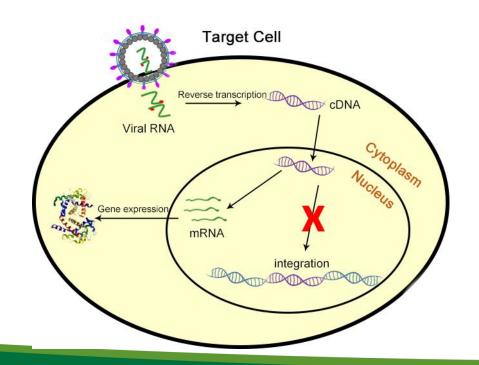
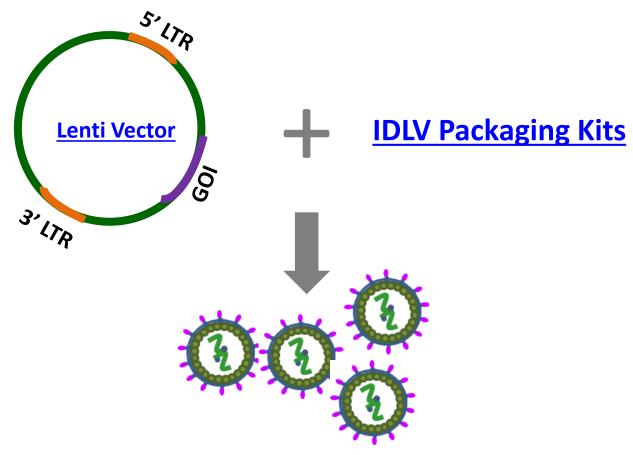
#### What is IDLV?

- ➤ IDLV is Integration Deficient Lentivirus
- ➤ No integration into the host Genome
- Genes/shRNA expressed transiently
- > Same cell spectrum & efficiency as regular lentivirus





### **How To Produce Integration Deficient Viral Particles?**

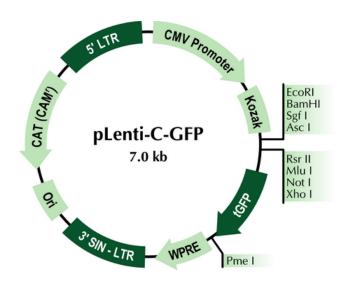


**Integration Deficient Lentivirus (IDLV)** 



### Titer of IDLV is Same as Regular Virus

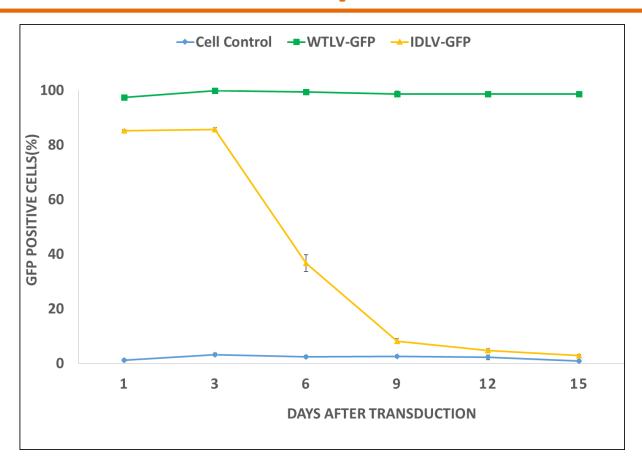
Type of Lentivirus	Titer (TU/ml)
Regular Lenti-GFP	2.9x10 <sup>7</sup>
IDLV-GFP (Integration Deficient)	2.7x10 <sup>7</sup>



Lenti-C-GFP vector was used to package into lentivirus using regular lenti packaging kit (cat# TR30037) or integration-deficient (IDLV) lenti packaging kit (cat# TR30036). Viral titer was the same.



#### **IDLV-GFP Mediated Expression is Transient**



Regular lenti-GFP virus (WTLV-GFP) and IDLV-GFP virus transduced HT1080 cells at 20 moi, GFP positive cells sorted at time points indicated.



### **Applications Using IDLV**

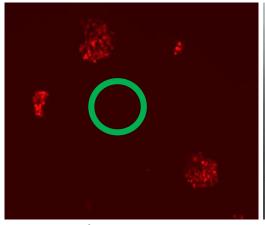
- √ Transient expression preferred
- ✓ Long-term or stable integration not wanted

CRISPR genome editing, Cas9 and sgRNA only needed to cut the genome temporally. Once genome editing is achieved, no need for Cas9 and sgRNA.

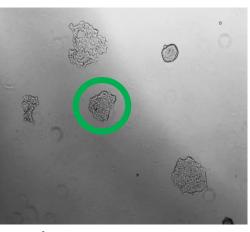


# **Functional Delivery of CRISPR by IDLV**

#### IDLV (Cas9 + gRNA-tRFP)

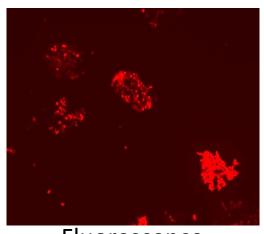




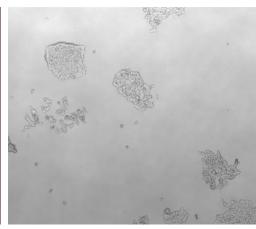


Phase contrast

#### IDLV (Cas9+ gRNA-Scramble)



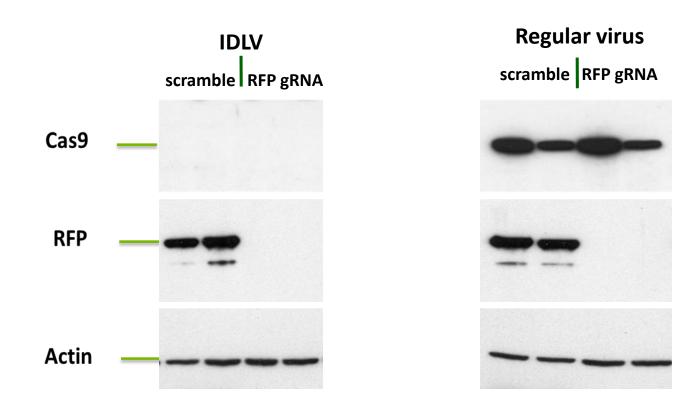
Fluorescence



Phase contrast

IDLV Cas9 and gRNA targeting tRFP were transduced into HEK293-RFP cells; scramble gRNA was used as negative control. Single cell colonies were isolated two weeks after transduction. RFP was knocked out in cells in green circle.

## **Effective KO of RFP W/O Footprint of Cas9**



3 weeks after transduction, WB was analyzed to check the RFP expression in single cell colonies; scramble control cells (RFP positive), RFP gRNA cells (not red). IDLV worked similarly as regular lentivirus (LV); RFP knocked out, but no Cas9 in cells.

