

# Andrew G Polson

## List of Publications by Year in descending order

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Version: 2024-02-01

32  
papers

1,915  
citations

361045

20  
h-index

454577

30  
g-index

32  
all docs

32  
docs citations

32  
times ranked

2502  
citing authors

#	ARTICLE	IF	CITATIONS
1	Antibody-Drug Conjugates for the Treatment of Non-Hodgkin's Lymphoma: Target and Linker-Drug Selection. <i>Cancer Research</i> , 2009, 69, 2358-2364.	0.4	229
2	Therapeutic potential of an anti-CD79b antibody-drug conjugate, anti-CD79b-vc-MMAE, for the treatment of non-Hodgkin lymphoma. <i>Blood</i> , 2009, 114, 2721-2729.	0.6	205
3	An anti-CD3/anti-CLL-1 bispecific antibody for the treatment of acute myeloid leukemia. <i>Blood</i> , 2017, 129, 609-618.	0.6	136
4	Antibody-drug conjugates targeted to CD79 for the treatment of non-Hodgkin lymphoma. <i>Blood</i> , 2007, 110, 616-623.	0.6	135
5	Antitumor Efficacy of a Bispecific Antibody That Targets HER2 and Activates T Cells. <i>Cancer Research</i> , 2014, 74, 5561-5571.	0.4	135
6	Targeting LGR5 <sup>+</sup> cells with an antibody-drug conjugate for the treatment of colon cancer. <i>Science Translational Medicine</i> , 2015, 7, 314ra186.	5.8	131
7	A Novel Anti-CD22 Anthracycline-Based Antibody-Drug Conjugate (ADC) That Overcomes Resistance to Auristatin-Based ADCs. <i>Clinical Cancer Research</i> , 2015, 21, 3298-3306.	3.2	124
8	Expression pattern of the human FcRH/IRTA receptors in normal tissue and in B-chronic lymphocytic leukemia. <i>International Immunology</i> , 2006, 18, 1363-1373.	1.8	100
9	Cathepsin B Is Dispensable for Cellular Processing of Cathepsin B-Cleavable Antibody-Drug Conjugates. <i>Cancer Research</i> , 2017, 77, 7027-7037.	0.4	99
10	DCDT2980S, an Anti-CD22-Monomethyl Auristatin E Antibody-Drug Conjugate, Is a Potential Treatment for Non-Hodgkin Lymphoma. <i>Molecular Cancer Therapeutics</i> , 2013, 12, 1255-1265.	1.9	72
11	Investigational antibody-drug conjugates for hematological malignancies. <i>Expert Opinion on Investigational Drugs</i> , 2011, 20, 75-85.	1.9	64
12	FcRL5 as a Target of Antibody-Drug Conjugates for the Treatment of Multiple Myeloma. <i>Molecular Cancer Therapeutics</i> , 2012, 11, 2222-2232.	1.9	63
13	An Anti-CLL-1 Antibody-Drug Conjugate for the Treatment of Acute Myeloid Leukemia. <i>Clinical Cancer Research</i> , 2019, 25, 1358-1368.	3.2	53
14	<i>In vivo</i> effects of targeting CD79b with antibodies and antibody-drug conjugates. <i>Molecular Cancer Therapeutics</i> , 2009, 8, 2937-2946.	1.9	45
15	The successes and limitations of preclinical studies in predicting the pharmacodynamics and safety of cell-surface-targeted biological agents in patients. <i>British Journal of Pharmacology</i> , 2012, 166, 1600-1602.	2.7	37
16	Phase I study of the anti-FcRH5 antibody-drug conjugate DFRF4539A in relapsed or refractory multiple myeloma. <i>Blood Cancer Journal</i> , 2019, 9, 17.	2.8	35
17	Intratumoral Payload Concentration Correlates with the Activity of Antibody-Drug Conjugates. <i>Molecular Cancer Therapeutics</i> , 2018, 17, 677-685.	1.9	30
18	Antibody-drug conjugates for the treatment of B-cell non-Hodgkin's lymphoma and leukemia. <i>Future Oncology</i> , 2013, 9, 355-368.	1.1	27

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19	Antibody Drug Conjugates Differentiate Uptake and DNA Alkylation of Pyrrolobenzodiazepines in Tumors from Organs of Xenograft Mice. <i>Drug Metabolism and Disposition</i> , 2016, 44, 1958-1962.	1.7	23
20	A homogeneous high-DAR antibody-drug conjugate platform combining THIOMAB antibodies and XTEN polypeptides. <i>Chemical Science</i> , 2022, 13, 3147-3160.	3.7	23
21	Anti-CD22 and anti-CD79b antibody-drug conjugates preferentially target proliferating B cells. <i>British Journal of Pharmacology</i> , 2017, 174, 628-640.	2.7	22
22	Exposure-Efficacy Analysis of Antibody-Drug Conjugates Delivering an Excessive Level of Payload to Tissues. <i>Drug Metabolism and Disposition</i> , 2019, 47, 1146-1155.	1.7	20
23	Stabilizing a Tubulysin Antibody-Drug Conjugate To Enable Activity Against Multidrug-Resistant Tumors. <i>ACS Medicinal Chemistry Letters</i> , 2017, 8, 1037-1041.	1.3	19
24	Evaluation and use of an anti-cynomolgus monkey CD79b surrogate antibody-drug conjugate to enable clinical development of polatuzumab vedotin. <i>British Journal of Pharmacology</i> , 2019, 176, 3805-3818.	2.7	18
25	A BCMA/CD16A bispecific innate cell engager for the treatment of multiple myeloma. <i>Leukemia</i> , 2022, 36, 1006-1014.	3.3	17
26	Antibody-Drug Conjugates Derived from Cytotoxic seco-CBI-Dimer Payloads Are Highly Efficacious in Xenograft Models and Form Protein Adducts In Vivo. <i>Bioconjugate Chemistry</i> , 2019, 30, 1356-1370.	1.8	15
27	Preclinical pharmacokinetics and pharmacodynamics of DCLL9718A: An antibody-drug conjugate for the treatment of acute myeloid leukemia. <i>MAbs</i> , 2018, 10, 1312-1321.	2.6	13
28	Toward an Effective Targeted Chemotherapy for Multiple Myeloma. <i>Clinical Cancer Research</i> , 2009, 15, 3906-3907.	3.2	11
29	An Anti-CD22-seco-CBI-Dimer Antibody-Drug Conjugate (ADC) for the Treatment of Non-Hodgkin Lymphoma That Provides a Longer Duration of Response than Auristatin-Based ADCs in Preclinical Models. <i>Molecular Cancer Therapeutics</i> , 2021, 20, 340-346.	1.9	9
30	Novel Anti-LY6G6D/CD3 T-Cell-Dependent Bispecific Antibody for the Treatment of Colorectal Cancer. <i>Molecular Cancer Therapeutics</i> , 2022, 21, 974-985.	1.9	5
31	Antibody-Drug Conjugates for the Treatment of B-Cell Malignancies. , 2013, , 139-147.		0
32	Antibody-Drug Conjugates Targeted to CD79 for the Treatment of Non-Hodgkin's Lymphoma. <i>Blood</i> , 2006, 108, 2524-2524.	0.6	0