

Monika Pepek

List of Publications by Year in descending order

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Version: 2024-02-01

12
papers

66
citations

2257263

3
h-index

1719596

7
g-index

14
all docs

14
docs citations

14
times ranked

149
citing authors

#	ARTICLE	IF	CITATIONS
1	Nanoparticle-mediated decrease of lamin B1 pools promotes a TRF protein-based adaptive response in cultured cells. <i>Biomaterials</i> , 2015, 53, 107-116.	5.7	33
2	Differential Regulation of Telomeric Complex by BCR-ABL1 Kinase in Human Cellular Models of Chronic Myeloid Leukemia—From Single Cell Analysis to Next-Generation Sequencing. <i>Genes</i> , 2020, 11, 1145.	1.0	10
3	Epithelial Cells of Deep Infiltrating Endometriosis Harbor Mutations in Cancer Driver Genes. <i>Cells</i> , 2021, 10, 749.	1.8	8
4	Germline missense NF1 mutation in an elderly patient with a blastic plasmacytoid dendritic cell neoplasm. <i>International Journal of Hematology</i> , 2019, 110, 102-106.	0.7	5
5	ASXL1 Mutations Detectable at Diagnosis May Predict Response to Imatinib in Patients with Chronic Myeloid Leukemia. <i>Blood</i> , 2019, 134, 4148-4148.	0.6	4
6	Genomic landscape of human erythroleukemia K562 cell line, as determined by next-generation sequencing and cytogenetics. <i>Acta Haematologica Polonica</i> , 2017, 48, 343-349.	0.1	3
7	Predictive significance of selected gene mutations in relapsed and refractory chronic lymphocytic leukemia patients treated with ibrutinib. <i>European Journal of Haematology</i> , 2021, 106, 320-326.	1.1	2
8	Co-occurrence of unclassified myeloproliferative neoplasm and giant cell arteritis in a patient treated with allogeneic hematopoietic stem cell transplantation: a case report and literature review. <i>Central-European Journal of Immunology</i> , 2021, 46, 121-126.	0.4	1
9	Predictive Significance of Selected Gene Mutations Identified Using Next Generation Sequencing in Relapsed and Refractory Chronic Lymphocytic Leukemia Patients Treated with Ibrutinib. <i>Blood</i> , 2019, 134, 5456-5456.	0.6	0
10	First familial cases of type 2 congenital erythrocytosis (ECYT2) with a Chuvash pathogenic variant in VHL gene in Poland: example of the clinical utility of next-generation sequencing in diagnostics of orphan diseases. <i>Acta Haematologica Polonica</i> , 2020, 51, 220-225.	0.1	0
11	Wenetoklaks w monoterapii przewlekłej białaczki limfocytowej przed powtórnyim przeszczepieniem allogenicznym krwiotwórczych komórek macierzystych. <i>Hematologia</i> , 2020, 11, 95-100.	0.0	0
12	Clonal Hematopoiesis with Somatic Mutations in "AYA" Generation of Patients with Chronic Myeloid Leukemia. <i>Blood</i> , 2020, 136, 23-24.	0.6	0