Ozgur Mete

List of Publications by Year
in descending order

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233 papers	9,896 citations	53660 45 h-index	46693 89 g-index

242 all docs	242 docs citations	242 times ranked	8105 citing authors

#	Article	IF	Citations
1	Prognostic Significance of Pulmonary Multifocal Neuroendocrine Proliferation With Typical Carcinoid. Annals of Thoracic Surgery, 2022, 113, 966-974.	0.7	6
2	Letter to the Editor From Asa and Mete: â€œHypophysitis, the Growing Spectrum of a Rare Pituitary Diseaseâ€• Journal of Clinical Endocrinology and Metabolism, 2022, 107, e2649-e2649.	1.8	3
3	Endoscopic Treatment of Sellar Arachnoid Cysts via a Simple Cyst-Opening Technique: Long-Term Outcomes From a Single Center. World Neurosurgery, 2022, 161, e625-e634.	0.7	3
4	Overview of the 2022 WHO Classification of Parathyroid Tumors. Endocrine Pathology, 2022, 33, 64-89.	5.2	96
5	The Next Steps for Endocrine Pathology. Endocrine Pathology, 2022, 33, 228-230.	5.2	2
6	VHL mosaicism: the added value of multi-tissue analysis. Npj Genomic Medicine, 2022, 7, 21.	1.7	7
7	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.	1.3	41
8	Overview of the 2022 WHO Classification of Thyroid Neoplasms. Endocrine Pathology, 2022, 33, 27-63.	5.2	388
9	Clinical Application of Next-Generation Sequencing in Advanced Thyroid Cancers. Thyroid, 2022, 32, 657-666.	2.4	16
10	Overview of the 2022 WHO Classification of Pituitary Tumors. Endocrine Pathology, 2022, 33, 6-26.	5.2	174
11	Overview of the 2022 WHO Classification of Neuroendocrine Neoplasms. Endocrine Pathology, 2022, 33, 115-154.	5.2	227
12	Overview of the 2022 WHO Classification of Adrenal Cortical Tumors. Endocrine Pathology, 2022, 33, 155-196.	5.2	87
13	Overview of the 2022 WHO Classification of Paragangliomas and Pheochromocytomas. Endocrine Pathology, 2022, 33, 90-114.	5.2	115
14	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.	5.2	8
15	Is there a role for surgery after chemotherapy in recurrent/metastatic adrenal cortical cancer (ACC)?. Journal of Clinical Oncology, 2022, 40, 5092-5092.	0.8	0
16	International Histopathology Consensus for Unilateral Primary Aldosteronism. Journal of Clinical Endocrinology and Metabolism, 2021, 106, 42-54.	1.8	127
17	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.	1.1	18
18	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

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19	Cytokeratin profiles in pituitary neuroendocrine tumors. Human Pathology, 2021, 107, 87-95.	1.1	21
20	Inherited Follicular Epithelial-Derived Thyroid Carcinomas: From Molecular Biology to Histological Correlates. Endocrine Pathology, 2021, 32, 77-101.	5.2	21
21	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.	5.2	41
22	Special Issue on Molecular Pathology of Endocrine Neoplasms: Understanding the Basis of Endocrine Pathology Practice. Endocrine Pathology, 2021, 32, 1-2.	5.2	0
23	SILENT CORTICOTROPH TUMOR WITH ADRENOCORTICAL CHORISTOMA IN AN ELEVEN-YEAR-OLD BOY. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, .	0.4	3
24	Molecular Pathology of Well-Differentiated Pulmonary and Thymic Neuroendocrine Tumors: What Do Pathologists Need to Know?. Endocrine Pathology, 2021, 32, 154-168.	5.2	25
25	What Did We Learn from the Molecular Biology of Adrenal Cortical Neoplasia? From Histopathology to Translational Genomics. Endocrine Pathology, 2021, 32, 102-133.	5.2	33
26	The Pangenomic Classification of Pituitary Neuroendocrine Tumors: Quality Histopathology is Required for Accurate Translational Research. Endocrine Pathology, 2021, 32, 415-417.	5.2	9
27	P36.04 Molecular Prognostic Factors in Neuroendocrine Thymic Tumors: a Retrospective Multicentre Study. Journal of Thoracic Oncology, 2021, 16, S435-S436.	0.5	0
28	Significance of Crooke's Hyaline Change in Nontumorous Corticotrophs of Patients With Cushing Disease. Frontiers in Endocrinology, 2021, 12, 620005.	1.5	6
29	Significance of Alpha-inhibin Expression in Pheochromocytomas and Paragangliomas. American Journal of Surgical Pathology, 2021, 45, 1264-1273.	2.1	19
30	Oncocytic Change in Thyroid Pathology. Frontiers in Endocrinology, 2021, 12, 678119.	1.5	18
31	A Holistic Approach to Pathology Education During the Coronavirus Disease 2019 (COVID-19) Pandemic. Archives of Pathology and Laboratory Medicine, 2021, 145, 923-924.	1.2	3
32	Pituitary neuroendocrine tumors: a model for neuroendocrine tumor classification. Modern Pathology, 2021, 34, 1634-1650.	2.9	44
33	Middle Ear â€œAdenomaâ€• a Neuroendocrine Tumor with Predominant L Cell Differentiation. Endocrine Pathology, 2021, 32, 433-441.	5.2	15
34	Cribriform-Morular Thyroid Carcinoma Is a Distinct Thyroid Malignancy of Uncertain Cytogenesis. Endocrine Pathology, 2021, 32, 327-335.	5.2	25
35	Follicular cells in pituitary neuroendocrine tumors. Human Pathology, 2021, 114, 1-8.	1.1	4
36	Clinicopathological variables that correlate with sestamibi positivity in uniglandular parathyroid disease: a retrospective analysis of 378 parathyroid adenomas. Annals of Nuclear Medicine, 2021, , 1.	1.2	5

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37	Single-cell transcriptome and genome analysis: A much-needed tool for pituitary neuroendocrine tumor studies. Neuro-Oncology, 2021, 23, 1803-1804.	0.6	0
38	Diverse Oncogenic Fusions and Distinct Gene Expression Patterns Define the Genomic Landscape of Pediatric Papillary Thyroid Carcinoma. Cancer Research, 2021, 81, 5625-5637.	0.4	15
39	GHRH-producing tumors and other neuroendocrine neoplasms associated with acromegaly and/or gigantism. , 2021, , 259-274.		0
40	Familial Isolated Pituitary Adenoma (FIPA) Syndrome. Encyclopedia of Pathology, 2021, , 1-4.	0.0	0
41	Genomics and Epigenomics of Pituitary Tumors: What Do Pathologists Need to Know?. Endocrine Pathology, 2021, 32, 3-16.	5.2	15
42	Metastatic Neuroendocrine Neoplasms of Unknown Primary Site. , 2021, , 357-387.		13
43	Inherited Neuroendocrine Neoplasms. , 2021, , 409-459.		7
44	Multiple Endocrine Tumors Associated with Germline MAX Mutations: Multiple Endocrine Neoplasia Type 5?. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e1163-e1182.	1.8	43
45	Genetic and epigenetic characterization of posterior pituitary tumors. Acta Neuropathologica, 2021, 142, 1025-1043.	3.9	7
46	Recipient of the 2021 Endocrine Pathology Society Lifetime Achievement Award: Dr. Ronald A. DeLellis. Endocrine Pathology, 2021, 32, 429-431.	5.2	2
47	Paragangliomas and Pheochromocytomas. , 2021, , 263-285.		1
48	Pituitary Neuroendocrine Neoplasms. , 2021, , 55-83.		2
49	Thyroid Neuroendocrine Neoplasms. , 2021, , 119-136.		2
50	Parathyroid Neuroendocrine Neoplasms. , 2021, , 137-150.		0
51	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.	1.5	11
52	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.	5.2	21
53	Severe Primary Hyperparathyroidism Caused by Parathyroid Carcinoma in a 13â€Yearâ€Old Child; Novel Findings From HRpQCT. JBMR Plus, 2020, 4, e10324.	1.3	9
54	Pituitary neuroendocrine tumors (PitNETs): nomenclature evolution, not clinical revolution. Pituitary, 2020, 23, 322-325.	1.6	34

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55	Mixed Sparsely Granulated Lactotroph and Densely Granulated Somatotroph Pituitary Neuroendocrine Tumor Expands the Spectrum of Neuroendocrine Neoplasms in Ovarian Teratomas: the Role of Pituitary Neuroendocrine Cell Lineage Biomarkers. Endocrine Pathology, 2020, 31, 315-319.	5.2	4
56	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
57	Immunohistochemical Analysis of the Metabolic Phenotype of Adrenal Cortical Carcinoma. Endocrine Pathology, 2020, 31, 231-238.	5.2	7
58	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.	5.2	34
59	Book Review on â€œThe Survival Guide to Endocrine Pathology (Pathology Survival Guides Series 1,) Tj ETQq1 1 0.784314 rgBT /Over	5.2	0
60	Thyroid Tumor Capsular Invasion: the Bottom Line or Much Ado About Nothing?. Endocrine Pathology, 2020, 31, 141-142.	5.2	4
61	DGCR8 microprocessor defect characterizes familial multinodular goiter with schwannomatosis. Journal of Clinical Investigation, 2020, 130, 1479-1490.	3.9	31
62	Carney Complex. Encyclopedia of Pathology, 2020, , 1-4.	0.0	0
63	Marfan Syndrome. Encyclopedia of Pathology, 2020, , 1-5.	0.0	0
64	Multiple Endocrine Neoplasia Type 1 (MEN1). Encyclopedia of Pathology, 2020, , 1-6.	0.0	0
65	Multiple Endocrine Neoplasia Type 2 (MEN2). Encyclopedia of Pathology, 2020, , 1-5.	0.0	0
66	SUN-930 A Case of Acromegaly Secondary to Ectopic Growth Hormone-Releasing Hormone (GHRH) Secretion from a Bronchial Neuroendocrine Tumour. Journal of the Endocrine Society, 2020, 4, .	0.1	0
67	RARE-22. GERMLINE PATHOGENIC VARIANT c.1552G>A;p.E518K IN DGCR8 CONFERS SUSCEPTIBILITY FOR SCHWANNOMATOSIS AND THYROID TUMORS. Neuro-Oncology, 2020, 22, iii447-iii447.	0.6	0
68	DICER1 Syndrome. Encyclopedia of Pathology, 2020, , 1-5.	0.0	0
69	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.	1.6	28
70	Comprehensive characterization of a Canadian cohort of von Hippelâ€Lindau disease patients. Clinical Genetics, 2019, 96, 461-467.	1.0	16
71	VEGFRâ€2 is downregulated in sestamibiâ€negative parathyroid adenomas. Head and Neck, 2019, 41, 3564-3569.	0.9	4
72	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

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73	The Clinicopathological Spectrum of Parathyroid Carcinoma. Frontiers in Endocrinology, 2019, 10, 731.	1.5	25
74	A Diagnostic Approach to Adrenocortical Tumors. Surgical Pathology Clinics, 2019, 12, 967-995.	0.7	25
75	Hypothalamic Endocrine Tumors: An Update. Journal of Clinical Medicine, 2019, 8, 1741.	1.0	15
76	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
77	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
78	Adrenal cortical neoplasia: from histology to molecular biology. Diagnostic Histopathology, 2019, 25, 178-189.	0.2	1
79	The Role of Disease Label in Patient Perceptions and Treatment Decisions in the Setting of Low-Risk Malignant Neoplasms. JAMA Oncology, 2019, 5, 817.	3.4	29
80	Diagnostic Pitfall: Parathyroid Carcinoma Expands the Spectrum of Calcitonin and Calcitonin Gene-Related Peptide Expressing Neuroendocrine Neoplasms. Endocrine Pathology, 2019, 30, 168-172.	5.2	12
81	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
82	Hypothalamic Vasopressin-Producing Tumors. American Journal of Surgical Pathology, 2019, 43, 251-260.	2.1	24
83	Neuroendocrine Neoplasms Associated with Germline Pathogenic Variants in the Homologous Recombination Pathway. Endocrine Pathology, 2019, 30, 237-245.	5.2	14
84	Evolving concepts in prostatic neuroendocrine manifestations: from focal divergent differentiation to amphicrine carcinoma. Human Pathology, 2019, 85, 313-327.	1.1	24
85	GATA3 immunoreactivity expands the transcription factor profile of pituitary neuroendocrine tumors. Modern Pathology, 2019, 32, 484-489.	2.9	48
86	Ki67 Quantitative Interpretation: Insights using Image Analysis. Journal of Pathology Informatics, 2019, 10, 8.	0.8	23
87	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
88	DICER1 Mutations Are Frequent in Adolescent-Onset Papillary Thyroid Carcinoma. Journal of Clinical Endocrinology and Metabolism, 2018, 103, 2009-2015.	1.8	79
89	Immunohistochemical Biomarkers in Pituitary Pathology. Endocrine Pathology, 2018, 29, 130-136.	5.2	26
90	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

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91	Epidemiology and biomarker profile of pituitary adenohypophysial tumors. Modern Pathology, 2018, 31, 900-909.	2.9	120
92	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.	5.2	16
93	Immunohistochemical Biomarkers of Adrenal Cortical Neoplasms. Endocrine Pathology, 2018, 29, 137-149.	5.2	45
94	Noninvasive follicular thyroid neoplasm with papillary-like nuclear features: a review for pathologists. Modern Pathology, 2018, 31, 39-55.	2.9	107
95	Diagnostic and Prognostic Biomarkers of Adrenal Cortical Carcinoma. American Journal of Surgical Pathology, 2018, 42, 201-213.	2.1	56
96	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
97	What's new in pituitary pathology?. Histopathology, 2018, 72, 133-141.	1.6	24
98	Endocrine pathology: past, present and future. Pathology, 2018, 50, 111-118.	0.3	23
99	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.	5.2	12
100	Healing of Oral Lichenoid Lesions following Replacement of Dental Amalgam Restorations with Feldspathic Ceramic Inlay-Onlay Restorations: Clinical Results of a Follow-Up Period Varied from Three Months up to Five Years. BioMed Research International, 2018, 2018, 1-7.	0.9	4
101	Autoinfarction of Giant Parathyroid Adenoma after Preoperative Withdrawal of Anticoagulants. Case Reports in Surgery, 2018, 2018, 1-5.	0.2	3
102	A Young Male with Parafibromin-Deficient Parathyroid Carcinoma Due to a Rare Germline HRPT2/CDC73 Mutation. Endocrine Pathology, 2018, 29, 374-379.	5.2	6
103	An Unusual Salivary Gland Tumor Mimicking Papillary Thyroid Carcinoma: Mammary Analog Secretory Carcinoma. Frontiers in Endocrinology, 2018, 9, 555.	1.5	9
104	The Diagnosis and Clinical Significance of Paragangliomas in Unusual Locations. Journal of Clinical Medicine, 2018, 7, 280.	1.0	104
105	Synchronous Multiple Pituitary Neuroendocrine Tumors of Different Cell Lineages. Endocrine Pathology, 2018, 29, 332-338.	5.2	28
106	Mixed Adenoma Well-differentiated Neuroendocrine Tumor (MANET) of the Digestive System. American Journal of Surgical Pathology, 2018, 42, 1503-1512.	2.1	38
107	The Many Faces of Primary Aldosteronism and Cushing Syndrome: A Reflection of Adrenocortical Tumor Heterogeneity. Frontiers in Medicine, 2018, 5, 54.	1.2	19
108	Immunohistochemistry in Diagnostic Parathyroid Pathology. Endocrine Pathology, 2018, 29, 113-129.	5.2	78

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109	Immunohistochemical Biomarkers in Thyroid Pathology. Endocrine Pathology, 2018, 29, 91-112.	5.2	48
110	Editorial: Special Issue on Immunohistochemical Biomarkers in Endocrine Pathology. Endocrine Pathology, 2018, 29, 89-90.	5.2	1
111	Liver Transplantation in a Young Patient with Severe and Refractory Carcinoid Syndrome. AACE Clinical Case Reports, 2018, 4, e289-e293.	0.4	0
112	Comprehensive Molecular Characterization of Pheochromocytoma and Paraganglioma. Cancer Cell, 2017, 31, 181-193.	7.7	532
113	From pituitary adenoma to pituitary neuroendocrine tumor (PitNET): an International Pituitary Pathology Club proposal. Endocrine-Related Cancer, 2017, 24, C5-C8.	1.6	262
114	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.	2.9	161
115	Xanthomatous Hypophysitis Is Associated with Ruptured Rathkeâ€™s Cleft Cyst. Endocrine Pathology, 2017, 28, 83-90.	5.2	31
116	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.	5.2	8
117	Pituitary Adenomas Presenting as Sinonasal or Nasopharyngeal Masses. American Journal of Surgical Pathology, 2017, 41, 525-534.	2.1	26
118	Pathologic Reporting of Tall-Cell Variant of Papillary Thyroid Cancer: Have We Reached a Consensus?. Thyroid, 2017, 27, 1498-1504.	2.4	32
119	Pancreatic endocrine neoplasia: familial syndromes. Diagnostic Histopathology, 2017, 23, 378-385.	0.2	6
120	Prostate carcinoma with amphicrine features: further refining the spectrum of neuroendocrine differentiation in tumours of primary prostatic origin?. Histopathology, 2017, 71, 926-933.	1.6	23
121	Overview of the 2017 WHO Classification of Pituitary Tumors. Endocrine Pathology, 2017, 28, 228-243.	5.2	319
122	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
123	TFE3-Expressing Perivascular Epithelioid Cell Neoplasm (PEComa) of the Sella Turcica. Endocrine Pathology, 2017, 28, 22-26.	5.2	9
124	Hereditary Endocrine Tumor Syndromes: The Clinical and Predictive Role of Molecular Histopathology. AJSP Review and Reports, 2017, 22, 246-268.	0.0	11
125	An International Ki67 Reproducibility Study in Adrenal Cortical Carcinoma. American Journal of Surgical Pathology, 2016, 40, 569-576.	2.1	75
126	Familial hyperparathyroidism syndromes. Diagnostic Histopathology, 2016, 22, 92-100.	0.2	12

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127	Diagnosis and management of gastrointestinal neuroendocrine tumors: An evidence-based Canadian consensus. Cancer Treatment Reviews, 2016, 47, 32-45.	3.4	74
128	Nomenclature Revision for Encapsulated Follicular Variant of Papillary Thyroid Carcinoma. JAMA Oncology, 2016, 2, 1023.	3.4	1,192
129	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.	5.2	20
130	Comprehensive Pan-Genomic Characterization of Adrenocortical Carcinoma. Cancer Cell, 2016, 29, 723-736.	7.7	482
131	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
132	Cytology and Pathology: Pitfalls and Challenges. , 2016, , 33-46.		3
133	Algorithmic approach to neuroendocrine tumors in targeted biopsies: Practical applications of immunohistochemical markers. Cancer Cytopathology, 2016, 124, 871-884.	1.4	72
134	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.	5.2	7
135	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.	5.2	18
136	Parathyroid Lipoadenoma: a Clinicopathological Diagnosis and Possible Trap for the Unaware Pathologist. Endocrine Pathology, 2016, 27, 34-41.	5.2	27
137	Inter-Observer Variation in the Pathologic Identification of Minimal Extrathyroidal Extension in Papillary Thyroid Carcinoma. Thyroid, 2016, 26, 512-517.	2.4	56
138	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.	5.2	42
139	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
140	Monomorphous Plurihormonal Pituitary Adenoma of Pit-1 Lineage in a Giant Adolescent with Central Hyperthyroidism. Endocrine Pathology, 2016, 27, 25-33.	5.2	26
141	Not All Post-FNA Spindle Cell Proliferations in the Thyroid Are of Myofibroblastic Origin: Follicular Adenoma with Spindle Cell Metaplasia. Endocrine Pathology, 2015, 26, 374-376.	5.2	3
142	Hobnailâ€variant of papillary thyroid carcinoma in liquidâ€based cytology. Diagnostic Cytopathology, 2015, 43, 990-992.	0.5	18
143	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
144	Parathyroid carcinoma: diagnosis and clinical implications. Turk Patoloji Dergisi, 2015, 31 Suppl 1, 80-97.	0.1	46

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145	Update in endocrine pathology. Turk Patoloji Dergisi, 2015, 31 Suppl 1, 1-3.	0.1	0
146	Pancreatic Neuroendocrine Tumors Producing GHRH, GH, Ghrelin, PTH, or PTHrP. , 2015, , 125-139.		3
147	Thyroglossal Duct Cyst Associated with Xanthogranulomatous Inflammation. Head and Neck Pathology, 2015, 9, 530-533.	1.3	7
148	Republished: Clinicopathological correlates of adrenal Cushing's syndrome. Postgraduate Medical Journal, 2015, 91, 331-342.	0.9	6
149	Endocrine Manifestations of von Hippelâ€“Lindau Disease. Archives of Pathology and Laboratory Medicine, 2015, 139, 263-268.	1.2	48
150	Images in Endocrine Pathology: Papillary Variant of Medullary Thyroid Carcinoma with Cystic Change. Endocrine Pathology, 2015, 26, 87-89.	5.2	9
151	Null Cell Adenomas of the Pituitary Gland: an Institutional Review of Their Clinical Imaging and Behavioral Characteristics. Endocrine Pathology, 2015, 26, 63-70.	5.2	59
152	Clear Cell Sarcomaâ€“Like Tumor of the Gastrointestinal Tract. International Journal of Surgical Pathology, 2015, 23, 61-67.	0.4	18
153	Clinicopathological correlates of hyperparathyroidism. Journal of Clinical Pathology, 2015, 68, 771-787.	1.0	88
154	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
155	An Unusual Adrenal Cortical Nodule: Composite Adrenal Cortical Adenoma and Adenomatoid Tumor. Endocrine Pathology, 2015, 26, 370-373.	5.2	7
156	The Complementary Role of Transcription Factors in the Accurate Diagnosis of Clinically Nonfunctioning Pituitary Adenomas. Endocrine Pathology, 2015, 26, 349-355.	5.2	167
157	Clinicopathologic Correlates of Primary Aldosteronism. Archives of Pathology and Laboratory Medicine, 2015, 139, 948-954.	1.2	18
158	Clinicopathological correlates of adrenal Cushing's syndrome. Journal of Clinical Pathology, 2015, 68, 175-186.	1.0	30
159	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
160	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
161	A Mimic of Sarcomatoid Adrenal Cortical Carcinoma: Epithelioid Angiosarcoma Occurring in Adrenal Cortical Adenoma. Endocrine Pathology, 2014, 25, 404-409.	5.2	18
162	Metastatic Thyroid Carcinoma to the Gastric Body. Journal of Clinical Endocrinology and Metabolism, 2014, 99, 3958-3959.	1.8	9

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163	Non-pheochromocytoma (PCC)/paraganglioma (PGL) tumors in patients with succinate dehydrogenase-related PCCâ€“PGL syndromes: a clinicopathological and molecular analysis. European Journal of Endocrinology, 2014, 170, 1-12.	1.9	219
164	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
165	Multiple Endocrine Neoplasia Type 1. , 2014, 19, 85-89.		6
166	Endobronchial Gangliocytic Paraganglioma: Not All Keratin-Positive Endobronchial Neuroendocrine Neoplasms are Pulmonary Carcinoids. Endocrine Pathology, 2014, 25, 356-358.	5.2	14
167	An Unusual Malignant Thyroid Nodule: Coexistence of Epithelioid Angiosarcoma and Follicular Variant Papillary Thyroid Carcinoma. Endocrine Pathology, 2014, 25, 350-352.	5.2	14
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