Andrew L Folpe

List of Publications by Year
in descending order

173 papers	13,217 citations	22153 59 h-index	24258 110 g-index
173 all docs	173 docs citations	173 times ranked	7234 citing authors

#	Article	IF	Citations
1	Rapidly fatal SMARCA4-deficient undifferentiated sarcoma originating from hybrid hemosiderotic fibrolipomatous tumor/pleomorphic hyalinizing angiectatic tumor of the foot. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2022, 480, 1115-1120.	2.8	6
2	Head and Neck Low-Grade Fibromyxoid Sarcoma: A Clinicopathologic Study of 15 Cases. Head and Neck Pathology, 2022, 16, 434-443.	2.6	4
3	Sternoclavicular joint pseudotumor: a clinicopathologic and radiologic study of 25 cases occurring in patients without prior neck dissection. Human Pathology, 2022, 122, 32-39.	2.0	0
4	Cellular variant of kaposiform lymphangiomatosis: a report of three cases, expanding the morphologic and molecular genetic spectrum of this rare entity. Human Pathology, 2022, 122, 72-81.	2.0	9
5	â€˜I Canâ€™t Keep Up!â€™: an update on advances in soft tissue pathology occurring after the publication of the 2020 World Health Organization classification of soft tissue and bone tumours. Histopathology, 2022, 80, 54-75.	2.9	13
6	Overlapping morphological, immunohistochemical and genetic features of superficial CD34-positive fibroblastic tumor and PRDM10-rearranged soft tissue tumor. Modern Pathology, 2022, 35, 767-776.	5.5	14
7	Loss of dimethylated H3K27 (H3K27me2) expression is not a specific marker of malignant peripheral nerve sheath tumor (MPNST): An immunohistochemical study of 137 cases, with emphasis on MPNST and melanocytic tumors. Annals of Diagnostic Pathology, 2022, 59, 151967.	1.3	3
8	Glomangiomatosis of the Upper Extremity. Journal of Hand Surgery, 2021, 46, 716.e1-716.e3.	1.6	2
9	Primary intraâ€abdominal melanoma arising in association with extracutaneous blue naevus: a report of two cases. Histopathology, 2021, 78, 281-289.	2.9	3
10	Mesenchymal tumors of the gastrointestinal tract with NTRK rearrangements: a clinicopathological, immunophenotypic, and molecular study of eight cases, emphasizing their distinction from gastrointestinal stromal tumor (GIST). Modern Pathology, 2021, 34, 95-103.	5.5	52
11	Lipoblastomas presenting in older children and adults: analysis of 22 cases with identification of novel PLAG1 fusion partners. Modern Pathology, 2021, 34, 584-591.	5.5	29
12	Update on <scp>SWI</scp>/<scp>SNF</scp>â€related gynecologic mesenchymal neoplasms: <scp>SMARCA4</scp>â€deficient uterine sarcoma and <scp>SMARCB1</scp>â€deficient vulvar neoplasms. Genes Chromosomes and Cancer, 2021, 60, 190-209.	2.8	13
13	â€œInflammatory Leiomyosarcomaâ€and â€œHistiocyte-rich Rhabdomyoblastic Tumorâ€: a clinicopathological, immunohistochemical and genetic study of 13 cases, with a proposal for reclassification as â€œInflammatory Rhabdomyoblastic Tumorâ€. Modern Pathology, 2021, 34, 758-769.	5.5	27
14	Recurrent novel HMGA2-NCOR2 fusions characterize a subset of keratin-positive giant cell-rich soft tissue tumors. Modern Pathology, 2021, 34, 1507-1520.	5.5	22
15	Inflammatory rhabdomyoblastic tumor with progression to high-grade rhabdomyosarcoma. Modern Pathology, 2021, 34, 1035-1036.	5.5	13
16	NUTM1-rearranged colorectal sarcoma: a clinicopathologically and genetically distinctive malignant neoplasm with a poor prognosis. Modern Pathology, 2021, 34, 1547-1557.	5.5	24
17	CD10 (neprilysin) expression: a potential adjunct in the distinction of hibernoma from morphologic mimics. Human Pathology, 2021, 110, 12-19.	2.0	3
18	Response to Lee et al: Toward a unifying entity that encompasses most, but perhaps not all, inflammatory leiomyosarcomas and histiocyte-rich rhabdomyoblastic tumors. Modern Pathology, 2021, 34, 1439.	5.5	3

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19	Myxoid pleomorphic liposarcomaâ€"a clinicopathologic, immunohistochemical, molecular genetic and epigenetic study of 12 cases, suggesting a possible relationship with conventional pleomorphic liposarcoma. Modern Pathology, 2021, 34, 2043-2049.	5.5	24
20	EWSR1-WT1 gene fusions in neoplasms other than desmoplastic small round cell tumor: a report of three unusual tumors involving the female genital tract and review of the literature. Modern Pathology, 2021, 34, 1912-1920.	5.5	17
21	Hiding in plain sight: Gene panel and genetic markers reveal 26-year undiagnosed tumor-induced osteomalacia of the rib concurrently misdiagnosed as X-linked hypophosphatemia. Bone Reports, 2021, 14, 100744.	0.4	6
22	RNAscope CSF1 chromogenic in situ hybridization: a potentially useful tool in the differential diagnosis of tenosynovial giant cell tumors. Human Pathology, 2021, 115, 1-9.	2.0	5
23	Radiation Therapy for Treatment of Soft Tissue Sarcoma in Adults: Executive Summary of an ASTRO Clinical Practice Guideline. Practical Radiation Oncology, 2021, 11, 339-351.	2.1	65
24	PRRX1â€"NCOA1â€•rearranged fibroblastic tumour: aÂclinicopathological, immunohistochemical and molecular genetic study of six cases of a potentially underâ€•recognised, distinctive mesenchymal tumour. Histopathology, 2021, 79, 997-1003.	2.9	11
25	Eccrine angiomatous hamartoma: First case in the cytology literature. Annals of Diagnostic Pathology, 2021, 54, 151796.	1.3	0
26	â€œHey! Whatever happened to hemangiopericytoma and fibrosarcoma?â€•An update on selected conceptual advances in soft tissue pathology which have occurred over the past 50 years. Human Pathology, 2020, 95, 113-136.	2.0	11
27	Extraneural perineurioma: CT and MRI imaging characteristics. Skeletal Radiology, 2020, 49, 109-114.	2.0	7
28	PIK3CA mutations in lipomatosis of nerve with or without nerve territory overgrowth. Modern Pathology, 2020, 33, 420-430.	5.5	33
29	Update on selected advances in the immunohistochemical and molecular genetic analysis of soft tissue tumors. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2020, 476, 3-15.	2.8	14
30	Contemporary approaches to soft tissue and bone pathology. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2020, 476, 1-2.	2.8	0
31	Frequent overexpression of klotho in fusion-negative phosphaturic mesenchymal tumors with tumorigenic implications. Modern Pathology, 2020, 33, 858-870.	5.5	17
32	Lymphatic-type â€œAngiosarcomaâ€•With Prominent Lymphocytic Infiltrate. American Journal of Surgical Pathology, 2020, 44, 271-279.	3.7	9
33	Paraspinal pseudoneoplasms: a series of 58 consultation cases emphasizing the importance of pathology-radiology correlation. Human Pathology, 2020, 103, 14-24.	2.0	2
34	Well-Differentiated/Dedifferentiated Liposarcoma Arising in the Upper Aerodigestive Tract: 8 Cases Mimicking Non-adipocytic Lesions. Head and Neck Pathology, 2020, 14, 974-981.	2.6	15
35	Xanthogranulomatous epithelial tumor: report of 6 cases of a novel, potentially deceptive lesion with a predilection for young women. Modern Pathology, 2020, 33, 1889-1895.	5.5	13
36	Colonic Angiosarcoma Arising in Association with Amyloid Deposits. Case Reports in Gastrointestinal Medicine, 2020, 2020, 1-6.	0.3	2

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37	MyoD1 expression in fibroepithelial stromal polyps. Human Pathology, 2020, 99, 75-79.	2.0	6
38	Juvenile Hyaline Fibromatosis. Mayo Clinic Proceedings, 2020, 95, 328-329.	3.0	5
39	Head and Neck Mesenchymal Neoplasms With GLI1 Gene Alterations. American Journal of Surgical Pathology, 2020, 44, 729-737.	3.7	46
40	Imaging features of phosphaturic mesenchymal tumors. Skeletal Radiology, 2019, 48, 119-127.	2.0	26
41	Spindle Epithelial Tumor with Thymus-Like Differentiation (SETTLE): A Next-Generation Sequencing Study. Head and Neck Pathology, 2019, 13, 162-168.	2.6	6
42	HepaticYAP1-TFE3Rearranged Epithelioid Hemangioendothelioma. Case Reports in Gastrointestinal Medicine, 2019, 2019, 1-5.	0.3	5
43	OLIG2 is a marker of the fusion protein-driven neurodevelopmental transcriptional signature in alveolar rhabdomyosarcoma. Human Pathology, 2019, 91, 77-85.	2.0	20
44	Phosphaturic mesenchymal tumors: A review and update. Seminars in Diagnostic Pathology, 2019, 36, 260-268.	1.5	77
45	Atypical lipomatous tumour/wellâ€differentiated liposarcoma and deâ€differentiated liposarcoma in patients agedÂâ‰Â40Âyears: a study of 116 patients. Histopathology, 2019, 75, 833-842.	2.9	16
46	Loss of succinate dehydrogenase B immunohistochemical expression distinguishes pulmonary chondromas from hamartomas. Histopathology, 2019, 75, 825-832.	2.9	6
47	Immunohistochemistry for TFE3 lacks specificity and sensitivity in the diagnosis of TFE3-rearranged neoplasms: a comparative, 2-laboratory study. Human Pathology, 2019, 87, 65-74.	2.0	41
48	Perinephric myxoid pseudotumor of fat: a distinctive pseudoneoplasm most often associated with non-neoplastic renal disease. Human Pathology, 2019, 87, 37-43.	2.0	15
49	A comparison of adult rhabdomyosarcoma and high-grade neuroendocrine carcinoma of the urinary bladder reveals novel PPP1R12A fusions in rhabdomyosarcoma. Human Pathology, 2019, 88, 48-59.	2.0	2
50	Tenosynovitis With Psammomatous Calcifications. American Journal of Surgical Pathology, 2019, 43, 261-267.	3.7	5
51	Malignant Tenosynovial Giant Cell Tumor: The True â€œSynovial Sarcoma?â€•A Clinicopathologic, Immunohistochemical, and Molecular Cytogenetic Study of 10 Cases, Supporting Origin from Synoviocytes. Modern Pathology, 2019, 32, 242-251.	5.5	29
52	Ancillary Diagnostic Tests in the Diagnosis of Cutaneous Soft Tissue Neoplasms. , 2019, , 15-56.		0
53	Aberrant receptor tyrosine kinase signaling in lipofibromatosis: a clinicopathological and molecular genetic study of 20 cases. Modern Pathology, 2019, 32, 423-434.	5.5	49
54	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.	5.5	29

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55	Phosphaturic mesenchymal tumor without osteomalacia: additional confirmation of the â€œnonphosphaturicâ€•variant, with emphasis on the roles of FGF23 chromogenic in situ hybridization and FN1-FGFR1 fluorescence in situ hybridization. Human Pathology, 2018, 80, 94-98.	2.0	18
56	Mediastinal Synovial Sarcoma. American Journal of Surgical Pathology, 2018, 42, 761-766.	3.7	16
57	Lowâ€grade fibromyxoid sarcoma arising within the median nerve. Neuropathology, 2018, 38, 309-314.	1.2	3
58	Recurrent <scp>GNA</scp>14 mutations in anastomosing haemangiomas. Histopathology, 2018, 73, 354-357.	2.9	33
59	Abdominopelvic and Retroperitoneal Low-Grade Fibromyxoid Sarcoma. American Journal of Clinical Pathology, 2018, 149, 128-134.	0.7	19
60	Mesenchymal chondrosarcomas showing immunohistochemical evidence of rhabdomyoblastic differentiation: a potential diagnostic pitfall. Human Pathology, 2018, 77, 28-34.	2.0	34
61	Malignant Peripheral Nerve Sheath Tumor in a Patient With BAP1 Tumor Predisposition Syndrome. World Neurosurgery, 2018, 109, 362-364.	1.3	3
62	Polypoid fibroadipose tumors of the esophagus: â€˜giant fibrovascular polypâ€™ or liposarcoma? A clinicopathological and molecular cytogenetic study of 13 cases. Modern Pathology, 2018, 31, 337-342.	5.5	37
63	â€œChondroblastomaâ€likeâ€•epithelioid fibrous histiocytoma: A previously undescribed and potentially confusing variant. Journal of Cutaneous Pathology, 2018, 45, 99-103.	1.3	6
64	Radiation Therapy for Retroperitoneal Sarcomas: Influences of Histology, Grade, and Size. Sarcoma, 2018, 2018, 1-8.	1.3	12
65	Spindle cell rhabdomyosarcoma of bone with <scp>FUS</scp>â€“<scp>TFCP</scp>2 fusion: confirmation of a very recently described rhabdomyosarcoma subtype. Histopathology, 2018, 73, 514-520.	2.9	63
66	Lipoblastoma-like tumor of the vulva: a clinicopathologic, immunohistochemical, fluorescence in situ hybridization and genomic copy number profiling study of seven cases. Modern Pathology, 2018, 31, 1862-1868.	5.5	25
67	Clinicopathologic features and outcomes of gastrointestinal stromal tumors arising from the esophagus and gastroesophageal junction. Journal of Gastrointestinal Oncology, 2018, 9, 718-727.	1.4	7
68	Hypoxia-related microRNA-210 is a diagnostic marker for discriminating osteoblastoma and osteosarcoma. Journal of Orthopaedic Research, 2017, 35, 1137-1146.	2.3	13
69	Fibrous hamartoma of infancy: a clinicopathologic study of 145 cases, including 2 with sarcomatous features. Modern Pathology, 2017, 30, 474-485.	5.5	61
70	Recurrent GNAQ mutations in anastomosing hemangiomas. Modern Pathology, 2017, 30, 722-727.	5.5	59
71	Hemosiderotic Fibrolipomatous Tumor, Pleomorphic Hyalinizing Angiectatic Tumor, and Myxoinflammatory Fibroblastic Sarcoma: Related or Not?. Advances in Anatomic Pathology, 2017, 24, 268-277.	4.3	32
72	BRAF V600E Mutations Occur in a Subset of Glomus Tumors, and Are Associated With Malignant Histologic Characteristics. American Journal of Surgical Pathology, 2017, 41, 1532-1541.	3.7	49

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73	Gastroblastoma harbors a recurrent somatic MALAT1â€“GLI1 fusion gene. Modern Pathology, 2017, 30, 1443-1452.	5.5	93
74	Composite hemangioendothelioma with neuroendocrine marker expression: an aggressive variant. Modern Pathology, 2017, 30, 1589-1602.	5.5	38
75	Comparision of New Diagnostic Tools for Malignant Peripheral Nerve Sheath Tumors. Pathology and Oncology Research, 2017, 23, 393-398.	1.9	6
76	Anastomosing Hemangiomas Arising in Unusual Locations. American Journal of Surgical Pathology, 2016, 40, 1084-1089.	3.7	47
77	Pseudolipoblastic perineurioma: an unusual morphological variant of perineurioma that may simulate liposarcoma. Human Pathology, 2016, 57, 22-27.	2.0	5
78	TGFBR3 and MGEA5 rearrangements are much more common in â€œhybridâ€•hemosiderotic fibrolipomatous tumor-myxoinflammatory fibroblastic sarcomas than in classical myxoinflammatory fibroblastic sarcomas: a morphological and fluorescence in situ hybridization study. Human Pathology, 2016, 53, 14-24.	2.0	36
79	Voluntary Second Opinions in Pediatric Bone and Soft Tissue Pathology. International Journal of Surgical Pathology, 2016, 24, 685-691.	0.8	16
80	Primary angiomatoid fibrous histiocytoma of the lung with mediastinal lymph node metastasis. Human Pathology, 2016, 58, 134-137.	2.0	15
81	Characterization of FN1â€“FGFR1 and novel FN1â€“FGF1 fusion genes in a large series of phosphaturic mesenchymal tumors. Modern Pathology, 2016, 29, 1335-1346.	5.5	139
82	Renal Leiomyoma and Leiomyosarcoma. American Journal of Surgical Pathology, 2016, 40, 1557-1563.	3.7	11
83	Oncocytic variant of malignant gastrointestinal neuroectodermal tumor: a potential diagnostic pitfall. Human Pathology, 2016, 57, 13-16.	2.0	25
84	SMARCB1-deficient Vulvar Neoplasms. American Journal of Surgical Pathology, 2015, 39, 836-849.	3.7	44
85	Aberrant intermediate filament and synaptophysin expression is a frequent event in malignant melanoma: an immunohistochemical study of 73 cases. Modern Pathology, 2015, 28, 1033-1042.	5.5	50
86	Solitary (juvenile) xanthogranuloma: a comprehensive immunohistochemical study emphasizing recently developed markers of histiocytic lineage. Human Pathology, 2015, 46, 1390-1397.	2.0	41
87	Intrathoracic peripheral nerve sheath tumorsâ€”a clinicopathological study of 75 cases. Human Pathology, 2015, 46, 419-425.	2.0	45
88	A Novel Chromogenic In Situ Hybridization Assay for FGF23 mRNA in Phosphaturic Mesenchymal Tumors. American Journal of Surgical Pathology, 2015, 39, 75-83.	3.7	61
89	Identification of a novelFN1-FGFR1genetic fusion as a frequent event in phosphaturic mesenchymal tumour. Journal of Pathology, 2015, 235, 539-545.	4.5	120
90	ERG expression in chondrogenic bone and soft tissue tumours. Journal of Clinical Pathology, 2015, 68, 125-129.	2.0	44

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91	Tumor-Induced Osteomalacia. Translational Endocrinology & Metabolism, 2015, 7, .	0.2	12
92	MYC amplification and overexpression in primary cutaneous angiosarcoma: a fluorescence in-situ hybridization and immunohistochemical study. Modern Pathology, 2014, 27, 509-515.	5.5	116
93	SMARCB1 deletion by a complex three-way chromosomal translocation in an extrarenal malignant rhabdoid tumor. Cancer Genetics, 2014, 207, 437-440.	0.4	4
94	TGFBR3 and MGEA5 Rearrangements in Pleomorphic Hyalinizing Angiectatic Tumors and the Spectrum of Related Neoplasms. American Journal of Surgical Pathology, 2014, 38, 1182-1992.	3.7	74
95	Myofibromas With Atypical Features. American Journal of Surgical Pathology, 2014, 38, 1649-1654.	3.7	36
96	Malignant Melanotic Schwannian Tumor. American Journal of Surgical Pathology, 2014, 38, 94-105.	3.7	169
97	Fibrosarcoma: a review and update. Histopathology, 2014, 64, 12-25.	2.9	100
98	Intraneural fibroma of the median nerve at the wrist. Journal of Clinical Neuroscience, 2014, 21, 1054-1056.	1.5	1
99	Selected topics in the pathology of epithelioid soft tissue tumors. Modern Pathology, 2014, 27, S64-S79.	5.5	40
100	Diagnostic utility of SOX10 to distinguish malignant peripheral nerve sheath tumor from synovial sarcoma, including intraneural synovial sarcoma. Modern Pathology, 2014, 27, 55-61.	5.5	79
101	Superficial CD34-positive fibroblastic tumor: report of 18 cases of a distinctive low-grade mesenchymal neoplasm of intermediate (borderline) malignancy. Modern Pathology, 2014, 27, 294-302.	5.5	82
102	Aberrant expression of neuroendocrine markers in angiosarcoma: a potential diagnostic pitfall. Human Pathology, 2014, 45, 1618-1624.	2.0	31
103	Tumorâ€Induced Osteomalacia Resulting from Primary Cutaneous Phosphaturic Mesenchymal Tumor: A Case and Review of the Medical Literature. Journal of Cutaneous Pathology, 2013, 40, 780-784.	1.3	18
104	Epithelioid sarcoma is associated with a high percentage of SMARCB1 deletions. Modern Pathology, 2013, 26, 385-392.	5.5	129
105	Cutaneous Neoplasms Showing EWSR1 Rearrangement. Advances in Anatomic Pathology, 2013, 20, 75-85.	4.3	33
106	Hemosiderotic Fibrolipomatous Tumor, Not an Entirely Benign Entity. American Journal of Surgical Pathology, 2013, 37, 1627-1630.	3.7	27
107	Myxochondroid metaplasia of the plantar foot: a distinctive pseudoneoplastic lesion resembling nuchal fibrocartilaginous pseudotumor and the equine digital cushion. Modern Pathology, 2013, 26, 1561-1567.	5.5	9
108	Benign Notochordal Cell Tumor of the Sacrum with Atypical Imaging Features: The Value of CT Guided Biopsy for Diagnosis. Open Neuroimaging Journal, 2013, 7, 36-40.	0.2	13

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109	â€œMalignantâ€•Perivascular Epithelioid Cell Neoplasm: Risk Stratification and Treatment Strategies. Sarcoma, 2012, 2012, 1-12.	1.3	150
110	Epithelioid Malignant Peripheral Nerve Sheath Tumor Arising in a Schwannoma, in a Patient With â€œNeuroblastoma-likeâ€•Schwannomatosis and a Novel Germline SMARCB1 Mutation. American Journal of Surgical Pathology, 2012, 36, 154-160.	3.7	102
111	Cellular Spindled Histiocytic Pseudotumor Complicating Mammary Fat Necrosis. American Journal of Surgical Pathology, 2012, 36, 1571-1578.	3.7	21
112	Liposarcomas of the Mediastinum and Thorax. American Journal of Surgical Pathology, 2012, 36, 1395-1403.	3.7	83
113	Melanotic Xp11.2 Neoplasm of the Ovary. American Journal of Surgical Pathology, 2012, 36, 1410-1414.	3.7	21
114	Merkel cell carcinoma with heterologous rhabdomyoblastic differentiation: the role of immunohistochemistry for Merkel cell polyomavirus large Tâ€antigen in confirmation. Journal of Cutaneous Pathology, 2012, 39, 47-51.	1.3	28
115	CD1a immunopositivity in perivascular epithelioid cell neoplasms: true expression or technical artifact? A streptavidin-biotin and polymer-based detection system immunohistochemical study of perivascular epithelioid cell neoplasms and their morphologic mimics. Human Pathology, 2011, 42, 369-374.	2.0	15
116	Intra-articular Epithelioid Sarcoma Showing Mixed Classic and Proximal-type Features. American Journal of Surgical Pathology, 2011, 35, 891-897.	3.7	25
117	Low-grade Fibromyxoid Sarcoma of the Small Intestine. American Journal of Surgical Pathology, 2011, 35, 1069-1073.	3.7	31
118	Cutaneous angiosarcoma arising in massive localized lymphedema of the morbidly obese: a report of five cases and review of the literature. Journal of Cutaneous Pathology, 2011, 38, 560-564.	1.3	68
119	Angiosarcoma: a study of 98 cases with immunohistochemical evaluation of TLE3, a recently described marker of potential taxane responsiveness. Journal of Cutaneous Pathology, 2011, 38, 961-966.	1.3	23
120	Sclerosing Epithelioid Fibrosarcomaâ€“A Report of Two Cases with Cytogenetic Analysis of FUS Gene Rearrangement by FISH Technique. Pathology and Oncology Research, 2011, 17, 145-148.	1.9	40
121	Angiomatoid fibrous histiocytoma: unusual sites and unusual morphology. Modern Pathology, 2011, 24, 1560-1570.	5.5	134
122	Ossifying Fibromyxoid Tumor of Soft Parts. American Journal of Surgical Pathology, 2011, 35, 1615-1625.	3.7	110
123	Adult-type Fibrosarcoma: A Reevaluation of 163 Putative Cases Diagnosed at a Single Institution Over a 48-year Period. American Journal of Surgical Pathology, 2010, 34, 1504-1513.	3.7	127
124	Primary Vascular Tumors and Tumor-like Lesions of the Kidney: A Clinicopathologic Analysis of 25 Cases. American Journal of Surgical Pathology, 2010, 34, 942-949.	3.7	109
125	The Impact of Advances in Molecular Genetic Pathology on the Classification, Diagnosis and Treatment of Selected Soft Tissue Tumors of the Head and Neck. Head and Neck Pathology, 2010, 4, 70-76.	2.6	25
126	Perivascular epithelioid cell neoplasms: pathology and pathogenesis. Human Pathology, 2010, 41, 1-15.	2.0	332

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127	Perivascular epithelioid cell neoplasm of the uterine cervix: an unusual tumor in an unusual location. Rare Tumors, 2010, 2, 56.	0.6	13
128	TLE1 expression is not specific for synovial sarcoma: a whole section study of 163 soft tissue and bone neoplasms. Modern Pathology, 2009, 22, 872-878.	5.5	221
129	Liposarcomas in Young Patients. American Journal of Surgical Pathology, 2009, 33, 645-658.	3.7	184
130	Primary Epithelioid Sarcoma of Bone. American Journal of Surgical Pathology, 2009, 33, 954-958.	3.7	35
131	Clusterin is Expressed in Normal Synoviocytes and in Tenosynovial Giant Cell Tumors of Localized and Diffuse Types. American Journal of Surgical Pathology, 2009, 33, 1225-1229.	3.7	67
132	INI1 and GLUT-1 Expression in Epithelioid Sarcoma and Its Cutaneous Neoplastic and Nonneoplastic Mimics. American Journal of Dermatopathology, 2009, 31, 152-156.	0.6	40
133	RT-PCR Analysis for FGF23 Using Paraffin Sections in the Diagnosis of Phosphaturic Mesenchymal Tumors With and Without Known Tumor Induced Osteomalacia. American Journal of Surgical Pathology, 2009, 33, 1348-1354.	3.7	100
134	Spindle Epithelial Tumor With Thymus-like Differentiation: A Morphologic, Immunohistochemical, and Molecular Genetic Study of 11 Cases. American Journal of Surgical Pathology, 2009, 33, 1179-1186.	3.7	61
135	Aberrant expression of epithelial and neuroendocrine markers in alveolar rhabdomyosarcoma: a potentially serious diagnostic pitfall. Modern Pathology, 2008, 21, 795-806.	5.5	156
136	GLUT-1 expression in mesenchymal tumors: an immunohistochemical study of 247 soft tissue and bone neoplasms. Human Pathology, 2008, 39, 1519-1526.	2.0	72
137	Dermatofibrosarcoma Protuberans Presenting as a Subcutaneous Mass: A Clinicopathological Study of 15 Cases With Exclusive or Near-Exclusive Subcutaneous Involvement. American Journal of Dermatopathology, 2008, 30, 327-332.	0.6	50
138	The Utility of Fluorescence In Situ Hybridization (FISH) in the Diagnosis of Myxoid Soft Tissue Neoplasms. American Journal of Surgical Pathology, 2008, 32, 8-13.	3.7	119
139	Cutaneous CD30-Positive Epithelioid Angiosarcoma Following Breast-Conserving Therapy and Irradiation: A Potential Diagnostic Pitfall. American Journal of Dermatopathology, 2008, 30, 370-372.	0.6	30
140	Sporadic Cutaneous Angiosarcomas: A Proposal for Risk Stratification Based on 69 Cases. American Journal of Surgical Pathology, 2008, 32, 72-77.	3.7	109
141	Activation of the mTOR pathway in sporadic angiomyolipomas and other perivascular epithelioid cell neoplasms. Human Pathology, 2007, 38, 1361-1371.	2.0	213
142	Tenosynovial giant cell tumor and pigmented villonodular synovitis. Skeletal Radiology, 2007, 36, 899-900.	2.0	2
143	Best Practices in Diagnostic Immunohistochemistry: Pleomorphic Cutaneous Spindle Cell Tumors. Archives of Pathology and Laboratory Medicine, 2007, 131, 1517-1524.	2.5	59
144	Perivascular Epithelioid Cell Neoplasms of Soft Tissue and Gynecologic Origin. American Journal of Surgical Pathology, 2005, 29, 1558-1575.	3.7	820

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145	Latency-associated nuclear antigen expression and human herpesvirus-8 polymerase chain reaction in the evaluation of Kaposi sarcoma and other vascular tumors in HIV-positive patients. Modern Pathology, 2005, 18, 463-468.	5.5	60
146	Morphologic and Immunophenotypic Diversity in Ewing Family Tumors. American Journal of Surgical Pathology, 2005, 29, 1025-1033.	3.7	376
147	Morphologic and immunophenotypic diversity in Ewing family tumors: a study of 66 genetically confirmed cases. American Journal of Surgical Pathology, 2005, 29, 1025-33.	3.7	267
148	Cutaneous Angiosarcoma Following Breast-conserving Surgery and Radiation: An Analysis of 27 Cases. American Journal of Surgical Pathology, 2004, 28, 781-788.	3.7	216
149	Most Osteomalacia-associated Mesenchymal Tumors Are a Single Histopathologic Entity. American Journal of Surgical Pathology, 2004, 28, 1-30.	3.7	587
150	Pleomorphic Hyalinizing Angiectatic Tumor. American Journal of Surgical Pathology, 2004, 28, 1417-1425.	3.7	148
151	Cutaneous and Subcutaneous Fibrohistiocytic Tumors of Intermediate Malignancy. American Journal of Dermatopathology, 2004, 26, 141-155.	0.6	92
152	Kaposiform Hemangioendothelioma. American Journal of Surgical Pathology, 2004, 28, 559-568.	3.7	361
153	Ossifying Fibromyxoid Tumor of Soft Parts. American Journal of Surgical Pathology, 2003, 27, 421-431.	3.7	204
154	Epithelioid Sarcoma-Like Hemangioendothelioma. American Journal of Surgical Pathology, 2003, 27, 48-57.	3.7	209
155	MyoD1 and Myogenin Expression in Human Neoplasia: A Review and Update. Advances in Anatomic Pathology, 2002, 9, 198-203.	4.3	76
156	Expression of Claudin-1, a Recently Described Tight Junction-Associated Protein, Distinguishes Soft Tissue Perineurioma From Potential Mimics. American Journal of Surgical Pathology, 2002, 26, 1620-1626.	3.7	188
157	Sclerosing Rhabdomyosarcoma in Adults. American Journal of Surgical Pathology, 2002, 26, 1175-1183.	3.7	164
158	Lipoleiomyosarcoma (Well-Differentiated Liposarcoma With Leiomyosarcomatous Differentiation). American Journal of Surgical Pathology, 2002, 26, 742-749.	3.7	77
159	Clear Cell Myomelanocytic Tumor of the Thigh. American Journal of Surgical Pathology, 2002, 26, 809-812.	3.7	81
160	Microphthalmia Transcription Factor and Melanoma Cell Adhesion Molecule Expression Distinguish Desmoplastic/Spindle Cell Melanoma From Morphologic Mimics. American Journal of Surgical Pathology, 2001, 25, 58-64.	3.7	92
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