

# Cliona M Rooney

## List of Publications by Year in descending order

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148  
papers

15,613  
citations

36271

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19169

118  
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148  
docs citations

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times ranked

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citing authors

#	ARTICLE	IF	CITATIONS
1	Epstein Barr virus- positive B-cell lymphoma is highly vulnerable to MDM2 inhibitors in vivo. <i>Blood Advances</i> , 2022, 6, 891-901.	2.5	2
2	Donor-derived multiple leukemia antigen-specific T-cell therapy to prevent relapse after transplant in patients with ALL. <i>Blood</i> , 2022, 139, 2706-2711.	0.6	13
3	A Costimulatory CAR Improves TCR-based Cancer Immunotherapy. <i>Cancer Immunology Research</i> , 2022, 10, 512-524.	1.6	12
4	Long-term follow-up for the development of subsequent malignancies in patients treated with genetically modified IECs. <i>Blood</i> , 2022, 140, 16-24.	0.6	14
5	Multi-antigen-targeted T-cell therapy to treat patients with relapsed/refractory breast cancer. <i>Therapeutic Advances in Medical Oncology</i> , 2022, 14, 175883592211071.	1.4	6
6	Adoptive T-Cell Therapy for Epstein-Barr Virus-Related Lymphomas. <i>Journal of Clinical Oncology</i> , 2021, 39, 514-524.	0.8	18
7	CD70-specific CAR T cells have potent activity against acute myeloid leukemia without HSC toxicity. <i>Blood</i> , 2021, 138, 318-330.	0.6	98
8	T-Cell Therapy for Lymphoma Using Nonengineered Multiantigen-Targeted T Cells Is Safe and Produces Durable Clinical Effects. <i>Journal of Clinical Oncology</i> , 2021, 39, 1415-1425.	0.8	30
9	The National Heart, Lung, and Blood Institute-funded Production Assistance for Cellular Therapies (PACT) program: Eighteen years of cell therapy. <i>Clinical and Translational Science</i> , 2021, 14, 2099-2110.	1.5	1
10	Autologous EBV-specific T cell treatment results in sustained responses in patients with advanced extranodal NK/T lymphoma: results of a multicenter study. <i>Annals of Hematology</i> , 2021, 100, 2529-2539.	0.8	12
11	Donor-Derived Adoptive T-Cell Therapy Targeting Multiple Tumor Associated Antigens to Prevent Post-Transplant Relapse in Patients with ALL. <i>Blood</i> , 2021, 138, 471-471.	0.6	0
12	Safety and Efficacy of Off-the-Shelf CD30.CAR-Modified Epstein-Barr Virus-Specific T Cells in Patients with CD30-Positive Lymphoma. <i>Blood</i> , 2021, 138, 1763-1763.	0.6	6
13	Anti-CD30 CAR-T Cell Therapy in Relapsed and Refractory Hodgkin Lymphoma. <i>Journal of Clinical Oncology</i> , 2020, 38, 3794-3804.	0.8	235
14	Identification of protective T-cell antigens for smallpox vaccines. <i>Cytotherapy</i> , 2020, 22, 642-652.	0.3	10
15	Oncolytic measles virus therapy enhances tumor antigen-specific T-cell responses in patients with multiple myeloma. <i>Leukemia</i> , 2020, 34, 3310-3322.	3.3	64
16	A Bank of CD30.CAR-Modified, Epstein-Barr Virus-Specific T Cells That Lacks Host Reactivity and Resists Graft Rejection for Patients with CD30-Positive Lymphoma. <i>Blood</i> , 2020, 136, 16-16.	0.6	6
17	A strategy to protect off-the-shelf cell therapy products using virus-specific T-cells engineered to eliminate alloreactive T-cells. <i>Journal of Translational Medicine</i> , 2019, 17, 240.	1.8	18
18	T-Cell Receptor Stimulation Enhances the Expansion and Function of CD19 Chimeric Antigen Receptor-Expressing T Cells. <i>Clinical Cancer Research</i> , 2019, 25, 7340-7350.	3.2	32

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19	Tâ€cell receptor sequencing demonstrates persistence of virusâ€specific T cells after antiviral immunotherapy. <i>British Journal of Haematology</i> , 2019, 187, 206-218.	1.2	29
20	Current challenges for CAR Tâ€cell therapy of acute myeloid leukemia. <i>Transfusion</i> , 2019, 59, 1171-1173.	0.8	7
21	Epstein-Barr Virus (EBV)-derived BART1 encodes CD4- and CD8-restricted epitopes as targets for T-cell immunotherapy. <i>Cytotherapy</i> , 2019, 21, 212-223.	0.3	16
22	NK Cells Expressing a Chimeric Activating Receptor Eliminate MDSCs and Rescue Impaired CAR-T Cell Activity against Solid Tumors. <i>Cancer Immunology Research</i> , 2019, 7, 363-375.	1.6	180
23	CD70-Specific CAR T Cells Have Potent Activity Against Acute Myeloid Leukemia (AML) without HSC Toxicity. <i>Blood</i> , 2019, 134, 1932-1932.	0.6	3
24	Transposon-modified antigen-specific T lymphocytes for sustained therapeutic protein delivery in vivo. <i>Nature Communications</i> , 2018, 9, 1325.	5.8	16
25	Enhanced Expression of Anti-CD19 Chimeric Antigen Receptor in piggyBac Transposon-Engineered T Cells. <i>Molecular Therapy - Methods and Clinical Development</i> , 2018, 8, 131-140.	1.8	49
26	Tumor-Specific T-Cells Engineered to Overcome Tumor Immune Evasion Induce Clinical Responses in Patients With Relapsed Hodgkin Lymphoma. <i>Journal of Clinical Oncology</i> , 2018, 36, 1128-1139.	0.8	137
27	Chimeric Antigen Receptor Signaling Domains Differentially Regulate Proliferation and Native T Cell Receptor Function in Virus-Specific T Cells. <i>Frontiers in Medicine</i> , 2018, 5, 343.	1.2	12
28	EBV/LMP-specific T cells maintain remissions of T- and B-cell EBV lymphomas after allogeneic bone marrow transplantation. <i>Blood</i> , 2018, 132, 2351-2361.	0.6	49
29	Inâ€Vivo Fate and Activity of Second- versus Third-Generation CD19-Specific CAR-T Cells in B Cell Non-Hodgkinâ€™s Lymphomas. <i>Molecular Therapy</i> , 2018, 26, 2727-2737.	3.7	180
30	Modeling cytokine release syndrome. <i>Nature Medicine</i> , 2018, 24, 705-706.	15.2	18
31	A New Method for Reactivating and Expanding T Cells Specific for <i>Rhizopus oryzae</i> . <i>Molecular Therapy - Methods and Clinical Development</i> , 2018, 9, 305-312.	1.8	24
32	Strategies for enhancing adoptive T-cell immunotherapy against solid tumors using engineered cytokine signaling and other modalities. <i>Expert Opinion on Biological Therapy</i> , 2018, 18, 653-664.	1.4	26
33	CD30-Chimeric Antigen Receptor (CAR) T Cells for Therapy of Hodgkin Lymphoma (HL). <i>Blood</i> , 2018, 132, 680-680.	0.6	20
34	Improving Chimeric Antigen Receptor-Modified T Cell Function by Reversing the Immunosuppressive Tumor Microenvironment of Pancreatic Cancer. <i>Molecular Therapy</i> , 2017, 25, 249-258.	3.7	217
35	Vaccination Targeting Native Receptors to Enhance the Function and Proliferation of Chimeric Antigen Receptor (CAR)-Modified T Cells. <i>Clinical Cancer Research</i> , 2017, 23, 3499-3509.	3.2	76
36	HER2-Specific Chimeric Antigen Receptorâ€Modified Virus-Specific T Cells for Progressive Glioblastoma. <i>JAMA Oncology</i> , 2017, 3, 1094.	3.4	608

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37	Comprehensive Approach for Identifying the T Cell Subset Origin of CD3 and CD28 Antibody-Activated Chimeric Antigen Receptor-Modified T Cells. <i>Journal of Immunology</i> , 2017, 199, 348-362.	0.4	41
38	CAR T Cells Administered in Combination with Lymphodepletion and PD-1 Inhibition to Patients with Neuroblastoma. <i>Molecular Therapy</i> , 2017, 25, 2214-2224.	3.7	378
39	Tonic 4-1BB Costimulation in Chimeric Antigen Receptors Impedes T Cell Survival and Is Vector-Dependent. <i>Cell Reports</i> , 2017, 21, 17-26.	2.9	203
40	Constitutive Signaling from an Engineered IL7 Receptor Promotes Durable Tumor Elimination by Tumor-Redirected T Cells. <i>Cancer Discovery</i> , 2017, 7, 1238-1247.	7.7	204
41	Phase 1 clinical trial of adoptive immunotherapy using "off-the-shelf" activated natural killer cells in patients with refractory and relapsed acute myeloid leukemia. <i>Cytotherapy</i> , 2017, 19, 1225-1232.	0.3	117
42	Treatment of Acute Myeloid Leukemia with T Cells Expressing Chimeric Antigen Receptors Directed to C-type Lectin-like Molecule 1. <i>Molecular Therapy</i> , 2017, 25, 2202-2213.	3.7	109
43	EBV-Directed T Cell Therapeutics for EBV-Associated Lymphomas. <i>Methods in Molecular Biology</i> , 2017, 1532, 255-265.	0.4	16
44	Clinical and immunological responses after CD30-specific chimeric antigen receptor-redirected lymphocytes. <i>Journal of Clinical Investigation</i> , 2017, 127, 3462-3471.	3.9	301
45	Off-the-Shelf Virus-Specific T Cells to Treat BK Virus, Human Herpesvirus 6, Cytomegalovirus, Epstein-Barr Virus, and Adenovirus Infections After Allogeneic Hematopoietic Stem-Cell Transplantation. <i>Journal of Clinical Oncology</i> , 2017, 35, 3547-3557.	0.8	367
46	Expansion of HER2-CAR T cells after lymphodepletion and clinical responses in patients with advanced sarcoma. <i>Journal of Clinical Oncology</i> , 2017, 35, 10508-10508.	0.8	32
47	Fine-tuning the CAR spacer improves T-cell potency. <i>Oncotarget</i> , 2016, 5, e1253656.	2.1	137
48	Highly Efficient Genome Editing of Murine and Human Hematopoietic Progenitor Cells by CRISPR/Cas9. <i>Cell Reports</i> , 2016, 17, 1453-1461.	2.9	223
49	Large-Scale Culture and Genetic Modification of Human Natural Killer Cells for Cellular Therapy. <i>Methods in Molecular Biology</i> , 2016, 1441, 195-202.	0.4	20
50	Adoptive immunotherapy for primary immunodeficiency disorders with virus-specific T lymphocytes. <i>Journal of Allergy and Clinical Immunology</i> , 2016, 137, 1498-1505.e1.	1.5	117
51	Direct Comparison of In Vivo Fate of Second and Third-Generation CD19-Specific Chimeric Antigen Receptor (CAR)-T Cells in Patients with B-Cell Lymphoma: Reversal of Toxicity from Tonic Signaling. <i>Blood</i> , 2016, 128, 1851-1851.	0.6	22
52	Administration of Most Closely HLA-Matched Multivirus-Specific T Cells for the Treatment of EBV, CMV, AdV, HHV6, and BKV Post Allogeneic Hematopoietic Stem Cell Transplant. <i>Blood</i> , 2016, 128, 501-501.	0.6	2
53	Phase 1 Clinical Trial of Adoptive Immunotherapy Using "Off-the-Shelf" Activated Natural Killer Cells (aNK) in Patients with Refractory/Relapsed Acute Myeloid Leukemia. <i>Blood</i> , 2016, 128, 1649-1649.	0.6	1
54	Fast and Efficient Gene Editing in Human Hematopoietic Cells. <i>Blood</i> , 2016, 128, 4704-4704.	0.6	0

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55	Tumor indoleamine 2,3-dioxygenase (IDO) inhibits CD19-CAR T cells and is downregulated by lymphodepleting drugs. <i>Blood</i> , 2015, 125, 3905-3916.	0.6	260
56	Anti-Tumor Effects after Adoptive Transfer of IL-12 Transposon-Modified Murine Splenocytes in the OT-I-Melanoma Mouse Model. <i>PLoS ONE</i> , 2015, 10, e0140744.	1.1	11
57	Expanded Cytotoxic T-cell Lymphocytes Target the Latent HIV Reservoir. <i>Journal of Infectious Diseases</i> , 2015, 212, 258-263.	1.9	86
58	Antigen-specific T cell therapies for cancer: Figure 1.. <i>Human Molecular Genetics</i> , 2015, 24, R67-R73.	1.4	32
59	CMV-specific T cells generated from naïve T cells recognize atypical epitopes and may be protective in vivo. <i>Science Translational Medicine</i> , 2015, 7, 285ra63.	5.8	93
60	Inducible caspase-9 suicide gene controls adverse effects from alloplete T cells after haploidentical stem cell transplantation. <i>Blood</i> , 2015, 125, 4103-4113.	0.6	188
61	Peripheral Blood-Derived Virus-Specific Memory Stem T Cells Mature to Functional Effector Memory Subsets with Self-Renewal Potency. <i>Journal of Immunology</i> , 2015, 194, 5559-5567.	0.4	36
62	Human Epidermal Growth Factor Receptor 2 (HER2) -Specific Chimeric Antigen Receptor-Modified T Cells for the Immunotherapy of HER2-Positive Sarcoma. <i>Journal of Clinical Oncology</i> , 2015, 33, 1688-1696.	0.8	778
63	Evaluating the potential for undesired genomic effects of the <i>piggyBac</i> transposon system in human cells. <i>Nucleic Acids Research</i> , 2015, 43, 1770-1782.	6.5	44
64	Broadly-specific Cytotoxic T Cells Targeting Multiple HIV Antigens Are Expanded From HIV+ Patients: Implications for Immunotherapy. <i>Molecular Therapy</i> , 2015, 23, 387-395.	3.7	46
65	Adoptive T-Cell Immunotherapy. <i>Current Topics in Microbiology and Immunology</i> , 2015, 391, 427-454.	0.7	48
66	Phase 1 Study of Intratumoral Pexa-Vec (JX-594), an Oncolytic and Immunotherapeutic Vaccinia Virus, in Pediatric Cancer Patients. <i>Molecular Therapy</i> , 2015, 23, 602-608.	3.7	132
67	Graft Versus Leukemia Response Without Graft-versus-host Disease Elicited By Adoptively Transferred Multivirus-specific T-cells. <i>Molecular Therapy</i> , 2015, 23, 179-183.	3.7	28
68	Safety of Multiple Doses of CAR T Cells. <i>Blood</i> , 2015, 126, 4425-4425.	0.6	5
69	Autologous HER2 CMV bispecific CAR T cells for progressive glioblastoma: Results from a phase I clinical trial.. <i>Journal of Clinical Oncology</i> , 2015, 33, 3008-3008.	0.8	44
70	Adoptively-Transferred EBV-Specific T Cells to Prevent or Treat EBV-Related Lymphoproliferative Disease in Allogeneic HSCT Recipients - a Single Center Experience Spanning 22 Years. <i>Blood</i> , 2015, 126, 1926-1926.	0.6	0
71	Optimizing the production of suspension cells using the G-Rex -series. <i>Molecular Therapy - Methods and Clinical Development</i> , 2014, 1, 14015.	1.8	71
72	Reversal of Tumor Immune Inhibition Using a Chimeric Cytokine Receptor. <i>Molecular Therapy</i> , 2014, 22, 1211-1220.	3.7	145

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73	Systemic Inflammatory Response Syndrome After Administration of Unmodified T Lymphocytes. <i>Molecular Therapy</i> , 2014, 22, 1134-1138.	3.7	28
74	Harnessing the Immune System to Potentiate Oncolytics. <i>Molecular Therapy</i> , 2014, 22, 239-240.	3.7	1
75	Can Treg elimination enhance NK cell therapy for AML?. <i>Blood</i> , 2014, 123, 3848-3849.	0.6	7
76	T lymphocytes targeting native receptors. <i>Immunological Reviews</i> , 2014, 257, 39-55.	2.8	34
77	Reply to S. Yuan et al. <i>Journal of Clinical Oncology</i> , 2014, 32, 2820-2821.	0.8	0
78	Activity of Broad-Spectrum T Cells as Treatment for AdV, EBV, CMV, BKV, and HHV6 Infections after HSCT. <i>Science Translational Medicine</i> , 2014, 6, 242ra83.	5.8	357
79	Sustained Complete Responses in Patients With Lymphoma Receiving Autologous Cytotoxic T Lymphocytes Targeting Epstein-Barr Virus Latent Membrane Proteins. <i>Journal of Clinical Oncology</i> , 2014, 32, 798-808.	0.8	433
80	Anti-leukemic potency of piggyBac-mediated CD19-specific T cells against refractory Philadelphia chromosome-positive acute lymphoblastic leukemia. <i>Cytotherapy</i> , 2014, 16, 1257-1269.	0.3	42
81	Kinetics of Tumor Destruction by Chimeric Antigen Receptor-modified T Cells. <i>Molecular Therapy</i> , 2014, 22, 623-633.	3.7	113
82	Expansion and Homing of Adoptively Transferred Human Natural Killer Cells in Immunodeficient Mice Varies with Product Preparation and In Vivo Cytokine Administration: Implications for Clinical Therapy. <i>Biology of Blood and Marrow Transplantation</i> , 2014, 20, 1252-1257.	2.0	71
83	Closely related T-memory stem cells correlate with in vivo expansion of CAR-CD19-T cells and are preserved by IL-7 and IL-15. <i>Blood</i> , 2014, 123, 3750-3759.	0.6	534
84	Combinatorial treatment with oncolytic adenovirus and helper-dependent adenovirus augments adenoviral cancer gene therapy. <i>Molecular Therapy - Oncolytics</i> , 2014, 1, 14008.	2.0	19
85	Long-term outcome after haploidentical stem cell transplant and infusion of T cells expressing the inducible caspase 9 safety transgene. <i>Blood</i> , 2014, 123, 3895-3905.	0.6	161
86	Clinical Grade Purification and Expansion of Natural Killer Cells. <i>Critical Reviews in Oncogenesis</i> , 2014, 19, 121-132.	0.2	56
87	T Cells Expressing CD19-Specific Chimeric Antigen Receptors Are Inhibited By Indoleamine 2,3-Dioxygenase in Tumors. <i>Blood</i> , 2014, 124, 2434-2434.	0.6	0
88	Safety and clinical efficacy of rapidly-generated trivirus-directed T cells as treatment for adenovirus, EBV, and CMV infections after allogeneic hematopoietic stem cell transplant. <i>Molecular Therapy</i> , 2013, 21, 2113-2121.	3.7	200
89	Multicenter study of banked third-party virus-specific T cells to treat severe viral infections after hematopoietic stem cell transplantation. <i>Blood</i> , 2013, 121, 5113-5123.	0.6	507
90	Infusion of donor-derived CD19-redirected virus-specific T cells for B-cell malignancies relapsed after allogeneic stem cell transplant: a phase 1 study. <i>Blood</i> , 2013, 122, 2965-2973.	0.6	470

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91	Clinical Responses In Patients Infused With T Lymphocytes Redirected To Target Î² <sub>2</sub> -Light Immunoglobulin Chain. <i>Blood</i> , 2013, 122, 506-506.	0.6	6
92	T-cell therapy in the treatment of post-transplant lymphoproliferative disease. <i>Nature Reviews Clinical Oncology</i> , 2012, 9, 510-519.	12.5	230
93	Moving Successful Virus-specific T-cell Therapy for Hematopoietic Stem Cell Recipients to Late Phase Clinical Trials. <i>Molecular Therapy - Nucleic Acids</i> , 2012, 1, e55.	2.3	10
94	Adoptive transfer of virus-directed T cells: will this fly for flu?. <i>Cytotherapy</i> , 2012, 14, 133-134.	0.3	2
95	Optimal Xenogeneic Adoptive Transfer of Human NK Cells: Fresh NK Cells and IL-15 Administration Are Superior to Frozen NK Cells and IL-2. <i>Blood</i> , 2012, 120, 346-346.	0.6	1
96	Multicenter Study of "off-the-Shelf" Third Party Virus-Specific T Cells (VSTs) to Treat Adenovirus (Adv), Cytomegalovirus (CMV) or Epstein Barr Virus (EBV) Infection After Hemopoietic Stem Cell Transplantation (HSCT). <i>Blood</i> , 2012, 120, 457-457.	0.6	2
97	Fresh Ex Vivo Expanded Natural Killer Cells Demonstrate Robust Proliferation in Vivo in High-Risk Relapsed Multiple Myeloma (MM) Patients. <i>Blood</i> , 2012, 120, 579-579.	0.6	2
98	Human papillomavirus type 16 (HPV16) E6/E7-specific cytotoxic T lymphocytes (CTL) for immunotherapy of HPV-associated cancer (Ca).. <i>Journal of Clinical Oncology</i> , 2012, 30, 2558-2558.	0.8	0
99	Expanded Natural Killer (NK) Cells for Immunotherapy: Fresh and Made to Order. <i>Blood</i> , 2012, 120, 1912-1912.	0.6	0
100	Safety and Clinical Efficacy of Rapidly-Generated Trivirus-Directed T Cells After Allogeneic Hematopoietic Stem Cell Transplant. <i>Blood</i> , 2012, 120, 223-223.	0.6	25
101	Cytotoxic T Lymphocytes Simultaneously Targeting Multiple Tumor-associated Antigens to Treat EBV Negative Lymphoma. <i>Molecular Therapy</i> , 2011, 19, 2258-2268.	3.7	80
102	CD28 costimulation improves expansion and persistence of chimeric antigen receptor-modified T cells in lymphoma patients. <i>Journal of Clinical Investigation</i> , 2011, 121, 1822-1826.	3.9	876
103	Combining Oncolytic Vaccinia Virotherapy with Adoptive T Cell Therapy,. <i>Blood</i> , 2011, 118, 4042-4042.	0.6	1
104	Complete Tumor Responses in Lymphoma Patients Receiving Autologous Cytotoxic T Lymphocytes Targeting Epstein Barr Virus (EBV) - Latent Membrane Proteins. <i>Blood</i> , 2011, 118, 956-956.	0.6	1
105	Towards Phase 2/3 Trials for Epstein - Barr Virus (EBV)-Associated Malignancies,. <i>Blood</i> , 2011, 118, 4043-4043.	0.6	0
106	Human Papillomavirus Type 16 (HPV16) E6/E7-Specific Cytotoxic T Lymphocytes (CTLs) for Immunotherapy of HPV-Associated Malignancies. <i>Blood</i> , 2011, 118, 1913-1913.	0.6	0
107	EBV-Induced Lymphoproliferation. <i>Blood</i> , 2011, 118, SCI-9-SCI-9.	0.6	0
108	Vaccination with Î³CD40L or Flagellin Gene-Modified T Cells Activates Dendritic Cells In Vivo and Induces a Potent Anti-Tumor Immune Response. <i>Blood</i> , 2011, 118, 1909-1909.	0.6	0



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109	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.	0.6	721
110	Generation of Epstein-Barr virus-specific cytotoxic T lymphocytes resistant to the immunosuppressive drug tacrolimus (FK506). <i>Blood</i> , 2009, 114, 4784-4791.	0.6	86
111	Cytotoxic T lymphocyte therapy with donor T cells prevents and treats adenovirus and Epstein-Barr virus infections after haploidentical and matched unrelated stem cell transplantation. <i>Blood</i> , 2009, 114, 4283-4292.	0.6	311
112	IL15 Enhances Proliferation and Effector Function of Antigen-Specific Cytotoxic T Lymphocytes (CTLs) and Mitigates the Suppressive Action of Regulatory T Cells (Tregs).. <i>Blood</i> , 2009, 114, 4088-4088.	0.6	1
113	Cytotoxic T Lymphocytes (CTL) Specific for CMV, Adenovirus, and EBV Can Be Generated From Naive T Cells for Adoptive Immunotherapy.. <i>Blood</i> , 2009, 114, 504-504.	0.6	0
114	An Inducible Caspase 9 Suicide Gene to Improve the Safety of Mesenchymal Stromal Cell Therapies.. <i>Blood</i> , 2009, 114, 1444-1444.	0.6	0
115	Adverse Events Following Infusion of T Cells for Adoptive Immunotherapy: A 10 Year Experience.. <i>Blood</i> , 2009, 114, 3212-3212.	0.6	0
116	Immune-Based Therapies Targeting Mage-A4 for Relapsed/Refractory Hodgkin's Lymphoma After Stem Cell Transplant.. <i>Blood</i> , 2009, 114, 4089-4089.	0.6	7
117	Monoculture-Derived T Lymphocytes Providing Multiple Virus Specificity and Anti-Leukemia Activity for Recipients of Hematopoietic Stem Cells or Umbilical Cord Blood Transplants. <i>Blood</i> , 2008, 112, 3909-3909.	0.6	1
118	Generation of Epstein Barr Virus Specific Cytotoxic T Lymphocytes (EBVCTLs) Resistant to the Immunosuppressive Drug Tacrolimus (FK506). <i>Blood</i> , 2008, 112, 3536-3536.	0.6	0
119	Complete Tumor Responses in Lymphoma Patients Who Receive Autologous Cytotoxic T Lymphocytes Targeting EBV Latent Membrane Proteins. <i>Blood</i> , 2008, 112, 230-230.	0.6	0
120	The "Side-Population" of Human Lymphoma Cells Have Increased Chemo-Resistance, Stem-Cell Like Properties and Are Potential Targets for Immunotherapy. <i>Blood</i> , 2008, 112, 2620-2620.	0.6	0
121	Selective Loss of a Putative Precursor Population of B-Chronic Lymphocytic Leukemia Cells Following Immunization with hCD40L/IL-2 Expressing Autologous Tumor Cells. <i>Blood</i> , 2008, 112, 3172-3172.	0.6	0
122	Exploiting Cytokine Secretion to Rapidly Produce Multivirus-Specific T Cells for Adoptive Immunotherapy. <i>Blood</i> , 2008, 112, 4594-4594.	0.6	0
123	Polyclonal PRAME-Specific Cytotoxic T Lymphocytes Generated Using Protein-Spanning Pools of Overlapping Pentadecapeptides Target Chronic Myeloid Leukemia. <i>Blood</i> , 2008, 112, 3899-3899.	0.6	0
124	Complete responses of relapsed lymphoma following genetic modification of tumor-antigen presenting cells and T-lymphocyte transfer. <i>Blood</i> , 2007, 110, 2838-2845.	0.6	266
125	Adoptive immunotherapy for herpesviruses. , 2007, , 1318-1331.		2
126	T-Cells Redirected Against CD70 for the Immunotherapy of Hematological Malignancies.. <i>Blood</i> , 2007, 110, 2757-2757.	0.6	3



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127	Adoptive cellular immunotherapy. , 2006, , 648-660.		0
128	Treatment of solid organ transplant recipients with autologous Epstein Barr virus-specific cytotoxic T lymphocytes (CTLs). Blood, 2006, 108, 2942-2949.	0.6	241
129	Monoculture-derived T lymphocytes specific for multiple viruses expand and produce clinically relevant effects in immunocompromised individuals. Nature Medicine, 2006, 12, 1160-1166.	15.2	536
130	Generation and Expansion of PRAME-Specific Cytotoxic T-Lymphocytes for Adoptive T-Cell Therapy of Hematological Malignancies.. Blood, 2006, 108, 2205-2205.	0.6	0
131	Immune Responses Are Induced Against Side-Population B-CLL Stem Cells by Patient Vaccination with hCD40L/IL2 Gene Modified Tumor Cells.. Blood, 2006, 108, 2552-2552.	0.6	0
132	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.	3.7	426
133	Genetically Modified Her2-Specific T Cells Recognize Low and High Her2 Expressing Breast Cancer Cells.. Blood, 2005, 106, 5540-5540.	0.6	0
134	The Clinical Use of Donor-Derived Virus-Specific Cytotoxic T Lymphocytes Reactive Against Cytomegalovirus (CMV), Adenovirus and Epstein Barr Virus (EBV).. Blood, 2005, 106, 81-81.	0.6	0
135	The Use of Autologous LMP2-Specific Cytotoxic T Lymphocytes (CTL) for the Treatment of Relapsed EBV-Positive Hodgkin Disease and Non-Hodgkin Lymphoma.. Blood, 2005, 106, 773-773.	0.6	0
136	Cytotoxic T Lymphocyte Therapy for Epstein-Barr Virus+ Hodgkin's Disease. Journal of Experimental Medicine, 2004, 200, 1623-1633.	4.2	371
137	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.	0.6	2
138	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.	0.6	0
139	Retrovirus-Transduced T Cell Blasts Have Not Only Antigen-Presenting Capabilities but Also Suppressor Regulatory T Cell-Inducing Capability.. Blood, 2004, 104, 3855-3855.	0.6	0
140	New insights into EBV-associated post-transplant lymphoproliferative disease. Lancet, The, 2003, 361, 192-193.	6.3	6
141	Counting EBV and T cells to predict PTLD. Blood, 2003, 101, 4227-4228.	0.6	2
142	Adapting a transforming growth factor Î²-related tumor protection strategy to enhance antitumor immunity. Blood, 2002, 99, 3179-3187.	0.6	310
143	An Epstein-Barr virus deletion mutant associated with fatal lymphoproliferative disease unresponsive to therapy with virus-specific CTLs. Blood, 2001, 97, 835-843.	0.6	249
144	Adenoviral gene transfer into dendritic cells efficiently amplifies the immune response to LMP2A antigen: A potential treatment strategy for Epstein-Barr virus-positive Hodgkin's lymphoma. International Journal of Cancer, 2001, 93, 706-713.	2.3	80

#	ARTICLE	IF	CITATIONS
145	Transfer of EBV-specific CTL to prevent EBV lymphoma post bone marrow transplant. , 1999, 14, 154-156.		29
146	Administration of Neomycin Resistance Gene Marked EBV Specific Cytotoxic T-Lymphocytes to Patients with Relapsed EBV-Positive Hodgkin Disease. Center for Cell and Gene Therapy, Baylor College of Medicine, Houston, Texas. Human Gene Therapy, 1998, 9, 1237-1250.	1.4	24
147	Infusion of Cytotoxic T Cells for the Prevention and Treatment of Epstein-Barr Virus-Induced Lymphoma in Allogeneic Transplant Recipients. Blood, 1998, 92, 1549-1555.	0.6	269
148	Long-term restoration of immunity against Epstein-Barr virus infection by adoptive transfer of gene-modified virus-specific T lymphocytes. Nature Medicine, 1996, 2, 551-555.	15.2	820