Antonio Di Stasi

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/6169432/publications.pdf

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22 2,772 11 papers citations h-index

23 23 23 3507 all docs docs citations times ranked citing authors

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g-index

#	Article	IF	CITATIONS
1	Inducible Apoptosis as a Safety Switch for Adoptive Cell Therapy. New England Journal of Medicine, 2011, 365, 1673-1683.	27.0	1,264
2	T lymphocytes coexpressing CCR4 and a chimeric antigen receptor targeting CD30 have improved homing and antitumor activity in a Hodgkin tumor model. Blood, 2009, 113, 6392-6402.	1.4	458
3	Epstein Barr virus–specific cytotoxic T lymphocytes expressing the anti-CD30ζ artificial chimeric T-cell receptor for immunotherapy of Hodgkin disease. Blood, 2007, 110, 2620-2630.	1.4	227
4	Inducible caspase-9 suicide gene controls adverse effects from alloreplete T cells after haploidentical stem cell transplantation. Blood, 2015, 125, 4103-4113.	1.4	188
5	Improving the safety of cell therapy products by suicide gene transfer. Frontiers in Pharmacology, 2014, 5, 254.	3.5	165
6	Long-term outcome after haploidentical stem cell transplant and infusion of T cells expressing the inducible caspase 9 safety transgene. Blood, 2014, 123, 3895-3905.	1.4	161
7	Review of the Results of WT1 Peptide Vaccination Strategies for Myelodysplastic Syndromes and Acute Myeloid Leukemia from Nine Different Studies. Frontiers in Immunology, 2015, 6, 36.	4.8	88
8	In Vitro Pre-Clinical Validation of Suicide Gene Modified Anti-CD33 Redirected Chimeric Antigen Receptor T-Cells for Acute Myeloid Leukemia. PLoS ONE, 2016, 11, e0166891.	2.5	72
9	Seatbelts in CAR therapy: How Safe Are CARS?. Pharmaceuticals, 2015, 8, 230-249.	3.8	42
10	Generation of Suicide Gene-Modified Chimeric Antigen Receptor-Redirected T-Cells for Cancer Immunotherapy. Methods in Molecular Biology, 2019, 1895, 57-73.	0.9	29
11	Regulated Apoptosis of Genetically Modified Hematopoietic Stem and Progenitor Cells Via an Inducible Caspase-9 Suicide Gene in Rhesus Macaques. Stem Cells, 2015, 33, 91-100.	3.2	28
12	Gene Therapy to Improve Migration of T Cells to the Tumor Site. Methods in Molecular Biology, 2010, 651, 103-118.	0.9	9
13	Clinical-scale manufacturing of $\hat{I}^3\hat{I}$ cells for protection against infection and disease recurrence following haploidentical peripheral blood stem cell transplantation and cyclophosphamide gyhd prophylaxis. Bone Marrow Transplantation, 2018, 53, 766-769.	2.4	8
14	Tumor lysis syndrome and infectious complications during treatment with venetoclax combined with azacitidine or decitabine in patients with acute myeloid leukemia. Leukemia Research, 2022, 117, 106844.	0.8	7
15	Exploiting Cell Death Pathways for Inducible Cell Elimination to Modulate Graft-versus-Host-Disease. Biomedicines, 2017, 5, 30.	3.2	5
16	Impact of high-dose steroid premedication on the outcome of myeloablative T-cell replete haploidentical peripheral blood stem cell transplant. Bone Marrow Transplantation, 2018, 53, 1345-1348.	2.4	4
17	Impact of access to care on 1-year mortality following allogeneic blood or marrow transplantation. Bone Marrow Transplantation, 2021, 56, 1364-1372.	2.4	4
18	Phase <scp>II</scp> clinical trial of one dose of postâ€transplant cyclophosphamide for graft versus host disease prevention following myeloablative, peripheral blood stem cell, matchedâ€unrelated donor transplantation. American Journal of Hematology, 2021, 96, E396-E398.	4.1	4

#	Article	IF	CITATIONS
19	Observation Versus Immediate Reinduction for Acute Myeloid Leukemia Patients With Indeterminate Day 14 Bone Marrow Results. Clinical Lymphoma, Myeloma and Leukemia, 2020, 20, 31-38.	0.4	3
20	IDH1 inhibitor-induced neutrophilic dermatosis in a patient with acute myeloid leukemia. Cancer Treatment and Research Communications, 2022, 31, 100560.	1.7	2
21	Passenger Lymphocyte Syndrome and Autoimmune Hypothyroidism Following Hematopoietic Stem Cell Transplantation. Case Reports in Immunology, 2022, 2022, 1-6.	0.4	1
22	Survival outcomes of patients with relapsed or refractory acute myeloid leukemia after venetoclax combined with hypomethylating agents Journal of Clinical Oncology, 2022, 40, e18808-e18808.	1.6	0