Andrew L Folpe

List of Publications by Citations

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	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
169	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
168	Atypical and malignant glomus tumors: analysis of 52 cases, with a proposal for the reclassification of glomus tumors. *American Journal of Surgical Pathology*, **2001**, 25, 1-12	6.7	455
167	Morphologic and Immunophenotypic Diversity in Ewing Family Tumors. *American Journal of Surgical Pathology*, **2005**, 29, 1025-1033	6.7	313
166	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
165	Thyroid transcription factor-1 is expressed in extrapulmonary small cell carcinomas but not in other extrapulmonary neuroendocrine tumors. *Modern Pathology*, **2000**, 13, 238-42	9.8	298
164	Immunohistochemical detection of FLI-1 protein expression: a study of 132 round cell tumors with emphasis on CD99-positive mimics of Ewing© sarcoma/primitive neuroectodermal tumor. *American Journal of Surgical Pathology*, **2000**, 24, 1657-62	6.7	266
163	Perivascular epithelioid cell neoplasms: pathology and pathogenesis. *Human Pathology*, **2010**, 41, 1-15	3.7	264
162	Low-grade fibromyxoid sarcoma and hyalinizing spindle cell tumor with giant rosettes: a clinicopathologic study of 73 cases supporting their identity and assessing the impact of high-grade areas. *American Journal of Surgical Pathology*, **2000**, 24, 1353-60	6.7	257
161	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
160	Expression of Fli-1, a nuclear transcription factor, distinguishes vascular neoplasms from potential mimics. *American Journal of Surgical Pathology*, **2001**, 25, 1061-6	6.7	243
159	Clear cell myomelanocytic tumor of the falciform ligament/ligamentum teres: a novel member of the perivascular epithelioid clear cell family of tumors with a predilection for children and young adults. *American Journal of Surgical Pathology*, **2000**, 24, 1239-46	6.7	224
158	Poorly differentiated synovial sarcoma: immunohistochemical distinction from primitive neuroectodermal tumors and high-grade malignant peripheral nerve sheath tumors. *American Journal of Surgical Pathology*, **1998**, 22, 673-82	6.7	205
157	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
156	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
155	Activation of the mTOR pathway in sporadic angiomyolipomas and other perivascular epithelioid cell neoplasms. *Human Pathology*, **2007**, 38, 1361-71	3.7	181
154	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

153	Epithelioid sarcoma-like hemangioendothelioma. *American Journal of Surgical Pathology*, **2003**, 27, 48-57	6.7	170
152	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
151	Vascular endothelial growth factor receptor-3 (VEGFR-3): a marker of vascular tumors with presumed lymphatic differentiation, including Kaposi© sarcoma, kaposiform and Dabska-type hemangioendotheliomas, and a subset of angiosarcomas. *Modern Pathology*, **2000**, 13, 180-5	9.8	156
150	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
149	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
148	CD31 expression in intratumoral macrophages: a potential diagnostic pitfall. *American Journal of Surgical Pathology*, **2001**, 25, 1167-73	6.7	137
147	Aberrant expression of epithelial and neuroendocrine markers in alveolar rhabdomyosarcoma: a potentially serious diagnostic pitfall. *Modern Pathology*, **2008**, 21, 795-806	9.8	129
146	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
145	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
144	"Malignant" perivascular epithelioid cell neoplasm: risk stratification and treatment strategies. *Sarcoma*, **2012**, 2012, 541626	3.1	111
143	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
142	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
141	Angiomatoid fibrous histiocytoma: unusual sites and unusual morphology. *Modern Pathology*, **2011**, 24, 1560-70	9.8	103
140	Epithelioid sarcoma is associated with a high percentage of SMARCB1 deletions. *Modern Pathology*, **2013**, 26, 385-92	9.8	96
139	Cytokeratin 8 immunostaining pattern and E-cadherin expression distinguish lobular from ductal breast carcinoma. *American Journal of Clinical Pathology*, **2000**, 114, 190-6	1.9	96
138	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
137	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
136	Ossifying fibromyxoid tumor of soft parts: a clinicopathologic, proteomic, and genomic study. *American Journal of Surgical Pathology*, **2011**, 35, 1615-25	6.7	92

135	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
134	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
133	Sporadic cutaneous angiosarcomas: a proposal for risk stratification based on 69 cases. *American Journal of Surgical Pathology*, **2008**, 32, 72-7	6.7	88
132	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
131	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
130	Consultative (expert) second opinions in soft tissue pathology. Analysis of problem-prone diagnostic situations. *American Journal of Clinical Pathology*, **2001**, 116, 473-6	1.9	83
129	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	6.7	83
128	Immunohistochemical study of microphthalmia transcription factor and tyrosinase in angiomyolipoma of the kidney, renal cell carcinoma, and renal and retroperitoneal sarcomas: comparative evaluation with traditional diagnostic markers. *American Journal of Surgical Pathology*, **2001**, 25, 65-70	6.7	81
127	Cutaneous and subcutaneous fibrohistiocytic tumors of intermediate malignancy: an update. *American Journal of Dermatopathology*, **2004**, 26, 141-55	0.9	77
126	Lipomatous hemangiopericytoma: a rare variant of hemangiopericytoma that may be confused with liposarcoma. *American Journal of Surgical Pathology*, **1999**, 23, 1201-7	6.7	73
125	Clear cell myomelanocytic tumor of the thigh: report of a unique case. *American Journal of Surgical Pathology*, **2002**, 26, 809-12	6.7	67
124	MyoD1 and myogenin expression in human neoplasia: a review and update. *Advances in Anatomic Pathology*, **2002**, 9, 198-203	5.1	65
123	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
122	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
121	Fibrosarcoma: a review and update. *Histopathology*, **2014**, 64, 12-25	7.3	61
120	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
119	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
118	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

117	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
116	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
115	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
114	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
113	Gastroblastoma harbors a recurrent somatic MALAT1-GLI1 fusion gene. *Modern Pathology*, **2017**, 30, 1443-1452	9.8	49
112	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
111	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
110	Best practices in diagnostic immunohistochemistry: pleomorphic cutaneous spindle cell tumors. *Archives of Pathology and Laboratory Medicine*, **2007**, 131, 1517-24	5	44
109	Intrathoracic peripheral nerve sheath tumors-a clinicopathological study of 75 cases. *Human Pathology*, **2015**, 46, 419-25	3.7	40
108	Phosphaturic mesenchymal tumors: A review and update. *Seminars in Diagnostic Pathology*, **2019**, 36, 260-268	4.3	40
107	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
106	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
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	Solitary (juvenile) xanthogranuloma: a comprehensive immunohistochemical study emphasizing recently developed markers of histiocytic lineage. *Human Pathology*, **2015**, 46, 1390-7	3.7	36
102	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
101	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
100	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

99	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
98	Selected topics in the pathology of epithelioid soft tissue tumors. *Modern Pathology*, **2014**, 27 Suppl 1, S64-79	9.8	32
97	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
96	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
95	ERG expression in chondrogenic bone and soft tissue tumours. *Journal of Clinical Pathology*, **2015**, 68, 125-9	3.9	29
94	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
93	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 study. *Human Pathology*, **2016**, 53, 14-24	3.7	28
92	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
91	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
90	Myofibromas with atypical features: expanding the morphologic spectrum of a benign entity. *American Journal of Surgical Pathology*, **2014**, 38, 1649-54	6.7	27
89	Cutaneous neoplasms showing EWSR1 rearrangement. *Advances in Anatomic Pathology*, **2013**, 20, 75-85	5.1	27
88	Aberrant expression of neuroendocrine markers in angiosarcoma: a potential diagnostic pitfall. *Human Pathology*, **2014**, 45, 1618-24	3.7	26
87	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
86	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
85	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
84	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
83	Mesenchymal chondrosarcomas showing immunohistochemical evidence of rhabdomyoblastic differentiation: a potential diagnostic pitfall. *Human Pathology*, **2018**, 77, 28-34	3.7	21
82	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

81	Hemosiderotic fibrolipomatous tumor, not an entirely benign entity. *American Journal of Surgical Pathology*, **2013**, 37, 1627-30	6.7	21
80	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
79	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
78	Recurrent GNA14 mutations in anastomosing haemangiomas. *Histopathology*, **2018**, 73, 354-357	7.3	20
77	Melanotic xp11.2 neoplasm of the ovary: report of a unique case. *American Journal of Surgical Pathology*, **2012**, 36, 1410-4	6.7	20
76	Oncocytic variant of malignant gastrointestinal neuroectodermal tumor: a potential diagnostic pitfall. *Human Pathology*, **2016**, 57, 13-16	3.7	19
75	Composite hemangioendothelioma with neuroendocrine marker expression: an aggressive variant. *Modern Pathology*, **2017**, 30, 1589-1602	9.8	19
74	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
73	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
72	Cellular spindled histiocytic pseudotumor complicating mammary fat necrosis: a potential diagnostic pitfall. *American Journal of Surgical Pathology*, **2012**, 36, 1571-8	6.7	17
71	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
70	PIK3CA mutations in lipomatosis of nerve with or without nerve territory overgrowth. *Modern Pathology*, **2020**, 33, 420-430	9.8	17
69	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
68	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
67	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
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	9.8	13
65	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
64	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

63	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
62	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
61	Mediastinal Synovial Sarcoma: Clinicopathologic Analysis of 21 Cases With Molecular Confirmation. *American Journal of Surgical Pathology*, **2018**, 42, 761-766	6.7	11
60	Primary angiomatoid fibrous histiocytoma of the lung with mediastinal lymph node metastasis. *Human Pathology*, **2016**, 58, 134-137	3.7	11
59	Imaging features of phosphaturic mesenchymal tumors. *Skeletal Radiology*, **2019**, 48, 119-127	2.7	11
58	OLIG2 is a marker of the fusion protein-driven neurodevelopmental transcriptional signature in alveolar rhabdomyosarcoma. *Human Pathology*, **2019**, 91, 77-85	3.7	11
57	Tumor-Induced Osteomalacia. *Translational Endocrinology & Metabolism*, **2015**, 7,		11
56	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
55	Renal Leiomyoma and Leiomyosarcoma: A Study of 57 Cases. *American Journal of Surgical Pathology*, **2016**, 40, 1557-1563	6.7	10
54	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
53	Perivascular epithelioid cell neoplasm of the uterine cervix: an unusual tumor in an unusual location. *Rare Tumors*, **2010**, 2, e56	1.1	10
52	"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
51	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
50	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
49	"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
48	Radiation Therapy for Retroperitoneal Sarcomas: Influences of Histology, Grade, and Size. *Sarcoma*, **2018**, 2018, 7972389	3.1	9
47	Hypoxia-related microRNA-210 is a diagnostic marker for discriminating osteoblastoma and osteosarcoma. *Journal of Orthopaedic Research*, **2017**, 35, 1137-1146	3.8	8
46	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

45	Frequent overexpression of klotho in fusion-negative phosphaturic mesenchymal tumors with tumorigenic implications. *Modern Pathology*, **2020**, 33, 858-870	9.8	7
44	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
43	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
42	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
41	Spindle Epithelial Tumor with Thymus-Like Differentiation (SETTLE): A Next-Generation Sequencing Study. *Head and Neck Pathology*, **2019**, 13, 162-168	3.3	5
40	Lymphatic-type "Angiosarcoma" With Prominent Lymphocytic Infiltrate. *American Journal of Surgical Pathology*, **2020**, 44, 271-279	6.7	5
39	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
38	NUTM1-rearranged colorectal sarcoma: a clinicopathologically and genetically distinctive malignant neoplasm with a poor prognosis. *Modern Pathology*, **2021**, 34, 1547-1557	9.8	5
37	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
36	Loss of succinate dehydrogenase B immunohistochemical expression distinguishes pulmonary chondromas from hamartomas. *Histopathology*, **2019**, 75, 825-832	7.3	4
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	9.8	4
34	Comparision of New Diagnostic Tools for Malignant Peripheral Nerve Sheath Tumors. *Pathology and Oncology Research*, **2017**, 23, 393-398	2.6	4
33	Pseudolipoblastic perineurioma: an unusual morphological variant of perineurioma that may simulate liposarcoma. *Human Pathology*, **2016**, 57, 22-27	3.7	4
32	"Chondroblastoma-like" epithelioid fibrous histiocytoma: A previously undescribed and potentially confusing variant. *Journal of Cutaneous Pathology*, **2018**, 45, 99-103	1.7	4
31	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
30	Low-grade fibromyxoid sarcoma arising within the median nerve. *Neuropathology*, **2018**, 38, 309-314	2	3
29	Hepatic Rearranged Epithelioid Hemangioendothelioma. *Case Reports in Gastrointestinal Medicine*, **2019**, 2019, 7530845	0.6	3
28	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

27	Juvenile Hyaline Fibromatosis. *Mayo Clinic Proceedings*, **2020**, 95, 328-329	6.4	3
26	Inflammatory rhabdomyoblastic tumor with progression to high-grade rhabdomyosarcoma. *Modern Pathology*, **2021**, 34, 1035-1036	9.8	3
25	Extraneural perineurioma: CT and MRI imaging characteristics. *Skeletal Radiology*, **2020**, 49, 109-114	2.7	3
24	Colonic Angiosarcoma Arising in Association with Amyloid Deposits. *Case Reports in Gastrointestinal Medicine*, **2020**, 2020, 3780763	0.6	2
23	MyoD1 expression in fibroepithelial stromal polyps. *Human Pathology*, **2020**, 99, 75-79	3.7	2
22	Malignant Peripheral Nerve Sheath Tumor in a Patient With BAP1 Tumor Predisposition Syndrome. *World Neurosurgery*, **2018**, 109, 362-364	2.1	2
21	SMARCB1 deletion by a complex three-way chromosomal translocation in an extrarenal malignant rhabdoid tumor. *Cancer Genetics*, **2014**, 207, 437-40	2.3	2
20	Tenosynovial giant cell tumor and pigmented villonodular synovitis. *Skeletal Radiology*, **2007**, 36, 899-900; author reply 901	2.7	2
19	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
18	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
17	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
16	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
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