Andrew L Folpe

List of Publications by Year in Descending Order

170 papers	10,351 citations	54 h-index	99 g-index
173 ext. papers	11,900 ext. citations	5.6 avg, IF	6.37 L-index

#	Paper	IF	Citations
170	ΦCanΦ Keep Up!ΦAn 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	7.3	0
169	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	2.2	0
168	Glomangiomatosis of the Upper Extremity. *Journal of Hand Surgery*, **2021**, 46, 716.e1-716.e3	2.6	1
167	Recurrent novel HMGA2-NCOR2 fusions characterize a subset of keratin-positive giant cell-rich soft tissue tumors. *Modern Pathology*, **2021**, 34, 1507-1520	9.8	5
166	Inflammatory rhabdomyoblastic tumor with progression to high-grade rhabdomyosarcoma. *Modern Pathology*, **2021**, 34, 1035-1036	9.8	3
165	NUTM1-rearranged colorectal sarcoma: a clinicopathologically and genetically distinctive malignant neoplasm with a poor prognosis. *Modern Pathology*, **2021**, 34, 1547-1557	9.8	5
164	CD10 (neprilysin) expression: a potential adjunct in the distinction of hibernoma from morphologic mimics. *Human Pathology*, **2021**, 110, 12-19	3.7	1
163	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	9.8	0
162	Myxoid pleomorphic liposarcoma-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	9.8	2
161	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	9.8	7
160	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	2.6	2
159	Primary intra-abdominal melanoma arising in association with extracutaneous blue naevus: a report of two cases. *Histopathology*, **2021**, 78, 281-289	7.3	0
158	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	9.8	16
157	Lipoblastomas presenting in older children and adults: analysis of 22 cases with identification of novel PLAG1 fusion partners. *Modern Pathology*, **2021**, 34, 584-591	9.8	11
156	Update on SWI/SNF-related gynecologic mesenchymal neoplasms: SMARCA4-deficient uterine sarcoma and SMARCB1-deficient vulvar neoplasms. *Genes Chromosomes and Cancer*, **2021**, 60, 190-209	5	5
155	"Inflammatory Leiomyosarcoma" and "Histiocyte-rich Rhabdomyoblastic Tumor": a clinicopathological, immunohistochemical and genetic study of 13 cases, with a proposal for reclassification as "Inflammatory Rhabdomyoblastic Tumor". *Modern Pathology*, **2021**, 34, 758-769	9.8	10
154	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*, **2021**, 1	5.1	0

153	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	3.7	2
152	Head and Neck Low-Grade Fibromyxoid Sarcoma: A Clinicopathologic Study of 15 Cases. *Head and Neck Pathology*, **2021**, 1	3.3	1
151	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	2.8	2
150	PRRX1-NCOA1-rearranged fibroblastic tumour: a clinicopathological, immunohistochemical and molecular genetic study of six cases of a potentially under-recognised, distinctive mesenchymal tumour. *Histopathology*, **2021**, 79, 997-1003	7.3	4
149	Eccrine angiomatous hamartoma: First case in the cytology literature. *Annals of Diagnostic Pathology*, **2021**, 54, 151796	2.2	
148	Well-Differentiated/Dedifferentiated Liposarcoma Arising in the Upper Aerodigestive Tract: 8 Cases Mimicking Non-adipocytic Lesions. *Head and Neck Pathology*, **2020**, 14, 974-981	3.3	9
147	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	9.8	4
146	Colonic Angiosarcoma Arising in Association with Amyloid Deposits. *Case Reports in Gastrointestinal Medicine*, **2020**, 2020, 3780763	0.6	2
145	MyoD1 expression in fibroepithelial stromal polyps. *Human Pathology*, **2020**, 99, 75-79	3.7	2
144	Juvenile Hyaline Fibromatosis. *Mayo Clinic Proceedings*, **2020**, 95, 328-329	6.4	3
143	Head and Neck Mesenchymal Neoplasms With GLI1 Gene Alterations: A Pathologic Entity With Distinct Histologic Features and Potential for Distant Metastasis. *American Journal of Surgical Pathology*, **2020**, 44, 729-737	6.7	17
142	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	5.1	9
141	Contemporary approaches to soft tissue and bone pathology. *Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin*, **2020**, 476, 1-2	5.1	
140	Frequent overexpression of klotho in fusion-negative phosphaturic mesenchymal tumors with tumorigenic implications. *Modern Pathology*, **2020**, 33, 858-870	9.8	7
139	Lymphatic-type "Angiosarcoma" With Prominent Lymphocytic Infiltrate. *American Journal of Surgical Pathology*, **2020**, 44, 271-279	6.7	5
138	Paraspinal pseudoneoplasms: a series of 58 consultation cases emphasizing the importance of pathology-radiology correlation. *Human Pathology*, **2020**, 103, 14-24	3.7	0
137	"Hey! Whatever happened to hemangiopericytoma and fibrosarcoma?" An update on selected conceptual advances in soft tissue pathology which have occurred over the past 50 years. *Human Pathology*, **2020**, 95, 113-136	3.7	9
136	Extraneural perineurioma: CT and MRI imaging characteristics. *Skeletal Radiology*, **2020**, 49, 109-114	2.7	3

135	PIK3CA mutations in lipomatosis of nerve with or without nerve territory overgrowth. *Modern Pathology*, **2020**, 33, 420-430	9.8	17
134	Loss of succinate dehydrogenase B immunohistochemical expression distinguishes pulmonary chondromas from hamartomas. *Histopathology*, **2019**, 75, 825-832	7.3	4
133	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	3.7	22
132	Perinephric myxoid pseudotumor of fat: a distinctive pseudoneoplasm most often associated with non-neoplastic renal disease. *Human Pathology*, **2019**, 87, 37-43	3.7	5
131	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	3.7	1
130	Tenosynovitis With Psammomatous Calcifications: A Distinctive Trauma-Associated Subtype of Idiopathic Calcifying Tenosynovitis With a Predilection for the Distal extremities of Middle-Aged Women-A Report of 23 Cases. *American Journal of Surgical Pathology*, **2019**, 43, 261-267	6.7	1
129	Imaging features of phosphaturic mesenchymal tumors. *Skeletal Radiology*, **2019**, 48, 119-127	2.7	11
128	Spindle Epithelial Tumor with Thymus-Like Differentiation (SETTLE): A Next-Generation Sequencing Study. *Head and Neck Pathology*, **2019**, 13, 162-168	3.3	5
127	Hepatic Rearranged Epithelioid Hemangioendothelioma. *Case Reports in Gastrointestinal Medicine*, **2019**, 2019, 7530845	0.6	3
126	OLIG2 is a marker of the fusion protein-driven neurodevelopmental transcriptional signature in alveolar rhabdomyosarcoma. *Human Pathology*, **2019**, 91, 77-85	3.7	11
125	Phosphaturic mesenchymal tumors: A review and update. *Seminars in Diagnostic Pathology*, **2019**, 36, 260-268	4.3	40
124	Atypical lipomatous tumour/well-differentiated liposarcoma and de-differentiated liposarcoma in patients aged ⩾40 years: a study of 116 patients. *Histopathology*, **2019**, 75, 833-842	7.3	12
123	Malignant Tenosynovial Giant Cell Tumor: The True "Synovial Sarcoma?" A Clinicopathologic, Immunohistochemical, and Molecular Cytogenetic Study of 10 Cases, Supporting Origin from Synoviocytes. *Modern Pathology*, **2019**, 32, 242-251	9.8	18
122	Ancillary Diagnostic Tests in the Diagnosis of Cutaneous Soft Tissue Neoplasms **2019**, 15-56		
121	Aberrant receptor tyrosine kinase signaling in lipofibromatosis: a clinicopathological and molecular genetic study of 20 cases. *Modern Pathology*, **2019**, 32, 423-434	9.8	29
120	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	9.8	14
119	Phosphaturic mesenchymal tumor without osteomalacia: additional confirmation of the "nonphosphaturic" 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	3.7	11
118	Mediastinal Synovial Sarcoma: Clinicopathologic Analysis of 21 Cases With Molecular Confirmation. *American Journal of Surgical Pathology*, **2018**, 42, 761-766	6.7	11

117	Low-grade fibromyxoid sarcoma arising within the median nerve. *Neuropathology*, **2018**, 38, 309-314	2	3
116	Recurrent GNA14 mutations in anastomosing haemangiomas. *Histopathology*, **2018**, 73, 354-357	7.3	20
115	Abdominopelvic and Retroperitoneal Low-Grade Fibromyxoid Sarcoma: A Clinicopathologic Study of 13 Cases. *American Journal of Clinical Pathology*, **2018**, 149, 128-134	1.9	12
114	Mesenchymal chondrosarcomas showing immunohistochemical evidence of rhabdomyoblastic differentiation: a potential diagnostic pitfall. *Human Pathology*, **2018**, 77, 28-34	3.7	21
113	Malignant Peripheral Nerve Sheath Tumor in a Patient With BAP1 Tumor Predisposition Syndrome. *World Neurosurgery*, **2018**, 109, 362-364	2.1	2
112	Polypoid fibroadipose tumors of the esophagus: Giant fibrovascular polyp or liposarcoma? A clinicopathological and molecular cytogenetic study of 13 cases. *Modern Pathology*, **2018**, 31, 337-342	9.8	18
111	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	9.8	13
110	Clinicopathologic features and outcomes of gastrointestinal stromal tumors arising from the esophagus and gastroesophageal junction. *Journal of Gastrointestinal Oncology*, **2018**, 9, 718-727	2.8	3
109	"Chondroblastoma-like" epithelioid fibrous histiocytoma: A previously undescribed and potentially confusing variant. *Journal of Cutaneous Pathology*, **2018**, 45, 99-103	1.7	4
108	Radiation Therapy for Retroperitoneal Sarcomas: Influences of Histology, Grade, and Size. *Sarcoma*, **2018**, 2018, 7972389	3.1	9
107	Spindle cell rhabdomyosarcoma of bone with FUS-TFCP2 fusion: confirmation of a very recently described rhabdomyosarcoma subtype. *Histopathology*, **2018**, 73, 514-520	7.3	34
106	Hypoxia-related microRNA-210 is a diagnostic marker for discriminating osteoblastoma and osteosarcoma. *Journal of Orthopaedic Research*, **2017**, 35, 1137-1146	3.8	8
105	Fibrous hamartoma of infancy: a clinicopathologic study of 145 cases, including 2 with sarcomatous features. *Modern Pathology*, **2017**, 30, 474-485	9.8	36
104	Recurrent GNAQ mutations in anastomosing hemangiomas. *Modern Pathology*, **2017**, 30, 722-727	9.8	36
103	Hemosiderotic Fibrolipomatous Tumor, Pleomorphic Hyalinizing Angiectatic Tumor, and Myxoinflammatory Fibroblastic Sarcoma: Related or Not?. *Advances in Anatomic Pathology*, **2017**, 24, 268-277	5.1	27
102	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	6.7	27
101	Gastroblastoma harbors a recurrent somatic MALAT1-GLI1 fusion gene. *Modern Pathology*, **2017**, 30, 1443-1452	9.8	49
100	Composite hemangioendothelioma with neuroendocrine marker expression: an aggressive variant. *Modern Pathology*, **2017**, 30, 1589-1602	9.8	19

99	Comparision of New Diagnostic Tools for Malignant Peripheral Nerve Sheath Tumors. *Pathology and Oncology Research*, **2017**, 23, 393-398	2.6	4
98	Primary angiomatoid fibrous histiocytoma of the lung with mediastinal lymph node metastasis. *Human Pathology*, **2016**, 58, 134-137	3.7	11
97	Characterization of FN1-FGFR1 and novel FN1-FGF1 fusion genes in a large series of phosphaturic mesenchymal tumors. *Modern Pathology*, **2016**, 29, 1335-1346	9.8	95
96	Renal Leiomyoma and Leiomyosarcoma: A Study of 57 Cases. *American Journal of Surgical Pathology*, **2016**, 40, 1557-1563	6.7	10
95	Oncocytic variant of malignant gastrointestinal neuroectodermal tumor: a potential diagnostic pitfall. *Human Pathology*, **2016**, 57, 13-16	3.7	19
94	Anastomosing Hemangiomas Arising in Unusual Locations: A Clinicopathologic Study of 17 Soft Tissue Cases Showing a Predilection for the Paraspinal Region. *American Journal of Surgical Pathology*, **2016**, 40, 1084-9	6.7	31
93	Pseudolipoblastic perineurioma: an unusual morphological variant of perineurioma that may simulate liposarcoma. *Human Pathology*, **2016**, 57, 22-27	3.7	4
92	TGFBR3 and MGEA5 rearrangements are much more common in "hybrid" hemosiderotic fibrolipomatous tumor-myxoinflammatory fibroblastic sarcomas than in classical myxoinflammatory fibroblastic sarcomas: a morphological and fluorescence in situ hybridization	3.7	28
91	Voluntary Second Opinions in Pediatric Bone and Soft Tissue Pathology: A Retrospective Review of 1601 Cases From a Single Mesenchymal Tumor Consultation Service. *International Journal of Surgical Pathology*, **2016**, 24, 685-691	1.2	8
90	Solitary (juvenile) xanthogranuloma: a comprehensive immunohistochemical study emphasizing recently developed markers of histiocytic lineage. *Human Pathology*, **2015**, 46, 1390-7	3.7	36
89	Intrathoracic peripheral nerve sheath tumors-a clinicopathological study of 75 cases. *Human Pathology*, **2015**, 46, 419-25	3.7	40
88	A novel chromogenic in situ hybridization assay for FGF23 mRNA in phosphaturic mesenchymal tumors. *American Journal of Surgical Pathology*, **2015**, 39, 75-83	6.7	51
87	Identification of a novel FN1-FGFR1 genetic fusion as a frequent event in phosphaturic mesenchymal tumour. *Journal of Pathology*, **2015**, 235, 539-45	9.4	90
86	ERG expression in chondrogenic bone and soft tissue tumours. *Journal of Clinical Pathology*, **2015**, 68, 125-9	3.9	29
85	SMARCB1-deficient Vulvar Neoplasms: A Clinicopathologic, Immunohistochemical, and Molecular Genetic Study of 14 Cases. *American Journal of Surgical Pathology*, **2015**, 39, 836-49	6.7	34
84	Aberrant intermediate filament and synaptophysin expression is a frequent event in malignant melanoma: an immunohistochemical study of 73 cases. *Modern Pathology*, **2015**, 28, 1033-42	9.8	40
83	Tumor-Induced Osteomalacia. *Translational Endocrinology & Metabolism*, **2015**, 7,		11
82	Selected topics in the pathology of epithelioid soft tissue tumors. *Modern Pathology*, **2014**, 27 Suppl 1, S64-79	9.8	32

81	Diagnostic utility of SOX10 to distinguish malignant peripheral nerve sheath tumor from synovial sarcoma, including intraneural synovial sarcoma. *Modern Pathology*, **2014**, 27, 55-61	9.8	62
80	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	9.8	53
79	Aberrant expression of neuroendocrine markers in angiosarcoma: a potential diagnostic pitfall. *Human Pathology*, **2014**, 45, 1618-24	3.7	26
78	MYC amplification and overexpression in primary cutaneous angiosarcoma: a fluorescence in-situ hybridization and immunohistochemical study. *Modern Pathology*, **2014**, 27, 509-15	9.8	91
77	SMARCB1 deletion by a complex three-way chromosomal translocation in an extrarenal malignant rhabdoid tumor. *Cancer Genetics*, **2014**, 207, 437-40	2.3	2
76	TGFBR3 and MGEA5 rearrangements in pleomorphic hyalinizing angiectatic tumors and the spectrum of related neoplasms. *American Journal of Surgical Pathology*, **2014**, 38, 1182-992	6.7	56
75	Myofibromas with atypical features: expanding the morphologic spectrum of a benign entity. *American Journal of Surgical Pathology*, **2014**, 38, 1649-54	6.7	27
74	Malignant melanotic schwannian tumor: a clinicopathologic, immunohistochemical, and gene expression profiling study of 40 cases, with a proposal for the reclassification of "melanotic schwannoma". *American Journal of Surgical Pathology*, **2014**, 38, 94-105	6.7	121
73	Fibrosarcoma: a review and update. *Histopathology*, **2014**, 64, 12-25	7.3	61
72	Intraneural fibroma of the median nerve at the wrist. *Journal of Clinical Neuroscience*, **2014**, 21, 1054-6	2.2	1
71	Tumor-induced osteomalacia resulting from primary cutaneous phosphaturic mesenchymal tumor: a case and review of the medical literature. *Journal of Cutaneous Pathology*, **2013**, 40, 780-4; quiz 779	1.7	14
70	Epithelioid sarcoma is associated with a high percentage of SMARCB1 deletions. *Modern Pathology*, **2013**, 26, 385-92	9.8	96
69	Cutaneous neoplasms showing EWSR1 rearrangement. *Advances in Anatomic Pathology*, **2013**, 20, 75-85	5.1	27
68	Hemosiderotic fibrolipomatous tumor, not an entirely benign entity. *American Journal of Surgical Pathology*, **2013**, 37, 1627-30	6.7	21
67	Myxochondroid metaplasia of the plantar foot: a distinctive pseudoneoplastic lesion resembling nuchal fibrocartilaginous pseudotumor and the equine digital cushion. *Modern Pathology*, **2013**, 26, 1561-7	9.8	6
66	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	0.1	10
65	Merkel cell carcinoma with heterologous rhabdomyoblastic differentiation: the role of immunohistochemistry for Merkel cell polyomavirus large T-antigen in confirmation. *Journal of Cutaneous Pathology*, **2012**, 39, 47-51	1.7	21
64	"Malignant" perivascular epithelioid cell neoplasm: risk stratification and treatment strategies. *Sarcoma*, **2012**, 2012, 541626	3.1	111

63	Epithelioid malignant peripheral nerve sheath tumor arising in a schwannoma, in a patient with "neuroblastoma-like" schwannomatosis and a novel germline SMARCB1 mutation. *American Journal of Surgical Pathology*, **2012**, 36, 154-60	6.7	85
62	Cellular spindled histiocytic pseudotumor complicating mammary fat necrosis: a potential diagnostic pitfall. *American Journal of Surgical Pathology*, **2012**, 36, 1571-8	6.7	17
61	Liposarcomas of the mediastinum and thorax: a clinicopathologic and molecular cytogenetic study of 24 cases, emphasizing unusual and diverse histologic features. *American Journal of Surgical Pathology*, **2012**, 36, 1395-403	6.7	59
60	Melanotic xp11.2 neoplasm of the ovary: report of a unique case. *American Journal of Surgical Pathology*, **2012**, 36, 1410-4	6.7	20
59	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, 369-74	3.7	13
58	Intra-articular epithelioid sarcoma showing mixed classic and proximal-type features: report of 2 cases, with immunohistochemical and molecular cytogenetic INI-1 study. *American Journal of Surgical Pathology*, **2011**, 35, 891-7	6.7	21
57	Low-grade fibromyxoid sarcoma of the small intestine: report of 4 cases with molecular cytogenetic confirmation. *American Journal of Surgical Pathology*, **2011**, 35, 1069-73	6.7	22
56	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, 560-4	1.7	54
55	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, 961-6	1.7	21
54	Sclerosing epithelioid fibrosarcoma - a report of two cases with cytogenetic analysis of FUS gene rearrangement by FISH technique. *Pathology and Oncology Research*, **2011**, 17, 145-8	2.6	30
53	Angiomatoid fibrous histiocytoma: unusual sites and unusual morphology. *Modern Pathology*, **2011**, 24, 1560-70	9.8	103
52	Ossifying fibromyxoid tumor of soft parts: a clinicopathologic, proteomic, and genomic study. *American Journal of Surgical Pathology*, **2011**, 35, 1615-25	6.7	92
51	Perivascular epithelioid cell neoplasms: pathology and pathogenesis. *Human Pathology*, **2010**, 41, 1-15	3.7	264
50	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, 1504-13	6.7	104
49	Primary vascular tumors and tumor-like lesions of the kidney: a clinicopathologic analysis of 25 cases. *American Journal of Surgical Pathology*, **2010**, 34, 942-9	6.7	93
48	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, 70-6	3.3	24
47	Perivascular epithelioid cell neoplasm of the uterine cervix: an unusual tumor in an unusual location. *Rare Tumors*, **2010**, 2, e56	1.1	10
46	TLE1 expression is not specific for synovial sarcoma: a whole section study of 163 soft tissue and bone neoplasms. *Modern Pathology*, **2009**, 22, 872-8	9.8	193

45	Liposarcomas in young patients: a study of 82 cases occurring in patients younger than 22 years of age. *American Journal of Surgical Pathology*, **2009**, 33, 645-58	6.7	140
44	Primary epithelioid sarcoma of bone: report of a unique case, with immunohistochemical and fluorescent in situ hybridization confirmation of INI1 deletion. *American Journal of Surgical Pathology*, **2009**, 33, 954-8	6.7	33
43	Clusterin is expressed in normal synoviocytes and in tenosynovial giant cell tumors of localized and diffuse types: diagnostic and histogenetic implications. *American Journal of Surgical Pathology*, **2009**, 33, 1225-9	6.7	44
42	INI1 and GLUT-1 expression in epithelioid sarcoma and its cutaneous neoplastic and nonneoplastic mimics. *American Journal of Dermatopathology*, **2009**, 31, 152-6	0.9	35
41	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, 1348-54	6.7	85
40	Spindle epithelial tumor with thymus-like differentiation: a morphologic, immunohistochemical, and molecular genetic study of 11 cases. *American Journal of Surgical Pathology*, **2009**, 33, 1179-86	6.7	54
39	Aberrant expression of epithelial and neuroendocrine markers in alveolar rhabdomyosarcoma: a potentially serious diagnostic pitfall. *Modern Pathology*, **2008**, 21, 795-806	9.8	129
38	GLUT-1 expression in mesenchymal tumors: an immunohistochemical study of 247 soft tissue and bone neoplasms. *Human Pathology*, **2008**, 39, 1519-26	3.7	60
37	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, 327-32	0.9	37
36	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	6.7	107
35	Cutaneous CD30-positive epithelioid angiosarcoma following breast-conserving therapy and irradiation: a potential diagnostic pitfall. *American Journal of Dermatopathology*, **2008**, 30, 370-2	0.9	24
34	Sporadic cutaneous angiosarcomas: a proposal for risk stratification based on 69 cases. *American Journal of Surgical Pathology*, **2008**, 32, 72-7	6.7	88
33	Tenosynovial giant cell tumor and pigmented villonodular synovitis. *Skeletal Radiology*, **2007**, 36, 899-900; author reply 901	2.7	2
32	Activation of the mTOR pathway in sporadic angiomyolipomas and other perivascular epithelioid cell neoplasms. *Human Pathology*, **2007**, 38, 1361-71	3.7	181
31	Best practices in diagnostic immunohistochemistry: pleomorphic cutaneous spindle cell tumors. *Archives of Pathology and Laboratory Medicine*, **2007**, 131, 1517-24	5	44
30	Perivascular epithelioid cell neoplasms of soft tissue and gynecologic origin: a clinicopathologic study of 26 cases and review of the literature. *American Journal of Surgical Pathology*, **2005**, 29, 1558-75	6.7	666
29	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, 463-8	9.8	48
28	Morphologic and Immunophenotypic Diversity in Ewing Family Tumors. *American Journal of Surgical Pathology*, **2005**, 29, 1025-1033	6.7	313

27	Morphologic and immunophenotypic diversity in Ewing family tumors: a study of 66 genetically confirmed cases. *American Journal of Surgical Pathology*, **2005**, 29, 1025-33	6.7	244
26	Cutaneous angiosarcoma following breast-conserving surgery and radiation: an analysis of 27 cases. *American Journal of Surgical Pathology*, **2004**, 28, 781-8	6.7	192
25	Most osteomalacia-associated mesenchymal tumors are a single histopathologic entity: an analysis of 32 cases and a comprehensive review of the literature. *American Journal of Surgical Pathology*, **2004**, 28, 1-30	6.7	498
24	Pleomorphic hyalinizing angiectatic tumor: analysis of 41 cases supporting evolution from a distinctive precursor lesion. *American Journal of Surgical Pathology*, **2004**, 28, 1417-25	6.7	128
23	Cutaneous and subcutaneous fibrohistiocytic tumors of intermediate malignancy: an update. *American Journal of Dermatopathology*, **2004**, 26, 141-55	0.9	77
22	Kaposiform hemangioendothelioma: a study of 33 cases emphasizing its pathologic, immunophenotypic, and biologic uniqueness from juvenile hemangioma. *American Journal of Surgical Pathology*, **2004**, 28, 559-68	6.7	302
21	Ossifying fibromyxoid tumor of soft parts: a clinicopathologic study of 70 cases with emphasis on atypical and malignant variants. *American Journal of Surgical Pathology*, **2003**, 27, 421-31	6.7	181
20	Epithelioid sarcoma-like hemangioendothelioma. *American Journal of Surgical Pathology*, **2003**, 27, 48-57	6.7	170
19	MyoD1 and myogenin expression in human neoplasia: a review and update. *Advances in Anatomic Pathology*, **2002**, 9, 198-203	5.1	65
18	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, 1620-6	6.7	160
17	Sclerosing rhabdomyosarcoma in adults: report of four cases of a hyalinizing, matrix-rich variant of rhabdomyosarcoma that may be confused with osteosarcoma, chondrosarcoma, or angiosarcoma. *American Journal of Surgical Pathology*, **2002**, 26, 1175-83	6.7	148
16	Lipoleiomyosarcoma (well-differentiated liposarcoma with leiomyosarcomatous differentiation): a clinicopathologic study of nine cases including one with dedifferentiation. *American Journal of Surgical Pathology*, **2002**, 26, 742-9	6.7	64
15	Clear cell myomelanocytic tumor of the thigh: report of a unique case. *American Journal of Surgical Pathology*, **2002**, 26, 809-12	6.7	67
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