

Paul A Beavis

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

61
papers

5,046
citations

32
h-index

71
g-index

72
ext. papers

6,850
ext. citations

14.2
avg, IF

5.55
L-index

#	Paper	IF	Citations
61	Differential location of NKT and MAIT cells within lymphoid tissue.. <i>Scientific Reports</i> , 2022 , 12, 4034	4.9	0
60	Challenges of Creating New Tumor-Infiltrating Lymphocyte for Combating Breast Cancer.. <i>Journal of Clinical Oncology</i> , 2022 , JCO2200284	2.2	
59	Adoptive transfer of tumor-specific Th9 cells eradicates heterogeneous antigen-expressing tumor cells. <i>Cancer Cell</i> , 2021 ,	24.3	3
58	PTP1B is an intracellular checkpoint that limits T cell and CAR T cell anti-tumor immunity. <i>Cancer Discovery</i> , 2021 ,	24.4	3
57	Augmenting Adoptive T-cell Immunotherapy by Targeting the PD-1/PD-L1 Axis. <i>Cancer Research</i> , 2021 , 81, 5803-5805	10.1	0
56	Cellular networks controlling T cell persistence in adoptive cell therapy. <i>Nature Reviews Immunology</i> , 2021 , 21, 769-784	36.5	17
55	CRISPR/Cas9 mediated deletion of the adenosine A2A receptor enhances CAR T cell efficacy. <i>Nature Communications</i> , 2021 , 12, 3236	17.4	22
54	Myeloma natural killer cells are exhausted and have impaired regulation of activation. <i>Haematologica</i> , 2021 , 106, 2522-2526	6.6	2
53	CDK4/6 Inhibition Promotes Antitumor Immunity through the Induction of T-cell Memory. <i>Cancer Discovery</i> , 2021 , 11, 2582-2601	24.4	12
52	The role of exhaustion in CAR T cell therapy. <i>Cancer Cell</i> , 2021 , 39, 885-888	24.3	5
51	MAIT cells regulate NK cell-mediated tumor immunity. <i>Nature Communications</i> , 2021 , 12, 4746	17.4	8
50	Adoptive cellular therapy with T cells expressing the dendritic cell growth factor Flt3L drives epitope spreading and antitumor immunity. <i>Nature Immunology</i> , 2020 , 21, 914-926	19.1	53
49	p38 Kinase: A Key Target for Driving Potent T Cells for Adoptive Immunotherapy. <i>Cancer Cell</i> , 2020 , 37, 756-758	24.3	0
48	Promising Immuno-Oncology Options for the Future: Cellular Therapies and Personalized Cancer Vaccines. <i>American Society of Clinical Oncology Educational Book / ASCO American Society of Clinical Oncology Meeting</i> , 2020 , 40, 1-6	7.1	5
47	Efficient CRISPR/Cas9 Gene Editing in Uncultured Naive Mouse T Cells for In Vivo Studies. <i>Journal of Immunology</i> , 2020 , 204, 2308-2315	5.3	12
46	Sex-specific adipose tissue imprinting of regulatory T cells. <i>Nature</i> , 2020 , 579, 581-585	50.4	72
45	Tissue-resident memory T cells in breast cancer control and immunotherapy responses. <i>Nature Reviews Clinical Oncology</i> , 2020 , 17, 341-348	19.4	70

44	A New Safety Approach Allowing Reversible Control of CAR T Cell Responses. <i>Molecular Therapy</i> , 2020 , 28, 1563-1566	11.7	
43	PTPN2 phosphatase deletion in T cells promotes anti-tumour immunity and CAR T-cell efficacy in solid tumours. <i>EMBO Journal</i> , 2020 , 39, e103637	13	36
42	IL-15 Preconditioning Augments CAR T Cell Responses to Checkpoint Blockade for Improved Treatment of Solid Tumors. <i>Molecular Therapy</i> , 2020 , 28, 2379-2393	11.7	19
41	Pharmacological and genetic strategies for targeting adenosine to enhance adoptive T cell therapy of cancer. <i>Current Opinion in Pharmacology</i> , 2020 , 53, 91-97	5.1	1
40	Targeting the epigenetic regulation of antitumour immunity. <i>Nature Reviews Drug Discovery</i> , 2020 , 19, 776-800	64.1	81
39	Intratumoral Copper Modulates PD-L1 Expression and Influences Tumor Immune Evasion. <i>Cancer Research</i> , 2020 , 80, 4129-4144	10.1	19
38	Macrophage-Derived CXCL9 and CXCL10 Are Required for Antitumor Immune Responses Following Immune Checkpoint Blockade. <i>Clinical Cancer Research</i> , 2020 , 26, 487-504	12.9	138
37	An Evolutionarily Conserved Function of Polycomb Silences the MHC Class I Antigen Presentation Pathway and Enables Immune Evasion in Cancer. <i>Cancer Cell</i> , 2019 , 36, 385-401.e8	24.3	169
36	Supercharging adoptive T cell therapy to overcome solid tumor-induced immunosuppression. <i>Science Translational Medicine</i> , 2019 , 11,	17.5	65
35	Switching on the green light for chimeric antigen receptor T-cell therapy. <i>Clinical and Translational Immunology</i> , 2019 , 8, e1046	6.8	9
34	Tissue-specific tumor microenvironments influence responses to immunotherapies. <i>Clinical and Translational Immunology</i> , 2019 , 8, e1094	6.8	11
33	Antagonism of IAPs Enhances CAR T-cell Efficacy. <i>Cancer Immunology Research</i> , 2019 , 7, 183-192	12.5	33
32	Tumor-derived exosomes modulate T cell function through transfer of RNA. <i>FEBS Journal</i> , 2018 , 285, 1030-1032	5.7	4
31	Chimeric antigen receptor T cells form nonclassical and potent immune synapses driving rapid cytotoxicity. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2018 , 115, E2068-E2076	11.5	129
30	Dual PD-1 and CTLA-4 Checkpoint Blockade Promotes Antitumor Immune Responses through CD4Foxp3 Cell-Mediated Modulation of CD103 Dendritic Cells. <i>Cancer Immunology Research</i> , 2018 , 6, 1069-1081	12.5	38
29	Single-cell profiling of breast cancer T cells reveals a tissue-resident memory subset associated with improved prognosis. <i>Nature Medicine</i> , 2018 , 24, 986-993	50.5	420
28	Targeting Adenosine Receptor Signaling in Cancer Immunotherapy. <i>International Journal of Molecular Sciences</i> , 2018 , 19,	6.3	90
27	Tumor immune evasion arises through loss of TNF sensitivity. <i>Science Immunology</i> , 2018 , 3,	28	119

26	A Multifunctional Role for Adjuvant Anti-4-1BB Therapy in Augmenting Antitumor Response by Chimeric Antigen Receptor T Cells. <i>Cancer Research</i> , 2017 , 77, 1296-1309	10.1	46
25	BET-Bromodomain Inhibitors Engage the Host Immune System and Regulate Expression of the Immune Checkpoint Ligand PD-L1. <i>Cell Reports</i> , 2017 , 18, 2162-2174	10.6	170
24	A novel combination strategy for effectively targeting cancer stem-like cells. <i>Immunology and Cell Biology</i> , 2017 , 95, 573-574	5	3
23	Dual-specific Chimeric Antigen Receptor T Cells and an Indirect Vaccine Eradicate a Variety of Large Solid Tumors in an Immunocompetent, Self-antigen Setting. <i>Clinical Cancer Research</i> , 2017 , 23, 2478-2490	12.9	71
22	T cell inhibitory mechanisms in a model of aggressive Non-Hodgkin's Lymphoma. <i>Oncolmmunology</i> , 2017 , 7, e1365997	7.2	1
21	Agonist immunotherapy restores T cell function following MEK inhibition improving efficacy in breast cancer. <i>Nature Communications</i> , 2017 , 8, 606	17.4	60
20	A Novel Target Antigen for the Treatment of Acute Myeloid Leukemia by CAR T Cells. <i>Molecular Therapy</i> , 2017 , 25, 1997-1998	11.7	2
19	CMTM6 maintains the expression of PD-L1 and regulates anti-tumour immunity. <i>Nature</i> , 2017 , 549, 101-105	10.4	375
18	Targeting the adenosine 2A receptor enhances chimeric antigen receptor T cell efficacy. <i>Journal of Clinical Investigation</i> , 2017 , 127, 929-941	15.9	183
17	Reprogramming the tumor microenvironment to enhance adoptive cellular therapy. <i>Seminars in Immunology</i> , 2016 , 28, 64-72	10.7	41
16	RAS/MAPK Activation Is Associated with Reduced Tumor-Infiltrating Lymphocytes in Triple-Negative Breast Cancer: Therapeutic Cooperation Between MEK and PD-1/PD-L1 Immune Checkpoint Inhibitors. <i>Clinical Cancer Research</i> , 2016 , 22, 1499-509	12.9	311
15	Immunosuppressive activities of adenosine in cancer. <i>Current Opinion in Pharmacology</i> , 2016 , 29, 7-16	5.1	156
14	Enhancing the efficacy of adoptive cellular therapy by targeting tumor-induced immunosuppression. <i>Immunotherapy</i> , 2015 , 7, 499-512	3.8	16
13	Relevance of tumor-infiltrating lymphocytes in breast cancer. <i>BMC Medicine</i> , 2015 , 13, 202	11.4	131
12	CD73: A potential biomarker for anti-PD-1 therapy. <i>Oncolmmunology</i> , 2015 , 4, e1046675	7.2	23
11	Adenosine Receptor 2A Blockade Increases the Efficacy of Anti-PD-1 through Enhanced Antitumor T-cell Responses. <i>Cancer Immunology Research</i> , 2015 , 3, 506-17	12.5	198
10	CD3bright signals on T cells identify IL-17A-producing VβVα+ T cells. <i>Immunology and Cell Biology</i> , 2015 , 93, 198-212	5	50
9	Cross-talk between tumors can affect responses to therapy. <i>Oncolmmunology</i> , 2015 , 4, e975572	7.2	7

8	CD73 promotes anthracycline resistance and poor prognosis in triple negative breast cancer. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013 , 110, 11091-6	11.5	303
7	A blockade enhances anti-metastatic immune responses. <i>Oncotmunology</i> , 2013 , 2, e26705	7.2	13
6	Blockade of A2A receptors potently suppresses the metastasis of CD73+ tumors. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013 , 110, 14711-6	11.5	244
5	Anti-PD-1 antibody therapy potently enhances the eradication of established tumors by gene-modified T cells. <i>Clinical Cancer Research</i> , 2013 , 19, 5636-46	12.9	485
4	CD73-deficient mice are resistant to carcinogenesis. <i>Cancer Research</i> , 2012 , 72, 2190-6	10.1	156
3	CD73: a potent suppressor of antitumor immune responses. <i>Trends in Immunology</i> , 2012 , 33, 231-7	14.4	253
2	Activation of p38 mitogen-activated protein kinase is critical step for acquisition of effector function in cytokine-activated T cells, but acts as a negative regulator in T cells activated through the T-cell receptor. <i>Immunology</i> , 2011 , 132, 104-10	7.8	18
1	Resistance to regulatory T cell-mediated suppression in rheumatoid arthritis can be bypassed by ectopic foxp3 expression in pathogenic synovial T cells. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011 , 108, 16717-22	11.5	43