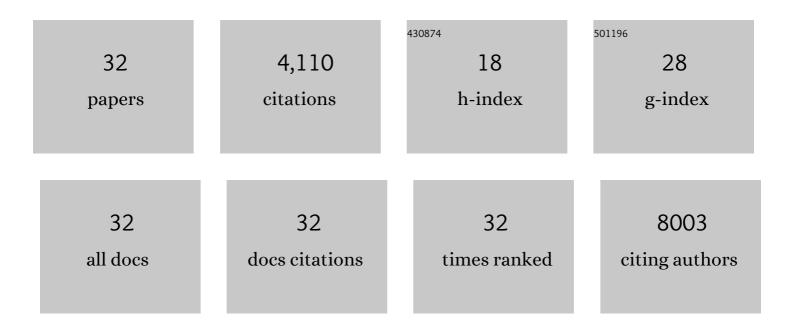
Young Rock Chung

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Early type I IFN blockade improves the efficacy of viral vaccines. Journal of Experimental Medicine, 2020, 217, .	8.5	38
2	Interrogating Adaptive Immunity Using LCMV. Current Protocols in Immunology, 2020, 130, e99.	3.6	19
3	Muscarinic acetylcholine receptor regulates self-renewal of early erythroid progenitors. Science Translational Medicine, 2019, 11, .	12.4	12
4	TLR4 signaling improves PD-1 blockade therapy during chronic viral infection. PLoS Pathogens, 2019, 15, e1007583.	4.7	17
5	KMT2C mediates the estrogen dependence of breast cancer through regulation of $ER\hat{I}\pm$ enhancer function. Oncogene, 2018, 37, 4692-4710.	5.9	102
6	Synthetic Lethal and Convergent Biological Effects of Cancer-Associated Spliceosomal Gene Mutations. Cancer Cell, 2018, 34, 225-241.e8.	16.8	162
7	ASXL2 is essential for haematopoiesis and acts as a haploinsufficient tumour suppressor in leukemia. Nature Communications, 2017, 8, 15429.	12.8	55
8	Robust patient-derived xenografts of MDS/MPN overlap syndromes capture the unique characteristics of CMML and JMML. Blood, 2017, 130, 397-407.	1.4	112
9	A somatic mutation in erythro-myeloid progenitors causes neurodegenerative disease. Nature, 2017, 549, 389-393.	27.8	144
10	Oncogenic Mutations in <i>XPO1</i> Promote Lymphoid Transformation By Altering Nuclear/Cytoplasmic Localization of NFI®B Signaling Intermediates. Blood, 2017, 130, 879-879.	1.4	0
11	Characterization of Ntrk fusions and Therapeutic Response to Ntrk Inhibition in Hematologic Malignancies. Blood, 2017, 130, 794-794.	1.4	0
12	Modulation of splicing catalysis for therapeutic targeting of leukemia with mutations in genes encoding spliceosomal proteins. Nature Medicine, 2016, 22, 672-678.	30.7	301
13	Reply to "Uveal melanoma cells are resistant to EZH2 inhibition regardless of BAP1 status". Nature Medicine, 2016, 22, 578-579.	30.7	7
14	Frequent Clinical Overlap of Histiocytic Neoplasms and WHO-Classified Myeloid Malignancies Leads to Functional Insights into the Cell-of-Origin of Histiocytoses. Blood, 2016, 128, 951-951.	1.4	3
15	Synthetic Lethal Interactions of MDS-Associated Spliceosomal Gene Mutations Identifies the Basis for Their Mutual Exclusivity. Blood, 2016, 128, 961-961.	1.4	6
16	Serine/Arginine-Rich Splicing Factor 1 (SRSF1) Is Required for Adult and Embryonic Hematopoiesis and Has Non-Overlapping Roles with SRSF2 in Hematopoiesis. Blood, 2016, 128, 1478-1478.	1.4	1
17	SRSF2 Mutations Contribute to Myelodysplasia by Mutant-Specific Effects on Exon Recognition. Cancer Cell, 2015, 27, 617-630.	16.8	449
18	Loss of BAP1 function leads to EZH2-dependent transformation. Nature Medicine, 2015, 21, 1344-1349.	30.7	297

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#	Article	IF	CITATIONS
19	Prospective Blinded Study of <i>BRAF</i> V600E Mutation Detection in Cell-Free DNA of Patients with Systemic Histiocytic Disorders. Cancer Discovery, 2015, 5, 64-71.	9.4	115
20	ASXL2 Is a Novel Mediator of RUNX1-ETO Transcriptional Function and Collaborates with RUNX1-ETO to Promote Leukemogenesis. Blood, 2015, 126, 302-302.	1.4	2
21	Therapeutic Targeting of Spliceosomal Mutant Myeloid Leukemias through Modulation of Splicing Catalysis. Blood, 2015, 126, 4-4.	1.4	4
22	Diverse Mechanisms of Vemurafenib Resistance in BRAF-Mutant Hairy Cell Leukemia. Blood, 2015, 126, 449-449.	1.4	3
23	Genomic and functional analysis of leukemic transformation of myeloproliferative neoplasms. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, E5401-10.	7.1	238
24	Hematopoietic Stem Cell Origin of <i>BRAF</i> V600E Mutations in Hairy Cell Leukemia. Science Translational Medicine, 2014, 6, 238ra71.	12.4	102
25	Femoral Bone Marrow Aspiration in Live Mice. Journal of Visualized Experiments, 2014, , .	0.3	12
26	EZH2 Is Required for Germinal Center Formation and Somatic EZH2 Mutations Promote Lymphoid Transformation. Cancer Cell, 2013, 23, 677-692.	16.8	706
27	Direct migration of follicular melanocyte stem cells to the epidermis after wounding or UVB irradiation is dependent on Mc1r signaling. Nature Medicine, 2013, 19, 924-929.	30.7	151
28	Deletion of Asxl1 results in myelodysplasia and severe developmental defects in vivo. Journal of Experimental Medicine, 2013, 210, 2641-2659.	8.5	278
29	ASXL1 Mutations Promote Myeloid Transformation through Loss of PRC2-Mediated Gene Repression. Cancer Cell, 2012, 22, 180-193.	16.8	504
30	Progression of RAS-Mutant Leukemia during RAF Inhibitor Treatment. New England Journal of Medicine, 2012, 367, 2316-2321.	27.0	222
31	Epigenetic alterations in hematopoietic malignancies. International Journal of Hematology, 2012, 96, 413-427.	1.6	48
32	Conditional Deletion of Asxl1 Results in Myelodysplasia. Blood, 2012, 120, 308-308.	1.4	0