Abbas Agaimy

List of Publications by Year
in descending order

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330 papers	11,569 citations	28190 55 h-index	48187 88 g-index

342 all docs	342 docs citations	342 times ranked	10438 citing authors

#	Article	IF	Citations
1	Minute Gastric Sclerosing Stromal Tumors (GIST Tumorlets) Are Common in Adults and Frequently Show c-KIT Mutations. American Journal of Surgical Pathology, 2007, 31, 113-120.	2.1	338
2	Numerous IgG4-positive plasma cells are ubiquitous in diverse localised non-specific chronic inflammatory conditions and need to be distinguished from IgG4-related systemic disorders. Journal of Clinical Pathology, 2011, 64, 237-243.	1.0	307
3	Succinate Dehydrogenase (SDH)-deficient Renal Carcinoma. American Journal of Surgical Pathology, 2014, 38, 1588-1602.	2.1	282
4	Solitary Fibrous Tumors/Hemangiopericytomas with Different Variants of the NAB2-STAT6 Gene Fusion Are Characterized by Specific Histomorphology and Distinct Clinicopathological Features. American Journal of Pathology, 2014, 184, 1209-1218.	1.9	198
5	SMARCB1 (INI-1)-deficient Sinonasal Carcinoma. American Journal of Surgical Pathology, 2017, 41, 458-471.	2.1	198
6	Occurrence of other malignancies in patients with gastrointestinal stromal tumors. Seminars in Diagnostic Pathology, 2006, 23, 120-129.	1.0	187
7	Fumarate Hydrataseâ€“deficient Renal Cell Carcinoma Is Strongly Correlated With Fumarate Hydratase Mutation and Hereditary Leiomyomatosis and Renal Cell Carcinoma Syndrome. American Journal of Surgical Pathology, 2016, 40, 865-875.	2.1	182
8	SWI/SNF Complexâ€“deficient Undifferentiated/Rhabdoid Carcinomas of the Gastrointestinal Tract. American Journal of Surgical Pathology, 2016, 40, 544-553.	2.1	175
9	Somatostatin receptor expression related to TP53 and RB1 alterations in pancreatic and extrapancreatic neuroendocrine neoplasms with a Ki67-index above 20%. Modern Pathology, 2017, 30, 587-598.	2.9	162
10	Aberrant DNA hypermethylation of SDHC: a novel mechanism of tumor development in Carney triad. Endocrine-Related Cancer, 2014, 21, 567-577.	1.6	161
11	Enhancer hijacking activates oncogenic transcription factor NR4A3 in acinic cell carcinomas of the salivary glands. Nature Communications, 2019, 10, 368.	5.8	153
12	Clear Cell Myoepithelial Carcinoma of Salivary Glands Showing EWSR1 Rearrangement. American Journal of Surgical Pathology, 2015, 39, 338-348.	2.1	141
13	SMARCB1(INI1)-deficient Sinonasal Basaloid Carcinoma. American Journal of Surgical Pathology, 2014, 38, 1274-1281.	2.1	140
14	The Expanding Family of SMARCB1(INI1)-deficient Neoplasia. Advances in Anatomic Pathology, 2014, 21, 394-410.	2.4	140
15	New developments in existing WHO entities and evolving molecular concepts: The Genitourinary Pathology Society (GUPS) update on renal neoplasia. Modern Pathology, 2021, 34, 1392-1424.	2.9	138
16	Spectrum of KIT/PDGFRA/BRAF mutations and Phosphatidylinositol-3-Kinase pathway gene alterations in gastrointestinal stromal tumors (GIST). Cancer Letters, 2011, 312, 43-54.	3.2	125
17	SMARCA4 loss is synthetic lethal with CDK4/6 inhibition in non-small cell lung cancer. Nature Communications, 2019, 10, 557.	5.8	125
18	Gastrointestinal stromal tumours: a regular origin in the muscularis propria, but an extremely diverse gross presentation. Langenbeck's Archives of Surgery, 2006, 391, 322-329.	0.8	119

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19	SMARCA4 and SMARCA2 deficiency in nonâ€“small cell lung cancer: immunohistochemical survey of 316 consecutive specimens. Annals of Diagnostic Pathology, 2017, 26, 47-51.	0.6	118
20	Novel, emerging and provisional renal entities: The Genitourinary Pathology Society (GUPS) update on renal neoplasia. Modern Pathology, 2021, 34, 1167-1184.	2.9	118
21	Tubulocystic Carcinoma of the Kidney With Poorly Differentiated Foci. American Journal of Surgical Pathology, 2016, 40, 1457-1472.	2.1	112
22	Disentangling inflammatory from fibrotic disease activity by fibroblast activation protein imaging. Annals of the Rheumatic Diseases, 2020, 79, 1485-1491.	0.5	111
23	Paediatric and adult soft tissue sarcomas with NTRK1 gene fusions: a subset of spindle cell sarcomas unified by a prominent myopericytic/haemangiopericytic pattern. Journal of Pathology, 2016, 238, 700-710.	2.1	108
24	ISL1 expression is not restricted to pancreatic well-differentiated neuroendocrine neoplasms, but is also commonly found in well and poorly differentiated neuroendocrine neoplasms of extrapancreatic origin. Modern Pathology, 2013, 26, 995-1003.	2.9	107
25	Synthetic vulnerabilities of mesenchymal subpopulations in pancreatic cancer. Nature, 2017, 542, 362-366.	13.7	105
26	Pattern of SMARCB1 (INI1) and SMARCA4 (BRG1) in poorly differentiated endometrioid adenocarcinoma of the uterus: analysis of a series with emphasis on a novel SMARCA4-deficient dedifferentiated rhabdoid variant. Annals of Diagnostic Pathology, 2015, 19, 198-202.	0.6	102
27	Reappraisal of Morphologic Differences Between Renal Medullary Carcinoma, Collecting Duct Carcinoma, and Fumarate Hydrataseâ€“deficient Renal Cell Carcinoma. American Journal of Surgical Pathology, 2018, 42, 279-292.	2.1	101
28	Metastatic Malignant Melanoma With Complete Loss of Differentiation Markers (Undifferentiated/Dedifferentiated Melanoma). American Journal of Surgical Pathology, 2016, 40, 181-191.	2.1	100
29	SMARCB1 (INI1)-negative Rhabdoid Carcinomas of the Gastrointestinal Tract. American Journal of Surgical Pathology, 2014, 38, 910-920.	2.1	96
30	Multimodal analysis of cell-free DNA whole-genome sequencing for pediatric cancers with low mutational burden. Nature Communications, 2021, 12, 3230.	5.8	95
31	Nuclear NR4A3 Immunostaining Is a Specific and Sensitive Novel Marker for Acinic Cell Carcinoma of the Salivary Glands. American Journal of Surgical Pathology, 2019, 43, 1264-1272.	2.1	94
32	SMARCA4-deficient pulmonary adenocarcinoma: clinicopathological, immunohistochemical, and molecular characteristics of a novel aggressive neoplasm with a consistent TTF1neg/CK7pos/HepPar-1pos immunophenotype. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2017, 471, 599-609.	1.4	94
33	Peripheral nerve sheath tumors of the gastrointestinal tract: a multicenter study of 58 patients including NF1-associated gastric schwannoma and unusual morphologic variants. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2010, 456, 411-422.	1.4	92
34	Gastrointestinal manifestations of neurofibromatosis type 1 (Recklinghausen's disease): clinicopathological spectrum with pathogenetic considerations. International Journal of Clinical and Experimental Pathology, 2012, 5, 852-62.	0.5	91
35	Collecting Duct Carcinoma Versus Renal Medullary Carcinoma. American Journal of Surgical Pathology, 2014, 38, 871-874.	2.1	90
36	Pancreatic undifferentiated rhabdoid carcinoma: KRAS alterations and SMARCB1 expression status define two subtypes. Modern Pathology, 2015, 28, 248-260.	2.9	90

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37	SMARCA4-deficient Sinonasal Carcinoma. American Journal of Surgical Pathology, 2020, 44, 703-710.	2.1	90
38	Influence of low FODMAP and gluten-free diets on disease activity and intestinal microbiota in patients with non-celiac gluten sensitivity. Clinical Nutrition, 2019, 38, 697-707.	2.3	89
39	Sclerosing nodular lesions of the gastrointestinal tract containing large numbers of IgG4 plasma cells. Pathology, 2011, 43, 31-35.	0.3	86
40	Calcifying Fibrous Tumor of the Stomach: Clinicopathologic and Molecular Study of Seven Cases With Literature Review and Reappraisal of Histogenesis. American Journal of Surgical Pathology, 2010, 34, 271-278.	2.1	83
41	NCOA4-RET and TRIM27-RET Are Characteristic Gene Fusions in Salivary Intraductal Carcinoma, Including Invasive and Metastatic Tumors. American Journal of Surgical Pathology, 2019, 43, 1303-1313.	2.1	82
42	Gastrointestinal stromal tumors (GIST) from risk stratification systems to the new TNM proposal: more questions than answers? A review emphasizing the need for a standardized GIST reporting. International Journal of Clinical and Experimental Pathology, 2010, 3, 461-71.	0.5	80
43	Phosphaturic Mesenchymal Tumors. American Journal of Surgical Pathology, 2017, 41, 1371-1380.	2.1	77
44	Recurrent Somatic PDGFRB Mutations in Sporadic Infantile/Solitary Adult Myofibromas But Not in Angioleiomyomas and Myopericytomas. American Journal of Surgical Pathology, 2017, 41, 195-203.	2.1	76
45	True smooth muscle neoplasms of the gastrointestinal tract: morphological spectrum and classification in a series of 85 cases from a single institute. Langenbeck's Archives of Surgery, 2007, 392, 75-81.	0.8	75
46	Microscopic Gastrointestinal Stromal Tumors in Esophageal and Intestinal Surgical Resection Specimens. American Journal of Surgical Pathology, 2008, 32, 867-873.	2.1	74
47	Epithelial-Myoepithelial Carcinoma. American Journal of Surgical Pathology, 2018, 42, 18-27.	2.1	71
48	Salivary gland mucoepidermoid carcinoma is a clinically, morphologically and genetically heterogeneous entity: a clinicopathological study of 40 cases with emphasis on grading, histological variants and presence of the t(11;19) translocation. Histopathology, 2011, 58, 557-570.	1.6	70
49	Lymph node metastasis in gastrointestinal stromal tumours (GIST) occurs preferentially in young patients â‰40Âyears: an overview based on our case material and the literature. Langenbeck's Archives of Surgery, 2009, 394, 375-381.	0.8	69
50	Claudin-4 expression distinguishes SWI/SNF complex-deficient undifferentiated carcinomas from sarcomas. Modern Pathology, 2017, 30, 539-548.	2.9	69
51	Genomic EWSR1 Fusion Sequence as Highly Sensitive and Dynamic Plasma Tumor Marker in Ewing Sarcoma. Clinical Cancer Research, 2016, 22, 4356-4365.	3.2	68
52	Primary and metastatic cardiac tumors: imaging characteristics, surgical treatment, and histopathological spectrum: a 10-year-experience at a German heart center. Cardiovascular Pathology, 2012, 21, 436-443.	0.7	65
53	Recurrent Loss of SMARCA4 in Sinonasal Teratocarcinosarcoma. American Journal of Surgical Pathology, 2020, 44, 1331-1339.	2.1	64
54	ETV6 Gene Rearrangements Characterize a Morphologically Distinct Subset of Sinonasal Low-grade Nonâ€“intestinal-type Adenocarcinoma. American Journal of Surgical Pathology, 2017, 41, 1552-1560.	2.1	61

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55	Loss of expression of the SWI/SNF complex is a frequent event in undifferentiated/dedifferentiated urothelial carcinoma of the urinary tract. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2016, 469, 321-330.	1.4	58
56	SMARCA4-deficient Sinonasal Carcinoma. Head and Neck Pathology, 2017, 11, 541-545.	1.3	58
57	Rhabdoid and Undifferentiated Phenotype in Renal Cell Carcinoma. American Journal of Surgical Pathology, 2017, 41, 253-262.	2.1	56
58	Colonic Adenocarcinomas Harboring NTRK Fusion Genes. American Journal of Surgical Pathology, 2020, 44, 162-173.	2.1	56
59	Defining Ewing and Ewing-like small round cell tumors (SRCT): The need for molecular techniques in their categorization and differential diagnosis. A study of 200 cases. Annals of Diagnostic Pathology, 2016, 22, 25-32.	0.6	55
60	Molecular Profiling of Clear Cell Myoepithelial Carcinoma of Salivary Glands With EWSR1 Rearrangement Identifies Frequent PLAG1 Gene Fusions But No EWSR1 Fusion Transcripts. American Journal of Surgical Pathology, 2021, 45, 1-13.	2.1	54
61	Lipomatous Salivary Gland Tumors. American Journal of Surgical Pathology, 2013, 37, 128-137.	2.1	52
62	Reappraisal of sinonasal undifferentiated carcinoma: SMARCB1 (INI1)-deficient sinonasal carcinoma: a single-institution experience. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2015, 467, 649-656.	1.4	52
63	Recurrent Fusions Between YAP1 and KMT2A in Morphologically Distinct Neoplasms Within the Spectrum of Low-grade Fibromyxoid Sarcoma and Sclerosing Epithelioid Fibrosarcoma. American Journal of Surgical Pathology, 2020, 44, 594-606.	2.1	52
64	Multiple Sporadic Gastrointestinal Stromal Tumors (GISTs) of the Proximal Stomach are Caused by Different Somatic KIT Mutations Suggesting a Field Effect. American Journal of Surgical Pathology, 2008, 32, 1553-1559.	2.1	51
65	Hereditary SWI/SNF complex deficiency syndromes. Seminars in Diagnostic Pathology, 2018, 35, 193-198.	1.0	51
66	Primary and metastatic cardiac sarcomas: a 12-year experience at a German heart center. International Journal of Clinical and Experimental Pathology, 2012, 5, 928-38.	0.5	51
67	Cytomegalovirus infection presenting as isolated inflammatory polyps of the gastrointestinal tract. Pathology, 2011, 43, 440-446.	0.3	50
68	Recurrent Mutations within the Amino-Terminal Region of Î²-Catenin Are Probable Key Molecular Driver Events in Sinonasal Hemangiopericytoma. American Journal of Pathology, 2015, 185, 563-571.	1.9	49
69	ALK rearranged renal cell carcinoma (ALK-RCC): a multi-institutional study of twelve cases with identification of novel partner genes CLIP1, KIF5B and KIAA1217. Modern Pathology, 2020, 33, 2564-2579.	2.9	49
70	Biphasic Squamoid Alveolar Renal Cell Carcinoma. American Journal of Surgical Pathology, 2016, 40, 664-675.	2.1	48
71	Combination of 5-fluorouracil and thymoquinone targets stem cell gene signature in colorectal cancer cells. Cell Death and Disease, 2019, 10, 379.	2.7	48
72	Benign Serrated Colorectal Fibroblastic Polyps/Intramucosal Perineuriomas Are True Mixed Epithelial-stromal Polyps (Hybrid Hyperplastic Polyp/Mucosal Perineurioma) With Frequent BRAF Mutations. American Journal of Surgical Pathology, 2010, 34, 1663-1671.	2.1	48

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73	DEK-AFF2 Carcinoma of the Sinonasal Region and Skull Base. American Journal of Surgical Pathology, 2021, 45, 1682-1693.	2.1	47
74	Coexistence of gastrointestinal stromal tumours (GIST) and malignant neoplasms of different origin: Prognostic implications. International Journal of Surgery, 2014, 12, 371-377.	1.1	46
75	Adenomatoid tumors of the female and male genital tract. A comparative clinicopathologic and immunohistochemical analysis of 47 cases emphasizing their site-specific morphologic diversity. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2011, 458, 593-602.	1.4	45
76	TRIM28 haploinsufficiency predisposes to Wilms tumor. International Journal of Cancer, 2019, 145, 941-951.	2.3	45
77	Malignant teratoid tumor of the thyroid gland: an aggressive primitive multiphenotypic malignancy showing organotypical elements and frequent DICER1 alterationsâ€"is the term â€œthyroblastomaâ€•more appropriate?. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2020, 477, 787-798.	1.4	45
78	NUT Carcinoma of the Salivary Glands. American Journal of Surgical Pathology, 2018, 42, 877-884.	2.1	44
79	SMARCB1 (INI-1)-Deficient Adenocarcinoma of the Sinonasal Tract: A Potentially Under-Recognized form of Sinonasal Adenocarcinoma with Occasional Yolk Sac Tumor-Like Features. Head and Neck Pathology, 2020, 14, 465-472.	1.3	44
80	Surgery with Radical Intent: Is There an Indication for G3 Neuroendocrine Neoplasms?. Annals of Surgical Oncology, 2020, 27, 1348-1355.	0.7	44
81	High-grade Transformation/Dedifferentiation in Salivary Gland Carcinomas: Occurrence Across Subtypes and Clinical Significance. Advances in Anatomic Pathology, 2021, 28, 107-118.	2.4	44
82	Sellar Region Atypical Teratoid/Rhabdoid Tumors (ATRT) in Adults Display DNA Methylation Profiles of the ATRT-MYC Subgroup. American Journal of Surgical Pathology, 2018, 42, 506-511.	2.1	43
83	Dedifferentiated and Undifferentiated Melanomas. American Journal of Surgical Pathology, 2021, 45, 240-254.	2.1	43
84	Fat-Containing Salivary Gland Tumors: A Review. Head and Neck Pathology, 2013, 7, 90-96.	1.3	42
85	Fumarate hydratase (FH) deficiency in uterine leiomyomas: recognition by histological features versus blind immunoscreening. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2018, 472, 789-796.	1.4	42
86	SWI/SNF Complex-Deficient Soft Tissue Neoplasms. Surgical Pathology Clinics, 2019, 12, 149-163.	0.7	42
87	Sinonasal Undifferentiated Carcinoma (SNUC): From an Entity to Morphologic Pattern and Back Againâ€"A Historical Perspective. Advances in Anatomic Pathology, 2020, 27, 51-60.	2.4	42
88	EWSR1-SMAD3â€“rearranged Fibroblastic Tumor. American Journal of Surgical Pathology, 2018, 42, 1325-1333.	2.1	40
89	Eosinophilic vacuolated tumor (EVT) of kidney demonstrates sporadic TSC/MTOR mutations: next-generation sequencing multi-institutional study of 19 cases. Modern Pathology, 2022, 35, 344-351.	2.9	40
90	Sporadic Cajal cell hyperplasia is common in resection specimens for distal oesophageal carcinoma. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2006, 448, 288-294.	1.4	39

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91	Phenotypical and molecular distinctness of sinonasal haemangiopericytoma compared to solitary fibrous tumour of the sinonasal tract. Histopathology, 2014, 65, 667-673.	1.6	39
92	Comparative study of soft tissue perineurioma and meningioma using a fiveâ€marker immunohistochemical panel. Histopathology, 2014, 65, 60-70.	1.6	39
93	Impact of age and gender on tumor related prognosis in gastrointestinal stromal tumors (GIST). BMC Cancer, 2015, 15, 57.	1.1	39
94	Sinonasal Tract Alveolar Rhabdomyosarcoma in Adults: A Clinicopathologic and Immunophenotypic Study of Fifty-Two Cases with Emphasis on Epithelial Immunoreactivity. Head and Neck Pathology, 2018, 12, 181-192.	1.3	39
95	Expression of Neuroendocrine Markers in Different Molecular Subtypes of Breast Carcinoma. BioMed Research International, 2014, 2014, 1-9.	0.9	38
96	Defined morphological criteria allow reliable diagnosis of colorectal serrated polyps and predict polyp genetics. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2014, 464, 663-672.	1.4	38
97	Pancreatic panniculitis in a patient with pancreatic-type acinar cell carcinoma of the liver â€“ case report and review of literature. BMC Cancer, 2016, 16, 130.	1.1	38
98	Superficial acral fibromyxoma: clinicopathological, immunohistochemical, and molecular study of 11 cases highlighting frequent Rb1 loss/deletions. Human Pathology, 2017, 60, 192-198.	1.1	38
99	YAP1-NUTM1 Gene Fusion in Porocarcinoma of the External Auditory Canal. Head and Neck Pathology, 2020, 14, 982-990.	1.3	38
100	CTNNB1 (Î²-Catenin)-altered Neoplasia. Advances in Anatomic Pathology, 2016, 23, 1-12.	2.4	37
101	Follicular dendritic cell sarcoma: clinicopathologic study of 15 cases with emphasis on novel expression of MDM2, somatostatin receptor 2A, and PD-L1. Annals of Diagnostic Pathology, 2016, 23, 21-28.	0.6	37
102	PD-L1 expression in tumor tissue and peripheral blood of patients with oral squamous cell carcinoma. Oncotarget, 2017, 8, 112584-112597.	0.8	37
103	Dysplastic Lipoma. American Journal of Surgical Pathology, 2018, 42, 1530-1540.	2.1	36
104	Intraductal Papillary Mucinous Neoplasms of Minor Salivary Glands With AKT1 p.Glu17Lys Mutation. American Journal of Surgical Pathology, 2018, 42, 1076-1082.	2.1	36
105	Desmoplastic myxoid tumor, SMARCB1-mutant: clinical, histopathological and molecular characterization of a pineal region tumor encountered in adolescents and adults. Acta Neuropathologica, 2020, 139, 277-286.	3.9	36
106	Misses and near misses in diagnosing nodular fasciitis and morphologically related reactive myofibroblastic proliferations: experience of a referral center with emphasis on frequency of USP6 gene rearrangements. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2018, 473, 351-360.	1.4	35
107	Distinct genetic alterations and luminal molecular subtype in nested variant of urothelial carcinoma. Histopathology, 2019, 75, 865-875.	1.6	35
108	Malignant transformation of oral leukoplakia is associated with macrophage polarization. Journal of Translational Medicine, 2020, 18, 11.	1.8	34

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109	Mammary Analog Secretory Carcinoma of the Nasal Cavity. American Journal of Surgical Pathology, 2018, 42, 735-743.	2.1	32
110	Multiple sporadic gastrointestinal stromal tumours arising at different gastrointestinal sites: pattern of involvement of the muscularis propria as a clue to independent primary GISTs. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2009, 455, 101-108.	1.4	31
111	Angioleiomyoma of the Sinonasal Tract: Analysis of 16 Cases and Review of the Literature. Head and Neck Pathology, 2015, 9, 463-473.	1.3	31
112	Biphasic papillary renal cell carcinoma is a rare morphological variant with frequent multifocality: a study of 28 cases. Histopathology, 2018, 72, 777-785.	1.6	31
113	Escalation in mucus cystatin 2, pappalysinâ€A, and periostin levels over time predict need for recurrent surgery in chronic rhinosinusitis with nasal polyps. International Forum of Allergy and Rhinology, 2019, 9, 1212-1219.	1.5	31
114	Inflammatory leiomyosarcoma shows frequent co-expression of smooth and skeletal muscle markers supporting a primitive myogenic phenotype: a report of 9 cases with a proposal for reclassification as low-grade inflammatory myogenic tumor. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2020, 477, 219-230.	1.4	30
115	Gene Expression in Solitary Fibrous Tumors (SFTs) Correlates with Anatomic Localization and NAB2-STAT6 Gene Fusion Variants. American Journal of Pathology, 2021, 191, 602-617.	1.9	30
116	Programmed death-1 (PD-1) receptor/PD-1 ligand (PD-L1) expression in fumarate hydratase-deficient renal cell carcinoma. Annals of Diagnostic Pathology, 2017, 29, 17-22.	0.6	29
117	Histiocyte-rich rhabdomyoblastic tumor: rhabdomyosarcoma, rhabdomyoma, or rhabdomyoblastic tumor of uncertain malignant potential? A histologically distinctive rhabdomyoblastic tumor in search of a place in the classification of skeletal muscle neoplasms. Modern Pathology, 2019, 32, 446-457.	2.9	29
118	Microsecretory Adenocarcinoma of Salivary Glands: An Expanded Series of 24 Cases. Head and Neck Pathology, 2021, 15, 1192-1201.	1.3	29
119	Epithelioid gastric stromal tumours of the antrum in young females with the Carney triad: a report of three new cases with mutational analysis and comparative genomic hybridization. Oncology Reports, 2007, 18, 9-15.	1.2	29
120	Primary and metastatic high-grade pleomorphic sarcoma/malignant fibrous histiocytoma of the gastrointestinal tract: an approach to the differential diagnosis in a series of five cases with emphasis on myofibroblastic differentiation. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2007, 451, 949-957.	1.4	28
121	Impact of postoperative radiotherapy and HER2/new overexpression in salivary duct carcinoma. Strahlentherapie Und Onkologie, 2017, 193, 961-970.	1.0	28
122	Inflamed benign tumors of the parotid gland: Diagnostic pitfalls from a potentially misleading entity. Head and Neck, 2015, 37, 23-29.	0.9	27
123	Angiosarcoma arising in association with vascular Dacron grafts and orthopedic joint prostheses: clinicopathologic, immunohistochemical, and molecular study. Annals of Diagnostic Pathology, 2016, 21, 21-28.	0.6	27
124	Dual Functional States of R406W-Desmin Assembly Complexes Cause Cardiomyopathy With Severe Intercalated Disc Derangement in Humans and in Knock-In Mice. Circulation, 2020, 142, 2155-2171.	1.6	27
125	Sclerosing Microcystic Adenocarcinoma of the Head and Neck Mucosa: A Neoplasm Closely Resembling Microcystic Adnexal Carcinoma. Head and Neck Pathology, 2016, 10, 501-508.	1.3	26
126	Mammary Analogue Secretory Carcinoma of Salivary Glands: Diagnostic Pitfall with Distinct Immunohistochemical Profile and Molecular Features. Rare Tumors, 2017, 9, 89-92.	0.3	26

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127	IDH2 R172 Mutations Across Poorly Differentiated Sinonasal Tract Malignancies. American Journal of Surgical Pathology, 2021, 45, 1190-1204.	2.1	26
128	Anorectal gastrointestinal stromal tumors: a retrospective multicenter analysis of 15 cases emphasizing their high local recurrence rate and the need for standardized therapeutic approach. International Journal of Colorectal Disease, 2013, 28, 1057-1064.	1.0	25
129	Primary signet ring stromal tumor of the testis: a study of 13 cases indicating their phenotypic and genotypic analogy to pancreatic solid pseudopapillary neoplasm. Human Pathology, 2017, 67, 85-93.	1.1	25
130	Head and Neck Kaposi Sarcoma: Clinicopathological Analysis of 11 Cases. Head and Neck Pathology, 2018, 12, 511-516.	1.3	25
131	Importance of the PD-1/PD-L1 Axis for Malignant Transformation and Risk Assessment of Oral Leukoplakia. Biomedicines, 2021, 9, 194.	1.4	25
132	Perineurioma of the stomach. Pathology Research and Practice, 2005, 201, 463-467.	1.0	24
133	Intestinal-type adenocarcinoma arising in a congenital sublingual teratoid cyst. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2007, 450, 479-481.	1.4	24
134	Dermatofibrosarcoma protuberans: surgical management of a challenging mesenchymal tumor. World Journal of Surgical Oncology, 2019, 17, 90.	0.8	24
135	Sinonasal papillomas: A single centre experience on 137 cases with emphasis on malignant transformation and EGFR/KRAS status in â€œcarcinoma ex papillomaâ€• Annals of Diagnostic Pathology, 2020, 46, 151504.	0.6	24
136	EWSR1-PATZ1-rearranged sarcoma: a report of nine cases of spindle and round cell neoplasms with predilection for thoracoabdominal soft tissues and frequent expression of neural and skeletal muscle markers. Modern Pathology, 2021, 34, 770-785.	2.9	24
137	SWI/SNF-deficient head and neck neoplasms: An overview. Seminars in Diagnostic Pathology, 2021, 38, 175-182.	1.0	24
138	Pancreatic-type acinar cell carcinoma of the liver: a clinicopathologic study of four patients. Modern Pathology, 2011, 24, 1620-1626.	2.9	23
139	High proliferation rate and TNM stage but not histomorphological subtype are independent prognostic markers for overall survival in papillary renal cell carcinoma. Human Pathology, 2019, 83, 212-223.	1.1	23
140	Malignant ascites occurs most often in patients with high-grade serous papillary ovarian cancer at initial diagnosis: a retrospective analysis of 191 women treated at Bayreuth Hospital, 2006â€“2015. Archives of Gynecology and Obstetrics, 2019, 299, 515-523.	0.8	23
141	Hepatic angiomyolipoma: a series of six cases with emphasis on pathological-radiological correlations and unusual variants diagnosed by core needle biopsy. International Journal of Clinical and Experimental Pathology, 2012, 5, 512-21.	0.5	23
142	Multifocal gastric gastrointestinal stromal tumors (GISTs) with lymph node metastases in children and young adults: A comparative clinical and histomorphological study of three cases including a new case of Carney triad. Diagnostic Pathology, 2011, 6, 52.	0.9	22
143	Juvenileâ€like (inflammatory/hyperplastic) mucosal polyps of the gastrointestinal tract in neurofibromatosis type 1. Histopathology, 2014, 64, 777-786.	1.6	22
144	â€˜Neuroendocrineâ€™ middle ear adenomas: consistent expression of the transcription factor <scp>ISL</scp>1 further supports their neuroendocrine derivation. Histopathology, 2015, 66, 182-191.	1.6	22

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145	MUC4 is a valuable marker for distinguishing secretory carcinoma of the salivary glands from its mimics. Histopathology, 2021, 79, 315-324.	1.6	22
146	Recurrent novel HMGA2-NCOR2 fusions characterize a subset of keratin-positive giant cell-rich soft tissue tumors. Modern Pathology, 2021, 34, 1507-1520.	2.9	22
147	Morphological heterogeneity of oral salivary gland carcinomas: a clinicopathologic study of 41 cases with long term follow-up emphasizing the overlapping spectrum of adenoid cystic carcinoma and polymorphous low-grade adenocarcinoma. International Journal of Clinical and Experimental Pathology, 2011, 4, 336-48.	0.5	22
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