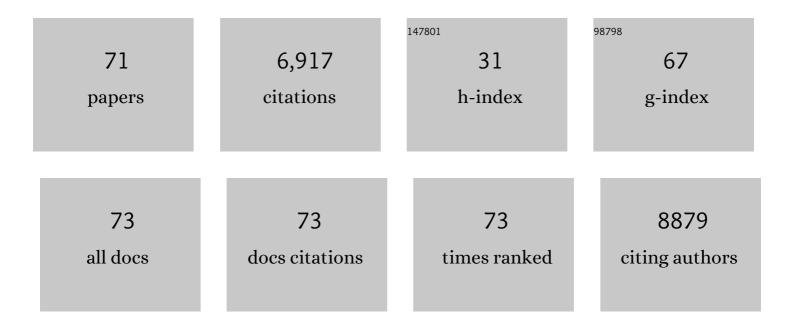
## Hanno Glimm

List of Publications by Year in descending order

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HANNO CLIMM

#	Article	IF	CITATIONS
1	Correction of X-linked chronic granulomatous disease by gene therapy, augmented by insertional activation of MDS1-EVI1, PRDM16 or SETBP1. Nature Medicine, 2006, 12, 401-409.	30.7	1,129
2	Genomic instability and myelodysplasia with monosomy 7 consequent to EVI1 activation after gene therapy for chronic granulomatous disease. Nature Medicine, 2010, 16, 198-204.	30.7	727
3	Stem-Cell Gene Therapy for the Wiskott–Aldrich Syndrome. New England Journal of Medicine, 2010, 363, 1918-1927.	27.0	505
4	An unbiased genome-wide analysis of zinc-finger nuclease specificity. Nature Biotechnology, 2011, 29, 816-823.	17.5	488
5	Gene Therapy for Wiskott-Aldrich Syndrome—Long-Term Efficacy and Genotoxicity. Science Translational Medicine, 2014, 6, 227ra33.	12.4	460
6	High-resolution insertion-site analysis by linear amplification–mediated PCR (LAM-PCR). Nature Methods, 2007, 4, 1051-1057.	19.0	281
7	Distinct Types of Tumor-Initiating Cells Form Human Colon Cancer Tumors and Metastases. Cell Stem Cell, 2011, 9, 357-365.	11.1	276
8	Vector integration is nonrandom and clustered and influences the fate of lymphopoiesis in SCID-X1 gene therapy. Journal of Clinical Investigation, 2007, 117, 2225-2232.	8.2	221
9	Integrative genomic and transcriptomic analysis of leiomyosarcoma. Nature Communications, 2018, 9, 144.	12.8	197
10	<i>NRG1</i> Fusions in <i>KRAS</i> Wild-Type Pancreatic Cancer. Cancer Discovery, 2018, 8, 1087-1095.	9.4	189
11	Gammaretrovirus-mediated correction of SCID-X1 is associated with skewed vector integration site distribution in vivo. Journal of Clinical Investigation, 2007, 117, 2241-2249.	8.2	185
12	Comprehensive genomic access to vector integration in clinical gene therapy. Nature Medicine, 2009, 15, 1431-1436.	30.7	173
13	Genome-wide high-throughput integrome analyses by nrLAM-PCR and next-generation sequencing. Nature Protocols, 2010, 5, 1379-1395.	12.0	161
14	The <scp>tRNA</scp> methyltransferase Dnmt2 is required forÂaccurate polypeptide synthesis duringÂhaematopoiesis. EMBO Journal, 2015, 34, 2350-2362.	7.8	154
15	Precision oncology based on omics data: The NCT Heidelberg experience. International Journal of Cancer, 2017, 141, 877-886.	5.1	133
16	Integrating next-generation sequencing into clinical oncology: strategies, promises and pitfalls. ESMO Open, 2016, 1, e000094.	4.5	126
17	Comprehensive Genomic and Transcriptomic Analysis for Guiding Therapeutic Decisions in Patients with Rare Cancers. Cancer Discovery, 2021, 11, 2780-2795.	9.4	125
18	High-resolution analysis of the human T-cell receptor repertoire. Nature Communications, 2015, 6, 8081.	12.8	123

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19	Recurrent CDKN1B (p27) mutations in hairy cell leukemia. Blood, 2015, 126, 1005-1008.	1.4	88
20	Harmonization and Standardization of Panel-Based Tumor Mutational Burden Measurement: Real-World Results and Recommendations ofÂtheÂQuality in Pathology Study. Journal of Thoracic Oncology, 2020, 15, 1177-1189.	1.1	81
21	The landscape of chromothripsis across adult cancer types. Nature Communications, 2020, 11, 2320.	12.8	75
22	Cell-of-Origin DNA Methylation Signatures Are Maintained during Colorectal Carcinogenesis. Cell Reports, 2018, 23, 3407-3418.	6.4	66
23	The AP-1 transcription factor JunB is essential for multiple myeloma cell proliferation and drug resistance in the bone marrow microenvironment. Leukemia, 2017, 31, 1570-1581.	7.2	60
24	Integration of genomics and histology revises diagnosis and enables effective therapy of refractory cancer of unknown primary with <i>PDL1</i> amplification. Journal of Physical Education and Sports Management, 2016, 2, a001180.	1.2	57
25	Cooperation of BRAFF595L and mutant HRAS in histiocytic sarcoma provides new insights into oncogenic BRAF signaling. Leukemia, 2016, 30, 937-946.	7.2	52
26	PD-L1 (CD274) copy number gain, expression, and immune cell infiltration as candidate predictors for response to immune checkpoint inhibitors in soft-tissue sarcoma. Oncolmmunology, 2017, 6, e1279777.	4.6	50
27	Asymmetric Centriole Numbers at Spindle Poles Cause Chromosome Missegregation in Cancer. Cell Reports, 2017, 20, 1906-1920.	6.4	49
28	Degradation of CCNK/CDK12 is a druggable vulnerability of colorectal cancer. Cell Reports, 2021, 36, 109394.	6.4	41
29	Succession of transiently active tumorâ€initiating cell clones in human pancreatic cancer xenografts. EMBO Molecular Medicine, 2017, 9, 918-932.	6.9	36
30	Efficient marking of human cells with rapid but transient repopulating activity in autografted recipients. Blood, 2005, 106, 893-898.	1.4	33
31	Genome-wide Specificity of Highly Efficient TALENs and CRISPR/Cas9 for T Cell Receptor Modification. Molecular Therapy - Methods and Clinical Development, 2017, 4, 213-224.	4.1	32
32	YAP Orchestrates Heterotypic Endothelial Cell Communication via HGF/c-MET Signaling in Liver Tumorigenesis. Cancer Research, 2020, 80, 5502-5514.	0.9	31
33	Genetic subclone architecture of tumor clone-initiating cells in colorectal cancer. Journal of Experimental Medicine, 2017, 214, 2073-2088.	8.5	30
34	Mutant KIT as imatinib-sensitive target in metastatic sinonasal carcinoma. Annals of Oncology, 2017, 28, 142-148.	1.2	30
35	Targeting Fibroblast Growth Factor Receptor 1 for Treatment of Soft-Tissue Sarcoma. Clinical Cancer Research, 2017, 23, 962-973.	7.0	29
36	Salinomycin: Anti-tumor activity in a pre-clinical colorectal cancer model. PLoS ONE, 2019, 14, e0211916.	2.5	27

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37	Patient-derived xenografts of gastrointestinal cancers are susceptible to rapid and delayed B-lymphoproliferation. International Journal of Cancer, 2017, 140, 1356-1363.	5.1	26
38	A Model for the Detection of Clonality in Marked Hematopoietic Stem Cells. Annals of the New York Academy of Sciences, 2001, 938, 146-156.	3.8	25
39	Conceptual framework for precision cancer medicine in Germany: Consensus statement of the Deutsche Krebshilfe working group â€~Molecular Diagnostics and Therapy'. European Journal of Cancer, 2020, 135, 1-7.	2.8	23
40	Targetable ERBB2 mutations identified in neurofibroma/schwannoma hybrid nerve sheath tumors. Journal of Clinical Investigation, 2020, 130, 2488-2495.	8.2	23
41	KIT-Dependent and KIT-Independent Genomic Heterogeneity of Resistance in Gastrointestinal Stromal Tumors — TORC1/2 Inhibition as Salvage Strategy. Molecular Cancer Therapeutics, 2019, 18, 1985-1996.	4.1	22
42	Validating Comprehensive Next-Generation Sequencing Results for Precision Oncology: The NCT/DKTK Molecularly Aided Stratification for Tumor Eradication Research Experience. JCO Precision Oncology, 2018, 2, 1-13.	3.0	20
43	So rare we need to hunt for them: reframing the ethical debate on incidental findings. Genome Medicine, 2015, 7, 83.	8.2	19
44	Identification and characterization of a BRAF fusion oncoprotein with retained autoinhibitory domains. Oncogene, 2020, 39, 814-832.	5.9	19
45	Identification of BCL-XL as highly active survival factor and promising therapeutic target in colorectal cancer. Cell Death and Disease, 2020, 11, 875.	6.3	17
46	YAPâ€induced Ccl2 expression is associated with a switch in hepatic macrophage identity and vascular remodelling in liver cancer. Liver International, 2021, 41, 3011-3023.	3.9	17
47	Pheno-seq – linking visual features and gene expression in 3D cell culture systems. Scientific Reports, 2019, 9, 12367.	3.3	16
48	Mapping Active Gene-Regulatory Regions in Human Repopulating Long-Term HSCs. Cell Stem Cell, 2018, 23, 132-146.e9.	11.1	14
49	Stk33 is required for spermatid differentiation and male fertility in mice. Developmental Biology, 2018, 433, 84-93.	2.0	13
50	Colorectal cancerâ€initiating cells caught in the act. EMBO Molecular Medicine, 2017, 9, 856-858.	6.9	12
51	Stable Long-Term Blood Formation by Stem Cells in Murine Steady-State Hematopoiesis. Stem Cells, 2012, 30, 1961-1970.	3.2	11
52	Phenotypic differentiation does not affect tumorigenicity of primary human colon cancer initiating cells. Cancer Letters, 2016, 371, 326-333.	7.2	11
53	Genomics of Immunotherapy-Associated Hyperprogressors—Letter. Clinical Cancer Research, 2017, 23, 6374-6375.	7.0	11
54	Metastatic adult pancreatoblastoma: Multimodal treatment and molecular characterization of a very rare disease. Pancreatology, 2020, 20, 425-432.	1.1	11

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#	Article	IF	CITATIONS
55	Functional States in Tumor-Initiating Cell Differentiation in Human Colorectal Cancer. Cancers, 2021, 13, 1097.	3.7	11
56	You Can Count on This: Barcoded Hematopoietic Stem Cells. Cell Stem Cell, 2011, 9, 390-392.	11.1	10
57	Induction of the proapoptotic tumor suppressor gene <i>Cell Adhesion Molecule 1</i> by chemotherapeutic agents is repressed in therapy resistant acute myeloid leukemia. Molecular Carcinogenesis, 2015, 54, 1815-1819.	2.7	9
58	Systematic Generation of Patient-Derived Tumor Models in Pancreatic Cancer. Cells, 2019, 8, 142.	4.1	9
59	High-throughput monitoring of integration site clonality in preclinical and clinical gene therapy studies. Molecular Therapy - Methods and Clinical Development, 2015, 2, 14061.	4.1	8
60	Detection of Structural Variants in Circulating Cell-Free DNA from Sarcoma Patients Using Next Generation Sequencing. Cancers, 2020, 12, 3627.	3.7	7
61	Sustained Polyclonal Hematopoietic Repopulation after Successful SCID-X1 Gene Therapy by Means of a Non Random Integrating Pseudotyped Gammaretrovector Blood, 2004, 104, 290-290.	1.4	7
62	High tumour mutational burden and EGFR/MAPK pathway activation are therapeutic targets in metastatic porocarcinoma. British Journal of Dermatology, 2021, , .	1.5	6
63	Rationale and design of the CRAFT (Continuous ReAssessment with Flexible ExTension in Rare) Tj ETQq1 1 0.784	314 rgBT 4.5	/Oyerlock 10
64	A perivascular niche in the bone marrow hosts quiescent and proliferating tumorigenic colorectal cancer cells. International Journal of Cancer, 2020, 147, 519-531.	5.1	5
65	Ruxolitinib is effective in the treatment of a patient with refractory Tâ€ALL. EJHaem, 2021, 2, 139-142.	1.0	4
66	Response to Cabozantinib Following Acquired Entrectinib Resistance in a Patient With <i>ETV6-NTRK3</i> Fusion-Positive Carcinoma Harboring the <i>NTRK3</i> <sup>G623R</sup> Solvent-Front Mutation. JCO Precision Oncology, 2021, 5, 687-694.	3.0	3
67	<scp>MGMT</scp> inactivation as a new biomarker in patients with advanced biliary tract cancers. Molecular Oncology, 2022, 16, 2733-2746.	4.6	2
68	Persistence of eGFP Marked Bone Marrow Cells in Long-Term Hematopoiesis Blood, 2004, 104, 2111-2111.	1.4	0
69	Insertional Activation of MDS1/EVI1, PRDM16 and SETBP1 in a Successful Chronic Granulomatous Disease (CGD) Gene Therapy Trial Blood, 2006, 108, 3274-3274.	1.4	0
70	Deregulated EVI1 Expression Leads to Genomic Instability and G1 Cell Cycle Arrest. Blood, 2011, 118, 2431-2431.	1.4	0
71	Recurrent Germline Variant in the Cohesin Complex Gene <i>RAD21</i> Predisposes Children to Lymphoblastic Leukemia and Lymphoma. Blood, 2021, 138, 3358-3358.	1.4	0