

Jennifer L Picarsic

List of Publications by Year in descending order

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Version: 2024-02-01

41
papers

4,422
citations

304743

22
h-index

289244

40
g-index

44
all docs

44
docs citations

44
times ranked

3978
citing authors

#	ARTICLE	IF	CITATIONS
1	ALK-positive histiocytosis: a new clinicopathologic spectrum highlighting neurologic involvement and responses to ALK inhibition. <i>Blood</i> , 2022, 139, 256-280.	1.4	60
2	International expert consensus recommendations for the diagnosis and treatment of Langerhans cell histiocytosis in adults. <i>Blood</i> , 2022, 139, 2601-2621.	1.4	63
3	The 5th edition of the World Health Organization Classification of Haematolymphoid Tumours: Myeloid and Histiocytic/Dendritic Neoplasms. <i>Leukemia</i> , 2022, 36, 1703-1719.	7.2	1,211
4	Discrepancies between 18 F-FDG PET/CT findings and conventional imaging in Langerhans cell histiocytosis. <i>Pediatric Blood and Cancer</i> , 2021, 68, e28891.	1.5	20
5	BRAF fusions in pediatric histiocytic neoplasms define distinct therapeutic responsiveness to RAF paradox breakers. <i>Pediatric Blood and Cancer</i> , 2021, 68, e28933.	1.5	9
6	BRAFV600E-induced senescence drives Langerhans cell histiocytosis pathophysiology. <i>Nature Medicine</i> , 2021, 27, 851-861.	30.7	38
7	A chronic eyelid lesion in a child: multi-disciplinary approach to diagnosis, treatment and management of a highly atypical histiocytic lesion. <i>Pediatric Hematology and Oncology</i> , 2021, , 1-7.	0.8	1
8	Histiocytic disorders. <i>Nature Reviews Disease Primers</i> , 2021, 7, 73.	30.5	46
9	New molecular insights into the pathogenesis of lipoblastomas: clinicopathologic, immunohistochemical, and molecular analysis in pediatric cases. <i>Human Pathology</i> , 2020, 104, 30-41.	2.0	20
10	Bone marrow-derived myeloid progenitors as driver mutation carriers in high- and low-risk Langerhans cell histiocytosis. <i>Blood</i> , 2020, 136, 2188-2199.	1.4	18
11	Dasatinib induces a dramatic response in a child with refractory juvenile xanthogranuloma with a novel MRC1-PDGFRB fusion. <i>Blood Advances</i> , 2020, 4, 2991-2995.	5.2	10
12	Interleukin-18 and cytotoxic impairment are independent and synergistic causes of murine virus-induced hyperinflammation. <i>Blood</i> , 2020, 136, 2162-2174.	1.4	20
13	Fibrous histiocytoma/dermatofibroma in children: the same as adults?. <i>Human Pathology</i> , 2020, 99, 107-115.	2.0	11
14	Erdheim-Chester disease: consensus recommendations for evaluation, diagnosis, and treatment in the molecular era. <i>Blood</i> , 2020, 135, 1929-1945.	1.4	191
15	Langerin staining identifies most littoral cell angiomas but not most other splenic angiomatous lesions. <i>Human Pathology</i> , 2019, 83, 43-49.	2.0	3
16	BRAF V600E mutation in Juvenile Xanthogranuloma family neoplasms of the central nervous system (CNS-JXG): a revised diagnostic algorithm to include pediatric Erdheim-Chester disease. <i>Acta Neuropathologica Communications</i> , 2019, 7, 168.	5.2	32
17	Mechanisms of action of ruxolitinib in murine models of hemophagocytic lymphohistiocytosis. <i>Blood</i> , 2019, 134, 147-159.	1.4	99
18	A Rare Case of Uterine Torsion With Juvenile Granulosa Cell Tumor in the Pediatric Patient. <i>Urology</i> , 2019, 128, 87-89.	1.0	8

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19	BRAF-V600E mutated Rosai-Dorfman-Destombes disease and Langerhans cell histiocytosis with response to BRAF inhibitor. <i>Blood Advances</i> , 2019, 3, 1848-1853.	5.2	28
20	Activating mutations in CSF1R and additional receptor tyrosine kinases in histiocytic neoplasms. <i>Nature Medicine</i> , 2019, 25, 1839-1842.	30.7	122
21	Improving medical students' understanding of pediatric diseases through an innovative and tailored web-based digital pathology program with philips pathology Tutor (Formerly PathXL). <i>Journal of Pathology Informatics</i> , 2019, 10, 18.	1.7	2
22	CNS Langerhans cell histiocytosis: Common hematopoietic origin for LCH-associated neurodegeneration and mass lesions. <i>Cancer</i> , 2018, 124, 2607-2620.	4.1	73
23	Interleukin-18 diagnostically distinguishes and pathogenically promotes human and murine macrophage activation syndrome. <i>Blood</i> , 2018, 131, 1442-1455.	1.4	288
24	Pediatric Testicular Hemangioma in a 10-Year-old: A Rare Entity That May Mimic Malignancy With Appraisal of the Literature. <i>Urology</i> , 2018, 114, 175-180.	1.0	6
25	Consensus recommendations for the diagnosis and clinical management of Rosai-Dorfman-Destombes disease. <i>Blood</i> , 2018, 131, 2877-2890.	1.4	335
26	Pathology of Histiocytic Disorders and Neoplasms and Related Disorders. , 2018, , 3-50.		11
27	Activating Mutations in CSF1R and Additional Receptor Tyrosine Kinases in Sporadic and Familial Histiocytic Neoplasms. <i>Blood</i> , 2018, 132, 49-49.	1.4	10
28	Novel <i>NLR4</i> Mutation Causes a Syndrome of Perinatal Autoinflammation With Hemophagocytic Lymphohistiocytosis, Hepatosplenomegaly, Fetal Thrombotic Vasculopathy, and Congenital Anemia and Ascites. <i>Pediatric and Developmental Pathology</i> , 2017, 20, 498-505.	1.0	62
29	Late graft dysfunction after pediatric heart transplantation is associated with fibrosis and microvasculopathy by automated, digital whole-slide analysis. <i>Journal of Heart and Lung Transplantation</i> , 2017, 36, 1336-1343.	0.6	15
30	Langerhans cell histiocytosis and Erdheim-Chester disease, both with cutaneous presentations, and papillary thyroid carcinoma all harboring the <i>BRAF</i> ^{V600E} mutation. <i>Journal of Cutaneous Pathology</i> , 2016, 43, 270-275.	1.3	27
31	Role of Epstein-Barr virus status and immunophenotypic studies in the evaluation of exfoliative cytology specimens from patients with post-transplant lymphoproliferative disorders. <i>Cancer Cytopathology</i> , 2016, 124, 425-435.	2.4	2
32	Revised classification of histiocytoses and neoplasms of the macrophage-dendritic cell lineages. <i>Blood</i> , 2016, 127, 2672-2681.	1.4	1,040
33	Mutations in the nuclear bile acid receptor FXR cause progressive familial intrahepatic cholestasis. <i>Nature Communications</i> , 2016, 7, 10713.	12.8	227
34	Molecular Characterization of Sporadic Pediatric Thyroid Carcinoma with the DNA/RNA ThyroSeq v2 Next-Generation Sequencing Assay. <i>Pediatric and Developmental Pathology</i> , 2016, 19, 115-122.	1.0	63
35	Phenotype and Immunophenotype of the Most Common Pediatric Tumors. <i>Applied Immunohistochemistry and Molecular Morphology</i> , 2015, 23, 313-326.	1.2	13
36	Can Malignant Thyroid Nodules Be Distinguished from Benign Thyroid Nodules in Children and Adolescents by Clinical Characteristics? A Review of 89 Pediatric Patients with Thyroid Nodules. <i>Thyroid</i> , 2015, 25, 392-400.	4.5	56

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37	Histologic Patterns of Thymic Involvement in Langerhans Cell Proliferations: A Clinicopathologic Study and Review of the Literature. <i>Pediatric and Developmental Pathology</i> , 2015, 18, 127-138.	1.0	9
38	Identification of Unique, Heterozygous Germline Mutation, <i>STK11</i> (p.F354L), in a Child with an Encapsulated Follicular Variant of Papillary Thyroid Carcinoma within Six Months of Completing Treatment for Neuroblastoma. <i>Pediatric and Developmental Pathology</i> , 2015, 18, 318-323.	1.0	16
39	Nosology and Pathology of Langerhans Cell Histiocytosis. <i>Hematology/Oncology Clinics of North America</i> , 2015, 29, 799-823.	2.2	52
40	Postâ€transplant Burkitt lymphoma is a more aggressive and distinct form of postâ€transplant lymphoproliferative disorder. <i>Cancer</i> , 2011, 117, 4540-4550.	4.1	46
41	Selfâ€Reported Napping and Duration and Quality of Sleep in the Lifestyle Interventions and Independence for Elders Pilot Study. <i>Journal of the American Geriatrics Society</i> , 2008, 56, 1674-1680.	2.6	58