

# Piers A Blombery

## List of Publications by Year in descending order

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Version: 2024-02-01

72  
papers

1,984  
citations

331538

21  
h-index

265120

42  
g-index

74  
all docs

74  
docs citations

74  
times ranked

3112  
citing authors

#	ARTICLE	IF	CITATIONS
1	Clonal hematopoiesis, myeloid disorders and <i>BAX</i> -mutated myelopoiesis in patients receiving venetoclax for CLL. <i>Blood</i> , 2022, 139, 1198-1207.	0.6	34
2	Single-cell sequencing demonstrates complex resistance landscape in CLL and MCL treated with BTK and BCL2 inhibitors. <i>Blood Advances</i> , 2022, 6, 503-508.	2.5	16
3	Findings from precision oncology in the clinic: rare, novel variants are a significant contributor to scaling molecular diagnostics. <i>BMC Medical Genomics</i> , 2022, 15, 70.	0.7	2
4	Health economic evidence for the use of molecular biomarker tests in hematological malignancies: A systematic review. <i>European Journal of Haematology</i> , 2022, 108, 469-485.	1.1	4
5	Methyl-CpG binding domain 4, DNA glycosylase ( <i>MBD4</i> )-associated neoplasia syndrome associated with a homozygous missense variant in <i>MBD4</i> : Expansion of an emerging phenotype. <i>British Journal of Haematology</i> , 2022, , .	1.2	2
6	Panel-based gene testing in myelodysplastic/myeloproliferative neoplasm- overlap syndromes: Australasian Leukaemia and Lymphoma Group (ALLG) consensus statement. <i>Pathology</i> , 2022, , .	0.3	2
7	Laboratory quality assessment of candidate gene panel testing for acute myeloid leukaemia: a joint ALLG / RCPAQAP initiative. <i>Pathology</i> , 2021, 53, 487-492.	0.3	0
8	Myeloid somatic mutation panel testing in myeloproliferative neoplasms. <i>Pathology</i> , 2021, 53, 339-348.	0.3	13
9	CAR-T cell therapy: practical guide to routine laboratory monitoring. <i>Pathology</i> , 2021, 53, 408-415.	0.3	10
10	T cell receptor beta locus sequencing early post-allogeneic stem cell transplant identifies patients at risk of initial and recurrent cytomegalovirus infection. <i>Bone Marrow Transplantation</i> , 2021, 56, 2582-2590.	1.3	2
11	Investigation of product-derived lymphoma following infusion of <i>piggyBac</i> -modified CD19 chimeric antigen receptor T cells. <i>Blood</i> , 2021, 138, 1391-1405.	0.6	87
12	Cryptic molecular lesion in acute promyelocytic leukemia with negative initial FISH. <i>Leukemia and Lymphoma</i> , 2021, 62, 3060-3062.	0.6	1
13	T-cell replete allogeneic stem cell transplant for mantle cell lymphoma achieves durable disease control, including against TP53-mutated disease. <i>Bone Marrow Transplantation</i> , 2021, 56, 2857-2859.	1.3	7
14	Transient, flexible gene editing in zebrafish neutrophils and macrophages for determination of cell-autonomous functions. <i>DMM Disease Models and Mechanisms</i> , 2021, 14, .	1.2	11
15	Response to everolimus in a patient with refractory HGBL-NOS harboring multiple genomic aberrations in PTEN. <i>Leukemia and Lymphoma</i> , 2021, , 1-5.	0.6	1
16	Characterisation of immune checkpoints in Richter syndrome identifies LAG3 as a potential therapeutic target. <i>British Journal of Haematology</i> , 2021, 195, 113-118.	1.2	8
17	Outcomes of patients with CLL sequentially resistant to both BCL2 and BTK inhibition. <i>Blood Advances</i> , 2021, 5, 4054-4058.	2.5	39
18	Expert Curation of Somatic FLT3 Variants By the Clingen Somatic Hematologic Cancer Taskforce (ClinGen HCT). <i>Blood</i> , 2021, 138, 4387-4387.	0.6	0

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19	Mechanisms of intrinsic and acquired resistance to venetoclax in B-cell lymphoproliferative disease. <i>Leukemia and Lymphoma</i> , 2020, 61, 257-262.	0.6	15
20	Immune recovery in patients with mantle cell lymphoma receiving long-term ibrutinib and venetoclax combination therapy. <i>Blood Advances</i> , 2020, 4, 4849-4859.	2.5	14
21	Clonal independence of <i>JAK2</i> and <i>CALR</i> or <i>MPL</i> mutations in comutated myeloproliferative neoplasms demonstrated by single cell DNA sequencing. <i>Haematologica</i> , 2020, 106, 313-315.	1.7	17
22	Recovery of natural killer cell cytotoxicity in a A91V perforin homozygous patient following severe haemophagocytic lymphohistiocytosis. <i>British Journal of Haematology</i> , 2020, 190, 458-461.	1.2	2
23	Utility of clinical comprehensive genomic characterization for diagnostic categorization in patients presenting with hypocellular bone marrow failure syndromes. <i>Haematologica</i> , 2020, 106, 64-73.	1.7	14
24	Diagnostic evaluation and considerations in hypocellular bone marrow failure—A focus on genomics. <i>International Journal of Laboratory Hematology</i> , 2020, 42, 82-89.	0.7	4
25	A synonymous <i>GATA2</i> variant underlying familial myeloid malignancy with striking intrafamilial phenotypic variability. <i>British Journal of Haematology</i> , 2020, 190, e297-e301.	1.2	14
26	High dose-rate brachytherapy of localized prostate cancer converts tumors from cold to hot. , 2020, 8, e000792.		45
27	Inotuzumab ozogamicin resistance associated with a novel <i>CD22</i> truncating mutation in a case of acute lymphoblastic leukaemia. <i>British Journal of Haematology</i> , 2020, 191, 123-126.	1.2	5
28	Severe chemotherapy toxicity in a 10-year-old with T-acute lymphoblastic lymphoma harboring biallelic <i>FANCM</i> variants. <i>Leukemia and Lymphoma</i> , 2020, 61, 1257-1259.	0.6	2
29	Detection of an <i>IGH-BRAF</i> fusion in a patient with <i>BRAF</i> Val600Glu negative hairy cell leukemia. <i>Leukemia and Lymphoma</i> , 2020, 61, 2024-2026.	0.6	6
30	Multiple <i>BCL2</i> mutations cooccurring with Gly101Val emerge in chronic lymphocytic leukemia progression on venetoclax. <i>Blood</i> , 2020, 135, 773-777.	0.6	115
31	Undetectable peripheral blood MRD should be the goal of venetoclax in CLL, but attainment plateaus after 24 months. <i>Blood Advances</i> , 2020, 4, 165-173.	2.5	34
32	Safety and Efficacy of Induction and Maintenance Avelumab Plus R-CHOP in Patients with Diffuse Large B-Cell Lymphoma (DLBCL): Analysis of the Phase II Avr-CHOP Study. <i>Blood</i> , 2020, 136, 43-44.	0.6	9
33	Acquired Mutations in <i>BAX</i> Confer Resistance to BH3 Mimetics in Acute Myeloid Leukemia. <i>Blood</i> , 2020, 136, 7-8.	0.6	13
34	<i>BTK</i> inhibitor therapy is effective in patients with CLL resistant to venetoclax. <i>Blood</i> , 2020, 135, 2266-2270.	0.6	67
35	A Phase II, Open-Label, Single Arm Trial to Assess the Efficacy and Safety of the Combination of Tisagenlecleucel and Ibrutinib in Mantle Cell Lymphoma (TARMAC). <i>Blood</i> , 2020, 136, 34-35.	0.6	4
36	Longitudinal Genomic Characterization Using Cell-Free DNA in Patients with Idiopathic Aplastic Anemia. <i>Blood</i> , 2020, 136, 5-6.	0.6	0

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37	High Clonal Complexity of Resistance Mechanisms Occurring at Progression after Single-Agent Targeted Therapy Strategies in Chronic Lymphocytic Leukemia. <i>Blood</i> , 2020, 136, 15-16.	0.6	2
38	Mutational and Copy Number Profiling of Circulating Tumor DNA in Acute Myeloid Leukemia Using Targeted Next Generation Sequencing. <i>Blood</i> , 2020, 136, 39-40.	0.6	2
39	Expert Curation of Somatic Variants in Hematological Malignancies By the Clingen Somatic Hematological Cancer Taskforce (ClinGen HCT). <i>Blood</i> , 2020, 136, 23-23.	0.6	0
40	Circulating tumor DNA for disease monitoring in the era of CAR T-cell therapy. <i>Leukemia and Lymphoma</i> , 2019, 60, 279-280.	0.6	0
41	Characterization of a novel venetoclax resistance mutation (BCL2 Phe104Ile) observed in follicular lymphoma. <i>British Journal of Haematology</i> , 2019, 186, e188-e191.	1.2	37
42	Structures of BCL-2 in complex with venetoclax reveal the molecular basis of resistance mutations. <i>Nature Communications</i> , 2019, 10, 2385.	5.8	139
43	First-in-Human RNA Polymerase I Transcription Inhibitor CX-5461 in Patients with Advanced Hematologic Cancers: Results of a Phase I Dose-Escalation Study. <i>Cancer Discovery</i> , 2019, 9, 1036-1049.	7.7	129
44	CNSpector: a web-based tool for visualisation and clinical diagnosis of copy number variation from next generation sequencing. <i>Scientific Reports</i> , 2019, 9, 6426.	1.6	17
45	Rapid and Durable Complete Remission of Refractory AITL with Azacitidine Treatment in Absence of TET2 Mutation or Concurrent MDS. <i>HemaSphere</i> , 2019, 3, e187.	1.2	14
46	Molecular Drivers of Breast Implant-Associated Anaplastic Large Cell Lymphoma. <i>Plastic and Reconstructive Surgery</i> , 2019, 143, 59S-64S.	0.7	28
47	What does good FISHing look like in MDS?. <i>Leukemia and Lymphoma</i> , 2019, 60, 571-572.	0.6	0
48	Dynamic molecular monitoring reveals that SWI539 mutations mediate resistance to ibrutinib plus venetoclax in mantle cell lymphoma. <i>Nature Medicine</i> , 2019, 25, 119-129.	15.2	147
49	Revisiting acquired aplastic anaemia: current concepts in diagnosis and management. <i>Internal Medicine Journal</i> , 2019, 49, 152-159.	0.5	15
50	Acquisition of the Recurrent Gly101Val Mutation in BCL2 Confers Resistance to Venetoclax in Patients with Progressive Chronic Lymphocytic Leukemia. <i>Cancer Discovery</i> , 2019, 9, 342-353.	7.7	306
51	BTK Leu528Trp - a Potential Secondary Resistance Mechanism Specific for Patients with Chronic Lymphocytic Leukemia Treated with the Next Generation BTK Inhibitor Zanubrutinib. <i>Blood</i> , 2019, 134, 170-170.	0.6	33
52	Characterization of the "Immune Evasion" Phenotype of Richter Syndrome and the Implications for Immune-Checkpoint Inhibitor Therapy. <i>Blood</i> , 2019, 134, 4290-4290.	0.6	0
53	Sensitive NPM1 Mutation Quantitation in Acute Myeloid Leukemia Using Ultradeep Next-Generation Sequencing in the Diagnostic Laboratory. <i>Archives of Pathology and Laboratory Medicine</i> , 2018, 142, 606-612.	1.2	8
54	Incidental detection of germline variants of potential clinical significance by massively parallel sequencing in haematological malignancies. <i>Journal of Clinical Pathology</i> , 2018, 71, 84-87.	1.0	18

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55	Detection of clinically relevant early genomic lesions in B-cell malignancies from circulating tumour DNA using a single hybridisation-based next generation sequencing assay. <i>British Journal of Haematology</i> , 2018, 183, 146-149.	1.2	8
56	Adaptive reprogramming of NK cells in X-linked lymphoproliferative syndrome. <i>Blood</i> , 2018, 131, 699-702.	0.6	5
57	The price of success-health economics of personalized diffuse large B-cell lymphoma treatment. <i>Leukemia and Lymphoma</i> , 2018, 59, 1517-1519.	0.6	1
58	Comprehensive genomic characterization dissects the complex biology of a case of synchronous Burkitt lymphoma and myeloid malignancy with shared hematopoietic ancestry. <i>Leukemia and Lymphoma</i> , 2018, 59, 992-995.	0.6	2
59	Natural killer receptor ligand expression on acute myeloid leukemia impacts survival and relapse after chemotherapy. <i>Blood Advances</i> , 2018, 2, 335-346.	2.5	47
60	Novel genomic findings in multiple myeloma identified through routine diagnostic sequencing. <i>Journal of Clinical Pathology</i> , 2018, 71, 895-899.	1.0	28
61	Frequent activating STAT3 mutations and novel recurrent genomic abnormalities detected in breast implant-associated anaplastic large cell lymphoma. <i>Oncotarget</i> , 2018, 9, 36126-36136.	0.8	62
62	Providing Diagnoses in Bone Marrow Failure Syndromes through Multimodal Comprehensive Genomic Evaluation and Multidisciplinary Care: The Melbourne Genomics Health Alliance Bone Marrow Failure Flagship. <i>Blood</i> , 2018, 132, 3867-3867.	0.6	0
63	Clinicopathological differences exist between CALR- and JAK2-mutated myeloproliferative neoplasms despite a similar molecular landscape: data from targeted next-generation sequencing in the diagnostic laboratory. <i>Annals of Hematology</i> , 2017, 96, 725-732.	0.8	17
64	Primary Breast Lymphoma—Population-Level Insights into an Infrequent but Increasingly Recognized Subtype of Lymphoma. <i>Journal of the National Cancer Institute</i> , 2017, 109, .	3.0	5
65	Canary: an atomic pipeline for clinical amplicon assays. <i>BMC Bioinformatics</i> , 2017, 18, 555.	1.2	4
66	Whole exome sequencing reveals activating JAK1 and STAT3 mutations in breast implant-associated anaplastic large cell lymphoma. <i>Haematologica</i> , 2016, 101, e387-e390.	1.7	124
67	Copper deficiency mimicking myelodysplastic syndrome. <i>Leukemia and Lymphoma</i> , 2016, 57, 1223-1226.	0.6	1
68	The molecular pathogenesis of B-cell non-Hodgkin lymphoma. <i>European Journal of Haematology</i> , 2015, 95, 280-293.	1.1	22
69	Molecular lesions in B-cell lymphoproliferative disorders: recent contributions from studies utilizing high-throughput sequencing techniques. <i>Leukemia and Lymphoma</i> , 2014, 55, 19-30.	0.6	5
70	Detection of BRAF mutations in patients with hairy cell leukemia and related lymphoproliferative disorders. <i>Haematologica</i> , 2012, 97, 780-783.	1.7	63
71	Erdheim-Chester Disease Harboring the BRAF V600E Mutation. <i>Journal of Clinical Oncology</i> , 2012, 30, e331-e332.	0.8	46
72	Prophylactic intravenous immunoglobulin during autologous haemopoietic stem cell transplantation for multiple myeloma is not associated with reduced infectious complications. <i>Annals of Hematology</i> , 2011, 90, 1167-1172.	0.8	27