

Martin A Pule

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/6390146/publications.pdf>

Version: 2024-02-01

48
papers

6,275
citations

279798

23
h-index

276875

41
g-index

49
all docs

49
docs citations

49
times ranked

6218
citing authors

#	ARTICLE	IF	CITATIONS
1	Chimeric antigen receptor T cells for gamma-δ T cell malignancies. <i>Leukemia</i> , 2022, 36, 577-579.	7.2	8
2	Paediatric Strategy Forum for medicinal product development of chimeric antigen receptor T-cells in children and adolescents with cancer. <i>European Journal of Cancer</i> , 2022, 160, 112-133.	2.8	24
3	A compact and simple method of achieving differential transgene expression by exploiting translational readthrough. <i>BioTechniques</i> , 2022, 72, 143-154.	1.8	2
4	Anti-CCR9 chimeric antigen receptor T cells for T-cell acute lymphoblastic leukemia. <i>Blood</i> , 2022, 140, 25-37.	1.4	29
5	Intratumoral IL-12 delivery empowers CAR-T cell immunotherapy in a pre-clinical model of glioblastoma. <i>Nature Communications</i> , 2021, 12, 444.	12.8	150
6	Clonal expansion of T memory stem cells determines early anti-leukemic responses and long-term CAR T cell persistence in patients. <i>Nature Cancer</i> , 2021, 2, 629-642.	13.2	59
7	Durable Responses and Low Toxicity After Fast Off-Rate CD19 Chimeric Antigen Receptor-T Therapy in Adults With Relapsed or Refractory B-Cell Acute Lymphoblastic Leukemia. <i>Journal of Clinical Oncology</i> , 2021, 39, 3352-3363.	1.6	59
8	CAR T cells with dual targeting of CD19 and CD22 in pediatric and young adult patients with relapsed or refractory B-cell acute lymphoblastic leukemia: a phase 1 trial. <i>Nature Medicine</i> , 2021, 27, 1797-1805.	30.7	125
9	Depletion of T cells via Inducible Caspase 9 Increases Safety of Adoptive T-Cell Therapy Against Chronic Hepatitis B. <i>Frontiers in Immunology</i> , 2021, 12, 734246.	4.8	15
10	Functional antibody and T cell immunity following SARS-CoV-2 infection, including by variants of concern, in patients with cancer: the CAPTURE study. <i>Nature Cancer</i> , 2021, 2, 1321-1337.	13.2	66
11	Tunable control of CAR T cell activity through tetracycline mediated disruption of protein-protein interaction. <i>Scientific Reports</i> , 2021, 11, 21902.	3.3	12
12	Siglec-6 CAR T: magic bullet for a moving target. <i>Blood</i> , 2021, 138, 1786-1787.	1.4	1
13	Antitumor activity without on-target off-tumor toxicity of GD2-chimeric antigen receptor T cells in patients with neuroblastoma. <i>Science Translational Medicine</i> , 2020, 12, .	12.4	108
14	A primer set for the rapid isolation of scFv fragments against cell surface antigens from immunised rats. <i>Scientific Reports</i> , 2020, 10, 19168.	3.3	4
15	Enhanced CAR T cell expansion and prolonged persistence in pediatric patients with ALL treated with a low-affinity CD19 CAR. <i>Nature Medicine</i> , 2019, 25, 1408-1414.	30.7	394
16	Anti-CD1a CAR T cells to selectively target T-ALL. <i>Blood</i> , 2019, 133, 2246-2247.	1.4	9
17	Imaging of X-Ray-Excited Emissions from Quantum Dots and Biological Tissue in Whole Mouse. <i>Scientific Reports</i> , 2019, 9, 19223.	3.3	10
18	Clonal Dynamics of Early Responder and Long-Term Surviving CAR-T Cells in Humans. <i>Blood</i> , 2019, 134, 52-52.	1.4	2

#	ARTICLE	IF	CITATIONS
19	An APRIL-based chimeric antigen receptor for dual targeting of BCMA and TACI in multiple myeloma. <i>Blood</i> , 2018, 131, 746-758.	1.4	131
20	Vaccination to improve the persistence of CD19CAR gene-modified T cells in relapsed pediatric acute lymphoblastic leukemia. <i>Leukemia</i> , 2017, 31, 1087-1095.	7.2	64
21	Targeting the T cell receptor β -chain constant region for immunotherapy of T cell malignancies. <i>Nature Medicine</i> , 2017, 23, 1416-1423.	30.7	196
22	An Optimized GD2-Targeting Retroviral Cassette for More Potent and Safer Cellular Therapy of Neuroblastoma and Other Cancers. <i>PLoS ONE</i> , 2016, 11, e0152196.	2.5	57
23	Automated manufacturing of chimeric antigen receptor T cells for adoptive immunotherapy using CliniMACS Prodigy. <i>Cytotherapy</i> , 2016, 18, 1002-1011.	0.7	174
24	TALEN-Mediated Inactivation of PD-1 in Tumor-Reactive Lymphocytes Promotes Intratumoral T-cell Persistence and Rejection of Established Tumors. <i>Cancer Research</i> , 2016, 76, 2087-2093.	0.9	67
25	A highly compact epitope-based marker/suicide gene for easier and safer T-cell therapy. <i>Blood</i> , 2014, 124, 1277-1287.	1.4	308
26	Towards gene therapy for EBV-associated posttransplant lymphoma with genetically modified EBV-specific cytotoxic T cells. <i>Blood</i> , 2014, 124, 2514-2522.	1.4	51
27	Comparison of Different Suicide-Gene Strategies for the Safety Improvement of Genetically Manipulated T Cells. <i>Human Gene Therapy Methods</i> , 2012, 23, 376-386.	2.1	102
28	Antitumor activity and long-term fate of chimeric antigen receptor ⁺ positive T cells in patients with neuroblastoma. <i>Blood</i> , 2011, 118, 6050-6056.	1.4	984
29	Long-term outcome of EBV-specific T-cell infusions to prevent or treat EBV-related lymphoproliferative disease in transplant recipients. <i>Blood</i> , 2010, 115, 925-935.	1.4	721
30	Cytotoxic T cells transduced with chimeric anti-CD19 receptors prevent engraftment of primary lymphoblastic leukemia in vivo. <i>Leukemia</i> , 2010, 24, 1080-1084.	7.2	16
31	Virus-specific T cells engineered to coexpress tumor-specific receptors: persistence and antitumor activity in individuals with neuroblastoma. <i>Nature Medicine</i> , 2008, 14, 1264-1270.	30.7	1,063
32	Flanking-sequence exponential anchored ⁺ polymerase chain reaction amplification: a sensitive and highly specific method for detecting retroviral integrant ⁺ host ⁺ junction sequences. <i>Cytotherapy</i> , 2008, 10, 526-539.	0.7	12
33	Co-expression of cytokine and suicide genes to enhance the activity and safety of tumor-specific cytotoxic T lymphocytes. <i>Blood</i> , 2007, 110, 2793-2802.	1.4	157
34	Gene transfer: methods and applications. , 2006, , 661-678.		0
35	Transgenic Expression of IL15 Selectively Expands Antigen Specific Cytotoxic T Cells (CTLs) Enhancing Their Anti-Tumor Effect In Vivo.. <i>Blood</i> , 2006, 108, 1721-1721.	1.4	0
36	Transgenic Expression of Inducible Caspase9 Suicide Gene for In Vivo Elimination of Antigen Specific Cytotoxic T Cells (CTLs) Engineered To Produce Cytokines.. <i>Blood</i> , 2006, 108, 137-137.	1.4	0

#	ARTICLE	IF	CITATIONS
37	An inducible caspase 9 safety switch for T-cell therapy. Blood, 2005, 105, 4247-4254.	1.4	607
38	A chimeric T cell antigen receptor that augments cytokine release and supports clonal expansion of primary human T cells. Molecular Therapy, 2005, 12, 933-941.	8.2	426
39	T-Cells Redirected Against the kappa Light Chain of Human Immunoglobulins Target Mature B Cell Derived Malignancies In Vitro and In Vivo.. Blood, 2005, 106, 612-612.	1.4	1
40	Genetically Modified Her2-Specific T Cells Recognize Low and High Her2 Expressing Breast Cancer Cells.. Blood, 2005, 106, 5540-5540.	1.4	0
41	Three-Module Signaling Endo-Domain Artificial T-Cell Receptor Which Transmits CD28, OX40 and CD3-Î¶ Signals Enhances IL-2 Release and Proliferative Response in Transduced Primary T-Cells.. Blood, 2004, 104, 1747-1747.	1.4	2
42	Fas Down-Modulation in Epstein Barr Virus (EBV)-Specific Cytotoxic T-Lymphocytes (CTLs) Reduces Their Sensitivity to Fas/FasL-Induced Apoptosis.. Blood, 2004, 104, 2647-2647.	1.4	0
43	Flanking-Sequence Exponential Anchored (FLEA) PCR - a Sensitive and Highly Specific Method for Detecting Retroviral Integrant-Host-Junction Sequences.. Blood, 2004, 104, 2112-2112.	1.4	1
44	Multiple Integration Events into Several Putative Oncogenes Was Required To Cause Leukemogenesis in Two Primate Recipients of RCR Contaminated Stem-Cells.. Blood, 2004, 104, 2102-2102.	1.4	3
45	Epstein Barr Virus (EBV)-Specific Cytotoxic T Lymphocytes (CTL) Expressing an Anti-CD30 Chimeric T Cell Receptor (CTCR) for the Treatment of Hodgkinâ€™s Disease (HD).. Blood, 2004, 104, 745-745.	1.4	0
46	Inducible Caspase 9 as a Safety Switch in Genetically Modified Cytotoxic T Cells.. Blood, 2004, 104, 1743-1743.	1.4	0
47	Artificial T-cell receptors. Cytotherapy, 2003, 5, 211-226.	0.7	53
48	Genetically engineered T-cells for adoptive immunotherapy. Current Opinion in Molecular Therapeutics, 2002, 4, 467-75.	2.8	2