

## DNA &/or siRNA Transfection Reagent

- ✦ Powerful
- ✦ Economical: low amounts of nucleic acid & reagent
- ✦ Gentle to cells
- ✦ Versatile (DNA & siRNA)
- ✦ Convenient protocol

jetPRIME<sup>®</sup> is a new versatile and powerful DNA and siRNA transfection reagent for day-to-day experiments. jetPRIME<sup>®</sup> ensures high DNA transfection efficiency and excellent gene silencing in a variety of adherent cells. jetPRIME<sup>®</sup> is ideal for DNA/siRNA co-transfection. jetPRIME<sup>®</sup> is very gentle to cells since it requires low amounts of nucleic acid and reagent during transfection. Effective and nontoxic DNA and siRNA delivery is essential for reliable scientific results.

### ✦ Superior DNA transfection efficiency

Superior transfection efficiencies ranging between 70 and 90% are obtained when using jetPRIME<sup>®</sup> reagent versus the top competitor's reagent for several commonly used cell lines (Fig. 1-2).

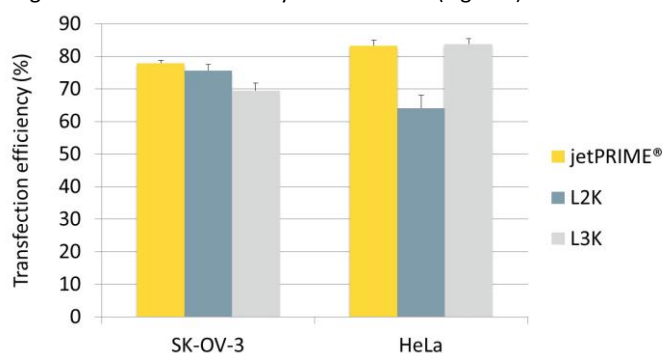


Fig. 1. Transfection efficiency assessed by FACS analysis in various cell lines 24 h following transfection in 24-well plates according to the manufacturer's recommendation for competitors L2K and L3K and 0.5 µg plasmid, 1 µl reagent per well for jetPRIME<sup>®</sup>.

Many other cell lines of various origins, as well as primary cells, are transfected with unusually high percentages (Table 1).

Cell types	Cell lines	Description	Transfection efficiency
Epithelial	B16-F10	Murine melanoma	70-80%
	BNL-C12	Murine normal embryonic hepatocyte	50-60%
	CaCO2	Human colon carcinoma epithelial	20%
	CHO-K1	Chinese hamster ovary	70%
	HCT-116	Human colon carcinoma	70%
	HeLa	Human cervix epitheloid carcinoma	70-90%
	HepG2	Human hepatocarcinoma	50-70%
	Huh-7	Human hepatocarcinoma	30-50%
	MCF-7	Human breast adenocarcinoma	50%
	MCF-10A	Human breast adenocarcinoma	40-50%
	MDCK	Canine kidney epithelial	20%
	PC-3	Human prostate carcinoma	70%
	Vero	African green monkey kidney	50%
Fibroblast	COS-7	African green monkey kidney	60-80%
	HEK-293	Human embryonic kidney fibroblast	80-90%
	MRC-5	Human lung fibroblast	50%
	NIH-3T3	Murine embryonic fibroblast	50-70%
Myeloblast	Raw 264.7	Murine monocyte/macrophage	40-50%
Myoblast	C2C12	Murine myoblast	70-90%
Neuronal	SH-SY5Y	Human neuroblastoma	70-80%
Primary Hepatocytes		Human primary hepatocyte cell	20-30%
Primary Melanocytes		Human primary melanocyte cell	40-50%

Table 1. Transfection efficiency of various cell types using jetPRIME<sup>®</sup>. The percentage of GFP-positive cells was determined by FACS analysis 24 h after transfection.

### ✦ Economical: less reagent and less DNA

jetPRIME<sup>®</sup> is such a powerful *in vitro* transfection reagent that it only requires a small amount of reagent and plasmid DNA (Table 2), making it very economical.

6-well plate			
Reagent	Volume of Reagent per well	Amount of DNA per well	Number of transfections per 1.5 ml vial
jetPRIME <sup>®</sup>	2 - 4 µl	1 - 2 µg	375 - 750
L2K	5 - 12.5 µl	2.5 µg	120 - 300
L3K	3.75 - 7.5 µl	2.5 µg	200 - 400

Table 2. Amounts of reagent and DNA (jetPRIME<sup>®</sup> and competitors) added per well in 6-well plate for transfection according to manufacturers' recommendations.

In addition to reducing costs, using less DNA also minimizes adverse cytotoxic effects triggered by transfection. Hence, jetPRIME<sup>®</sup> is the reagent of choice for high transfection efficiency with excellent cell viability.

### + Better cell viability

jetPRIME<sup>®</sup> is extremely gentle to cells during transfection leading to increased cell viability (Fig. 2) and improved transfection results. Cells transfected with jetPRIME<sup>®</sup> are healthy, while major cytotoxicity is observed with competitor.

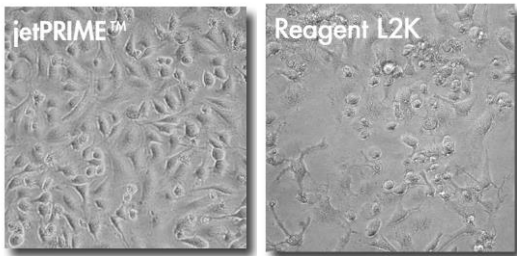


Fig. 2. Phase contrast microscopy of HeLa cells 24 h after transfections performed according to the manufacturer's recommendations for each reagent.

### + Co-transfection of DNA & siRNA

jetPRIME<sup>®</sup> can be used for DNA and siRNA co-transfection experiments. It shows highly efficient gene silencing in a variety of cell lines with very low toxicity. Over 90% silencing is achieved in adherent cells, using 10 nM siRNA (Fig. 3).

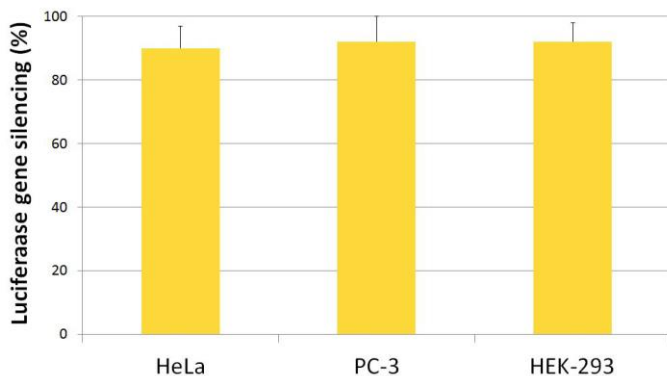


Fig. 3. Exogenous luciferase gene silencing in several cell lines after DNA & siRNA co-transfection using jetPRIME<sup>®</sup> performed with 400 ng pCMV-Luc and 10 nM of siRNA anti-Luc per well in 6-well plates.

### LISTE DES PRODUITS

Référence	Désignation
POL101000027	jetPRIME <sup>®</sup> 0.1 mL
POL101000015	jetPRIME <sup>®</sup> 0.75 mL
POL101000046	jetPRIME <sup>®</sup> 1.5 mL
POL101000001	jetPRIME <sup>®</sup> 5 x 1.5 mL
POL201000003	jetPRIME <sup>®</sup> buffer 60 ml

1.5 ml of jetPRIME<sup>®</sup> transfection reagent is sufficient to perform ca. 375 transfections in 6-well plates. Bulk quantities are available upon request.

### + Excellent gene silencing

jetPRIME<sup>®</sup> leads to over 90% knock-down of endogenous gene expression in a variety of cell lines. For example, jetPRIME<sup>®</sup>-mediated transfection of 10 nM siRNA duplexes targeting endogenous lamin A/C in HeLa cells drastically reduces gene expression to barely detectable level (Fig. 4).

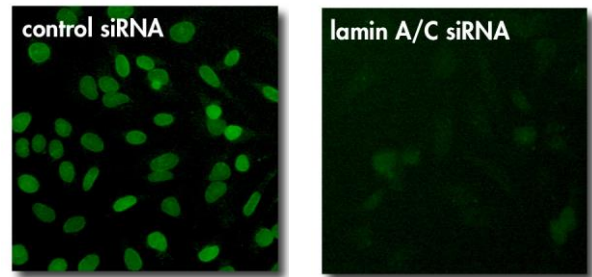


Fig. 4. Endogenous lamin A/C silencing using jetPRIME<sup>®</sup>. HeLa cells were transfected with 10 nM of 21-mer siRNA duplexes matching the lamin A/C sequence. After 48 h, lamin A/C silencing efficiency was determined by immunofluorescence microscopy using an antibody against lamin A/C.

### + Convenient protocol

jetPRIME<sup>®</sup> is an easy-to-use transfection reagent (Fig. 5):

- Fast and easy to scale up and down
- Compatible with serum and antibiotics

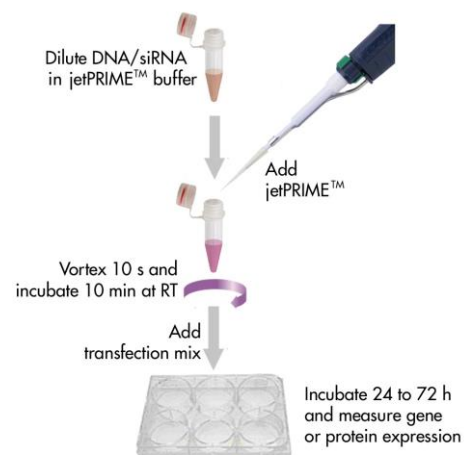


Fig. 5. jetPRIME<sup>®</sup> convenient protocol for DNA, siRNA and co-transfection of DNA and siRNA.

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Cell Transfection Database

