , 2016, 43, 1784-1791.	3.3	138
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362	Rodent models of pheochromocytoma, parallels in rodent and human tumorigenesis. <i>Cell and Tissue Research</i> , 2018, 372, 379-392.	1.5	16
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#	ARTICLE	IF	CITATIONS
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#	ARTICLE	IF	CITATIONS
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830	Pediatric adrenocortical tumor – review and management update. <i>Current Opinion in Endocrinology, Diabetes and Obesity</i> , 2020, 27, 177-186.	1.2	16
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1031	Phakomatoses and Endocrine Gland Tumors: Noteworthy and (Not so) Rare Associations. <i>Frontiers in Endocrinology</i> , 2021, 12, 678869.	1.5	3
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1074	Optimized procedures for testing plasma metanephrines in patients on hemodialysis. <i>Scientific Reports</i> , 2021, 11, 14706.	1.6	5
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1079	Tracheal Paraganglioma: A Case report and Review of the Pertinent Literature. <i>Internal Medicine</i> , 2021, 60, 2275-2283.	0.3	4
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1122	An Incidental Uptake of 18F-Choline in Paraganglioma. <i>Clinical Nuclear Medicine</i> , 2021, Publish Ahead of Print, .	0.7	1
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1138	Paraganglioma intraabdominal por mutaci3n de SDHB. A prop3sito de un caso. <i>Medicina Clinica Practica</i> , 2022, 5, 100273.	0.2	0
1139	Pheochromocytoma: Perioperative and Intraoperative Management. , 2022, , 143-154.		0
1140	Acute Adrenal Hypertensive Emergencies: Pheochromocytoma, Cushing's, Hyperaldosteronism. , 2022, , 127-142.		0
1141	Head and Neck Paragangliomas: Patterns of Otolaryngology Referrals for Genetic Testing Over 2 Decades. <i>OTO Open</i> , 2021, 5, 2473974X21995453.	0.6	2
1142	Targeting Loss of Heterozygosity: A Novel Paradigm for Cancer Therapy. <i>Pharmaceuticals</i> , 2021, 14, 57.	1.7	27

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1144	The 3PAs syndrome and succinate dehydrogenase deficiency in pituitary tumors. , 2021, , 127-155.		0
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1535	Case Report: Giant Paraganglioma of the Skull Base With Two Somatic Mutations in SDHB and PTEN Genes. <i>Frontiers in Endocrinology</i> , 2022, 13, 857504.	1.5	2
1536	Influence of Receptor Polymorphisms on the Response to $\alpha$ -Adrenergic Receptor Blockers in Pheochromocytoma Patients. <i>Biomedicines</i> , 2022, 10, 896.	1.4	1
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1553	A Giant Adrenal Mass in a Super Obese Patient. <i>Cureus</i> , 2017, 9, e1572.	0.2	2
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1617	Anesthetic management of a giant paraganglioma resection: a case report. <i>BMC Anesthesiology</i> , 2022, 22, .	0.7	1
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1627	Pheochromocytoma and Paraganglioma. , 2023, , 127-131.		0
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1633	Evolution of perioperative management of catecholamine-producing tumors. <i>Russian Journal of Anesthesiology and Reanimatology /Anesteziologiya I Reanimatologiya</i> , 2022, , 85.	0.2	1
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1640	Noradrenergic Pheochromocytoma: A Case Report. <i>Cureus</i> , 2022, , .	0.2	0
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1646	A Study of Paraganglioma Cases With Non-European Ancestry. <i>Cureus</i> , 2022, , .	0.2	0
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1650	A Clinicopathologic and Molecular Analysis of Fumarate Hydratase-deficient Pheochromocytoma and Paraganglioma. <i>American Journal of Surgical Pathology</i> , 2023, 47, 25-36.	2.1	1
1651	Surgical and postsurgical management of abdominal paragangliomas and pheochromocytomas. <i>Actas Urológicas Españolas (English Edition)</i> , 2022, , .	0.2	0
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1663	Neuroendocrine Tumors: Therapy with 131I-MIBG. , 2022, , 1461-1480.		1
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1668	Case report: Significant liver atrophy due to giant cystic pheochromocytoma. <i>Frontiers in Oncology</i> , 0, 12, .	1.3	0
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1670	Pheochromocytoma manifestation associated with acute infectious disease. <i>MÃ¼narodnij EndokrinologÃ½nÃ½j Å½urnal</i> , 2022, 18, 315-317.	0.1	0
1671	Unusually large paraganglioma complicated with successive catecholamine crises: A case report and review of the literature. <i>Frontiers in Surgery</i> , 0, 9, .	0.6	2
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1674	Favorable outcome in advanced pheochromocytoma and paraganglioma after hypofractionated intensity modulated radiotherapy. <i>Journal of Endocrinological Investigation</i> , 2023, 46, 477-485.	1.8	1
1675	A retrospective study on the association between urine metanephrines and cardiometabolic risk in patients with nonfunctioning adrenal incidentaloma. <i>Scientific Reports</i> , 2022, 12, .	1.6	1
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1678	New Biology of Pheochromocytoma and Paraganglioma. <i>Endocrine Practice</i> , 2022, 28, 1253-1269.	1.1	8
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1681	Incidence and risk factors for myocardial injury after laparoscopic adrenalectomy for pheochromocytoma: A retrospective cohort study. <i>Frontiers in Oncology</i> , 0, 12, .	1.3	2
1682	The Clinical Characteristics of Pheochromocytomas and Paragangliomas with Negative Catecholamines. <i>Journal of Clinical Medicine</i> , 2022, 11, 5583.	1.0	1
1683	Case report: Incidentally discovered case of pheochromocytoma as a cause of long COVID-19 syndrome. <i>Frontiers in Endocrinology</i> , 0, 13, .	1.5	2
1684	Causes, Evaluation, and Treatment of Secondary and Resistant Hypertension. <i>Nephrology Self-assessment Program: NephSAP</i> , 2022, 21, 296-310.	3.0	0
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1689	Diagnostic features and therapeutic strategies for malignant paraganglioma in a patient: A case report. <i>World Journal of Clinical Cases</i> , 2022, 10, 9834-9844.	0.3	1
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1924	Case Report: A 65-year-old man with paraganglioma accompanied by elevated interleukin-6 levels and KIF1B single gene mutation. <i>Frontiers in Endocrinology</i> , 0, 14, .	1.5	1
1925	Endocrine causes of hypertension: literature review and practical approach. <i>Hypertension Research</i> , 2023, 46, 2679-2692.	1.5	1
1935	Secondary diabetes mellitus in pheochromocytomas and paragangliomas. <i>Endocrine</i> , 0, , .	1.1	0
1938	Editorial: A year in review: discussions in adrenal endocrinology. <i>Frontiers in Endocrinology</i> , 0, 14, .	1.5	0
1943	Secondary Hypertension: Pheochromocytoma and Paraganglioma. , 2024, , 187-197.		0
1944	The Psychosocial Impact of Familial Endocrine Cancer Syndromes (FECS) on the Patient and Caregiver. , 2023, , 1-35.		0
1945	Genetics, Biology, Clinical Presentation, Laboratory Diagnostics, and Management of Pediatric and Adolescent Pheochromocytoma and Paraganglioma. , 2023, , 107-125.		0
1946	Imaging Approach to Pediatric and Adolescent Familial Cancer Syndromes. , 2023, , 127-148.		0
1985	VHL: Trends and Insight into a Multi-Modality, Interdisciplinary Approach for Management of Central Nervous System Hemangioblastoma. <i>Acta Neurochirurgica Supplementum</i> , 2023, , 81-88.	0.5	0
1997	Testing for Secondary Hypertension and Difficult to Control Patients. , 2023, , 217-227.		0
2001	Rare Endocrine Disorders. , 2024, , 315-331.		0
2009	Malignome endokriner Organe. , 2024, , 993-1051.		0