

# David Wu

## List of Publications by Year in descending order

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Version: 2024-02-01

54  
papers

2,883  
citations

331670

21  
h-index

214800

47  
g-index

55  
all docs

55  
docs citations

55  
times ranked

5250  
citing authors

#	ARTICLE	IF	CITATIONS
1	Acquisition of a CD19-negative myeloid phenotype allows immune escape of MLL-rearranged B-ALL from CD19 CAR-T-cell therapy. <i>Blood</i> , 2016, 127, 2406-2410.	1.4	622
2	International, evidence-based consensus diagnostic criteria for HHV-8“negative/idiopathic multicentric Castleman disease. <i>Blood</i> , 2017, 129, 1646-1657.	1.4	381
3	Validation and Implementation of Targeted Capture and Sequencing for the Detection of Actionable Mutation, Copy Number Variation, and Gene Rearrangement in Clinical Cancer Specimens. <i>Journal of Molecular Diagnostics</i> , 2014, 16, 56-67.	2.8	234
4	High-Throughput Sequencing Detects Minimal Residual Disease in Acute T Lymphoblastic Leukemia. <i>Science Translational Medicine</i> , 2012, 4, 134ra63.	12.4	207
5	Measurable residual disease detection by high-throughput sequencing improves risk stratification for pediatric B-ALL. <i>Blood</i> , 2018, 131, 1350-1359.	1.4	158
6	Detection of Minimal Residual Disease in B Lymphoblastic Leukemia by High-Throughput Sequencing of <i>IGH</i> . <i>Clinical Cancer Research</i> , 2014, 20, 4540-4548.	7.0	138
7	Mass cytometry of Hodgkin lymphoma reveals a CD4+ regulatory T-cell“rich and exhausted T-effector microenvironment. <i>Blood</i> , 2018, 132, 825-836.	1.4	121
8	ClinGen Myeloid Malignancy Variant Curation Expert Panel recommendations for germline RUNX1 variants. <i>Blood Advances</i> , 2019, 3, 2962-2979.	5.2	110
9	Genomic analyses of flow-sorted Hodgkin Reed-Sternberg cells reveal complementary mechanisms of immune evasion. <i>Blood Advances</i> , 2019, 3, 4065-4080.	5.2	99
10	International evidence-based consensus diagnostic and treatment guidelines for unicentric Castleman disease. <i>Blood Advances</i> , 2020, 4, 6039-6050.	5.2	94
11	Pathology of Castleman Disease. <i>Hematology/Oncology Clinics of North America</i> , 2018, 32, 37-52.	2.2	83
12	Clinicopathologic Features and Prognostic Impact of Lymph Node Involvement in Patients With Breast Implant-associated Anaplastic Large Cell Lymphoma. <i>American Journal of Surgical Pathology</i> , 2018, 42, 293-305.	3.7	80
13	Detection of minimal residual disease in NPM1-mutated acute myeloid leukemia by next-generation sequencing. <i>Modern Pathology</i> , 2014, 27, 1438-1446.	5.5	49
14	CADD score has limited clinical validity for the identification of pathogenic variants in noncoding regions in a hereditary cancer panel. <i>Genetics in Medicine</i> , 2016, 18, 1269-1275.	2.4	45
15	Pretransplantation Minimal Residual Disease Predicts Survival in Patients with Mantle Cell Lymphoma Undergoing Autologous Stem Cell Transplantation in Complete Remission. <i>Biology of Blood and Marrow Transplantation</i> , 2016, 22, 380-385.	2.0	37
16	Deep NPM1 Sequencing Following Allogeneic Hematopoietic Cell Transplantation Improves Risk Assessment in Adults with NPM1-Mutated AML. <i>Biology of Blood and Marrow Transplantation</i> , 2018, 24, 1615-1620.	2.0	29
17	Ultrasensitive detection of acute myeloid leukemia minimal residual disease using single molecule molecular inversion probes. <i>Haematologica</i> , 2017, 102, 1549-1557.	3.5	28
18	Pattern associated leukemia immunophenotypes and measurable disease detection in acute myeloid leukemia or myelodysplastic syndrome with mutated <i>NPM1</i> . <i>Cytometry Part B - Clinical Cytometry</i> , 2019, 96, 67-72.	1.5	26

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19	A novel disease-causing synonymous exonic mutation in GATA2 affecting RNA splicing. Blood, 2018, 132, 1211-1215.	1.4	25
20	Flow Cytometry for Non-Hodgkin and Classical Hodgkin Lymphoma. Methods in Molecular Biology, 2013, 971, 27-47.	0.9	23
21	Recurrent somatic loss of <scp><i>TNFRSF14</i></scp> in classical Hodgkin lymphoma. Genes Chromosomes and Cancer, 2016, 55, 278-287.	2.8	23
22	How I curate: applying American Society of Hematology-Clinical Genome Resource Myeloid Malignancy Variant Curation Expert Panel rules for RUNX1 variant curation for germline predisposition to myeloid malignancies. Haematologica, 2020, 105, 870-887.	3.5	23
23	Reactive T cells by flow cytometry distinguish Hodgkin lymphomas from T cell/histiocyteâ€rich large B cell lymphoma. Cytometry Part B - Clinical Cytometry, 2016, 90, 424-432.	1.5	22
24	Myeloid/lymphoid neoplasms with FLT3 rearrangement. Modern Pathology, 2021, 34, 1673-1685.	5.5	21
25	Flow cytometry of ALK-negative anaplastic large cell lymphoma of breast implant-associated effusion and capsular tissue. , 2014, , n/a-n/a.		19
26	On-Going Evolution Of IGH In B-Cell Precursor Acute Lymphoblastic Leukemia Does Not Substantially Affect Day 29, Post-Treatment MRD Quantification By High-Throughput Sequencing. Blood, 2013, 122, 1341-1341.	1.4	19
27	Insufficient evidence exists to use histopathologic subtype to guide treatment of idiopathic multicentric Castleman disease. American Journal of Hematology, 2020, 95, 1553-1561.	4.1	18
28	Flow cytometry of ALK-negative anaplastic large cell lymphoma of breast implant-associated effusion and capsular tissue. , 2015, 88, 58-63.		16
29	Computer-Aided Detection of Rare Tumor Populations in Flow Cytometry. American Journal of Clinical Pathology, 2015, 144, 517-524.	0.7	12
30	Flow cytometric features of incidental indolent T lymphoblastic proliferations. Cytometry Part B - Clinical Cytometry, 2020, 98, 282-287.	1.5	12
31	Robust Detection Of Minimal Residual Disease In Unselected Patients With B-Cell Precursor Acute Lymphoblastic Leukemia By High-Throughput Sequencing Of IGH. Blood, 2013, 122, 2550-2550.	1.4	12
32	Clinical Experience With Modified, Single-Tube T-Cell Receptor VÎ²2 Flow Cytometry Analysis for T-Cell Clonality. American Journal of Clinical Pathology, 2016, 145, 467-485.	0.7	11
33	Crossâ€Platform DNA Encoding for Singleâ€Cell Imaging of Gene Expression. Angewandte Chemie - International Edition, 2016, 55, 8975-8978.	13.8	10
34	Flow Cytometry for Non-Hodgkin and Hodgkin Lymphomas. Methods in Molecular Biology, 2019, 1956, 35-60.	0.9	9
35	Full-Length Isoforms of Kaposi's Sarcoma-Associated Herpesvirus Latency-Associated Nuclear Antigen Accumulate in the Cytoplasm of Cells Undergoing the Lytic Cycle of Replication. Journal of Virology, 2017, 91, .	3.4	8
36	De Novo Identification and Visualization of Important Cell Populations for Classic Hodgkin Lymphoma Using Flow Cytometry and Machine Learning. American Journal of Clinical Pathology, 2021, 156, 1092-1102.	0.7	8

#	ARTICLE	IF	CITATIONS
37	Cutaneous T-Cell Lymphoma in Sub-Saharan Africa. Journal of the National Comprehensive Cancer Network: JNCCN, 2013, 11, 275-280.	4.9	7
38	Jumping translocations in myelodysplastic syndromes. Cancer Genetics, 2016, 209, 395-402.	0.4	7
39	Bendamustine with rituximab, etoposide and carboplatin (T(R) <scp>EC</scp>) in relapsed or refractory aggressive lymphoma: a prospective multicentre phase 1/2 clinical trial. British Journal of Haematology, 2018, 183, 601-607.	2.5	7
40	Early Tâ€Cell Precursor Acute Lymphoblastic Leukemia in an Infant With an <i>NRAS</i> Q61R Mutation and Clinical Features of Juvenile Myelomonocytic Leukemia. Pediatric Blood and Cancer, 2016, 63, 1667-1670.	1.5	5
41	Potential for Process Improvement of Clinical Flow Cytometry by Incorporating Real-Time Automated Screening of Data to Expedite Addition of Antibody Panels. American Journal of Clinical Pathology, 2022, 157, 443-450.	0.7	5
42	Targeted Next-Generation Sequencing of Acute Leukemia. Methods in Molecular Biology, 2017, 1633, 163-184.	0.9	4
43	A Curriculum for Genomic Education of Molecular Genetic Pathology Fellows. Journal of Molecular Diagnostics, 2021, 23, 1218-1240.	2.8	4
44	Ultrasensitive Detection of Chimerism by Single-Molecule Molecular Inversion Probe Capture and High-Throughput Sequencing of Copy Number Deletion Polymorphisms. Clinical Chemistry, 2018, 64, 938-949.	3.2	3
45	Ultrasensitive Quantitation of Genomic Chimerism by Single-Molecule Molecular Inversion Probe Capture and High-Throughput Sequencing of Copy Number Deletion Polymorphisms. Journal of Molecular Diagnostics, 2021, , .	2.8	3
46	Residual Disease Monitoring By High Throughput Sequencing Provides Risk Stratification in Childhood B-ALL and Identifies a Novel Subset of Patients Having Poor Outcome. Blood, 2016, 128, 1086-1086.	1.4	2
47	Comprehensive Evaluation and Validation of a Next-Generation Sequencing Assay for Minimal Residual Disease Detection in T-Lymphoblastic Leukemia/Lymphoma. Blood, 2019, 134, 1475-1475.	1.4	2
48	Bone Marrow Involvement Detected By Multi-Parameter Flow Cytometry Predicts Poor Outcome after Autologous Stem Cell Transplantation for Peripheral T-Cell Lymphoma. Blood, 2015, 126, 1972-1972.	1.4	1
49	Crossâ€Platform DNA Encoding for Singleâ€Cell Imaging of Gene Expression. Angewandte Chemie, 2016, 128, 9121-9124.	2.0	0
50	Commentary on A Case of Rapid Deterioration with Marked Hypergammaglobulinemia. Clinical Chemistry, 2020, 66, 1379-1380.	3.2	0
51	Detection Of Recurrent/Persistent Disease By T-Cell Receptor Repertoire Profiling In Patients With Mature T-Cell Neoplasm. Blood, 2013, 122, 2614-2614.	1.4	0
52	Detection of Mutations in Inherited Bone Marrow Failure and Myelodysplastic Syndrome Genes Using Genomic Capture and Massively Parallel Sequencing in Clinical Diagnostics. Blood, 2016, 128, 1507-1507.	1.4	0
53	Comparative Genomic Analyses Defines Shared and Unique Features of cHL and PMBL and New Mechanisms of Sensitivity to PD-1 Blockade. Blood, 2019, 134, 1493-1493.	1.4	0
54	Expert Curation of Somatic Variants in Hematological Malignancies By the Clingen Somatic Hematological Cancer Taskforce (ClinGen HCT). Blood, 2020, 136, 23-23.	1.4	0