## Kimberly Lezon-Geyda

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/3333214/publications.pdf

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		1307366	1199470
15	296	7	12
papers	citations	h-index	g-index
15	15	15	448
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	Clinical spectrum of pyruvate kinase deficiency: data from the Pyruvate Kinase Deficiency Natural History Study. Blood, 2018, 131, 2183-2192.	0.6	121
2	Genotypeâ€phenotype correlation and molecular heterogeneity in pyruvate kinase deficiency. American Journal of Hematology, 2020, 95, 472-482.	2.0	47
3	A Unique Epigenomic Landscape Defines Human Erythropoiesis. Cell Reports, 2019, 28, 2996-3009.e7.	2.9	41
4	Histone Acetyltransferases p300 and CBP Coordinate Distinct Chromatin Remodeling Programs in Vascular Smooth Muscle Plasticity. Circulation, 2022, 145, 1720-1737.	1.6	27
5	Briefâ€exposure to preoperative bevacizumab reveals a TGFâ€Î² signature predictive of response in HER2â€negative breast cancers. International Journal of Cancer, 2016, 138, 747-757.	2.3	16
6	Phase 1b study of the mammalian target of rapamycin inhibitor sirolimus in combination with nanoparticle albumin–bound paclitaxel in patients with advanced solid tumors. Cancer, 2015, 121, 1817-1826.	2.0	11
7	Hemoglobin C trait accentuates erythrocyte dehydration in hereditary xerocytosis. Pediatric Blood and Cancer, 2017, 64, e26444.	0.8	10
8	CTCF and CohesinSA-1 Mark Active Promoters and Boundaries of Repressive Chromatin Domains in Primary Human Erythroid Cells. PLoS ONE, 2016, 11, e0155378.	1.1	7
9	A Ser725Arg mutation in Band 3 abolishes transport function and leads to anemia and renal tubular acidosis. Blood, 2018, 131, 1759-1763.	0.6	5
10	Pklr Intron Splicing-Associated Mutations and Alternate Diagnoses Are Common in Pyruvate Kinase Deficient Patients with Single or No Pklr Coding Mutations. Blood, 2018, 132, 3607-3607.	0.6	4
11	Molecular Characterization of 140 Patients in the Pyruvate Kinase Deficiency (PKD) Natural History Study (NHS): Report of 20 New Variants. Blood, 2015, 126, 3337-3337.	0.6	4
12	Terminal Erythroid Maturation Is Associated with Dynamic Changes in the Abundance of Histone Marks Associated with Active Transcription Elongation and RNA Polymerase II Pausing. Blood, 2019, 134, 154-154.	0.6	2
13	Altered Splicing from a Mutated Alternate Branch Point Is Common in Severe Alpha-Spectrin Linked Inherited Anemia. Blood, 2018, 132, 503-503.	0.6	1
14	Role of EVI1 in Cell Cycle Regulation: Relevance of Specific Target Genes Blood, 2005, 106, 1614-1614.	0.6	0
15	Next-generation RNA sequencing reveals transcriptomic changes after brief exposure to preoperative nab-paclitaxel, bevacizumab, and trastuzumab Journal of Clinical Oncology, 2012, 30, 10508-10508.	0.8	0