

VCF Order No.:

Virus No.:

User information:

Name:

Group leader:

Lab genetic engineering facility number:

Lab Bio Safety level of user:

S1 S2

Vector service: Shuttle vector plasmid name:

Shuttle vector properties
(Promoter, GOI, reporter gene)

Shuttle vector provided by the user:

No Yes

Shuttle vector cloned by VCF:

No Yes , Virus number BL-

Generation of new vector by VCF:

No Yes , cloning by:

Cloning description with dates:

Plasmid DNA preparation for transfection:

No Yes , preparation by:

Prep Date:

Virus request:

No Yes , done by:

HEK cell seeding date:

Transfection method:

Transfection date:

Shuttle vector:

Prep date:

Transfection (%),
evaluation HEK cells:

Packaging plasmid:

Prep date:

Envelope plasmid:

Prep date:

Harvest date:

Volume:

Virus quantification / efficiency test:

Quantification by reporter gene expression in cell cultures:

No Yes , done by:

Cell-culture:

#Cell / date:

Viruses (μl):

Transducing date:

Counting date:

Titer (IU/ml):

μl 100% recom.:

Quantification by qPCR of viral genome:

No Yes , done by:

qPCR date:

Titer:

(attached are the Roche lightcycler sheets)

Efficiency test by qPCR of transcript of GOI or shRNA kd:

No Yes , done by:

Cell-culture:

#Cell / date:

Viruses (μl):

Transducing date:

Cell lysis date:

cDNA date:

qPCR date:

Roche UPL #

% Expression from WT cells (per virus amount):

Primer seq. fw: 5'-3'

Primer seq. rev: 5'-3'

(attached are the Roche lightcycler sheets)

Efficiency test by Western Blotting:

No Yes , done by:

Cell-culture:

#Cell / date:

Viruses (μl):

Transducing date:

Cell lysis date:

Protein (μg/μl):

Proteins / WBlot
(date)

Prim. Antibody
(Name/company):

Dilution 1.AB:

Incubation date:

ECL date:

Sec. Antibody
(Name/company):

Dilution 2.AB:

% Expression from WT cells.

Date project start:

Date material released/shipped to user: