David M Parham

List of Publications by Year in Descending Order

270 papers	12,861 citations	64 h-index	104 g-index
310 ext. papers	14,240 ext. citations	4 avg, IF	5.82 L-index

#	Paper	IF	Citations
270	Pediatric Sarcomas: The Next Generation of Molecular Studies. *Cancers*, **2022**, 14, 2515	6.6	
269	Rhabdomyosarcoma: How Advanced Molecular Methods Are Shaping the Diagnostic and Therapeutic Paradigm. *Pediatric and Developmental Pathology*, **2021**, 24, 395-404	2.2	2
268	The Relevance of and Surgical Approach to the Suprahyoid Region in Thyroglossal Duct Surgery. *Laryngoscope*, **2021**, 131, 553-558	3.6	0
267	The prognostic significance of anaplasia in childhood rhabdomyosarcoma: A report from the Children's Oncology Group. *European Journal of Cancer*, **2021**, 143, 127-133	7.5	2
266	Alveolar rhabdomyosarcoma with regional nodal involvement: Results of a combined analysis from two cooperative groups. *Pediatric Blood and Cancer*, **2021**, 68, e28832	3	3
265	Histopathologic features of alveolar capillary dysplasia with misalignment of pulmonary veins with atypical clinical presentation. *Cardiovascular Pathology*, **2021**, 50, 107289	3.8	3
264	Epithelioid Sarcoma Arising in a Long-Term Survivor of an Atypical Teratoid/Rhabdoid Tumor in a Patient With Rhabdoid Tumor Predisposition Syndrome. *Pediatric and Developmental Pathology*, **2021**, 24, 164-168	2.2	0
263	Rhabdomyosarcoma: From Obscurity to Clarity in Diagnosis But With Ongoing Challenges in Management: The Farber-Landing Lecture of 2020. *Pediatric and Developmental Pathology*, **2021**, 24, 87-95	2.2	1
262	Clinicopathologic Characteristics of Late Acute Antibody-mediated Rejection in Pediatric Liver Transplantation. *Transplantation*, **2021**, 105, 2045-2053	1.8	6
261	A History of Board Certification in Pediatric Pathology on Its 30th Anniversary. *Pediatric and Developmental Pathology*, **2021**, 24, 3-9	2.2	
260	A risk-based treatment strategy for non-rhabdomyosarcoma soft-tissue sarcomas in patients younger than 30 years (ARST0332): a Children's Oncology Group prospective study. *Lancet Oncology, The*, **2020**, 21, 145-161	21.7	42
259	Transcriptome analysis of desmoplastic small round cell tumors identifies actionable therapeutic targets: a report from the Children's Oncology Group. *Scientific Reports*, **2020**, 10, 12318	4.9	14
258	Infantile Epithelioid Sarcoma with Genomic Segmental Amplification of as Double Minutes Plus Trisomy 2: A Case Report. *Fetal and Pediatric Pathology*, **2020**, 39, 51-61	1.7	1
257	Expanding the spectrum of dicer1-associated sarcomas. *Modern Pathology*, **2020**, 33, 164-174	9.8	30
256	Risk-based treatment for patients with first relapse or progression of rhabdomyosarcoma: A report from the Children's Oncology Group. *Cancer*, **2019**, 125, 2602-2609	6.4	11
255	Clinical features and outcomes of young patients with epithelioid sarcoma: an analysis from the Children's Oncology Group and the European paediatric soft tissue Sarcoma Study Group prospective clinical trials. *European Journal of Cancer*, **2019**, 112, 98-106	7.5	14
254	Magnetic resonance and computed tomography imaging features of epithelioid sarcoma in children and young adults with pathological and clinical correlation: a report from Children's Oncology Group study ARST0332. *Pediatric Radiology*, **2019**, 49, 922-932	2.8	7

253	Expanding the Spectrum of Pediatric NTRK-rearranged Mesenchymal Tumors. *American Journal of Surgical Pathology*, **2019**, 43, 435-445	6.7	67
252	The incidence of microscopic thyroglossal duct tissue superior to the hyoid bone. *Laryngoscope*, **2019**, 129, 1215-1217	3.6	11
251	English Translation of M. Bŕard: Tumeur Embryonnaire Du Muscle Stri: [Embryonal Tumor of Striated Muscle]. Lyon Med 1894; 77: 52. *Fetal and Pediatric Pathology*, **2019**, 38, 182-184	1.7	3
250	The addition of cixutumumab or temozolomide to intensive multiagent chemotherapy is feasible but does not improve outcome for patients with metastatic rhabdomyosarcoma: A report from the Children's Oncology Group. *Cancer*, **2019**, 125, 290-297	6.4	33
249	On the First Usage of the Term "Rhabdomyosarcoma". *Pediatric and Developmental Pathology*, **2019**, 22, 87	2.2	
248	Solitary Fibrous Tumors in Pediatric Patients: A Rare and Potentially Overdiagnosed Neoplasm, Confirmed by STAT6 Immunohistochemistry. *Pediatric and Developmental Pathology*, **2018**, 21, 389-400	2.2	6
247	Transmission of a germline mutation from unaffected male carrier associated with pediatric glioblastoma in his child and gestational choriocarcinoma in his female partner. *Journal of Physical Education and Sports Management*, **2018**, 4,	2.8	6
246	OncoKids: A Comprehensive Next-Generation Sequencing Panel for Pediatric Malignancies. *Journal of Molecular Diagnostics*, **2018**, 20, 765-776	5.1	39
245	Fibroblastic and myofibroblastic tumors of children: new genetic entities and new ancillary testing. *F1000Research*, **2018**, 7,	3.6	11
244	Risk group accurately predicts outcome in primary extremity non-rhabdomyosarcoma soft tissue sarcomas (NRSTS) in patients . *Journal of Clinical Oncology*, **2018**, 36, 10546-10546	2.2	
243	Addition of Vincristine and Irinotecan to Vincristine, Dactinomycin, and Cyclophosphamide Does Not Improve Outcome for Intermediate-Risk Rhabdomyosarcoma: A Report From the Children's Oncology Group. *Journal of Clinical Oncology*, **2018**, 36, 2770-2777	2.2	74
242	Profiling targetable immune checkpoints in osteosarcoma. *OncoImmunology*, **2018**, 7, e1475873	7.2	19
241	Fibrous hamartoma of infancy: a clinicopathologic study of 145 cases, including 2 with sarcomatous features. *Modern Pathology*, **2017**, 30, 474-485	9.8	36
240	Reduction of cyclophosphamide dose for patients with subset 2 low-risk rhabdomyosarcoma is associated with an increased risk of recurrence: A report from the Soft Tissue Sarcoma Committee of the Children's Oncology Group. *Cancer*, **2017**, 123, 2368-2375	6.4	46
239	Histology, fusion status, and outcome in metastatic rhabdomyosarcoma: A report from the Children's Oncology Group. *Pediatric Blood and Cancer*, **2017**, 64, e26645	3	44
238	45 Gy is not sufficient radiotherapy dose for Group III orbital embryonal rhabdomyosarcoma after less than complete response to 12 weeks of ARST0331 chemotherapy: A report from the Soft Tissue Sarcoma Committee of the Children's Oncology Group. *Pediatric Blood and Cancer*, **2017**, 64, e26540	3	19
237	Inherited germline ATRX mutation in two brothers with ATR-X syndrome and osteosarcoma. *American Journal of Medical Genetics, Part A*, **2017**, 173, 1390-1395	2.5	21
236	Opportunities for Improvement in Pathology Reporting of Childhood Nonrhabdomyosarcoma Soft Tissue Sarcomas: A Report From Children's Oncology Group (COG) Study ARST0332. *American Journal of Clinical Pathology*, **2016**, 146, 328-38	1.9	10

235	Congenital Cyst of the Umbilical Cord. *Fetal and Pediatric Pathology*, **2016**, 35, 344-347	1.7	4
234	Increased Wnt and Notch signaling: a clue to the renal disease in Schimke immuno-osseous dysplasia?. *Orphanet Journal of Rare Diseases*, **2016**, 11, 149	4.2	12
233	Histology, Fusion Status, and Outcome in Alveolar Rhabdomyosarcoma With Low-Risk Clinical Features: A Report From the Children's Oncology Group. *Pediatric Blood and Cancer*, **2016**, 63, 634-9	3	44
232	Intensive Multiagent Therapy, Including Dose-Compressed Cycles of Ifosfamide/Etoposide and Vincristine/Doxorubicin/Cyclophosphamide, Irinotecan, and Radiation, in Patients With High-Risk Rhabdomyosarcoma: A Report From the Children's Oncology Group. *Journal of Clinical Oncology*, **2016**, 34, 117-22	2.2	110
231	MRI and CT of Low-Grade Fibromyxoid Sarcoma in Children: A Report From Children's Oncology Group Study ARST0332. *American Journal of Roentgenology*, **2015**, 205, 414-20	5.4	17
230	Atrial Thrombus in a Neonate: A Diagnostic Challenge. *AJP Reports*, **2015**, 5, e18-21	1.2	2
229	The World Health Organization Classification of Skeletal Muscle Tumors in Pediatric Rhabdomyosarcoma: A Report From the Children's Oncology Group. *Archives of Pathology and Laboratory Medicine*, **2015**, 139, 1281-7	5	81
228	Transcriptional and posttranscriptional mechanisms contribute to the dysregulation of elastogenesis in Schimke immuno-osseous dysplasia. *Pediatric Research*, **2015**, 78, 609-17	3.2	6
227	Pediatric Cancer in the Head and Neck **2015**, 203-226		
226	Soft Tissue Sarcomas **2015**, 19-68		
225	Malignant Tumors of Peripheral Nerves **2015**, 399-414		
224	Malignancies of the Pediatric Lower Respiratory Tract **2015**, 227-243		
223	Quantification of glypican 3, β-catenin and claudin-1 protein expression in hepatoblastoma and paediatric hepatocellular carcinoma by colour deconvolution. *Histopathology*, **2015**, 67, 905-13	7.3	20
222	Risk-based treatment for synovial sarcoma in patients under 30 years of age: Children's Oncology Group study ARST0332.. *Journal of Clinical Oncology*, **2015**, 33, 10012-10012	2.2	1
221	Early results from Children's Oncology Group (COG) ARST08P1: Pilot studies of cixutumumab or temozolomide with intensive multiagent chemotherapy for patients with metastatic rhabdomyosarcoma (RMS).. *Journal of Clinical Oncology*, **2015**, 33, 10015-10015	2.2	5
220	Immunohistochemical Markers of Soft Tissue Tumors: Pathologic Diagnosis, Genetic Contributions, and Therapeutic Options. *Analytical Chemistry Insights*, **2015**, 10, 1-10		7
219	Shorter-duration therapy using vincristine, dactinomycin, and lower-dose cyclophosphamide with or without radiotherapy for patients with newly diagnosed low-risk rhabdomyosarcoma: a report from the Soft Tissue Sarcoma Committee of the Children's Oncology Group. *Journal of Clinical Oncology*, **2014**, 32, 3547-52	2.2	94
218	Early response as assessed by anatomic imaging does not predict failure-free survival among patients with Group III rhabdomyosarcoma: a report from the Children's Oncology Group. *European Journal of Cancer*, **2014**, 50, 816-23	7.5	28

217	Ectomesenchymoma with embryonal rhabdomyosarcoma and ganglioneuroma, arising in association with benign triton tumor of the tongue. *Pediatric and Developmental Pathology*, **2014**, 17, 226-30	2.2	3
216	Imaging features of alveolar soft-part sarcoma: a report from Children's Oncology Group Study ARST0332. *American Journal of Roentgenology*, **2014**, 203, 1345-52	5.4	29
215	Myogenin, AP2, NOS-1, and HMGA2 are surrogate markers of fusion status in rhabdomyosarcoma: a report from the soft tissue sarcoma committee of the children's oncology group. *American Journal of Surgical Pathology*, **2014**, 38, 654-9	6.7	41
214	Vincristine, dactinomycin, cyclophosphamide (VAC) versus VAC/V plus irinotecan (VI) for intermediate-risk rhabdomyosarcoma (IRRMS): A report from the Children Oncology Group Soft Tissue Sarcoma Committee.. *Journal of Clinical Oncology*, **2014**, 32, 10004-10004	2.2	16
213	Risk-based treatment for nonrhabdomyosarcoma soft tissue sarcomas (NRSTS) in patients under 30 years of age: Children Oncology Group study ARST0332.. *Journal of Clinical Oncology*, **2014**, 32, 10008-10008	2.2	21
212	PAX-FOXO1 fusion status drives unfavorable outcome for children with rhabdomyosarcoma: a children's oncology group report. *Pediatric Blood and Cancer*, **2013**, 60, 1411-7	3	157
211	Dense pattern of embryonal rhabdomyosarcoma, a lesion easily confused with alveolar rhabdomyosarcoma: a report from the Soft Tissue Sarcoma Committee of the Children's Oncology Group. *American Journal of Clinical Pathology*, **2013**, 140, 82-90	1.9	60
210	Classification of rhabdomyosarcoma and its molecular basis. *Advances in Anatomic Pathology*, **2013**, 20, 387-97	5.1	166
209	Histologic and clinical characteristics can guide staging evaluations for children and adolescents with rhabdomyosarcoma: a report from the Children's Oncology Group Soft Tissue Sarcoma Committee. *Journal of Clinical Oncology*, **2013**, 31, 3226-32	2.2	67
208	A novel algorithm for simplification of complex gene classifiers in cancer. *Cancer Research*, **2013**, 73, 5625-32	10.1	4
207	Patterns of toxicities and outcome in children versus adolescents/young adults (AYA) with metastatic rhabdomyosarcoma (RMS): A report from the Children Oncology Group (COG) Soft Tissue Sarcoma Committee.. *Journal of Clinical Oncology*, **2013**, 31, 10554-10554	2.2	
206	Evaluation of early response by anatomic imaging to predict survival among patients with group III rhabdomyosarcoma: A report from the Children Oncology Group.. *Journal of Clinical Oncology*, **2013**, 31, 10012-10012	2.2	
205	Reduced elastogenesis: a clue to the arteriosclerosis and emphysematous changes in Schimke immuno-osseous dysplasia?. *Orphanet Journal of Rare Diseases*, **2012**, 7, 70	4.2	20
204	An English Translation of Joseph Luc Riopelle, MD, (Hôtel-Dieu of Montréal), and Jean Paul Thériault (Hôpital Général of Verdun, Québec, Canada): Sur une forme méconnue de sarcome des parties molles: le rhabdomyosarcome alvéolaire (concerning an unrecognized form of sarcoma of the soft tissues: alveolar rhabdomyosarcoma). *Annales d'anatomie pathologique* 1956;1:88-111. *Pediatric and Developmental Pathology* **2012**, 15, 407-16	2.2	5
203	Myogenic tumors in children and adolescents. *Pediatric and Developmental Pathology*, **2012**, 15, 211-38	2.2	37
202	Clinical utility gene card for: Alveolar rhabdomyosarcoma. *European Journal of Human Genetics*, **2012**, 20,	5.3	2
201	Vincristine (V), dactinomycin (A), and lower doses of cyclophosphamide (C) with or without radiation therapy for patients with newly diagnosed low-risk embryonal rhabdomyosarcoma (ERMS): A report from the Children Oncology Group (COG).. *Journal of Clinical Oncology*, **2012**, 30, 9509-9509	2.2	6
200	Relationship of fusion protein status and outcome for children with intermediate-risk rhabdomyosarcoma: A Children's Oncology Group report.. *Journal of Clinical Oncology*, **2012**, 30, 9535-9535	2.2	3

199	Langerhans cell histiocytosis preceding the development of juvenile xanthogranuloma: a case and review of recent developments. *Pediatric and Developmental Pathology*, **2011**, 14, 480-4	2.2	21
198	A novel t(8;13)(q21;q22) translocation in a pediatric lipoma. *Cancer Genetics*, **2011**, 204, 629-33	2.3	2
197	Pathobiologic markers of the ewing sarcoma family of tumors: state of the art and prediction of behaviour. *Sarcoma*, **2011**, 2011, 856190	3.1	51
196	Pediatric cutaneous angiosarcomas: a clinicopathologic study of 10 cases. *American Journal of Surgical Pathology*, **2011**, 35, 70-5	6.7	41
195	Evidence for an unanticipated relationship between undifferentiated pleomorphic sarcoma and embryonal rhabdomyosarcoma. *Cancer Cell*, **2011**, 19, 177-91	24.3	142
194	Origin of an anomalous left coronary artery from a left hilar pulmonary artery. *Pediatric Cardiology*, **2011**, 32, 98-101	2.1	
193	Unusual presentation of congenital infantile fibrosarcoma in seven infants with molecular-genetic analysis. *Fetal and Pediatric Pathology*, **2011**, 30, 329-37	1.7	27
192	Prognostic significance and tumor biology of regional lymph node disease in patients with rhabdomyosarcoma: a report from the Children's Oncology Group. *Journal of Clinical Oncology*, **2011**, 29, 1304-11	2.2	79
191	Results of the Intergroup Rhabdomyosarcoma Study Group D9602 protocol, using vincristine and dactinomycin with or without cyclophosphamide and radiation therapy, for newly diagnosed patients with low-risk embryonal rhabdomyosarcoma: a report from the Soft Tissue Sarcoma Committee of the Children's Oncology Group. *Journal of Clinical Oncology*, **2011**, 29, 1312-8	2.2	150
190	Cytology testing and proficiency: survey findings of the Practice Committee of Society for Pediatric Pathology. *Pediatric and Developmental Pathology*, **2011**, 14, 153-4	2.2	2
189	Randomized phase II window trial of two schedules of irinotecan with vincristine in patients with first relapse or progression of rhabdomyosarcoma: a report from the Children's Oncology Group. *Journal of Clinical Oncology*, **2010**, 28, 4658-63	2.2	70
188	Fusion-negative alveolar rhabdomyosarcoma: modification of risk stratification is premature. *Journal of Clinical Oncology*, **2010**, 28, e587-8; author reply e589-90	2.2	17
187	Modern Diagnosis of Small Cell Malignancies of Children. *Surgical Pathology Clinics*, **2010**, 3, 515-51	3.9	10
186	A mouse model of lethal synergism between influenza virus and Haemophilus influenzae. *American Journal of Pathology*, **2010**, 176, 800-11	5.8	51
185	Impaired Wnt signaling in embryonal rhabdomyosarcoma cells from p53/c-fos double mutant mice. *American Journal of Pathology*, **2010**, 177, 2055-66	5.8	30
184	Grading of nonrhabdomyosarcoma soft tissue sarcoma in children and adolescents: a comparison of parameters used for the Fdration Nationale des Centers de Lutte Contre le Cancer and Pediatric Oncology Group Systems. *Cancer*, **2010**, 116, 2266-74	6.4	26
183	Treatment results for patients with localized, completely resected (Group I) alveolar rhabdomyosarcoma on Intergroup Rhabdomyosarcoma Study Group (IRSG) protocols III and IV, 1984-1997: a report from the Children's Oncology Group. *Pediatric Blood and Cancer*, **2010**, 55, 612-6	3	17
182	Jay Bernstein. *Pediatric and Developmental Pathology*, **2009**, 12, 311-311	2.2	

181	Steve Qualman. *Pediatric and Developmental Pathology*, **2009**, 12, 81-82	2.2	
180	Vincristine, actinomycin, and cyclophosphamide compared with vincristine, actinomycin, and cyclophosphamide alternating with vincristine, topotecan, and cyclophosphamide for intermediate-risk rhabdomyosarcoma: children's oncology group study D9803. *Journal of Clinical Oncology*, **2009**, 27, 5182-8	2.2	261
179	Lymphoma arising from a calcinotic lesion in a patient with juvenile dermatomyositis. *Pediatric Dermatology*, **2009**, 26, 159-61	1.9	10
178	Paediatric soft tissue tumours: from histology to molecular diagnosis. *Diagnostic Histopathology*, **2009**, 15, 524-530	0.7	3
177	Angiosarcomas arising in the viscera and soft tissue of children and young adults: a clinicopathologic study of 15 cases. *American Journal of Surgical Pathology*, **2009**, 33, 264-9	6.7	57
176	Response to Buccoliero: Letter to the Editor; The Problems and Promise of Central Pathology Review. *Pediatric and Developmental Pathology*, **2008**, 11, 328-328	2.2	
175	Fatal non-transplant-related epstein-barr virus-associated atypical lymphoid proliferations in infants and children: a clinicopathologic study. *Pediatric and Developmental Pathology*, **2008**, 11, 443-9	2.2	11
174	Prevalence and clinical impact of anaplasia in childhood rhabdomyosarcoma : a report from the Soft Tissue Sarcoma Committee of the Children's Oncology Group. *Cancer*, **2008**, 113, 3242-7	6.4	54
173	Randomized phase II window study of two schedules of irinotecan (CPT-11) and vincristine (VCR) in rhabdomyosarcoma (RMS) at first relapse/disease progression. *Journal of Clinical Oncology*, **2008**, 26, 10013-10013	2.2	10
172	Two consecutive phase II window trials of irinotecan alone or in combination with vincristine for the treatment of metastatic rhabdomyosarcoma: the Children's Oncology Group. *Journal of Clinical Oncology*, **2007**, 25, 362-9	2.2	159
171	A newborn with Poland anomaly and liver ectopy: an unusual association with important prognostic implications. *Pediatric and Developmental Pathology*, **2007**, 10, 134-7	2.2	7
170	Melanotic neuroectodermal tumor of infancy: report of a case with myogenic differentiation. *Pediatric and Developmental Pathology*, **2007**, 10, 157-60	2.2	10
169	The problems and promise of central pathology review: development of a standardized procedure for the Children's Oncology Group. *Pediatric and Developmental Pathology*, **2007**, 10, 199-207	2.2	30
168	Penile myointimoma in children and adolescents: a clinicopathologic study of 5 cases supporting a distinct entity. *American Journal of Surgical Pathology*, **2007**, 31, 1622-6	6.7	23
167	Correlation between histology and PAX/FKHR fusion status in alveolar rhabdomyosarcoma: a report from the Children's Oncology Group. *American Journal of Surgical Pathology*, **2007**, 31, 895-901	6.7	64
166	Successful management of rhabdoid tumor of the liver. *Journal of Pediatric Hematology/Oncology*, **2007**, 29, 406-8	1.2	24
165	Immunohistochemistry of primary malignant neuroepithelial tumors of the kidney: a potential source of confusion? A study of 30 cases from the National Wilms Tumor Study Pathology Center. *Human Pathology*, **2007**, 38, 205-11	3.7	36
164	Soft-tissue aneurysmal bone cyst: report of a case with t(5;17)(q33;p13). *Pediatric and Developmental Pathology*, **2007**, 10, 46-9	2.2	16

163	Randomized phase III trial comparing vincristine, actinomycin, cyclophosphamide (VAC) with VAC/V topotecan/cyclophosphamide (TC) for intermediate risk rhabdomyosarcoma (IRRMS). D9803, COG study. *Journal of Clinical Oncology*, **2007**, 25, 9509-9509	2.2	4
162	Examination of gene fusion status in archival samples of alveolar rhabdomyosarcoma entered on the Intergroup Rhabdomyosarcoma Study-III trial: a report from the Children's Oncology Group. *Journal of Molecular Diagnostics*, **2006**, 8, 202-8	5.1	80
161	Fatal pulmonary fat embolism following spinal fusion surgery. *Pediatric Critical Care Medicine*, **2006**, 7, 263-6	3	7
160	An immunohistochemical algorithm to facilitate diagnosis and subtyping of rhabdomyosarcoma: the Children's Oncology Group experience. *American Journal of Surgical Pathology*, **2006**, 30, 962-8	6.7	153
159	Use of a novel FISH assay on paraffin-embedded tissues as an adjunct to diagnosis of alveolar rhabdomyosarcoma. *Laboratory Investigation*, **2006**, 86, 547-56	5.9	72
158	Rhabdomyosarcomas in adults and children: an update. *Archives of Pathology and Laboratory Medicine*, **2006**, 130, 1454-65	5	220
157	Are external quality assurance (EQA) slide schemes a valid tool for the performance assessment of histopathologists?. *Pathology Research and Practice*, **2005**, 201, 117-21	3.4	5
156	Malignant triton tumor presenting as a rectal mass in an 11-month-old. *Pediatric and Developmental Pathology*, **2005**, 8, 235-9	2.2	15
155	USCAP Specialty Conference: case 4. *Pediatric and Developmental Pathology*, **2005**, 8, 85-7	2.2	1
154	Review of tonsillar lymphoma in pediatric patients from the pediatric oncology group: what can be learned about some indications for microscopic examination?. *Pediatric and Developmental Pathology*, **2005**, 8, 533-40	2.2	18
153	Cytogenetic findings in pediatric T-lymphoblastic lymphomas: one institution's experience and a review of the literature. *Pediatric and Developmental Pathology*, **2005**, 8, 550-6	2.2	5
152	Sclerosing rhabdomyosarcomas in children and adolescents: a clinicopathologic review of 13 cases from the Intergroup Rhabdomyosarcoma Study Group and Children's Oncology Group. *Pediatric and Developmental Pathology*, **2005**, 8, 141	2.2	7
151	Upstream CpG island methylation of the PAX3 gene in human rhabdomyosarcomas. *Pediatric Blood and Cancer*, **2005**, 44, 328-37	3	23
150	A novel t(X;2)(q13;q35) in clear cell sugar tumor of bone. *Cancer Genetics and Cytogenetics*, **2004**, 154, 77-80		6
149	Lymphocytic infiltration in pediatric thyroid carcinomas. *Pediatric and Developmental Pathology*, **2004**, 7, 487-92	2.2	9
148	Cytogenetic distinction among benign fibro-osseous lesions of bone in children and adolescents: value of karyotypic findings in differential diagnosis. *Pediatric and Developmental Pathology*, **2004**, 7, 148-58	2.2	25
147	Sclerosing rhabdomyosarcomas in children and adolescents: a clinicopathologic review of 13 cases from the Intergroup Rhabdomyosarcoma Study Group and Children's Oncology Group. *Pediatric and Developmental Pathology*, **2004**, 7, 583-94	2.2	60
146	Age is an independent prognostic factor in rhabdomyosarcoma: a report from the Soft Tissue Sarcoma Committee of the Children's Oncology Group. *Pediatric Blood and Cancer*, **2004**, 42, 64-73	3	166

145	Prevalence of Alveolar Rhabdomyosarcoma. *Oncology Times*, **2003**, 25, 5	3.5	
144	Incidence of autopsy findings in unexpected deaths of children and adolescents. *Pediatric and Developmental Pathology*, **2003**, 6, 142-55	2.2	9
143	Satellite DNA hypomethylation in karyotyped Wilms tumors. *Cancer Genetics and Cytogenetics*, **2003**, 141, 97-105		76
142	Prognostic factors and clinical outcomes in children and adolescents with metastatic rhabdomyosarcoma--a report from the Intergroup Rhabdomyosarcoma Study IV. *Journal of Clinical Oncology*, **2003**, 21, 78-84	2.2	356
141	Protocol for the examination of specimens from patients (children and young adults) with rhabdomyosarcoma. *Archives of Pathology and Laboratory Medicine*, **2003**, 127, 1290-7	5	37
140	Renal Tumors in Childhood **2003**, 143-151		
139	Recurring breakpoints of 1p13 approximately p22 in osteochondroma. *Cancer Genetics and Cytogenetics*, **2002**, 138, 102-6		14
138	Comparison of manual and automated leukocyte counts for determination of the absolute neutrophil count: application to a pediatric oncology clinic. *Medical and Pediatric Oncology*, **2002**, 38, 183-6		7
137	Limited reliability of lipid-laden macrophage index restricts its use as a test for pulmonary aspiration: comparison with a simple semiquantitative assay. *Pediatric and Developmental Pathology* , **2002**, 5, 551-8	2.2	22
136	Appearances may be deceiving: does aggressive histology correlate with behavior of infantile sarcomas?. *Pediatric and Developmental Pathology*, **2002**, 5, 513-4	2.2	5
135	Hypomethylation and hypermethylation of DNA in Wilms tumors. *Oncogene*, **2002**, 21, 6694-702	9.2	154
134	Pulmonary inflammatory pseudotumor with HHV-8. *American Journal of Surgical Pathology*, **2002**, 26, 1089-91; author reply 1091-2	6.7	11
133	Thallium bone imaging as an indicator of response and outcome in nonmetastatic primary extremity osteosarcoma. *Pediatric Radiology*, **2001**, 31, 251-6	2.8	18
132	Comparison of immunohistochemistry and silver stain for the diagnosis of pediatric Helicobacter pylori infection in urease-negative gastric biopsies. *Pediatric and Developmental Pathology*, **2001**, 4, 82-8	2.2	17
131	The frozen section yesterday and today: pediatric solid tumors--crucial issues. *Pediatric and Developmental Pathology*, **2001**, 4, 252-66	2.2	8
130	Efficacy of ifosfamide and doxorubicin given as a phase II "window" in children with newly diagnosed metastatic rhabdomyosarcoma: a report from the Intergroup Rhabdomyosarcoma Study Group. *Medical and Pediatric Oncology*, **2001**, 37, 442-8		63
129	Rhabdomyosarcoma of the parotid region occurring in childhood and adolescence. A report from the Intergroup Rhabdomyosarcoma Study Group. *Cancer*, **2001**, 92, 3135-46	6.4	22
128	Axillary lymph node scarring and the association with tumour response following neoadjuvant chemoendocrine therapy for breast cancer. *Breast*, **2001**, 10, 61-6	3.6	12

127	Correspondence re: Parham DM. Pathologic classification of rhabdomyosarcomas and correlations with molecular studies. Mod Pathol 2001;14:506-14. *Modern Pathology*, **2001**, 14, 1068	9.8	4
126	Pathologic classification of rhabdomyosarcomas and correlations with molecular studies. *Modern Pathology*, **2001**, 14, 506-14	9.8	183
125	Neuroectodermal and neuroendocrine tumors principally seen in children. *Pathology Patterns Reviews*, **2001**, 115 Suppl, S113-28		12
124	Carboplatin/ifosfamide window therapy for osteosarcoma: results of the St Jude Children's Research Hospital OS-91 trial. *Journal of Clinical Oncology*, **2001**, 19, 171-82	2.2	94
123	Primary malignant neuroepithelial tumors of the kidney: a clinicopathologic analysis of 146 adult and pediatric cases from the National Wilms' Tumor Study Group Pathology Center. *American Journal of Surgical Pathology*, **2001**, 25, 133-46	6.7	156
122	Pathological case of the month. Disseminated brucellosis. *JAMA Pediatrics*, **2000**, 154, 311-2		1
121	Immunohistochemical detection of FLI-1 protein expression: a study of 132 round cell tumors with emphasis on CD99-positive mimics of Ewing's sarcoma/primitive neuroectodermal tumor. *American Journal of Surgical Pathology*, **2000**, 24, 1657-62	6.7	266
120	Clinicopathologic study of ectomesenchymomas from Intergroup Rhabdomyosarcoma Study Groups III and IV. *Pediatric and Developmental Pathology*, **2000**, 3, 290-300	2.2	37
119	Neural differentiation and prognosis in peripheral primitive neuroectodermal tumor. *Journal of Clinical Oncology*, **2000**, 18, 2187-8	2.2	8
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