List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Nomenclature Revision for Encapsulated Follicular Variant of Papillary Thyroid Carcinoma. JAMA Oncology, 2016, 2, 1023.	7.1	1,192
2	Comprehensive Molecular Characterization of Pheochromocytoma and Paraganglioma. Cancer Cell, 2017, 31, 181-193.	16.8	532
3	Comprehensive Pan-Genomic Characterization of Adrenocortical Carcinoma. Cancer Cell, 2016, 29, 723-736.	16.8	482
4	Overview of the 2022 WHO Classification of Thyroid Neoplasms. Endocrine Pathology, 2022, 33, 27-63.	9.0	388
5	Overview of the 2017 WHO Classification of Pituitary Tumors. Endocrine Pathology, 2017, 28, 228-243.	9.0	319
6	From pituitary adenoma to pituitary neuroendocrine tumor (PitNET): an International Pituitary Pathology Club proposal. Endocrine-Related Cancer, 2017, 24, C5-C8.	3.1	262
7	Overview of the 2022 WHO Classification of Neuroendocrine Neoplasms. Endocrine Pathology, 2022, 33, 115-154.	9.0	227
8	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.	3.7	219
9	Pathological definition and clinical significance of vascular invasion in thyroid carcinomas of follicular epithelial derivation. Modern Pathology, 2011, 24, 1545-1552.	5.5	178
10	Overview of the 2022 WHO Classification of Pituitary Tumors. Endocrine Pathology, 2022, 33, 6-26.	9.0	174
11	The Complementary Role of Transcription Factors in the Accurate Diagnosis of Clinically Nonfunctioning Pituitary Adenomas. Endocrine Pathology, 2015, 26, 349-355.	9.0	167
12	Molecular correlates and rate of lymph node metastasis of non-invasive follicular thyroid neoplasm with papillary-like nuclear features and invasive follicular variant papillary thyroid carcinoma: the impact of rigid criteria to distinguish non-invasive follicular thyroid neoplasm with papillary-like nuclear features. Modern Pathology, 2017, 30, 810-825.	5.5	161
13	Spindle Cell Oncocytomas and Granular Cell Tumors of the Pituitary Are Variants of Pituicytoma. American Journal of Surgical Pathology, 2013, 37, 1694-1699.	3.7	151
14	International Histopathology Consensus for Unilateral Primary Aldosteronism. Journal of Clinical Endocrinology and Metabolism, 2021, 106, 42-54.	3.6	127
15	Biomarkers of aggressive pituitary adenomas. Journal of Molecular Endocrinology, 2012, 49, R69-R78.	2.5	123
16	Clinicopathological Correlations in Pituitary Adenomas. Brain Pathology, 2012, 22, 443-453.	4.1	120
17	Epidemiology and biomarker profile of pituitary adenohypophysial tumors. Modern Pathology, 2018, 31, 900-909.	5.5	120
18	Overview of the 2022 WHO Classification of Paragangliomas and Pheochromocytomas. Endocrine Pathology, 2022, 33, 90-114.	9.0	115

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19	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.	5.5	114
20	Clinical Safety of Renaming Encapsulated Follicular Variant of Papillary Thyroid Carcinoma: Is NIFTP Truly Benign?. World Journal of Surgery, 2018, 42, 321-326.	1.6	114
21	Noninvasive follicular thyroid neoplasm with papillary-like nuclear features: a review for pathologists. Modern Pathology, 2018, 31, 39-55.	5.5	107
22	The Diagnosis and Clinical Significance of Paragangliomas in Unusual Locations. Journal of Clinical Medicine, 2018, 7, 280.	2.4	104
23	Overview of the 2022 WHO Classification of Parathyroid Tumors. Endocrine Pathology, 2022, 33, 64-89.	9.0	96
24	Controversies in Thyroid Pathology: Thyroid Capsule Invasion and Extrathyroidal Extension. Annals of Surgical Oncology, 2010, 17, 386-391.	1.5	94
25	Clinicopathological correlates of hyperparathyroidism. Journal of Clinical Pathology, 2015, 68, 771-787.	2.0	88
26	Overview of the 2022 WHO Classification of Adrenal Cortical Tumors. Endocrine Pathology, 2022, 33, 155-196.	9.0	87
27	Precursor lesions of endocrine system neoplasms. Pathology, 2013, 45, 316-330.	0.6	84
28	DICER1 Mutations Are Frequent in Adolescent-Onset Papillary Thyroid Carcinoma. Journal of Clinical Endocrinology and Metabolism, 2018, 103, 2009-2015.	3.6	79
29	Immunohistochemistry in Diagnostic Parathyroid Pathology. Endocrine Pathology, 2018, 29, 113-129.	9.0	78
30	Retrospective study of 23 pathologically proven cases of central nervous system tuberculomas. Clinical Neurology and Neurosurgery, 2006, 108, 353-357.	1.4	76
31	An International Ki67 Reproducibility Study in Adrenal Cortical Carcinoma. American Journal of Surgical Pathology, 2016, 40, 569-576.	3.7	75
32	Diagnosis and management of gastrointestinal neuroendocrine tumors: An evidence-based Canadian consensus. Cancer Treatment Reviews, 2016, 47, 32-45.	7.7	74
33	Algorithmic approach to neuroendocrine tumors in targeted biopsies: Practical applications of immunohistochemical markers. Cancer Cytopathology, 2016, 124, 871-884.	2.4	72
34	Oncocytes, Oxyphils, Hürthle, and Askanazy Cells: Morphological and Molecular Features Of Oncocytic Thyroid Nodules. Endocrine Pathology, 2010, 21, 16-24.	9.0	65
35	Clinical features of silent corticotroph adenomas. Acta Neurochirurgica, 2012, 154, 1493-1498.	1.7	59
36	Null Cell Adenomas of the Pituitary Gland: an Institutional Review of Their Clinical Imaging and Behavioral Characteristics. Endocrine Pathology, 2015, 26, 63-70.	9.0	59

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37	Carney Complex with Adrenal Cortical Carcinoma. Journal of Clinical Endocrinology and Metabolism, 2012, 97, E202-E206.	3.6	57
38	Biomarkers of Parathyroid Carcinoma. Endocrine Pathology, 2012, 23, 221-231.	9.0	57
39	Inter-Observer Variation in the Pathologic Identification of Minimal Extrathyroidal Extension in Papillary Thyroid Carcinoma. Thyroid, 2016, 26, 512-517.	4.5	56
40	Diagnostic and Prognostic Biomarkers of Adrenal Cortical Carcinoma. American Journal of Surgical Pathology, 2018, 42, 201-213.	3.7	56
41	Epithelioid Angiomyolipoma: A Morphologically Distinct Variant That Mimics a Variety of Intra-abdominal Neoplasms. Archives of Pathology and Laboratory Medicine, 2011, 135, 665-670.	2.5	55
42	Protocol for the Examination of Specimens From Patients With Pheochromocytomas and Extra-Adrenal Paragangliomas. Archives of Pathology and Laboratory Medicine, 2014, 138, 182-188.	2.5	52
43	Parathyroid cancer: Outcome analysis of 16 patients treated at the princess margaret hospital. Head and Neck, 2013, 35, 35-39.	2.0	49
44	Follicular epithelial dysplasia of the thyroid: morphological and immunohistochemical characterization of a putative preneoplastic lesion to papillary thyroid carcinoma in chronic lymphocytic thyroiditis. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2013, 462, 557-563.	2.8	49
45	Therapeutic implications of accurate classification of pituitary adenomas. Seminars in Diagnostic Pathology, 2013, 30, 158-164.	1.5	48
46	Endocrine Manifestations of von Hippel–Lindau Disease. Archives of Pathology and Laboratory Medicine, 2015, 139, 263-268.	2.5	48
47	Immunohistochemical Biomarkers in Thyroid Pathology. Endocrine Pathology, 2018, 29, 91-112.	9.0	48
48	GATA3 immunoreactivity expands the transcription factor profile of pituitary neuroendocrine tumors. Modern Pathology, 2019, 32, 484-489.	5.5	48
49	Parathyroid carcinoma: diagnosis and clinical implications. Turk Patoloji Dergisi, 2015, 31 Suppl 1, 80-97.	0.3	46
50	Immunohistochemical Biomarkers of Adrenal Cortical Neoplasms. Endocrine Pathology, 2018, 29, 137-149.	9.0	45
51	Pituitary neuroendocrine tumors: a model for neuroendocrine tumor classification. Modern Pathology, 2021, 34, 1634-1650.	5.5	44
52	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.	9.0	43
53	Multiple Endocrine Tumors Associated with Germline <i>MAX</i> Mutations: Multiple Endocrine Neoplasia Type 5?. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e1163-e1182.	3.6	43
54	Classic Architecture with Multicentricity and Local Recurrence, and Absence of TERT Promoter Mutations are Correlates of BRAF V600E Harboring Pediatric Papillary Thyroid Carcinomas. Endocrine Pathology, 2016, 27, 153-161.	9.0	42

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55	Genomics of High-Grade Neuroendocrine Neoplasms: Well-Differentiated Neuroendocrine Tumor with High-Grade Features (G3 NET) and Neuroendocrine Carcinomas (NEC) of Various Anatomic Sites. Endocrine Pathology, 2021, 32, 192-210.	9.0	41
56	Update from the 5th Edition of the World Health Organization Classification of Head and Neck Tumors: Overview of the 2022 WHO Classification of Head and Neck Neuroendocrine Neoplasms. Head and Neck Pathology, 2022, 16, 123-142.	2.6	41
57	Can renal oncocytoma be differentiated from its renal mimics? The utility of anti-mitochondrial, caveolin 1, CD63 and cytokeratin 14 antibodies in the differential diagnosis. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2005, 447, 938-946.	2.8	40
58	Growth Patterns of Pituitary Adenomas and Histopathological Correlates. Journal of Clinical Endocrinology and Metabolism, 2014, 99, 1330-1338.	3.6	40
59	The Effect of Varicocele Repair on Experimental Varicoceleâ€Induced Testicular Germ Cell Apoptosis. Journal of Andrology, 2008, 29, 29-34.	2.0	38
60	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.	2.8	38
61	Mixed Adenoma Well-differentiated Neuroendocrine Tumor (MANET) of the Digestive System. American Journal of Surgical Pathology, 2018, 42, 1503-1512.	3.7	38
62	Prognostic and Predictive Markers in Medullary Thyroid Carcinoma. Endocrine Pathology, 2012, 23, 232-242.	9.0	35
63	Pituitary neuroendocrine tumors (PitNETs): nomenclature evolution, not clinical revolution. Pituitary, 2020, 23, 322-325.	2.9	34
64	Programmed Death-Ligand 1 (PD-L1) Is a Potential Biomarker of Disease-Free Survival in Papillary Thyroid Carcinoma: a Systematic Review and Meta-Analysis of PD-L1 Immunoexpression in Follicular Epithelial Derived Thyroid Carcinoma. Endocrine Pathology, 2020, 31, 291-300.	9.0	34
65	What Did We Learn from the Molecular Biology of Adrenal Cortical Neoplasia? From Histopathology to Translational Genomics. Endocrine Pathology, 2021, 32, 102-133.	9.0	33
66	Pitfalls in the Diagnosis of Follicular Epithelial Proliferations of the Thyroid. Advances in Anatomic Pathology, 2012, 19, 363-373.	4.3	32
67	Pathologic Reporting of Tall-Cell Variant of Papillary Thyroid Cancer: Have We Reached a Consensus?. Thyroid, 2017, 27, 1498-1504.	4.5	32
68	Xanthomatous Hypophysitis Is Associated with Ruptured Rathke's Cleft Cyst. Endocrine Pathology, 2017, 28, 83-90.	9.0	31
69	DGCR8 microprocessor defect characterizes familial multinodular goiter with schwannomatosis. Journal of Clinical Investigation, 2020, 130, 1479-1490.	8.2	31
70	Effect of Thyroid Gland Volume in Preoperative Detection of Suspected Malignant Thyroid Nodules in a Multinodular Goiter. Archives of Surgery, 2008, 143, 558.	2.2	30
71	The Role of Mediators of Cell Invasiveness, Motility, and Migration in the Pathogenesis of Silent Corticotroph Adenomas. Endocrine Pathology, 2013, 24, 191-198.	9.0	30
72	Clinicopathological correlates of adrenal Cushing's syndrome. Journal of Clinical Pathology, 2015, 68, 175-186.	2.0	30

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73	Hormone profiling, WHO 2010 grading, and AJCC / UICC staging in pancreatic neuroendocrine tumor behavior. Cancer Medicine, 2013, 2, 701-711.	2.8	29
74	The Role of Disease Label in Patient Perceptions and Treatment Decisions in the Setting of Low-Risk Malignant Neoplasms. JAMA Oncology, 2019, 5, 817.	7.1	29
75	Inhibin-expressing clear cell neuroendocrine tumor of the ampulla: an unusual presentation of von Hippel–Lindau disease. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2013, 463, 593-597.	2.8	28
76	Synchronous Multiple Pituitary Neuroendocrine Tumors of Different Cell Lineages. Endocrine Pathology, 2018, 29, 332-338.	9.0	28
77	Clinical, pathologic, and imaging characteristics of pituitary null cell adenomas as defined according to the 2017 World Health Organization criteria: a case series from two pituitary centers. Pituitary, 2019, 22, 514-519.	2.9	28
78	Parathyroid Lipoadenoma: a Clinicopathological Diagnosis and Possible Trap for the Unaware Pathologist. Endocrine Pathology, 2016, 27, 34-41.	9.0	27
79	Monomorphous Plurihormonal Pituitary Adenoma of Pit-1 Lineage in a Giant Adolescent with Central Hyperthyroidism. Endocrine Pathology, 2016, 27, 25-33.	9.0	26
80	Pituitary Adenomas Presenting as Sinonasal or Nasopharyngeal Masses. American Journal of Surgical Pathology, 2017, 41, 525-534.	3.7	26
81	Immunohistochemical Biomarkers in Pituitary Pathology. Endocrine Pathology, 2018, 29, 130-136.	9.0	26
82	Evaluation of the WHO 2010 Grading and AJCC/UICC Staging Systems in Prognostic Behavior of Intestinal Neuroendocrine Tumors. PLoS ONE, 2013, 8, e61538.	2.5	26
83	The Clinicopathological Spectrum of Parathyroid Carcinoma. Frontiers in Endocrinology, 2019, 10, 731.	3.5	25
84	A Diagnostic Approach to Adrenocortical Tumors. Surgical Pathology Clinics, 2019, 12, 967-995.	1.7	25
85	Molecular Pathology of Well-Differentiated Pulmonary and Thymic Neuroendocrine Tumors: What Do Pathologists Need to Know?. Endocrine Pathology, 2021, 32, 154-168.	9.0	25
86	Cribriform-Morular Thyroid Carcinoma Is a Distinct Thyroid Malignancy of Uncertain Cytogenesis. Endocrine Pathology, 2021, 32, 327-335.	9.0	25
87	Aldosterone-Producing Adrenal Cortical Adenoma with Oncocytic Change and Cytoplasmic Eosinophilic Globular Inclusions. Endocrine Pathology, 2009, 20, 182-185.	9.0	24
88	What's new in pituitary pathology?. Histopathology, 2018, 72, 133-141.	2.9	24
89	A Systematic Review and Meta-Analysis of the Diagnostic Performance of BRAF V600E Immunohistochemistry in Thyroid Histopathology. Endocrine Pathology, 2019, 30, 201-218.	9.0	24
90	Hypothalamic Vasopressin-Producing Tumors. American Journal of Surgical Pathology, 2019, 43, 251-260.	3.7	24

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91	Evolving concepts in prostatic neuroendocrine manifestations: from focal divergent differentiation to amphicrine carcinoma. Human Pathology, 2019, 85, 313-327.	2.0	24
92	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.	9.0	24
93	Prostate carcinoma with amphicrine features: further refining the spectrum of neuroendocrine differentiation in tumours of primary prostatic origin?. Histopathology, 2017, 71, 926-933.	2.9	23
94	Endocrine pathology: past, present and future. Pathology, 2018, 50, 111-118.	0.6	23
95	Ki67 Quantitative Interpretation: Insights using Image Analysis. Journal of Pathology Informatics, 2019, 10, 8.	1.7	23
96	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.	4.5	22
97	Neuroendocrine carcinoma of the skin—An updated review. Seminars in Diagnostic Pathology, 2013, 30, 234-244.	1.5	21
98	TTF-1 Expressing Sellar Neoplasm with Ependymal Rosettes and Oncocytic Change: Mixed Ependymal and Oncocytic Variant Pituicytoma. Endocrine Pathology, 2014, 25, 436-438.	9.0	21
99	Do You Know the Details of Your PAX8 Antibody? Monoclonal PAX8 (MRQ-50) Is Not Expressed in a Series of 45 Medullary Thyroid Carcinomas. Endocrine Pathology, 2020, 31, 33-38.	9.0	21
100	Cytokeratin profiles in pituitary neuroendocrine tumors. Human Pathology, 2021, 107, 87-95.	2.0	21
101	Inherited Follicular Epithelial-Derived Thyroid Carcinomas: From Molecular Biology to Histological Correlates. Endocrine Pathology, 2021, 32, 77-101.	9.0	21
102	The Impact of Phosphohistone-H3-Assisted Mitotic Count and Ki67 Score in the Determination of Tumor Grade and Prediction of Distant Metastasis in Well-Differentiated Pancreatic Neuroendocrine Tumors. Endocrine Pathology, 2016, 27, 162-170.	9.0	20
103	Leukocytoclastic Vasculitis due to Thalidomide in Multiple Myeloma. Japanese Journal of Clinical Oncology, 2007, 37, 704-707.	1.3	19
104	The Many Faces of Primary Aldosteronism and Cushing Syndrome: A Reflection of Adrenocortical Tumor Heterogeneity. Frontiers in Medicine, 2018, 5, 54.	2.6	19
105	Significance of Alpha-inhibin Expression in Pheochromocytomas and Paragangliomas. American Journal of Surgical Pathology, 2021, 45, 1264-1273.	3.7	19
106	A Mimic of Sarcomatoid Adrenal Cortical Carcinoma: Epithelioid Angiosarcoma Occurring in Adrenal Cortical Adenoma. Endocrine Pathology, 2014, 25, 404-409.	9.0	18
107	Hobnailâ€variant of papillary thyroid carcinoma in liquidâ€based cytology. Diagnostic Cytopathology, 2015, 43, 990-992.	1.0	18
108	Clear Cell Sarcoma–Like Tumor of the Gastrointestinal Tract. International Journal of Surgical Pathology, 2015, 23, 61-67.	0.8	18

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109	Clinicopathologic Correlates of Primary Aldosteronism. Archives of Pathology and Laboratory Medicine, 2015, 139, 948-954.	2.5	18
110	The Value of HBME-1 and Claudin-1 Expression Profile in the Distinction of BRAF-Like and RAS-Like Phenotypes in Papillary Thyroid Carcinoma. Endocrine Pathology, 2016, 27, 224-232.	9.0	18
111	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.	1.5	18
112	Data set for reporting of carcinoma of the adrenal cortex: explanations and recommendations of the guidelines from the International Collaboration on Cancer Reporting. Human Pathology, 2021, 110, 50-61.	2.0	18
113	Oncocytic Change in Thyroid Pathology. Frontiers in Endocrinology, 2021, 12, 678119.	3.5	18
114	Heme Oxygenase-1 Prevents Hyperthyroidism Induced Hepatic Damage via an Antioxidant and Antiapoptotic Pathway. Journal of Surgical Research, 2010, 164, 266-275.	1.6	16
115	Functional Cardiac Paraganglioma Associated with a Rare SDHC Mutation. Endocrine Pathology, 2014, 25, 315-320.	9.0	16
116	Positivity for GATA3 and TTF-1 (SPT24), and Negativity for Monoclonal PAX8 Expand the Biomarker Profile of the Solid Cell Nests of the Thyroid Gland. Endocrine Pathology, 2018, 29, 49-58.	9.0	16
117	Comprehensive characterization of a Canadian cohort of von Hippelâ€Lindau disease patients. Clinical Genetics, 2019, 96, 461-467.	2.0	16
118	Clinical Application of Next-Generation Sequencing in Advanced Thyroid Cancers. Thyroid, 2022, 32, 657-666.	4.5	16
119	Oral postinflammatory pigmentation: An analysis of 7 cases. Medicina Oral, Patologia Oral Y Cirugia Bucal, 2011, 16, e11-e14.	1.7	15
120	Malignant Pheochromocytoma Secreting Vasoactive Intestinal Peptide and Response to Sunitinib: A Case Report and Literature Review. Endocrine Practice, 2014, 20, e145-e150.	2.1	15
121	Hypothalamic Endocrine Tumors: An Update. Journal of Clinical Medicine, 2019, 8, 1741.	2.4	15
122	Middle Ear "Adenoma― a Neuroendocrine Tumor with Predominant L Cell Differentiation. Endocrine Pathology, 2021, 32, 433-441.	9.0	15
123	Diverse Oncogenic Fusions and Distinct Gene Expression Patterns Define the Genomic Landscape of Pediatric Papillary Thyroid Carcinoma. Cancer Research, 2021, 81, 5625-5637.	0.9	15
124	Genomics and Epigenomics of Pituitary Tumors: What Do Pathologists Need to Know?. Endocrine Pathology, 2021, 32, 3-16.	9.0	15
125	Solitary angiokeratoma of the tongue treated with diode laser. Lasers in Medical Science, 2009, 24, 123-125.	2.1	14
126	Primary Orbital Leiomyosarcoma. Ophthalmic Plastic and Reconstructive Surgery, 2009, 25, 154-155.	0.8	14

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127	Endobronchial Gangliocytic Paraganglioma: Not All Keratin-Positive Endobronchial Neuroendocrine Neoplasms are Pulmonary Carcinoids. Endocrine Pathology, 2014, 25, 356-358.	9.0	14
128	An Unusual Malignant Thyroid Nodule: Coexistence of Epithelioid Angiosarcoma and Follicular Variant Papillary Thyroid Carcinoma. Endocrine Pathology, 2014, 25, 350-352.	9.0	14
129	Clinical implications of accurate subtyping of pituitary adenomas: perspectives from the treating physician. Turk Patoloji Dergisi, 2015, 31 Suppl 1, 4-17.	0.3	14
130	Neuroendocrine Neoplasms Associated with Germline Pathogenic Variants in the Homologous Recombination Pathway. Endocrine Pathology, 2019, 30, 237-245.	9.0	14
131	Clinical and Radiological Features of Adrenal Cysts. Urologia Internationalis, 2008, 80, 31-36.	1.3	13
132	Head and neck paragangliomas: what does the pathologist need to know?. Diagnostic Histopathology, 2014, 20, 316-325.	0.4	13
133	Metastatic Neuroendocrine Neoplasms of Unknown Primary Site. , 2021, , 357-387.		13
134	Clear Cell Odontogenic Carcinoma of the Maxilla. Acta Medica (Hradec Kralove), 2011, 54, 122-124.	0.5	13
135	Morphological distinction of cortisolâ€producing and aldosteroneâ€producing adrenal cortical adenomas: not only possible but a critical clinical responsibility. Histopathology, 2012, 60, 1015-1016.	2.9	12
136	Familial hyperparathyroidism syndromes. Diagnostic Histopathology, 2016, 22, 92-100.	0.4	12
137	Inter-Observer Variation in the Pathologic Identification of Extranodal Extension in Nodal Metastasis from Papillary Thyroid Carcinoma. Thyroid, 2016, 26, 816-819.	4.5	12
138	Expanding the Spectrum of Colonic Manifestations in Tuberous Sclerosis: L-Cell Neuroendocrine Tumor Arising in the Background of Rectal PEComa. Endocrine Pathology, 2018, 29, 21-26.	9.0	12
139	Diagnostic Pitfall: Parathyroid Carcinoma Expands the Spectrum of Calcitonin and Calcitonin Gene-Related Peptide Expressing Neuroendocrine Neoplasms. Endocrine Pathology, 2019, 30, 168-172.	9.0	12
140	Does Hyperbaric Oxygen Administration Before or After Irradiation Decrease Side Effects of Irradiation on Implant Sites?. Annals of Plastic Surgery, 2011, 67, 62-67.	0.9	11
141	Hereditary Endocrine Tumor Syndromes: The Clinical and Predictive Role of Molecular Histopathology. AJSP Review and Reports, 2017, 22, 246-268.	0.1	11
142	Oncocytic Papillary Thyroid Carcinoma and Oncocytic Poorly Differentiated Thyroid Carcinoma: Clinical Features, Uptake, and Response to Radioactive Iodine Therapy, and Outcome. Frontiers in Endocrinology, 2021, 12, 795184.	3.5	11
143	Thyroid metastasis of endometrial carcinosarcoma associated with Graves' disease. Gynecological Endocrinology, 2007, 23, 562-566.	1.7	10
144	Interstitial granulomatous drug reaction due to thalidomide. Journal of the European Academy of Dermatology and Venereology, 2009, 23, 490-493.	2.4	10

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145	Composite Medullary and Papillary Thyroid Carcinoma In a Patient With MEN 2B. , 2009, 14, 208-213.		10
146	A Tumor With Many Faces: Metastatic Malignant Melanoma With Extensive Cartilaginous Differentiation. International Journal of Surgical Pathology, 2010, 18, 217-218.	0.8	10
147	Villous Papillary Thyroid Carcinoma: a Variant Associated with Marfan Syndrome. Endocrine Pathology, 2012, 23, 254-259.	9.0	10
148	Endoscopic Endonasal Pituitary Surgery For Nonfunctioning Pituitary Adenomas: Long-Term Outcomes and Management of Recurrent Tumors. World Neurosurgery, 2021, 146, e341-e350.	1.3	10
149	Metastatic Thyroid Carcinoma to the Gastric Body. Journal of Clinical Endocrinology and Metabolism, 2014, 99, 3958-3959.	3.6	9
150	Images in Endocrine Pathology: Papillary Variant of Medullary Thyroid Carcinoma with Cystic Change. Endocrine Pathology, 2015, 26, 87-89.	9.0	9
151	TFE3-Expressing Perivascular Epithelioid Cell Neoplasm (PEComa) of the Sella Turcica. Endocrine Pathology, 2017, 28, 22-26.	9.0	9
152	An Unusual Salivary Gland Tumor Mimicking Papillary Thyroid Carcinoma: Mammary Analog Secretory Carcinoma. Frontiers in Endocrinology, 2018, 9, 555.	3.5	9
153	Severe Primary Hyperparathyroidism Caused by Parathyroid Carcinoma in a 13‥earâ€Old Child; Novel Findings From HRpQCT. JBMR Plus, 2020, 4, e10324.	2.7	9
154	The Pangenomic Classification of Pituitary Neuroendocrine Tumors: Quality Histopathology is Required for Accurate Translational Research. Endocrine Pathology, 2021, 32, 415-417.	9.0	9
155	An Unusual Clinical Presentation of Pancreatic Solid Pseudopapillary Tumor With Ovarian Metastases: A Diagnostic Dilemma. International Journal of Surgical Pathology, 2011, 19, 342-345.	0.8	8
156	Thyroid neoplasms of follicular cell derivation: A simplified approach. Seminars in Diagnostic Pathology, 2013, 30, 178-185.	1.5	8
157	MEN2 Syndrome-Related Medullary Thyroid Carcinoma with Focal Tyrosine Hydroxylase Expression: Does It Represent a Hybrid Cellular Phenotype or Functional State of Tumor Cells?. Endocrine Pathology, 2017, 28, 362-366.	9.0	8
158	Special Issue On the 2022 WHO Classification of Endocrine and Neuroendocrine Tumors: a New Primer for Endocrine Pathology Practice. Endocrine Pathology, 2022, 33, 1-2.	9.0	8
159	Silent Corticotroph Adenoma with Adrenal Cortical Choristoma: a Rare but Distinct Morphological Entity. Endocrine Pathology, 2013, 24, 162-166.	9.0	7
160	Thyroglossal Duct Cyst Associated with Xanthogranulomatous Inflammation. Head and Neck Pathology, 2015, 9, 530-533.	2.6	7
161	An Unusual Adrenal Cortical Nodule: Composite Adrenal Cortical Adenoma and Adenomatoid Tumor. Endocrine Pathology, 2015, 26, 370-373.	9.0	7
162	An Unusual Solitary Thyroid Nodule with Bloody Follicles: Metastatic Renal Cell Carcinoma Within an Infiltrative Follicular Variant Papillary Carcinoma. Endocrine Pathology, 2016, 27, 171-174.	9.0	7

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163	Immunohistochemical Analysis of the Metabolic Phenotype of Adrenal Cortical Carcinoma. Endocrine Pathology, 2020, 31, 231-238.	9.0	7
164	Inherited Neuroendocrine Neoplasms. , 2021, , 409-459.		7
165	Genetic and epigenetic characterization of posterior pituitary tumors. Acta Neuropathologica, 2021, 142, 1025-1043.	7.7	7
166	Flat-type verruciform xanthoma of the tongue and its differential diagnosis. Dermatology Online Journal, 2009, 15, 5.	0.5	7
167	VHL mosaicism: the added value of multi-tissue analysis. Npj Genomic Medicine, 2022, 7, 21.	3.8	7
168	Anti-CD10 (56C6) is expressed variably in adrenocortical tumors and cannot be used to discriminate clear cell renal cell carcinomas. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2010, 456, 515-521.	2.8	6
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