Kevin Luk

List of Publications by Year in descending order

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KEVINLLIK

#	Article	IF	CITATIONS
1	LONP-1 and ATFS-1 sustain deleterious heteroplasmy by promoting mtDNA replication in dysfunctional mitochondria. Nature Cell Biology, 2022, 24, 181-193.	10.3	33
2	Optimization of Nuclear Localization Signal Composition Improves CRISPR-Cas12a Editing Rates in Human Primary Cells. , 2022, 1, 271-284.		5
3	ZNF410 represses fetal globin by singular control of CHD4. Nature Genetics, 2021, 53, 719-728.	21.4	35
4	Dissecting ELANE neutropenia pathogenicity by human HSC gene editing. Cell Stem Cell, 2021, 28, 833-845.e5.	11.1	23
5	CRISPR-enhanced human adipocyte browning as cell therapy for metabolic disease. Nature Communications, 2021, 12, 6931.	12.8	41
6	Therapeutic base editing of human hematopoietic stem cells. Nature Medicine, 2020, 26, 535-541.	30.7	196
7	Small-Molecule PAPD5 Inhibitors Restore Telomerase Activity in Patient Stem Cells. Cell Stem Cell, 2020, 26, 896-909.e8.	11.1	57
8	BCL11A enhancer–edited hematopoietic stem cells persist in rhesus monkeys without toxicity. Journal of Clinical Investigation, 2020, 130, 6677-6687.	8.2	54
9	Rational targeting of a NuRD subcomplex guided by comprehensive in situ mutagenesis. Nature Genetics, 2019, 51, 1149-1159.	21.4	83
10	Editing aberrant splice sites efficiently restores β-globin expression in β-thalassemia. Blood, 2019, 133, 2255-2262.	1.4	57
11	Evaluating and Enhancing Target Specificity of Gene-Editing Nucleases and Deaminases. Annual Review of Biochemistry, 2019, 88, 191-220.	11.1	120
12	Enhanced Cas12a editing in mammalian cells and zebrafish. Nucleic Acids Research, 2019, 47, 4169-4180.	14.5	85
13	Highly efficient therapeutic gene editing of human hematopoietic stem cells. Nature Medicine, 2019, 25, 776-783.	30.7	344
14	Precise therapeutic gene correction by a simple nuclease-induced double-stranded break. Nature, 2019, 568, 561-565.	27.8	86
15	Genome editing of HBG1 and HBG2 to induce fetal hemoglobin. Blood Advances, 2019, 3, 3379-3392.	5.2	121
16	Gene Editing ELANE in Human Hematopoietic Stem and Progenitor Cells Reveals Disease Mechanisms and Therapeutic Strategies for Severe Congenital Neutropenia. Blood, 2019, 134, 3-3.	1.4	8
17	Orthogonal Cas9–Cas9 chimeras provide a versatile platform for genome editing. Nature Communications, 2018, 9, 4856.	12.8	27
18	Highly Efficient Therapeutic Gene Editing of BCL11A enhancer in Human Hematopoietic Stem Cells from ß-Hemoglobinopathy Patients for Fetal Hemoglobin Induction. Blood, 2018, 132, 3482-3482.	1.4	2