Lukasz P Gondek

List of Publications by Year
in descending order

24 papers	807 citations	858243 12 h-index	721071 23 g-index
24 all docs	24 docs citations	24 times ranked	1288 citing authors

#	Article	IF	Citations
1	Nonmyeloablative Allogeneic Transplantation With Post-Transplant Cyclophosphamide for Acute Myeloid Leukemia With IDH Mutations: A Single Center Experience. Clinical Lymphoma, Myeloma and Leukemia, 2022, 22, 260-269.	0.2	4
2	Donor Clonal Hematopoiesis and Recipient Outcomes After Transplantation. Journal of Clinical Oncology, 2022, 40, 189-201.	0.8	79
3	Post-Transplantation Cyclophosphamide-Based Graft- versus-Host Disease Prophylaxis with Nonmyeloablative Conditioning for Blood or Marrow Transplantation for Myelofibrosis. Transplantation and Cellular Therapy, 2022, 28, 259.e1-259.e11.	0.6	11
4	Impact of diagnostic genetics on remission MRD and transplantation outcomes in older patients with AML. Blood, 2022, 139, 3546-3557.	0.6	37
5	Genomic landscape of myelodysplastic/myeloproliferative neoplasm can predict response to hypomethylating agent therapy. Leukemia and Lymphoma, 2022, 63, 1942-1948.	0.6	8
6	The role of the atypical chemokine receptor CCRL2 in myelodysplastic syndrome and secondary acute myeloid leukemia. Science Advances, 2022, 8, eabl8952.	4.7	7
7	Allelic complexity of KMT2A partial tandem duplications in acute myeloid leukemia and myelodysplastic syndromes. Blood Advances, 2022, 6, 4236-4240.	2.5	6
8	Clonal Hematopoiesis and the Risk of Hematologic Malignancies after Curative Therapies for Sickle Cell Disease. Journal of Clinical Medicine, 2022, 11, 3160.	1.0	2
9	Wholeâ€exome sequencing identifies functional classes of gene mutations associated with bone marrow failure in pediatric Fanconi Anemia patients. European Journal of Haematology, 2021, 107, 293-294.	1.1	0
10	Deep learning for diagnosis of acute promyelocytic leukemia via recognition of genomically imprinted morphologic features. Npj Precision Oncology, 2021, 5, 38.	2.3	18
11	Genderâ€related differences in the outcomes and genomic landscape of patients with myelodysplastic syndrome/myeloproliferative neoplasm overlap syndromes. British Journal of Haematology, 2021, 193, 1142-1150.	1.2	21
12	Clonal hematopoiesis and bone marrow failure syndromes. Best Practice and Research in Clinical Haematology, 2021, 34, 101273.	0.7	6
13	CHIP: is clonal hematopoiesis a surrogate for aging and other disease?. Hematology American Society of Hematology Education Program, 2021, 2021, 384-389.	0.9	8
14	Assessing clonal haematopoiesis: clinical burdens and benefits of diagnosing myelodysplastic syndrome precursor states. Lancet Haematology,the, 2020, 7, e73-e81.	2.2	45
15	Stem cell donors should be screened for CHIP. Blood Advances, 2020, 4, 784-788.	2.5	28
16	The diagnostic utility of targeted gene panel sequencing in discriminating etiologies of cytopenia. American Journal of Hematology, 2019, 94, 1141-1148.	2.0	33
17	Fibrinogen consumption and use of heparin are risk factors for delayed bleeding during acute promyelocytic leukemia induction. Leukemia Research, 2019, 83, 106174.	0.4	8
18	Hedgehog/GLI1 activation leads to leukemic transformation of myelodysplastic syndrome in vivo and GLI1 inhibition results in antitumor activity. Oncogene, 2019, 38, 687-698.	2.6	21

#	Article	IF	Citations
19	Hotspot SF3B1 mutations induce metabolic reprogramming and vulnerability to serine deprivation. Journal of Clinical Investigation, 2019, 129, 4708-4723.	3.9	41
20	Anthracyclineâ€induced acute myocarditis and ventricular fibrillation arrest. American Journal of Hematology, 2018, 93, 469-470.	2.0	4
21	Donor cell leukemia arising from clonal hematopoiesis after bone marrow transplantation. Leukemia, 2016, 30, 1916-1920.	3.3	79
22	Integration of Hedgehog and mutant FLT3 signaling in myeloid leukemia. Science Translational Medicine, 2015, 7, 291ra96.	5.8	50
23	I Walk the Line: How to Tell MDS From Other Bone Marrow Failure Conditions. Current Hematologic Malignancy Reports, 2014, 9, 389-399.	1.2	9
24	250K Single Nucleotide Polymorphism Array Karyotyping Identifies Acquired Uniparental Disomy and Homozygous Mutations, Including Novel Missense Substitutions of c-Cbl, in Myeloid Malignancies. Cancer Research, 2008, 68, 10349-10357.	0.4	282