

Catherine W Rehder

List of Publications by Year in descending order

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Version: 2024-02-01

33
papers

682
citations

840119

11
h-index

580395

25
g-index

33
all docs

33
docs citations

33
times ranked

1075
citing authors

#	ARTICLE	IF	CITATIONS
1	Chronic lymphocytic leukemia with t(6;14) (p21;q32) <i>CCND3</i> - <i>IGH</i> : <i>CCND3</i> rearrangement does not necessarily define a cyclin D1-negative mantle cell lymphoma. <i>Hematological Oncology</i> , 2022, 40, 112-115.	0.8	1
2	Circulating blastoid cells: acute leukemia, prolymphocytic leukemia, or something else?. <i>Annals of Hematology</i> , 2021, 100, 3047-3048.	0.8	0
3	Pseudo-“Pelger-Huët” anomaly in a 58-year-old woman treated with mycophenolate mofetil for flare of systemic lupus erythematosus. <i>International Journal of Laboratory Hematology</i> , 2021, 43, 3-4.	0.7	1
4	An Unusual Association: Total Anomalous Pulmonary Venous Return and Aortic Arch Obstruction in Patients with Cat Eye Syndrome. <i>Journal of Pediatric Genetics</i> , 2021, 10, 035-038.	0.3	5
5	Infantile leukemia—What factors determine its distinct biological nature? Clinicopathological study of 78 cases. <i>International Journal of Laboratory Hematology</i> , 2021, 43, 1117-1122.	0.7	0
6	Biclonal evolution of chronic lymphocytic leukaemia. <i>British Journal of Haematology</i> , 2021, 194, e64-e66.	1.2	1
7	Detection of a mosaic <i>CDKL5</i> deletion and inversion by optical genome mapping ends an exhaustive diagnostic odyssey. <i>Molecular Genetics & Genomic Medicine</i> , 2021, 9, e1665.	0.6	11
8	Interpretation of Incidental Genetic Findings Localizing to Genes Associated With Cardiac Channelopathies and Cardiomyopathies. <i>Circulation Genomic and Precision Medicine</i> , 2021, 14, e003200.	1.6	8
9	Conventional Cytogenetic Analysis of Hematologic Neoplasms: A 20-Year Review of Proficiency Test Results From the College of American Pathologists/American College of Medical Genetics and Genomics Cytogenetics Committee. <i>Archives of Pathology and Laboratory Medicine</i> , 2021, 145, 176-190.	1.2	3
10	Chronic lymphocytic leukemia/small lymphocytic lymphoma with secondary acquisition of t(11;14)(q13;q32)/ <i>CCND1</i> - <i>IGH</i> : a rare variant of Richter transformation to mantle cell lymphoma. <i>Clinical Lymphoma, Myeloma and Leukemia</i> , 2021, , .	0.2	0
11	Clinical and Molecular Disease Spectrum and Outcomes in Patients with Infantile-Onset Pompe Disease. <i>Journal of Pediatrics</i> , 2020, 216, 44-50.e5.	0.9	22
12	A comprehensive testing algorithm for the diagnosis of Fabry disease in males and females. <i>Molecular Genetics and Metabolism</i> , 2020, 130, 209-214.	0.5	26
13	Evaluation of X-Linked Adrenoleukodystrophy Newborn Screening in North Carolina. <i>JAMA Network Open</i> , 2020, 3, e1920356.	2.8	44
14	Sequential development of human herpes virus 8-positive diffuse large B-cell lymphoma and chronic myelomonocytic leukemia in a 59 year old female patient with hemoglobin SC disease. <i>Pathology Research and Practice</i> , 2019, 215, 152704.	1.0	4
15	The North Carolina Experience with Mucopolysaccharidosis Type I Newborn Screening. <i>Journal of Pediatrics</i> , 2019, 211, 193-200.e2.	0.9	22
16	Points to consider in the reevaluation and reanalysis of genomic test results: a statement of the American College of Medical Genetics and Genomics (ACMG). <i>Genetics in Medicine</i> , 2019, 21, 1267-1270.	1.1	147
17	Myeloid Neoplasms Following Solid Organ Transplantation. <i>American Journal of Clinical Pathology</i> , 2018, 149, 55-66.	0.4	11
18	Clinical and Molecular Variability in Patients with <i>PHKA2</i> Variants and Liver Phosphorylase b Kinase Deficiency. <i>JIMD Reports</i> , 2017, 37, 63-72.	0.7	9

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19	Sensitivity of whole exome sequencing in detecting infantile- and late-onset Pompe disease. <i>Molecular Genetics and Metabolism</i> , 2017, 122, 189-197.	0.5	21
20	Lineage Switch Between B-Lymphoblastic Leukemia and Acute Myeloid Leukemia Intermediated by "Occult" Myelodysplastic Neoplasm. <i>American Journal of Clinical Pathology</i> , 2017, 148, 136-147.	0.4	11
21	Sequential development of chronic myelogenous leukemia and primary myelofibrosis in a patient with history of large B-cell lymphoma treated with radiotherapy and chemotherapy: two myeloid neoplasms with distinct genotypic profiles suggestive of biclonality in a single individual. <i>Annals of Hematology</i> , 2016, 95, 1383-1386.	0.8	1
22	Severe Cardiomyopathy as the Isolated Presenting Feature in an Adult with Late-Onset Pompe Disease: A Case Report. <i>JIMD Reports</i> , 2016, 31, 79-83.	0.7	20
23	Chromosomal microarray analysis in clinical evaluation of neurodevelopmental disorders-reporting a novel deletion of SETDB1 and illustration of counseling challenge. <i>Pediatric Research</i> , 2016, 80, 371-381.	1.1	16
24	B-lymphoblastic leukemia in a patient with chronic lymphocytic leukemia: Sequential development of biclonal B-cell neoplasms over a 23-year period in a single individual. <i>Pathology Research and Practice</i> , 2016, 212, 1089-1093.	1.0	4
25	Epstein-Barr virus-negative diffuse large B cell lymphoma with aberrant expression of CD3 and other T cell-associated antigens: report of three cases with a review of the literature. <i>Annals of Hematology</i> , 2016, 95, 1671-1683.	0.8	9
26	Clinical laboratory experience of blood CRIM testing in infantile Pompe disease. <i>Molecular Genetics and Metabolism Reports</i> , 2015, 5, 76-79.	0.4	11
27	CRIM-negative infantile Pompe disease: characterization of immune responses in patients treated with ERT monotherapy. <i>Genetics in Medicine</i> , 2015, 17, 912-918.	1.1	54
28	T-cell prolymphocytic leukemia in a 63-year-old female with a pre-existing T-cell large granular lymphocytic leukemia: Metachronous T-cell leukemias with discordant subset restrictions (CD4) Tj ETQq0 0 0 rgBT 10verlock d0 Tf 50 3	1.0	10
29	Metachronous/concomitant B-cell neoplasms with discordant light-chain or heavy-chain isotype restrictions: evidence of distinct B-cell neoplasms rather than clonal evolutions. <i>Human Pathology</i> , 2014, 45, 2063-2076.	1.1	7
30	Algorithm for the Early Diagnosis and Treatment of Patients with Cross Reactive Immunologic Material-Negative Classic Infantile Pompe Disease: A Step towards Improving the Efficacy of ERT. <i>PLoS ONE</i> , 2013, 8, e67052.	1.1	93
31	Predicting cross-reactive immunological material (CRIM) status in Pompe disease using <i>GAA</i> mutations: Lessons learned from 10 years of clinical laboratory testing experience. <i>American Journal of Medical Genetics, Part C: Seminars in Medical Genetics</i> , 2012, 160C, 40-49.	0.7	110
32	Donor Cell Leukemia: A Clinicopathological Study of 9 Cases and a Comprehensive Review of Literature.. <i>Blood</i> , 2010, 116, 3466-3466.	0.6	0
33	Transformation of Indolent Small B-Cell Lymphoma to Histiocytic Sarcoma: Report of Two Cases with Molecular/Genetic Evidence of Clonal Relationship Between Two Morphologically and Immunophenotypically Distinctive Neoplasms.. <i>Blood</i> , 2009, 114, 3960-3960.	0.6	10