

Magnus Tobiasson

List of Publications by Year in descending order

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Version: 2024-02-01

32
papers

1,384
citations

686830

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580395

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docs citations

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times ranked

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citing authors

#	ARTICLE	IF	CITATIONS
1	Implications of TP53 allelic state for genome stability, clinical presentation and outcomes in myelodysplastic syndromes. <i>Nature Medicine</i> , 2020, 26, 1549-1556.	15.2	372
2	Myelodysplastic Syndromes Are Propagated by Rare and Distinct Human Cancer Stem Cells In Vivo. <i>Cancer Cell</i> , 2014, 25, 794-808.	7.7	272
3	Molecular International Prognostic Scoring System for Myelodysplastic Syndromes. , 2022, 1, .		259
4	Integrative Genomics Identifies the Molecular Basis of Resistance to Azacitidine Therapy in Myelodysplastic Syndromes. <i>Cell Reports</i> , 2017, 20, 572-585.	2.9	99
5	SF3B1-initiating mutations in MDS-RSs target lymphomyeloid hematopoietic stem cells. <i>Blood</i> , 2017, 130, 881-890.	0.6	66
6	Myelodysplastic syndromes: moving towards personalized management. <i>Haematologica</i> , 2020, 105, 1765-1779.	1.7	52
7	Limited clinical efficacy of azacitidine in transfusion-dependent, growth factor-resistant, low- and Int-1-risk MDS: Results from the nordic NMDSG08A phase II trial. <i>Blood Cancer Journal</i> , 2014, 4, e189-e189.	2.8	48
8	Comprehensive mapping of the effects of azacitidine on DNA methylation, repressive/permissive histone marks and gene expression in primary cells from patients with MDS and MDS-related disease. <i>Oncotarget</i> , 2017, 8, 28812-28825.	0.8	42
9	Mutations in histone modulators are associated with prolonged survival during azacitidine therapy. <i>Oncotarget</i> , 2016, 7, 22103-22115.	0.8	37
10	Azacitidine in Lower-Risk Myelodysplastic Syndromes: A Meta-Analysis of Data from Prospective Studies. <i>Oncologist</i> , 2018, 23, 159-170.	1.9	27
11	High prevalence of restless legs syndrome among patients with polycythemia vera treated with venesection. <i>Medical Oncology</i> , 2010, 27, 105-107.	1.2	17
12	Early detection of relapse in patients with myelodysplastic syndrome after allo-SCT. <i>Bone Marrow Transplantation</i> , 2011, 46, 719-726.	1.3	15
13	Azacitidine induces profound genome-wide hypomethylation in primary myelodysplastic bone marrow cultures but may also reduce histone acetylation. <i>Leukemia</i> , 2014, 28, 411-413.	3.3	14
14	Treatment of myelodysplastic syndrome in the era of next-generation sequencing. <i>Journal of Internal Medicine</i> , 2019, 286, 41-62.	2.7	13
15	Angioimmunoblastic T-cell lymphoma and myelodysplastic syndrome with mutations in <i>TET2</i> , <i>DNMT3A</i> and <i>CUX1</i> – azacitidine induces only lymphoma remission. <i>Leukemia and Lymphoma</i> , 2019, 60, 3316-3319.	0.6	11
16	Male sex and the pattern of recurrent myeloid mutations are strong independent predictors of blood transfusion intensity in patients with myelodysplastic syndromes. <i>Leukemia</i> , 2019, 33, 522-527.	3.3	7
17	Clinical Impacts of Germline <i>DDX41</i> Mutations on Myeloid Neoplasms. <i>Blood</i> , 2020, 136, 38-40.	0.6	7
18	Randomized phase II study of azacitidine ± lenalidomide in higher-risk myelodysplastic syndromes and acute myeloid leukemia with a karyotype including Del(5q). <i>Leukemia</i> , 2022, 36, 1436-1439.	3.3	6

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19	Absence of a common founder mutation in patients with cooccurring myelodysplastic syndrome and plasma cell disorder. <i>Blood</i> , 2021, 137, 1260-1263.	0.6	5
20	Limited benefit in patients with MDS receiving venetoclax and azacitidine as a bridge to allogeneic stem cell transplantation. <i>Leukemia and Lymphoma</i> , 2022, 63, 755-758.	0.6	3
21	Mutation Profiles Identify Distinct Clusters of Lower Risk Myelodysplastic Syndromes with Unique Clinical and Biological Features and Clinical Endpoints. <i>Blood</i> , 2020, 136, 29-29.	0.6	2
22	Failure to reach hematopoietic allogeneic stem cell transplantation in patients with myelodysplastic syndromes planned for transplantation: a population-based study. <i>Bone Marrow Transplantation</i> , 2022, 57, 598-606.	1.3	2
23	The extent of residual WT HSPCs is associated with the degree of anemia in patients with <i>SF3B1</i>-mutated MDS-RS. <i>Blood Advances</i> , 2022, 6, 4705-4709.	2.5	2
24	Multicenter Next-Generation Sequencing Studies between Theory and Practice. <i>Journal of Molecular Diagnostics</i> , 2021, 23, 347-357.	1.2	1
25	Evaluation of Azacitidine in Transfusion-Dependent, Epo-Refractory Patients with Lower-Risk Myelodysplastic Syndrome,. <i>Blood</i> , 2011, 118, 3798-3798.	0.6	0
26	Allelic Methylation Levels of VTRNA2-1 Predict Outcome in Higher Risk MDS Patients Not Treated by Azacitidine.. <i>Blood</i> , 2012, 120, 2394-2394.	0.6	0
27	Diverse Genetic Lesions In Myelodysplastic Syndromes Originate Exclusively In Rare MDS Stem Cells. <i>Blood</i> , 2013, 122, 4195-4195.	0.6	0
28	Identification of a Prognostic Gene Expression Signature for AZA Response in MDS and CMML Patients. <i>Blood</i> , 2014, 124, 4601-4601.	0.6	0
29	Mutations in Histone Modulators and HOXA5 Methylation Levels Affects Survival in Azacitidine Treated MDS Patients. <i>Blood</i> , 2014, 124, 4613-4613.	0.6	0
30	Mutations in Histone Modulators Are Associated with Prolonged Survival during Azacitidine Therapy. <i>Blood</i> , 2015, 126, 2839-2839.	0.6	0
31	Prediction of Relapse after Allogeneic Stem Cell Transplantation Using Individualized Minimal Residual Markers; The Prospective Nordic Study NMDSG14B. <i>Blood</i> , 2020, 136, 5-6.	0.6	0
32	<i>Post-Treatment Clone Size Predicts Survival Independently of IPSS-R and Response after Azacitidine Therapy for MDS.</i>. <i>Blood</i> , 2020, 136, 12-13.	0.6	0