Van Trung Chu

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/7816592/publications.pdf Version: 2024-02-01



VAN TRUNC CHIL

#	Article	IF	CITATIONS
1	Precise CRISPR-Cas–mediated gene repair with minimal off-target and unintended on-target mutations in human hematopoietic stem cells. Science Advances, 2022, 8, .	4.7	18
2	CRISPR-Cas9-Mediated ELANE Mutation Correction in Hematopoietic Stem and Progenitor Cells to Treat Severe Congenital Neutropenia. Molecular Therapy, 2020, 28, 2621-2634.	3.7	28
3	Protocol for Efficient CRISPR/Cas9/AAV-Mediated Homologous Recombination in Mouse Hematopoietic Stem and Progenitor Cells. STAR Protocols, 2020, 1, 100028.	0.5	6
4	Functional interplay of Epstein-Barr virus oncoproteins in a mouse model of B cell lymphomagenesis. Proceedings of the National Academy of Sciences of the United States of America, 2020, 117, 14421-14432.	3.3	17
5	Efficient CRISPR/Cas9-Mediated Gene Knockin in Mouse Hematopoietic Stem and Progenitor Cells. Cell Reports, 2019, 28, 3510-3522.e5.	2.9	19
6	sgRNA Sequence Motifs Blocking Efficient CRISPR/Cas9-Mediated Gene Editing. Cell Reports, 2019, 26, 1098-1103.e3.	2.9	92
7	BCR-dependent lineage plasticity in mature B cells. Science, 2019, 363, 748-753.	6.0	76
8	A novel allele for inducible Cre expression in germinal center BÂcells. European Journal of Immunology, 2019, 49, 192-194.	1.6	8
9	CRISPR/Cas9-Mediated In Vitro Mutagenesis in GC-Like B Cells. Methods in Molecular Biology, 2017, 1623, 135-145.	0.4	5
10	Efficient CRISPR-mediated mutagenesis in primary immune cells using CrispRGold and a C57BL/6 Cas9 transgenic mouse line. Proceedings of the National Academy of Sciences of the United States of America, 2016, 113, 12514-12519.	3.3	110
11	Efficient generation of Rosa26 knock-in mice using CRISPR/Cas9 in C57BL/6 zygotes. BMC Biotechnology, 2016, 16, 4.	1.7	222
12	Pop in, pop out: a novel gene-targeting strategy for use with CRISPR-Cas9. Genome Biology, 2015, 16, 244.	3.8	7
13	Increasing the efficiency of homology-directed repair for CRISPR-Cas9-induced precise gene editing in mammalian cells. Nature Biotechnology, 2015, 33, 543-548.	9.4	1,024