

Divya S Vinjamur

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/6878281/publications.pdf>

Version: 2024-02-01

17
papers

1,164
citations

933447

10
h-index

1125743

13
g-index

18
all docs

18
docs citations

18
times ranked

2307
citing authors

#	ARTICLE	IF	CITATIONS
1	BCL11A enhancer dissection by Cas9-mediated in situ saturating mutagenesis. <i>Nature</i> , 2015, 527, 192-197.	27.8	726
2	Recent progress in understanding and manipulating haemoglobin switching for the haemoglobinopathies. <i>British Journal of Haematology</i> , 2018, 180, 630-643.	2.5	107
3	Rational targeting of a NuRD subcomplex guided by comprehensive in situ mutagenesis. <i>Nature Genetics</i> , 2019, 51, 1149-1159.	21.4	83
4	Synthetic Lethality of Wnt Pathway Activation and Asparaginase in Drug-Resistant Acute Leukemias. <i>Cancer Cell</i> , 2019, 35, 664-676.e7.	16.8	70
5	Transcription Factors KLF1 and KLF2 Positively Regulate Embryonic and Fetal $\hat{1}^2$ -Globin Genes through Direct Promoter Binding. <i>Journal of Biological Chemistry</i> , 2011, 286, 24819-24827.	3.4	36
6	ZNF410 represses fetal globin by singular control of CHD4. <i>Nature Genetics</i> , 2021, 53, 719-728.	21.4	35
7	Growing and Genetically Manipulating Human Umbilical Cord Blood-Derived Erythroid Progenitor (HUDEP) Cell Lines. <i>Methods in Molecular Biology</i> , 2018, 1698, 275-284.	0.9	31
8	Common variants in signaling transcription-factor-binding sites drive phenotypic variability in red blood cell traits. <i>Nature Genetics</i> , 2020, 52, 1333-1345.	21.4	24
9	Kruppel-like transcription factors KLF1 and KLF2 have unique and coordinate roles in regulating embryonic erythroid precursor maturation. <i>Haematologica</i> , 2014, 99, 1565-1573.	3.5	16
10	Kruppel-Like Transcription Factor KLF1 Is Required for Optimal $\hat{1}^3$ - and $\hat{1}^2$ -Globin Expression in Human Fetal Erythroblasts. <i>PLoS ONE</i> , 2016, 11, e0146802.	2.5	11
11	Development of a double shmiR lentivirus effectively targeting both BCL11A and ZNF410 for enhanced induction of fetal hemoglobin to treat $\hat{1}^2$ -hemoglobinopathies. <i>Molecular Therapy</i> , 2022, 30, 2693-2708.	8.2	11
12	Molecular analysis of the erythroid phenotype of a patient with BCL11A haploinsufficiency. <i>Blood Advances</i> , 2021, 5, 2339-2349.	5.2	7
13	Crispr-Cas9 Saturating Mutagenesis Reveals an Achilles Heel in the BCL11A Erythroid Enhancer for Fetal Hemoglobin Induction (by Genome Editing). <i>Blood</i> , 2015, 126, 638-638.	1.4	5
14	Synthetic Lethality of Wnt Pathway Activation and Asparaginase in Drug-Resistant Acute Leukemias. <i>Blood</i> , 2018, 132, 891-891.	1.4	0
15	Transcriptional Signaling Centers Govern Human Erythropoiesis and Harbor Genetic Variations of Red Blood Cell Traits. <i>Blood</i> , 2018, 132, 1277-1277.	1.4	0
16	Comprehensive Integrated Genomic Perturbations Reveal Molecular Mechanisms of Red Blood Cell Trait Associations. <i>Blood</i> , 2018, 132, 532-532.	1.4	0
17	ZNF410 Represses Fetal Globin By Devoted Control of CHD4/NuRD. <i>Blood</i> , 2020, 136, 1-1.	1.4	0