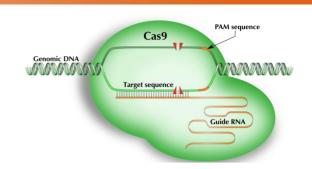
Genome Editing with CRISPR/Cas9

CRISPR/Cas9 is a RNA-guided, targeted genome editing tool which is *versatile*, *simple*, *and affordable*. Cas9 in complex with the guide RNA will lead to double-stranded breaks in a sequence-specific manner. Genome editing can be achieved via homology based and non-homology based repair mechanisms.



CRISPR/Cas9 Products

KN2.0 Gene knockout kits:

~50% bi-allelic knockout efficiency

CRISPR/Cas9 vectors (All-in-one, Lenti, Cas9)

Cas9 antibodies / Cas9 enzymes

AAVS1 / ROSA26 transgene insertion

Expression in genomic safe location

CRISPRa / CRISPRi kits

Functional Cas9 Protein

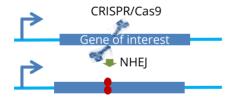
Custom Services

gRNA cloning service into CRISPR vectors

Donor vector of your own design

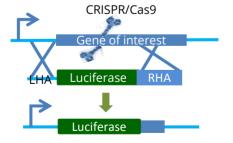
Applications using CRISPR/Cas9 as a genome editing tool

1. Indels (no donor template DNA)

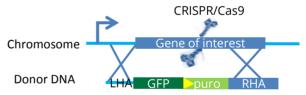


3. Native Promoter Study

Luciferase under the endogenous promoter

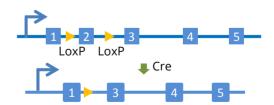


2. Gene knock-out with a reporter(s) knock-in



4. Conditional knockout

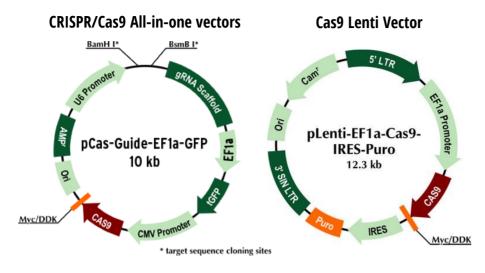
For essential genes or tissue-specific study inserting LoxP sites around the exon(s) to be knocked-out

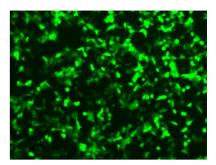


- 5. Specific mutations (with donor template DNA)
- SNPs—SNP associated disease, gene correction
- Desired deletions/insertions
- Tagging the endogenous genes



Genome Editing with CRISPR/Cas9





pCas-Guide-EF1a-GFP was transfected into HEK293 cells. The fluorescent picture shows the expression of GFP.

More CRISPR vectors: https://www.origene.com/products/vectors/crispr-vectors

Scheme of Genome-editing knockout: KN2.0

