

AnnaCarin Horne

List of Publications by Year in descending order

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Version: 2024-02-01

26
papers

9,230
citations

346980

22
h-index

620720

26
g-index

26
all docs

26
docs citations

26
times ranked

8316
citing authors

#	ARTICLE	IF	CITATIONS
1	Consensus-Based Guidelines for the Recognition, Diagnosis, and Management of Hemophagocytic Lymphohistiocytosis in Critically Ill Children and Adults. <i>Critical Care Medicine</i> , 2022, 50, 860-872.	0.4	29
2	Efficacy of Moderately Dosed Etoposide in Macrophage Activation Syndrome—Hemophagocytic Lymphohistiocytosis. <i>Journal of Rheumatology</i> , 2021, 48, 1596-1602.	1.0	26
3	Definition and validation of serum biomarkers for optimal differentiation of hyperferritinaemic cytokine storm conditions in children: a retrospective cohort study. <i>Lancet Rheumatology</i> , The, 2021, 3, e563-e573.	2.2	14
4	Diagnostic challenges for a novel SH2D1A mutation associated with X-linked lymphoproliferative disease. <i>Pediatric Blood and Cancer</i> , 2020, 67, e28184.	0.8	4
5	Development and initial validation of the MS score for diagnosis of macrophage activation syndrome in systemic juvenile idiopathic arthritis. <i>Annals of the Rheumatic Diseases</i> , 2019, 78, 1357-1362.	0.5	74
6	A novel disorder involving dyshematopoiesis, inflammation, and HLH due to aberrant CDC42 function. <i>Journal of Experimental Medicine</i> , 2019, 216, 2778-2799.	4.2	132
7	Recommendations for the management of hemophagocytic lymphohistiocytosis in adults. <i>Blood</i> , 2019, 133, 2465-2477.	0.6	587
8	Juvenile idiopathic arthritis and risk of cancer before and after the introduction of biological therapies. <i>RMD Open</i> , 2019, 5, e001055.	1.8	9
9	Recommendations for the Use of Etoposide-Based Therapy and Bone Marrow Transplantation for the Treatment of HLH: Consensus Statements by the HLH Steering Committee of the Histiocyte Society. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2018, 6, 1508-1517.	2.0	112
10	Confirmed efficacy of etoposide and dexamethasone in HLH treatment: long-term results of the cooperative HLH-2004 study. <i>Blood</i> , 2017, 130, 2728-2738.	0.6	418
11	Development and Initial Validation of the Macrophage Activation Syndrome/Primary Hemophagocytic Lymphohistiocytosis Score, a Diagnostic Tool that Differentiates Primary Hemophagocytic Lymphohistiocytosis from Macrophage Activation Syndrome. <i>Journal of Pediatrics</i> , 2017, 189, 72-78.e3.	0.9	50
12	Revised classification of histiocytoses and neoplasms of the macrophage-dendritic cell lineages. <i>Blood</i> , 2016, 127, 2672-2681.	0.6	1,040
13	2016 Classification Criteria for Macrophage Activation Syndrome Complicating Systemic Juvenile Idiopathic Arthritis: A European League Against Rheumatism/American College of Rheumatology/Paediatric Rheumatology International Trials Organisation Collaborative Initiative. <i>Arthritis and Rheumatology</i> , 2016, 68, 566-576.	2.9	427
14	Expert consensus on dynamics of laboratory tests for diagnosis of macrophage activation syndrome complicating systemic juvenile idiopathic arthritis. <i>RMD Open</i> , 2016, 2, e000161.	1.8	57
15	Macrophage activation syndrome in the era of biologic therapy. <i>Nature Reviews Rheumatology</i> , 2016, 12, 259-268.	3.5	323
16	2016 Classification Criteria for Macrophage Activation Syndrome Complicating Systemic Juvenile Idiopathic Arthritis. <i>Annals of the Rheumatic Diseases</i> , 2016, 75, 481-489.	0.5	338
17	Incidence and clinical presentation of primary hemophagocytic lymphohistiocytosis in Sweden. <i>Pediatric Blood and Cancer</i> , 2015, 62, 346-352.	0.8	63
18	Dissecting the Heterogeneity of Macrophage Activation Syndrome Complicating Systemic Juvenile Idiopathic Arthritis. <i>Journal of Rheumatology</i> , 2015, 42, 994-1001.	1.0	59

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19	Clinical Features, Treatment, and Outcome of Macrophage Activation Syndrome Complicating Systemic Juvenile Idiopathic Arthritis: A Multinational, Multicenter Study of 362 Patients. <i>Arthritis and Rheumatology</i> , 2014, 66, 3160-3169.	2.9	322
20	Performance of Current Guidelines for Diagnosis of Macrophage Activation Syndrome Complicating Systemic Juvenile Idiopathic Arthritis. <i>Arthritis and Rheumatology</i> , 2014, 66, 2871-2880.	2.9	101
21	Chemoimmunotherapy for hemophagocytic lymphohistiocytosis: long-term results of the HLH-94 treatment protocol. <i>Blood</i> , 2011, 118, 4577-4584.	0.6	493
22	Frequency and spectrum of central nervous system involvement in 193 children with haemophagocytic lymphohistiocytosis. <i>British Journal of Haematology</i> , 2008, 140, 327-335.	1.2	217
23	Characterization of <i>PRF1</i> , <i>STX11</i> and <i>UNC13D</i> genotype-phenotype correlations in familial hemophagocytic lymphohistiocytosis. <i>British Journal of Haematology</i> , 2008, 143, 75-83.	1.2	78
24	HLH-2004: Diagnostic and therapeutic guidelines for hemophagocytic lymphohistiocytosis. <i>Pediatric Blood and Cancer</i> , 2007, 48, 124-131.	0.8	4,018
25	Haematopoietic stem cell transplantation in haemophagocytic lymphohistiocytosis. <i>British Journal of Haematology</i> , 2005, 129, 622-630.	1.2	206
26	Subtyping of natural killer cell cytotoxicity deficiencies in haemophagocytic lymphohistiocytosis provides therapeutic guidance. <i>British Journal of Haematology</i> , 2005, 129, 658-666.	1.2	33