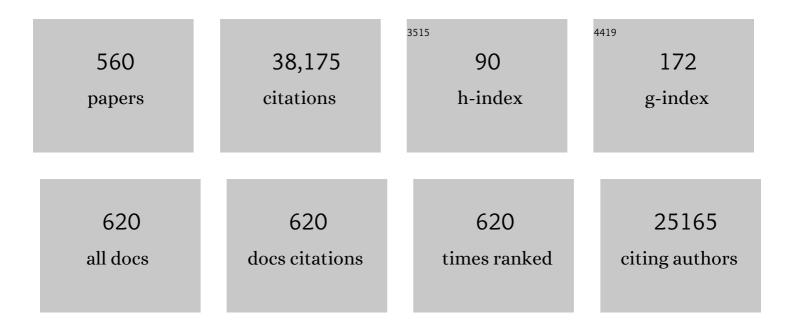
List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Pituitary corticotroph tumour with adrenocortical cells: A distinct clinicopathologic entity with unique morphology and methylation profile. Neuropathology and Applied Neurobiology, 2022, 48, .	1.8	0
2	Adrenal Gland. , 2022, , 461-486.		0
3	Letter to the Editor From Asa and Mete: "Hypophysitis, the Growing Spectrum of a Rare Pituitary Disease― Journal of Clinical Endocrinology and Metabolism, 2022, 107, e2649-e2649.	1.8	3
4	The Next Steps for Endocrine Pathology. Endocrine Pathology, 2022, 33, 228-230.	5.2	2
5	Overview of the 2022 WHO Classification of Thyroid Neoplasms. Endocrine Pathology, 2022, 33, 27-63.	5.2	388
6	Overview of the 2022 WHO Classification of Pituitary Tumors. Endocrine Pathology, 2022, 33, 6-26.	5.2	174
7	Overview of the 2022 WHO Classification of Neuroendocrine Neoplasms. Endocrine Pathology, 2022, 33, 115-154.	5.2	227
8	Pituitary carcinoma: reclassification and implications in the NET schema. Endocrine Oncology, 2022, 2, R14-R23.	0.1	3
9	Overview of the 2022 WHO Classification of Paragangliomas and Pheochromocytomas. Endocrine Pathology, 2022, 33, 90-114.	5.2	115
10	Ki-67 assessment of pancreatic neuroendocrine neoplasms: Systematic review and meta-analysis of manual vs. digital pathology scoring. Modern Pathology, 2022, 35, 712-720.	2.9	17
11	The Role of the Microbiome in Gastroentero-Pancreatic Neuroendocrine Neoplasms (GEP-NENs). Current Issues in Molecular Biology, 2022, 44, 2015-2028.	1.0	5
12	The diagnostic utility of BRAF VE1 mutation-specific immunohistochemistry in ameloblastoma. Modern Pathology, 2022, 35, 1570-1577.	2.9	6
13	Data set for the reporting of pheochromocytoma and paraganglioma: explanations and recommendations of the guidelines from the International Collaboration on Cancer Reporting. Human Pathology, 2021, 110, 83-97.	1.1	21
14	Perithyroidal Salivary Gland Acinic Cell Carcinoma: Morphological and Molecular Attributes of a Unique Lesion. Head and Neck Pathology, 2021, 15, 628-637.	1.3	1
15	Endoscopic Endonasal Pituitary Surgery For Nonfunctioning Pituitary Adenomas: Long-Term Outcomes and Management of Recurrent Tumors. World Neurosurgery, 2021, 146, e341-e350.	0.7	10
16	Cytokeratin profiles in pituitary neuroendocrine tumors. Human Pathology, 2021, 107, 87-95.	1.1	21
17	Inherited Follicular Epithelial-Derived Thyroid Carcinomas: From Molecular Biology to Histological Correlates. Endocrine Pathology, 2021, 32, 77-101.	5.2	21

Pathology of pituitary growth hormone excess., 2021, , 17-37.

#	Article	lF	CITATIONS
19	Molecular Pathology of Well-Differentiated Gastro-entero-pancreatic Neuroendocrine Tumors. Endocrine Pathology, 2021, 32, 169-191.	5.2	26
20	The Pangenomic Classification of Pituitary Neuroendocrine Tumors: Quality Histopathology is Required for Accurate Translational Research. Endocrine Pathology, 2021, 32, 415-417.	5.2	9
21	Significance of Crooke's Hyaline Change in Nontumorous Corticotrophs of Patients With Cushing Disease. Frontiers in Endocrinology, 2021, 12, 620005.	1.5	6
22	Significance of Alpha-inhibin Expression in Pheochromocytomas and Paragangliomas. American Journal of Surgical Pathology, 2021, 45, 1264-1273.	2.1	19
23	The North American Neuroendocrine Tumor Society Consensus Guidelines for Surveillance and Management of Metastatic and/or Unresectable Pheochromocytoma and Paraganglioma. Pancreas, 2021, 50, 469-493.	0.5	55
24	Challenges in the Diagnosis of Pituitary Neuroendocrine Tumors. Endocrine Pathology, 2021, 32, 222-227.	5.2	7
25	Oncocytic Change in Thyroid Pathology. Frontiers in Endocrinology, 2021, 12, 678119.	1.5	18
26	Pituitary neuroendocrine tumors: a model for neuroendocrine tumor classification. Modern Pathology, 2021, 34, 1634-1650.	2.9	44
27	Middle Ear "Adenomaâ€: a Neuroendocrine Tumor with Predominant L Cell Differentiation. Endocrine Pathology, 2021, 32, 433-441.	5.2	15
28	An Update on Pituitary Neuroendocrine Tumors Leading to Acromegaly and Gigantism. Journal of Clinical Medicine, 2021, 10, 2254.	1.0	15
29	Cribriform-Morular Thyroid Carcinoma Is a Distinct Thyroid Malignancy of Uncertain Cytogenesis. Endocrine Pathology, 2021, 32, 327-335.	5.2	25
30	Pendred Syndrome with C Cell Hyperplasia. Endocrine Pathology, 2021, 32, 427-428.	5.2	1
31	Follicular cells in pituitary neuroendocrine tumors. Human Pathology, 2021, 114, 1-8.	1.1	4
32	Nasopharyngeal neuroendocrine neoplasms: Systematic review of the literature and case presentation. Journal of Neuroendocrinology, 2021, 33, e13005.	1.2	4
33	Single-cell transcriptome and genome analysis: A much-needed tool for pituitary neuroendocrine tumor studies. Neuro-Oncology, 2021, 23, 1803-1804.	0.6	Ο
34	XB130 Deficiency Causes Congenital Hypothyroidism in Mice due to Disorganized Apical Membrane Structure and Function of Thyrocytes. Thyroid, 2021, 31, 1650-1661.	2.4	5
35	Hypothalamic hormone-producing tumors. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2021, 181, 67-74.	1.0	2
36	Genomics and Epigenomics of Pituitary Tumors: What Do Pathologists Need to Know?. Endocrine Pathology, 2021, 32, 3-16.	5.2	15

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37	Metastatic Neuroendocrine Neoplasms of Unknown Primary Site. , 2021, , 357-387.		13
38	Genetic and epigenetic characterization of posterior pituitary tumors. Acta Neuropathologica, 2021, 142, 1025-1043.	3.9	7
39	Neuroendocrine Neoplasms: Historical Background and Terminologies. , 2021, , 1-14.		2
40	Pancreatic Neuroendocrine Neoplasms. , 2021, , 245-261.		0
41	Paragangliomas and Pheochromocytomas. , 2021, , 263-285.		1
42	Pituitary Neuroendocrine Neoplasms. , 2021, , 55-83.		2
43	Thyroid Neuroendocrine Neoplasms. , 2021, , 119-136.		2
44	Parathyroid Neuroendocrine Neoplasms. , 2021, , 137-150.		0
45	Pathogenesis of multinodular goiter in elderly XB130 deficient mice: alteration of thyroperoxidase affinity with iodide and hydrogen peroxide. Thyroid, 2021, , .	2.4	2
46	Oncocytic Papillary Thyroid Carcinoma and Oncocytic Poorly Differentiated Thyroid Carcinoma: Clinical Features, Uptake, and Response to Radioactive lodine Therapy, and Outcome. Frontiers in Endocrinology, 2021, 12, 795184.	1.5	11
47	Pituitary neuroendocrine tumors (PitNETs): nomenclature evolution, not clinical revolution. Pituitary, 2020, 23, 322-325.	1.6	34
48	Syndrome of Inappropriate Antidiuresis in a Young Adult—Searching for the Causative Needle in the Proverbial Haystack. Kidney International Reports, 2020, 5, 231-234.	0.4	2
49	Characterization of pathological thyroid tissue using polarization-sensitive second harmonic generation microscopy. Laboratory Investigation, 2020, 100, 1280-1287.	1.7	19
50	Structure, Function, and Morphology in the Classification of Pituitary Neuroendocrine Tumors: the Importance of Routine Analysis of Pituitary Transcription Factors. Endocrine Pathology, 2020, 31, 330-336.	5.2	24
51	Immunohistochemical Analysis of the Metabolic Phenotype of Adrenal Cortical Carcinoma. Endocrine Pathology, 2020, 31, 231-238.	5.2	7
52	Images in Endocrine Pathology: Progressive Loss of Sustentacular Cells in a Case of Recurrent Jugulotympanic Paraganglioma over a Span of 5 years. Endocrine Pathology, 2020, 31, 310-314.	5.2	8
53	Images in Endocrine Pathology: High-Grade Intrathyroidal Parathyroid Carcinoma with Crooke's Hyalinization. Endocrine Pathology, 2020, 31, 190-194.	5.2	2
54	Centromeric cohesion failure invokes a conserved choreography of chromosomal mis-segregations in pancreatic neuroendocrine tumor. Genome Medicine, 2020, 12, 38.	3.6	9

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55	Thyroid Tumor Capsular Invasion: the Bottom Line or Much Ado About Nothing?. Endocrine Pathology, 2020, 31, 141-142.	5.2	4
56	Issues to Consider When Implementing Digital Pathology for Primary Diagnosis. Archives of Pathology and Laboratory Medicine, 2020, 144, 1297-1297.	1.2	4
57	Acidophil Stem Cell Tumor, Pituitary. Encyclopedia of Pathology, 2020, , 1-4.	0.0	Ο
58	Gonadotroph Tumor. Encyclopedia of Pathology, 2020, , 1-3.	0.0	0
59	Parathyroid Hyperplasia. Encyclopedia of Pathology, 2020, , 1-4.	0.0	0
60	Corticotroph Tumour. Encyclopedia of Pathology, 2020, , 1-6.	0.0	0
61	Pituitary Carcinoma. Encyclopedia of Pathology, 2020, , 1-3.	0.0	0
62	Null Cell Tumor. Encyclopedia of Pathology, 2020, , 1-3.	0.0	0
63	Mammosomatotroph Tumor. Encyclopedia of Pathology, 2020, , 1-3.	0.0	0
64	GATA2. Encyclopedia of Pathology, 2020, , 1-2.	0.0	0
65	Parathyroid Adenoma. Encyclopedia of Pathology, 2020, , 1-4.	0.0	0
66	C-Cell Hyperplasia. Encyclopedia of Pathology, 2020, , 1-4.	0.0	0
67	Lactotroph Tumor. Encyclopedia of Pathology, 2020, , 1-3.	0.0	0
68	Pituitary Neuroendocrine Tumor. Encyclopedia of Pathology, 2020, , 1-5.	0.0	0
69	Medullary Thyroid Carcinoma. Encyclopedia of Pathology, 2020, , 1-5.	0.0	0
70	Molecular profiling confirms historical immunohistochemistry in acromegaly. Endocrine-Related Cancer, 2020, 27, L1-L2.	1.6	0
71	Comprehensive characterization of a Canadian cohort of von Hippelâ€Lindau disease patients. Clinical Genetics, 2019, 96, 461-467.	1.0	16
72	VEGFRâ€2 is downregulated in sestamibiâ€negative parathyroid adenomas. Head and Neck, 2019, 41, 3564-3569	9.0.9	4

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73	A Systematic Review and Meta-Analysis of the Diagnostic Performance of BRAF V600E Immunohistochemistry in Thyroid Histopathology. Endocrine Pathology, 2019, 30, 201-218.	5.2	24
74	The Clinicopathological Spectrum of Parathyroid Carcinoma. Frontiers in Endocrinology, 2019, 10, 731.	1.5	25
75	Somatostatin Receptor Ligand Therapy—A Potential Therapy for Neurocytoma. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 2395-2402.	1.8	7
76	Diagnosis and Pathologic Characteristics of Medullary Thyroid Carcinoma—Review of Current Guidelines. Current Oncology, 2019, 26, 338-344.	0.9	65
77	Hypothalamic Endocrine Tumors: An Update. Journal of Clinical Medicine, 2019, 8, 1741.	1.0	15
78	An Institutional Experience of Tumor Progression to Pituitary Carcinoma in a 15-Year Cohort of 1055 Consecutive Pituitary Neuroendocrine Tumors. Endocrine Pathology, 2019, 30, 118-127.	5.2	43
79	The Current Histologic Classification of Thyroid Cancer. Endocrinology and Metabolism Clinics of North America, 2019, 48, 1-22.	1.2	66
80	Papillary Thyroid Cancers with Focal Tall Cell Change are as Aggressive as Tall Cell Variants and Should Not be Considered as Low-Risk Disease. Annals of Surgical Oncology, 2019, 26, 2533-2539.	0.7	18
81	Treatment Options for Pancreatic Neuroendocrine Tumors. Cancers, 2019, 11, 828.	1.7	55
82	A phase 2 trial of sunitinib in patients with progressive paraganglioma or pheochromocytoma: the SNIPP trial. British Journal of Cancer, 2019, 120, 1113-1119.	2.9	83
83	Interobserver Variability in the Histopathologic Assessment of Extrathyroidal Extension of Well Differentiated Thyroid Carcinoma Supports the New American Joint Committee on Cancer Eighth Edition Criteria for Tumor Staging. Thyroid, 2019, 29, 619-624.	2.4	22
84	Characterization of Pancreatic Cancer Tissue Using Multiphoton Excitation Fluorescence and Polarization-Sensitive Harmonic Generation Microscopy. Frontiers in Oncology, 2019, 9, 272.	1.3	32
85	Molecular Predictors of Clinical Behavior in Pituitary Adenohypophysial Tumors. Contemporary Endocrinology, 2019, , 155-172.	0.3	Ο
86	Hypothalamic Vasopressin-Producing Tumors. American Journal of Surgical Pathology, 2019, 43, 251-260.	2.1	24
87	The Clinicopathological Spectrum of Acromegaly. Journal of Clinical Medicine, 2019, 8, 1962.	1.0	42
88	Plurihormonal Pituitary Tumor of Pit-1 and SF-1 Lineages, with Synchronous Collision Corticotroph Tumor: a Possible Stem Cell Phenomenon. Endocrine Pathology, 2019, 30, 74-80.	5.2	25
89	GATA3 immunoreactivity expands the transcription factor profile of pituitary neuroendocrine tumors. Modern Pathology, 2019, 32, 484-489.	2.9	48
90	2020 Vision of Digital Pathology in Action. Journal of Pathology Informatics, 2019, 10, 27.	0.8	12

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91	Ki67 Quantitative Interpretation: Insights using Image Analysis. Journal of Pathology Informatics, 2019, 10, 8.	0.8	23
92	SUN-453 Absence of Crooke's Hyaline Changes May Predict Worse Outcomes in Patients with Cushing Disease. Journal of the Endocrine Society, 2019, 3, .	0.1	0
93	Immunohistochemical Biomarkers in Pituitary Pathology. Endocrine Pathology, 2018, 29, 130-136.	5.2	26
94	Noninvasive Follicular Thyroid Neoplasm with Papillaryâ€Like Nuclear Features (NIFTP): Trading Six for a Risky Half Dozen: Reply. World Journal of Surgery, 2018, 42, 2279-2279.	0.8	4
95	Epidemiology and biomarker profile of pituitary adenohypophysial tumors. Modern Pathology, 2018, 31, 900-909.	2.9	120
96	The evolving diagnosis of noninvasive follicular thyroid neoplasm with papillary-like nuclear features (NIFTP). Human Pathology, 2018, 74, 1-4.	1.1	45
97	Immunohistochemical Biomarkers of Adrenal Cortical Neoplasms. Endocrine Pathology, 2018, 29, 137-149.	5.2	45
98	Pancreatic Neuroendocrine Tumor Producing Insulin and Vasopressin. Endocrine Pathology, 2018, 29, 15-20.	5.2	9
99	Epigenetics of pituitary tumors: Pathogenetic and therapeutic implications. Molecular and Cellular Endocrinology, 2018, 469, 70-76.	1.6	27
100	The epigenetic landscape of differentiated thyroid cancer. Molecular and Cellular Endocrinology, 2018, 469, 3-10.	1.6	24
101	Diagnostic and Prognostic Biomarkers of Adrenal Cortical Carcinoma. American Journal of Surgical Pathology, 2018, 42, 201-213.	2.1	56
102	Clinical Safety of Renaming Encapsulated Follicular Variant of Papillary Thyroid Carcinoma: Is NIFTP Truly Benign?. World Journal of Surgery, 2018, 42, 321-326.	0.8	114
103	Integrated Pathology Informatics Enables High-Quality Personalized and Precision Medicine: Digital Pathology and Beyond. Archives of Pathology and Laboratory Medicine, 2018, 142, 369-382.	1.2	29
104	The retrotransposon gag domain containing protein Rgag4 is an Ikaros target in the pituitary. Molecular and Cellular Endocrinology, 2018, 461, 188-193.	1.6	4
105	What's new in pituitary pathology?. Histopathology, 2018, 72, 133-141.	1.6	24
106	Endocrine pathology: past, present and future. Pathology, 2018, 50, 111-118.	0.3	23
107	Intrathyroidal Parathyroid Carcinoma: An Atypical Thyroid Lesion. Frontiers in Endocrinology, 2018, 9, 641.	1.5	19

108 Pituitary Tumors; Diagnosis and Treatment. , 2018, , 257-257.

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109	An Unusual Salivary Gland Tumor Mimicking Papillary Thyroid Carcinoma: Mammary Analog Secretory Carcinoma. Frontiers in Endocrinology, 2018, 9, 555.	1.5	9
110	The Diagnosis and Clinical Significance of Paragangliomas in Unusual Locations. Journal of Clinical Medicine, 2018, 7, 280.	1.0	104
111	Synchronous Multiple Pituitary Neuroendocrine Tumors of Different Cell Lineages. Endocrine Pathology, 2018, 29, 332-338.	5.2	28
112	Immunohistochemical Biomarkers in Thyroid Pathology. Endocrine Pathology, 2018, 29, 91-112.	5.2	48
113	A common classification framework for neuroendocrine neoplasms: an International Agency for Research on Cancer (IARC) and World Health Organization (WHO) expert consensus proposal. Modern Pathology, 2018, 31, 1770-1786.	2.9	739
114	Liver Transplantation in a Young Patient with Severe and Refractory Carcinoid Syndrome. AACE Clinical Case Reports, 2018, 4, e289-e293.	0.4	0
115	Molecular Pathogenesis of Pituitary Tumors. , 2017, , 165-175.		Ο
116	Progressive epigenetic dysregulation in neuroendocrine tumour liver metastases. Endocrine-Related Cancer, 2017, 24, L21-L25.	1.6	37
117	Comprehensive Molecular Characterization of Pheochromocytoma and Paraganglioma. Cancer Cell, 2017, 31, 181-193.	7.7	532
118	Pituitary acromegaly: not one disease. Endocrine-Related Cancer, 2017, 24, C1-C4.	1.6	37
119	Pancreatic Struma with Papillary Thyroid Carcinoma: a Diagnostic Dilemma. Endocrine Pathology, 2017, 28, 91-94.	5.2	2
120	The evolution of differentiated thyroid cancer. Pathology, 2017, 49, 229-237.	0.3	20
121	From pituitary adenoma to pituitary neuroendocrine tumor (PitNET): an International Pituitary Pathology Club proposal. Endocrine-Related Cancer, 2017, 24, C5-C8.	1.6	262
122	The dangers of parathyroid biopsy. Journal of Otolaryngology - Head and Neck Surgery, 2017, 46, 4.	0.9	44
123	Xanthomatous Hypophysitis Is Associated with Ruptured Rathke's Cleft Cyst. Endocrine Pathology, 2017, 28, 83-90.	5.2	31
124	Pituitary Adenomas Presenting as Sinonasal or Nasopharyngeal Masses. American Journal of Surgical Pathology, 2017, 41, 525-534.	2.1	26
125	Pathologic Reporting of Tall-Cell Variant of Papillary Thyroid Cancer: Have We Reached a Consensus?. Thyroid, 2017, 27, 1498-1504.	2.4	32
126	Clinical Applications of Whole-slide Imaging in Anatomic Pathology. Advances in Anatomic Pathology, 2017, 24, 215-221.	2.4	23

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127	Malignant Ovarian Steroid Cell Tumor Causing Severe Hyperandrogenism: Case Report And Review Of The Literature. AACE Clinical Case Reports, 2017, 3, e269-e274.	0.4	1
128	Ikaros and its interacting partner CtBP target the metalloprotease ADAMTS10 to modulate pituitary cell function. Molecular and Cellular Endocrinology, 2017, 439, 126-132.	1.6	4
129	Template for Reporting Results of Biomarker Testing of Specimens From Patients With Thyroid Carcinoma. Archives of Pathology and Laboratory Medicine, 2017, 141, 559-563.	1.2	9
130	TFE3-Expressing Perivascular Epithelioid Cell Neoplasm (PEComa) of the Sella Turcica. Endocrine Pathology, 2017, 28, 22-26.	5.2	9
131	Hypothalamic Disease. , 2017, , 97-106.		0
132	FGFR4 polymorphic alleles modulate mitochondrial respiration: A novel target for somatostatin analog action in pituitary tumors. Oncotarget, 2017, 8, 3481-3494.	0.8	14
133	An International Ki67 Reproducibility Study in Adrenal Cortical Carcinoma. American Journal of Surgical Pathology, 2016, 40, 569-576.	2.1	75
134	Diagnosis and management of gastrointestinal neuroendocrine tumors: An evidence-based Canadian consensus. Cancer Treatment Reviews, 2016, 47, 32-45.	3.4	74
135	Nomenclature Revision for Encapsulated Follicular Variant of Papillary Thyroid Carcinoma. JAMA Oncology, 2016, 2, 1023.	3.4	1,192
136	Comprehensive Pan-Genomic Characterization of Adrenocortical Carcinoma. Cancer Cell, 2016, 29, 723-736.	7.7	482
137	Inter-Observer Variation in the Pathologic Identification of Extranodal Extension in Nodal Metastasis from Papillary Thyroid Carcinoma. Thyroid, 2016, 26, 816-819.	2.4	12
138	Establishment and Characterization of a Human Neuroendocrine Tumor Xenograft. Endocrine Pathology, 2016, 27, 97-103.	5.2	14
139	Cytology and Pathology: Pitfalls and Challenges. , 2016, , 33-46.		3
140	Synchronous Papillary Carcinoma of Thyroid and Lung. Endocrine Pathology, 2016, 27, 268-270.	5.2	1
141	The American Association of Endocrine Surgeons Guidelines for Definitive Management of Primary Hyperparathyroidism. JAMA Surgery, 2016, 151, 959.	2.2	840
142	Gonadotrope Tumors. Progress in Molecular Biology and Translational Science, 2016, 143, 187-210.	0.9	3
143	NG2 targets tumorigenic Rb inactivation in Pit1-lineage pituitary cells. Endocrine-Related Cancer, 2016, 23, 445-456.	1.6	8
144	Minichromosome maintenance protein 7 as prognostic marker of tumor aggressiveness in pituitary adenoma patients. European Journal of Endocrinology, 2016, 174, 307-314.	1.9	25

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145	Differential Clinicopathological Risk and Prognosis of Major Papillary Thyroid Cancer Variants. Journal of Clinical Endocrinology and Metabolism, 2016, 101, 264-274.	1.8	179
146	Inter-Observer Variation in the Pathologic Identification of Minimal Extrathyroidal Extension in Papillary Thyroid Carcinoma. Thyroid, 2016, 26, 512-517.	2.4	56
147	Aggressive Pituitary Tumors or Localized Pituitary Carcinomas: Defining Pituitary Tumors. Expert Review of Endocrinology and Metabolism, 2016, 11, 149-162.	1.2	42
148	Silent subtype 3 pituitary adenomas are not always silent and represent poorly differentiated monomorphous plurihormonal Pit-1 lineage adenomas. Modern Pathology, 2016, 29, 131-142.	2.9	114
149	Monomorphous Plurihormonal Pituitary Adenoma of Pit-1 Lineage in a Giant Adolescent with Central Hyperthyroidism. Endocrine Pathology, 2016, 27, 25-33.	5.2	26
150	Prognostic Impact of Novel Molecular Subtypes of Small Intestinal Neuroendocrine Tumor. Clinical Cancer Research, 2016, 22, 250-258.	3.2	149
151	High-throughput drug library screening identifies colchicine as a thyroid cancer inhibitor. Oncotarget, 2016, 7, 19948-19959.	0.8	15
152	Clinical implications of accurate subtyping of pituitary adenomas: perspectives from the treating physician. Turk Patoloji Dergisi, 2015, 31 Suppl 1, 4-17.	0.1	14
153	Pancreatic Neuroendocrine Tumors Producing GHRH, GH, Ghrelin, PTH, or PTHrP. , 2015, , 125-139.		3
154	Tumor tissue characterization using polarization-sensitive second harmonic generation microscopy. , 2015, , .		8
155	Modeling complexity in pathologist workload measurement: the Automatable Activity-Based Approach to Complexity Unit Scoring (AABACUS). Modern Pathology, 2015, 28, 324-339.	2.9	23
156	Revised American Thyroid Association Guidelines for the Management of Medullary Thyroid Carcinoma. Thyroid, 2015, 25, 567-610.	2.4	1,738
157	Familial pheochromocytoma and renal cell carcinoma syndrome: TMEM127 as a novel candidate gene for the association. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2015, 466, 727-732.	1.4	38
158	When Thyroid Carcinoma Goes Bad: A Morphological and Molecular Analysis. Head and Neck Pathology, 2015, 9, 16-23.	1.3	50
159	Ultrastructural features of collagen in thyroid carcinoma tissue observed by polarization second harmonic generation microscopy. Biomedical Optics Express, 2015, 6, 3475.	1.5	56
160	The Complementary Role of Transcription Factors in the Accurate Diagnosis of Clinically Nonfunctioning Pituitary Adenomas. Endocrine Pathology, 2015, 26, 349-355.	5.2	167
161	In Reply. Archives of Pathology and Laboratory Medicine, 2015, 139, 967-968.	1.2	0
162	Implications of the TCGA Genomic Characterization of Papillary Thyroid Carcinoma for Thyroid Pathology: Does Follicular Variant Papillary Thyroid Carcinoma Exist?. Thyroid, 2015, 25, 1-2.	2.4	54

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163	Targeted expression of a human pituitary tumor–derived isoform of FGF receptor-4 recapitulates pituitary tumorigenesis. Journal of Clinical Investigation, 2015, 125, 3303-3303.	3.9	1
164	NCIC CTG IND.206: A phase II umbrella trial of sunitinib (S) or temsirolimus (T) in advanced rare cancers Journal of Clinical Oncology, 2015, 33, 2594-2594.	0.8	4
165	An unusual case of an ACTH-secreting macroadenoma with a germline variant in the aryl hydrocarbon receptor-interacting protein (AIP) gene. Endocrinology, Diabetes and Metabolism Case Reports, 2015, 2015, 140105.	0.2	9
166	Malignant Pheochromocytoma Secreting Vasoactive Intestinal Peptide and Response to Sunitinib: A Case Report and Literature Review. Endocrine Practice, 2014, 20, e145-e150.	1.1	15
167	TTF-1 Expressing Sellar Neoplasm with Ependymal Rosettes and Oncocytic Change: Mixed Ependymal and Oncocytic Variant Pituicytoma. Endocrine Pathology, 2014, 25, 436-438.	5.2	21
168	The PI3K/AKT/mTOR pathway in the pathophysiology and treatment of pituitary adenomas. Endocrine-Related Cancer, 2014, 21, R331-R344.	1.6	61
169	Metastatic Thyroid Carcinoma to the Gastric Body. Journal of Clinical Endocrinology and Metabolism, 2014, 99, 3958-3959.	1.8	9
170	Non-pheochromocytoma (PCC)/paraganglioma (PGL) tumors in patients with succinate dehydrogenase-related PCC–PGL syndromes: a clinicopathological and molecular analysis. European Journal of Endocrinology, 2014, 170, 1-12.	1.9	219
171	Protocol for the Examination of Specimens From Patients With Pheochromocytomas and Extra-Adrenal Paragangliomas. Archives of Pathology and Laboratory Medicine, 2014, 138, 182-188.	1.2	52
172	Integrated Genomic Characterization of Papillary Thyroid Carcinoma. Cell, 2014, 159, 676-690.	13.5	2,318
173	Multiple Endocrine Neoplasia Type 1. , 2014, 19, 85-89.		6
174	Editorial: The Birth of Endocrine Pathology. Endocrine Pathology, 2014, 25, 2-2.	5.2	1
175	Functional Cardiac Paraganglioma Associated with a Rare SDHC Mutation. Endocrine Pathology, 2014, 25, 315-320.	5.2	16
176	A History of Pituitary Pathology. Endocrine Pathology, 2014, 25, 6-11.	5.2	3
177	Genomic Approaches to Problems in Pituitary Neoplasia. Endocrine Pathology, 2014, 25, 209-213.	5.2	10
178	FGFR4 Polymorphic Variants Modulate Phenotypic Features of Cushing Disease. Molecular Endocrinology, 2014, 28, 525-533.	3.7	18
179	Tyrosine kinase receptors as molecular targets in pheochromocytomas and paragangliomas. Modern Pathology, 2014, 27, 1050-1062.	2.9	17
180	Genetics and Epigenetics of Endocrine Neoplasia. Molecular and Cellular Endocrinology, 2014, 386, 1.	1.6	1

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181	A single-arm, phase II, multicenter trial of sunitinib (SU) in locally advanced or metastatic pheochromocytoma/paraganglioma (PC/PG): Updated interim results Journal of Clinical Oncology, 2014, 32, e15621-e15621.	0.8	0
182	Parathyroid cancer: Outcome analysis of 16 patients treated at the princess margaret hospital. Head and Neck, 2013, 35, 35-39.	0.9	49
183	Silent Corticotroph Adenoma with Adrenal Cortical Choristoma: a Rare but Distinct Morphological Entity. Endocrine Pathology, 2013, 24, 162-166.	5.2	7
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