

SÃ©bastien HÃ©ritier

List of Publications by Year in descending order

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Version: 2024-02-01

41
papers

2,402
citations

361045

20
h-index

288905

40
g-index

51
all docs

51
docs citations

51
times ranked

3668
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.	0.6	60
2	Determinants of long-term outcomes of splenectomy in pediatric autoimmune cytopenias. <i>Blood</i> , 2022, 140, 253-261.	0.6	6
3	Eye movement abnormalities in neurodegenerative langerhans cell histiocytosis. <i>Neurological Sciences</i> , 2022, 43, 6539-6546.	0.9	3
4	Multisystem inflammatory syndrome in children rose and fell with the first wave of the COVID-19 pandemic in France. <i>Acta Paediatrica, International Journal of Paediatrics</i> , 2021, 110, 922-932.	0.7	21
5	A circulating subset of BRAF ^{V600E} -positive cells in infants with high-risk Langerhans cell histiocytosis treated with BRAF inhibitors. <i>British Journal of Haematology</i> , 2021, 194, 745-749.	1.2	5
6	Post-COVID-19 severe neutropenia. <i>Pediatric Blood and Cancer</i> , 2021, 68, e28866.	0.8	9
7	Long-term follow-up of children with risk organ-negative Langerhans cell histiocytosis after 2-chlorodeoxyadenosine treatment. <i>British Journal of Haematology</i> , 2020, 191, 825-834.	1.2	14
8	Chest computed tomography findings for a cohort of children with pulmonary Langerhans cell histiocytosis. <i>Pediatric Blood and Cancer</i> , 2020, 67, e28496.	0.8	7
9	Childhood Langerhans cell histiocytosis with severe lung involvement: a nationwide cohort study. <i>Orphanet Journal of Rare Diseases</i> , 2020, 15, 241.	1.2	14
10	Chronic Granulomatous Disease with the McLeod Phenotype: a French National Retrospective Case Series. <i>Journal of Clinical Immunology</i> , 2020, 40, 752-762.	2.0	10
11	Progress towards molecular-based management of childhood Langerhans cell histiocytosis. <i>Archives De Pediatrie</i> , 2019, 26, 301-307.	0.4	24
12	Vemurafenib for Refractory Multisystem Langerhans Cell Histiocytosis in Children: An International Observational Study. <i>Journal of Clinical Oncology</i> , 2019, 37, 2857-2865.	0.8	132
13	Highly sensitive methods are required to detect mutations in histiocytoses. <i>Haematologica</i> , 2019, 104, e97-e99.	1.7	27
14	Lung involvement in childhood Langerhans cell histiocytosis, A multi-institutional study from the french LCH study group. , 2019, , .		1
15	Incidence and risk factors for clinical neurodegenerative Langerhans cell histiocytosis: a longitudinal cohort study. <i>British Journal of Haematology</i> , 2018, 183, 608-617.	1.2	54
16	Autoimmune and inflammatory manifestations occur frequently in patients with primary immunodeficiencies. <i>Journal of Allergy and Clinical Immunology</i> , 2017, 140, 1388-1393.e8.	1.5	222
17	Circulating cell-free BRAF ^{V600E} as a biomarker in children with Langerhans cell histiocytosis. <i>British Journal of Haematology</i> , 2017, 178, 457-467.	1.2	57
18	Disseminated Bacillus Calmette-GuÃ©rin Osteomyelitis in Twin Sisters Related to STAT1 Gene Deficiency. <i>Pediatric and Developmental Pathology</i> , 2017, 20, 255-261.	0.5	11

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19	New somatic BRAF splicing mutation in Langerhans cell histiocytosis. <i>Molecular Cancer</i> , 2017, 16, 115.	7.9	37
20	<i>BRAF</i> Mutation Correlates With High-Risk Langerhans Cell Histiocytosis and Increased Resistance to First-Line Therapy. <i>Journal of Clinical Oncology</i> , 2016, 34, 3023-3030.	0.8	233
21	Disseminated BCG osteomyelitis related to STAT 1 gene deficiency mimicking a metastatic neuroblastoma. <i>Pediatric and Developmental Pathology</i> , 2016, , .	0.5	3
22	Langerhans cell histiocytosis: therapeutic strategy and outcome in a 30-year nationwide cohort of 1478 patients under 18 years of age. <i>British Journal of Haematology</i> , 2016, 174, 887-898.	1.2	83
23	Specific T cells for the treatment of cytomegalovirus and/or adenovirus in the context of hematopoietic stem cell transplantation. <i>Journal of Allergy and Clinical Immunology</i> , 2016, 138, 920-924.e3.	1.5	21
24	Diverse and Targetable Kinase Alterations Drive Histiocytic Neoplasms. <i>Cancer Discovery</i> , 2016, 6, 154-165.	7.7	372
25	Spondyloenchondrodysplasia Due to Mutations in ACP5: A Comprehensive Survey. <i>Journal of Clinical Immunology</i> , 2016, 36, 220-234.	2.0	71
26	Common cancer-associated PIK3CA activating mutations rarely occur in Langerhans cell histiocytosis. <i>Blood</i> , 2015, 125, 2448-2449.	0.6	28
27	Vemurafenib Use in an Infant for High-Risk Langerhans Cell Histiocytosis. <i>JAMA Oncology</i> , 2015, 1, 836.	3.4	92
28	Outcomes Following Gene Therapy in Patients With Severe Wiskott-Aldrich Syndrome. <i>JAMA - Journal of the American Medical Association</i> , 2015, 313, 1550.	3.8	327
29	Inherited CARD9 Deficiency in 2 Unrelated Patients With Invasive <i>Exophiala</i> Infection. <i>Journal of Infectious Diseases</i> , 2015, 211, 1241-1250.	1.9	141
30	Langerhans cell histiocytosis in children: Correlation of <i>BRAF</i> status with clinical characteristic.. <i>Journal of Clinical Oncology</i> , 2015, 33, 10003-10003.	0.8	0
31	Diverse and Targetable Kinase Alterations Drive Histiocytic Neoplasms. <i>Blood</i> , 2015, 126, 481-481.	0.6	0
32	Immune deficiency-related enteropathy-lymphocytopenia-alopecia syndrome results from tetratricopeptide repeat domain 7A deficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2014, 134, 1354-1364.e6.	1.5	66
33	Safety of hematopoietic stem cell transplantation from hepatitis B core antibodies-positive donors with low/undetectable viremia in HBV-naïve children. <i>European Journal of Clinical Microbiology and Infectious Diseases</i> , 2014, 33, 545-550.	1.3	7
34	Syndromes d'activation lymphohistiocytaire constitutionnels. <i>Revue D'Oncologie Hématologie Pédiatrique</i> , 2013, 1, 104-110.	0.1	0
35	Temporal and Spatial Compartmentalization of Drug-Resistant Cytomegalovirus (CMV) in a Child with CMV Meningoencephalitis: Implications for Sampling in Molecular Diagnosis. <i>Journal of Clinical Microbiology</i> , 2013, 51, 4266-4269.	1.8	26
36	Circulating Endothelial Cells As a Reliable Marker Of Endothelial Damage In Children Undergoing Hematopoietic Stem Cell Transplantation. <i>Blood</i> , 2013, 122, 2049-2049.	0.6	0

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37	Prevalence and Clinical Impact of Norovirus Fecal Shedding in Children with Inherited Immune Deficiencies. <i>Journal of Infectious Diseases</i> , 2012, 206, 1269-1274.	1.9	65
38	Massive expansion of maternal T cells in response to EBV infection in a patient with SCID-XI. <i>Blood</i> , 2012, 120, 1957-1959.	0.6	21
39	Primary T-cell immunodeficiency with immunodysregulation caused by autosomal recessive LCK deficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2012, 130, 1144-1152.e11.	1.5	96
40	Retrospective French nationwide survey of childhood aggressive vascular anomalies of bone, 1988-2009. <i>Orphanet Journal of Rare Diseases</i> , 2010, 5, 3.	1.2	15
41	Immune control of tumors: host immune response and antibody-based immunotherapy. <i>Biomedicine and Pharmacotherapy</i> , 2008, 62, 516.	2.5	0