

jetOPTIMUS[®] transfection reagent

Short protocol - DNA Transfection

DAY 0: Cell seeding

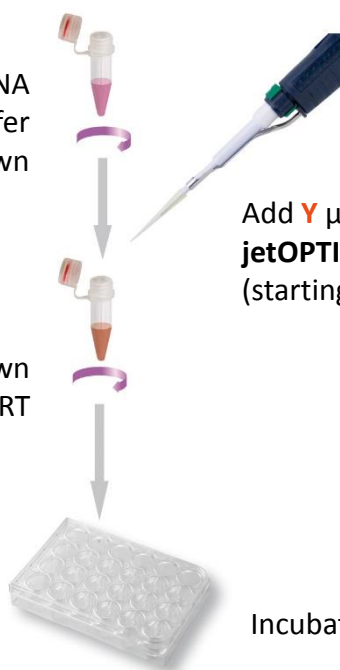
→ Seed cells in **V** mL of serum containing medium according to the table below

Quantities per well, dish or flask

Culture vessel	Number of cells	V = volume of growth medium for cell seeding
96-well	7500 - 25 000	0.125 mL
24-well	40 000 - 100 000	0.5 mL
6-well / 35 mm	150 000 - 400 000	2 mL
60 mm / flask 25 cm ²	200 000 - 850 000	5 mL
100 mm / flask 75 cm ²	1 x 10 ⁶ - 4 x 10 ⁶	10 mL

DAY 1: Transfection using jetOPTIMUS[®] reagent

Dilute **X** µg of DNA in **W** µL of jetOPTIMUS[®] buffer
Vortex 1 second and spin down



- Transfect cells at **60-80% confluency**
- Use **jetOPTIMUS[®] buffer only**
- Perform transfection **in the presence of serum**

Add **Y** µL of jetOPTIMUS[®] reagent (starting ratio 1:1)

Vortex 1 second, spin down and incubate 10 min at RT

Add transfection mix to cells in serum-containing medium

Incubate 24 to 48 h

Quantities per well, dish or flask

Culture vessel	W = volume of jetOPTIMUS [®] buffer	X = amount of DNA added	Y = volume of jetOPTIMUS [®] reagent
96-well	12.5 µL	0.13 µg	0.13 – 0.19 µL
24-well	50 µL	0.5 µg	0.5 – 0.75 µL
6-well / 35 mm	200 µL	2 µg	2 – 3 µL
60 mm / flask 25 cm ²	500 µL	4 µg	4 – 6 µL
100 mm / flask 75 cm ²	1000 µL	10 µg	10 – 15 µL

DAY 2-3: Measure gene expression

See back page for optimization tips

Download complete protocol on www.polyplus-transfection.com/resources/product-literature/

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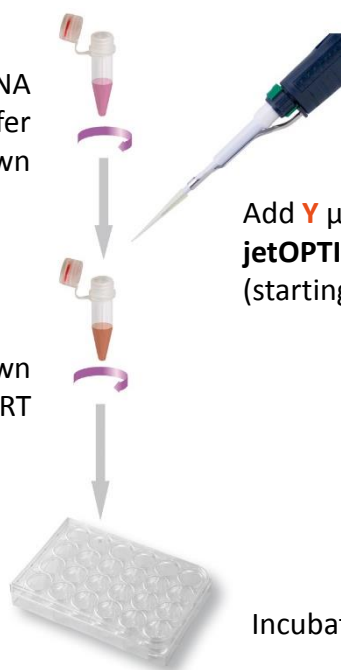
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Short protocol - Optimization Tips

+ Protocol Optimization

- + Test different DNA amounts: X, 0.5 X and 1.5 X
- + Test different DNA/jetOPTIMUS[®] ratios, 1:1 to 1:1.5

Quantities per well, dish or flask

Culture vessel	W = volume of jetOPTIMUS [®] buffer	X = amount of DNA added	Y = volume of jetOPTIMUS [®] reagent
96-well	12.5 µL	0.10 – 0.20 µg	0.10 – 0.30 µL
24-well	50 µL	0.25 – 0.75 µg	0.25 – 1 µL
6-well / 35 mm	200 µL	1 – 3 µg	1 – 4.5 µL
60 mm / flask 25 cm ²	500 µL	2 – 6 µg	2 – 9 µL
100 mm / flask 75 cm ²	1000 µL	5 – 15 µg	5 – 22 µL

For HEK-293 and HeLa cells, you may decrease the DNA amount to 0.5X and use the 1:1 DNA / jetOPTIMUS[®] ratio.

- + Check our online Transfection Database for cell specific protocols at:
www.polyplus-transfection.com/resources/cell-transfection-database/

+ Tips to increase cell viability of sensitive cells

- + Replace medium 4 h after transfection.
- + Decrease DNA amount to 0.5 X while maintaining the DNA/jetOPTIMUS[®] ratio previously used.
- + Analyze transfection at an earlier time point (24 h after transfection instead of 48 h for instance).
- + Perform transfection in reduced serum medium for sensitive cells.
- + Check that the target gene does not affect cell viability.
- + Ensure that cells have been passaged more than twice and less than 20 times prior to transfection. Discard overconfluent cells.

+ Good DNA Transfection Practices

- + Store appropriately jetOPTIMUS[®] (5 ± 3°C).
- + Regularly check for mycoplasma contamination.
- + Use a reporter gene to set up and optimize transfection conditions.
- + Serum quality may drastically affect transfection efficiency. When purchasing a new batch of serum or trypsin, check cell viability as well as transfection efficiency.

Note: Please refer to the complete protocol available online at:
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