Jay L Patel

List of Publications by Year
in descending order

18	247	1163117 8	996975 15
papers	citations	h-index	g-index

18	18	18	529
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	AML with germline DDX41 variants is a clinicopathologically distinct entity with an indolent clinical course and favorable outcome. Leukemia, 2022, 36, 664-674.	7.2	32
2	Near Complete Response to Trametinib Treatment in Histiocytic Sarcoma Harboring a Somatic KRAS Mutation. Journal of the National Comprehensive Cancer Network: JNCCN, 2022, 20, 618-621.	4.9	7
3	Realâ€world diagnostic testing patterns for assessment of ring sideroblasts and SF3B1 mutations in patients with newly diagnosed lowerâ€risk myelodysplastic syndromes. International Journal of Laboratory Hematology, 2021, 43, 426-432.	1.3	3
4	Diagnostic Challenge and Clinical Dilemma: The Long Reach of Clonal Hematopoiesis. Clinical Chemistry, 2021, 67, 1062-1070.	3.2	0
5	KLF1/EKLF expression in acute leukemia is correlated with chromosomal abnormalities. Blood Cells, Molecules, and Diseases, 2020, 83, 102434.	1.4	4
6	Molecular Pathology Economics 101: An Overview of Molecular Diagnostics Coding, Coverage, and Reimbursement. Journal of Molecular Diagnostics, 2020, 22, 975-993.	2.8	11
7	Current Aspects of Clonal Hematopoiesis: Implications for Clinical Diagnosis. Annals of Laboratory Medicine, 2019, 39, 509-514.	2.5	10
8	Chronic myelomonocytic leukemia with ETV6-ABL1 rearrangement and SMC1A mutation. Cancer Genetics, 2019, 238, 31-36.	0.4	9
9	Bone marrow findings in metastatic melanoma, including role of BRAF immunohistochemistry. International Journal of Laboratory Hematology, 2019, 41, 550-560.	1.3	3
10	Cdx Report Program: Heterogeneity Revealed in Current Reporting Practices for Hemato-Oncology Companion Diagnostic (CDx) Markers in Multiple Countries. Blood, 2019, 134, 5837-5837.	1.4	0
11	The Clinical and Laboratory Features of Clonal Hematopoiesis of Indeterminate Potential. Advances in Molecular Pathology, 2018, 1, 37-42.	0.4	1
12	Use of Next Generation Sequencing Panel for Routine Diagnosis of Hereditary Hemolytic Anemias. Blood, 2018, 132, 2325-2325.	1.4	2
13	Disease Characteristics and Treatment of Adult Langerhans Cell Histiocytosis: A Single Center Experience. Blood, 2018, 132, 4315-4315.	1.4	1
14	Coexisting and cooperating mutations in NPM1 -mutated acute myeloid leukemia. Leukemia Research, 2017, 56, 7-12.	0.8	51
15	Clinical utility of nextâ€generation sequencing in the diagnosis of hereditary haemolytic anaemias. British Journal of Haematology, 2016, 174, 806-814.	2.5	71
16	Concurrent detection of targeted copy number variants and mutations using a myeloid malignancy next generation sequencing panel allows comprehensive genetic analysis using a single testing strategy. British Journal of Haematology, 2016, 173, 49-58.	2.5	27
17	An Unusual Case of Cutaneous Blastic Plasmacytoid Dendritic Cell Neoplasm With Concomitant B-Cell Lymphoproliferative Disorder. American Journal of Dermatopathology, 2011, 33, e31-e36.	0.6	5
18	Performance characteristics of an automated assay for the quantitation of CYFRA 21-1 in human serum. Clinical Biochemistry, 2010, 43, 1449-1452.	1.9	10